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(54) **Title:** AMIDINO DERIVATIVES USEFUL AS NITRIC OXIDE SYNTHASE INHIBITORS

(57) **Abstract**

The current invention discloses amidino derivatives useful as nitric oxide synthase inhibitors.

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AMIDINO DERIVATIVES USEFUL AS NITRIC OXIDE SYNTHASE INHIBITORS

Background of the Invention

This application is a continuation-in-part of U.S. Patent Application Serial No. 08/139,970 filed October 21 1993.

Field of the Invention

The present invention relates to amidino derivatives and their use in therapy, in particular their use as nitric oxide synthase inhibitors.

Related Art

It has been known since the early 1980's that the vascular relaxation brought about by acetycholine is dependent on the presence of the endothelium and this activity was ascribed to a labile humoral factor termed endothelium-derived relaxing factor (EDRF). The activity of nitric oxide (NO) as a vasodilator has been known for well over 100 years and NO is the active component of amylnitrite, glyceryltrinitrite and other nitrovasodilators. The recent identification of EDRF as NO has coincided with the discovery of a biochemical pathway by which NO is synthesized from the amino acid L-arginine by the enzyme NO synthase.

NO is the endogenous stimulator of the soluble guanylate cyclase and is involved in a number of biological actions in addition to endothelium-dependent relaxation including cytotoxicity of phagocytic cells and cell-to-cell communication in the central nervous system (see Moncada et al, Biochemical Pharmacology, 38, 1709-1715 (1989) and Moncada et al, Pharmacological Reviews, 43, 109-142 (1991). It is now thought that excess NO

production may be involved in a number of conditions, particularly conditions which involve systemic hypotension such as toxic shock and therapy with certain cytokines.

The synthesis of NO from L-arginine can be inhibited by the L-arginine analogue, L-N-monomethyl-arginine (L-NMMA) and the therapeutic use of L-NMMA for the treatment of toxic shock and other types of systemic hypertension has been proposed (WO 91/04024 and GB-A-2240041). The therapeutic use of certain other NO synthase inhibitors apart from L-NMMA for the same purpose has also been proposed in WO 91/04024 and in EP-A-0446699.

It has recently become apparent that there are at least three types of NO synthase as follows:

(i) a constitutive, Ca^{++}/calmodulin dependent enzyme, located in the endothelium, that releases NO in response to receptor or physical stimulation.

(ii) a constitutive, Ca^{++}/calmodulin dependent enzyme, located in the brain, that releases NO in response to receptor or physical stimulation.

(iii) a Ca^{++} independent enzyme which is induced after activation of vascular smooth muscle, macrophages, endothelial cells, and a number of other cells by endotoxin and cytokines. Once expressed this inducible NO synthase synthesizes NO for long periods.

The NO released by the constitutive enzymes acts as a transduction mechanism underlying several physiological responses. The NO produced by the inducible enzyme is a cytotoxic molecule for tumor cells and invading microorganisms. It also appears that the adverse effects of excess NO production, in particular pathological vasodilation and tissue damage, may result largely from the effects of NO synthesized by the inducible NO synthase.

There is also a growing body of evidence that NO may

be involved in the degeneration of cartilage which takes place in certain conditions such as arthritis and it is also known that NO synthesis is increased in rheumatoid arthritis. Accordingly, further conditions in which there is an advantage in inhibiting NO production from L-arginine include autoimmune and/or inflammatory conditions affecting the joints, for example arthritis.

Conditions in which there is an advantage in inhibiting NO production from L-arginine include systemic hypotension associated with septic and/or toxic shock induced by a wide variety of agents; therapy with cytokines such as TNF, IL-1 and IL-2; and as an adjuvant to short term immunosuppression in transplant therapy. Further conditions in which there is an advantage in inhibiting NO production from L-arginine include autoimmune diseases and/or inflammatory conditions such as those affecting the joints, for example arthritis or inflammatory bowel disease, cardiovascular ischemia, diabetes, hyperalgesia (allodynia) cerebral ischemia (Both focal ischemia, thrombotic stroke and global ischemia, secondary to cardiac arrest) and other CNS disorders mediated by NO.

Some of the NO synthase inhibitors proposed for therapeutic use so far, and in particular L-NMMA, are non-selective in that they inhibit both the constitutive and the inducible NO synthase. Use of such a non-selective NO synthase inhibitor requires that great care be taken in order to avoid the potentially serious consequences of over-inhibition of the constitutive NO-synthase including hypertension and possible thrombosis and tissue damage. In particular, in the case of the therapeutic use of L-NMMA for the treatment of toxic shock it has been recommended that the patient must be subject to continuous blood pressure monitoring throughout the treatment. Thus, while non-selective NO synthase inhibitors have therapeutic utility provided

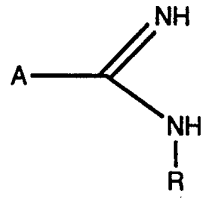
that appropriate precautions are taken, NO synthase inhibitors which are selective in the sense that they inhibit the inducible NO synthase to a considerably greater extent than the constitutive NO synthase would be of even greater therapeutic benefit and easier to use.

WO 94/12165, WO 94/14780, Wo93/13055, EP 0446699A1 and U.S. Patent No. 5,132,453 disclose compounds that inhibit nitric oxide synthesis and preferentially inhibit the inducible isoform of nitric oxide synthase. The disclosures of which are hereby incorporated by reference in their entirety as if written herein.

Summary of the Invention

In a broad aspect, the present invention is directed to inhibiting or modulating nitric oxide synthesis in a subject in need of such inhibition or modulation by administering a compound which preferentially inhibits or modulates the inducible isoform of nitric oxide synthase over the constitutive isoforms of nitric oxide synthase. It is also another object of the present invention to lower nitric oxide levels in a subject in need of such lowering.

The invention relates a method of inhibiting nitric oxide synthesis in a subject in need of such inhibition by administering a therapeutically effective amount of a compound having the formula:



(I)

and salts, and pharmaceutically acceptable ester and prodrugs thereof, wherein:

A is hydrogen, lower alkyl , lower alkenyl , lower alkynyl group, alkylthioalkyl group, alkyloxyalkyl group,

alkylsulfonylalkyl group, cycloalkyl group, bicycloalkyl group, cycloalkenyl group, cycloalkylalkyl group, phenylalkyl group, phenylalkenyl group, biphenylalkyl group, heterocyclic group, biaryl group, or aryl wherein each said radical may optionally be substituted by one or more of the following substituents such as alkyl, alkoxy, hydroxy, halogen, nitro, cyano, haloalkyl, carboxylic, carboxamide, amino, alkylamino or dialkylamino; and

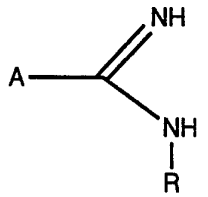
R is H, OH or lower alkyl group.

The invention further relates to pharmaceutical compositions comprising a compound of formula (I) for use in the above method. Such compounds and compositions have usefulness as inhibitors of nitric oxide synthase. Conditions in which there is an advantage in inhibiting NO production include systemic hypotension associated with septic and/or toxic shock induced by a wide variety of agents, therapy with cytokines such as TNF, IL-1 and IL-2; autoimmune and/or inflammatory diseases affecting the joints such as arthritis, diabetes and inflammatory bowel disease.

Compounds and compositions defined above have usefulness as inhibitors of nitric oxide synthase. These compounds also prefentially inhibit the inducible form over the constitutive form by at least 3 fold.

Detailed Description of the Invention

A preferred embodiment of the present invention is a method using a pharmaceutical composition including a compound of the formula (I)

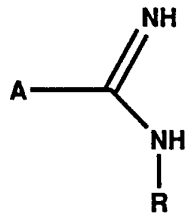


(I)

A is hydrogen, lower alkyl of 1 to about 10 carbon atoms, lower alkenyl group of 2 to about 6 carbon atoms, lower alkynyl group of 2 to about 6 carbon atoms, alkylthioalkyl group of 2 to about 6 carbon atoms, alkyloxyalkyl of 2 to about 6 carbon atoms, alkylsulfonylalkyl group of 2 to about 6 carbon atoms, cycloalkyl group of 3 to about 8 carbon atoms, bicycloalkyl group of 6 to about 10 carbon atoms, cycloalkenyl group of 3 to about 8 carbon atoms, cycloalkylalkyl group of 4 to about 10 carbon atoms, phenylalkyl group, phenylalkenyl group, biphenylalkyl group, heterocyclic group, or aryl substituted heterocyclic groups and which each group may be optionally be substituted by one or more of the following substituents: lower alkyl of 1 to about 4 carbon atoms, alkoxy of 1 to about 4 carbon atoms, hydroxy, halogen, nitro, cyano, haloalkyl, carboxyl, carboxamide, amino, monoalkylamino or dialkylamino; and

R is H, OH or lower alkyl group of 1 to about 6 carbon atoms.

Another preferred embodiment of the present invention is a compound of the formula (I)



(I)

A is lower alkyl of 2 to about 10 carbon atoms,

lower alkenyl of 2 to 6 carbon atoms, cycloalkyl group of 3 to 8 carbon atoms, bicycloalkyl group of 7 carbon atoms, alkylthioalkyl group of 2 to about 6 carbon atoms, alkyloxyalkyl of 2 to about 6 carbon atoms, alkylsulfonylalkyl group of 2 to about 6 carbon atoms , heterocyclic group, or aryl substituted heterocyclic group and which each may be optionally be substituted by one or more of the following lower alkyl, alkoxy, haloalkyl or halogen, nitro; and

R is H, OH or lower alkyl group of 1 to about 6 carbon atoms.

The present invention includes compounds of formula (I) in the form of salts, in particular acid addition salts. Suitable salts include those formed with both organic and inorganic acids. Such acid addition salts will normally be pharmaceutically acceptable although salts of non-pharmaceutically acceptable salts may be of utility in the preparation and purification of the compound in question. Thus, preferred salts include those formed from hydrochloric, hydrobromic, sulphuric, citric, tartaric, phosphoric, lactic, pyruvic, acetic, succinic, oxalic, fumaric, maleic, oxaloacetic, methanesulphonic, ethanesulphonic, p-toluenesulphonic, benzenesulphonic and isethionic acids. Salts of the compounds of formula (I) can be made by reacting the appropriate compound in the form of the free base with the appropriate acid.

While it may be possible for the compounds of formula (I) to be administered as the raw chemical, it is preferable to present them as a pharmaceutical formulation. According to a further aspect, the present invention provides a pharmaceutical formulation comprising a compound of formula (I) or a pharmaceutically acceptable salt or solvate thereof, together with one or more pharmaceutically acceptable

carriers thereof and optionally one or more other therapeutic ingredients. The carrier(s) must be "acceptable" in the sense of being compatible with the other ingredients of the formulation and not deleterious to the recipient thereof.

The formulations include those suitable for oral, parenteral (including subcutaneous, intradermal, intramuscular, intravenous and intraarticular), rectal and topical (including dermal, buccal, sublingual and intraocular) administration although the most suitable route may depend upon for example the condition and disorder of the recipient. The formulations may conveniently be presented in unit dosage form and may be prepared by any of the methods well known in the art of pharmacy. All methods include the step of bringing into association a compound of formula (I) or a pharmaceutically acceptable salt or solvate thereof ("active ingredient") with the carrier which constitutes one or more accessory ingredients. In general, the formulations are prepared by uniformly and intimately bringing into association the active ingredient with liquid carriers or finely divided solid carriers or both and then, if necessary, shaping the product into the desired formulation.

Formulations of the present invention suitable for oral administration may be presented as discrete units such as capsules, cachets or tablets each containing a predetermined amount of the active ingredient; as a powder or granules; as a solution or a suspension in an aqueous liquid or a non-aqueous liquid; or as an oil-in-water liquid emulsion or a water-in-oil liquid emulsion. The active ingredient may also be presented as a bolus, electuary or paste.

A tablet may be made by compression or moulding, optionally with one or more accessory ingredients. Compressed tablets may be prepared by compressing in a suitable machine the active ingredient in a free-flowing form such as a powder or granules, optionally mixed with a binder, lubricant, inert diluent, lubricating, surface active or dispersing agent. Moulded tablets may be made by moulding in a suitable machine a mixture of the

powdered compound moistened with an inert liquid diluent. The tablets may optionally be coated or scored and may be formulated so as to provide slow or controlled release of the active ingredient therein.

Formulations for parenteral administration include aqueous and non-aqueous sterile injection solutions which may contain anti-oxidants, buffers, bacteriostats and solutes which render the formulation isotonic with the blood of the intended recipient; and aqueous and non-aqueous sterile suspensions which may include suspending agents and thickening agents. The formulations may be presented in unit-dose or multi-dose containers, for example sealed ampoules and vials, and may be stored in a freeze-dried (lyophilized) condition requiring only the addition of the sterile liquid carrier, for example, saline, water-for-injection, immediately prior to use. Extemporaneous injection solutions and suspensions may be prepared from sterile powders, granules and tablets of the kind previously described.

Formulations for rectal administration may be presented as a suppository with the usual carriers such as cocoa butter or polyethylene glycol.

Formulations for topical administration in the mouth, for example buccally or sublingually, include lozenges comprising the active ingredient in a flavoured basis such as sucrose and acacia or tragacanth, and pastilles comprising the active ingredient in a basis such as gelatin and glycerin or sucrose and acacia.

Preferred unit dosage formulations are those containing an effective dose, as hereinbelow recited, or an appropriate fraction thereof, of the active ingredient.

It should be understood that in addition to the

ingredients particularly mentioned above, the formulations of this invention may include other agents conventional in the art having regard to the type of formulation in question, for example those suitable for oral administration may include flavouring agents.

The compounds of the invention may be administered orally or via injection at a dose of from 0.1 to 500 mg/kg per day. The dose range for adult humans is generally from 5mg to 2g/day. Tablets or other forms of presentation provided in discrete units may conveniently contain an amount of compound of the invention which is effective at such dosage or as a multiple of the same, for instance, units containing 5mg to 500mg, usually around 10mg to 200mg.

The compounds of formula (I) are preferably administered orally or by injection (intravenous or subcutaneous). The precise amount of compound administered to a patient will be the responsibility of the attendant physician. However, the dose employed will depend on a number of factors, including the age and sex of the patient, the precise disorder being treated, and its severity. Also, the route of administration may vary depending on the condition and its severity.

As utilized herein, the term "lower alkyl", alone or in combination, means an acyclic alkyl radical containing from 1 to about 20, preferably from 1 to about 10 carbon atoms and more preferably 1 to about 6 carbon atoms. Examples of such radicals include methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl, sec-butyl, tert-butyl, pentyl, iso-amyl, hexyl, octyl and the like.

The term "lower alkenyl" refers to an unsaturated acyclic hydrocarbon radical in so much as it contains at least one double bond. Such radicals containing from about 2 to about 20 carbon atoms, preferably from about 2

to about 10 carbon atoms and more preferably 2 to about 6 carbon atoms. Examples of suitable alkenyl radicals include propenyl, buten-1-yl, isobutenyl, penten-1-yl, 2-2-methylbuten-1-yl, 3-methylbuten-1-yl, hexen-1-yl, hepten-1-yl, and octen-1-yl, and the like.

The term "lower alkynyl" refers to an unsaturated acyclic hydrocarbon radical in so much as it contains one or more triple bonds, such radicals containing from about 2 to about 20 carbon atoms, preferably having from about 2 to about 10 carbon atoms and more preferably having 2 to about 6 carbon atoms. Examples of suitable alkynyl radicals include ethynyl, propynyl, butyn-1-yl, butyn-2-yl, pentyn-1-yl, pentyn-2-yl, 3-methylbutyn-1-yl, hexyn-1-yl, hexyn-2-yl, hexyn-3-yl, 3,3-dimethylbutyn-1-yl radicals and the like.

The term "heterocyclic radical" means a saturated or unsaturated cyclic hydrocarbon radical with 4 to about 10 carbon atoms, preferably about 5 to about 6; wherein 1 to about 3 carbon atoms are replaced by nitrogen, oxygen or sulfur. The "heterocyclic radical" may be fused to an aromatic hydrocarbon radical. Suitable examples include pyrrolyl, pyridinyl, pyrazolyl, triazolyl, pyrimidinyl, pyridazinyl, oxazolyl, thiazolyl, imidazolyl, indolyl, thiophenyl, furanyl, tetrazolyl, 2-pyrrolinyl, 3-pyrrolinyl, pyrrolindinyl, 1,3-dioxolanyl, 2-imidazolinyl, imidazolidinyl, 2-pyrazolinyl, pyrazolidinyl, isoxazolyl, isothiazolyl, 1,2,3-oxadiazolyl, 1,2,3-triazolyl, 1,3,4-thiadiazolyl, 2H-pyranyl, 4H-pyranyl, piperidinyl, 1,4-dioxanyl, morpholinyl, 1,4-dithianyl, thiomorpholinyl, pyrazinyl, piperazinyl, 1,3,5-triazinyl, 1,3,5-trithianyl, benzo(b)thiophenyl, benzimidazonyl, quinolinyl, and the like.

The term "Aryl" means an aromatic hydrocarbon radical of 4 to about 16 carbon atoms, preferably 6 to about 12 carbon atoms, more preferably 6 to about 10

carbon atoms. Examples of suitable aromatic hydrocarbon radicals include phenyl, naphthyl, and the like.

The terms "Cycloalkyl" or "cycloalkenyl" means an "alicyclic radical in a ring with 3 to about 10 carbon atoms, and preferably from 3 to about 6 carbon atoms. Examples of suitable alicyclic radicals include cyclopropyl, cyclopropylenyl, cyclobutyl, cyclopentyl, cyclohexyl, 2-cyclohexen-1-ylenyl, cyclohexenyl and the like.

The term "alkoxy", alone or in combination, means an alkyl ether radical wherein the term alkyl is as defined above and most preferably containing 1 to about 4 carbon atoms. Examples of suitable alkyl ether radicals include methoxy, ethoxy, n-propoxy, isopropoxy, n-butoxy, iso-butoxy, sec-butoxy, tert-butoxy and the like.

The term "halogen" means fluorine, chlorine, bromine or iodine.

The term "prodrug" refers to a compound that is made more active *in vivo*.

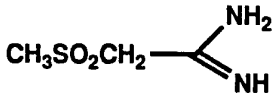
As used herein, reference to "treatment" of a patient is intended to include prophylaxis.

All references, patents or applications, U.S. or foreign, cited in the application are hereby incorporated by reference as if written herein.

The invention is illustrated by the following examples. Some of the compounds disclosed are publicly available from the source cited. Additional compounds of this invention have been described in publications as indicated or have been fully described herein.

Example 1

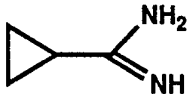
2-Methylsulfonylacetamidine hydrochloride



To a 250mL flask was added 5g (0.042mol) of 2-methylsulfonylacetonitrile and 75mL of anhydrous ethanol. This solution was cooled to 0°C in an ice bath while anhydrous HCl was bubbled in until saturated. This mixture was allowed to warm very slowly with constant stirring over a 24 hour period. The reaction mixture was concentrated to a reduced volume, diluted with ethyl ether and filtered to afford 5.5g of the ethyl imidate as a white solid. The imidate was added to 20mL of anhydrous ethanol and cooled to 0°C. To this reaction mixture was added 80mL of ethanol previously saturated with anhydrous ammonia. This mixture was capped and allowed to stir for three days. The reaction mixture was then concentrated to a reduced volume, diluted with ethyl ether and filtered to afford 4.7g (65%) of the 2-Methylsulfonylacetamidine Hydrochloride as a white solid, mp 186-194°C.

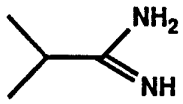
Example 2

Cyclopropylcarbamidine; Lancaster Synthesis Inc.



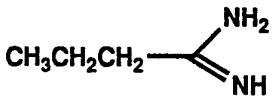
Example 3

Isopropylcarbamidine; C. R. Hauser and C. J. Eby, J. Am. Chem. Soc. 79, 725-727 (1957).



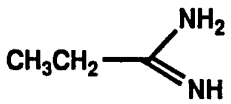
Example 4

1-Propylcarbamidine; R. Almquist, R. A. Huggins and R. A. Woodbury, J. Pharmacol. 89, 271-288 (1947).



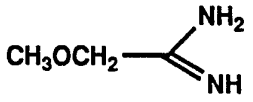
Example 5

Ethylcarbamidine; ibid.



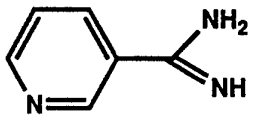
Example 6

2-Methoxyacetamidine; BELG. 645062, 1964.



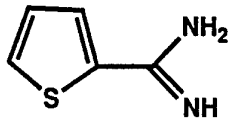
Example 7

3-Amidinopyridine; Ryan Scientific.



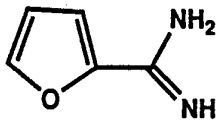
Example 8

2-Amidinothiophene; Ryan Scientific.



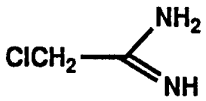
Example 9

2-Amidinofuran; T. J. Schwan and K.O. Ellis, J. Pharm. Sci. 64(2), 337-338 (1975).



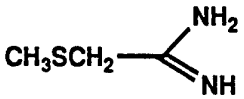
Example 10

2-Chloroacetamidine; Transworld Chemical Inc.

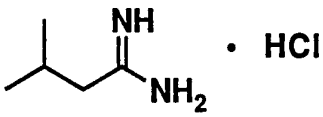


Example 11

2-Methylmercaptoacetamide; GER. 2,928,185, F. Maurer and I. Hammann (1901).



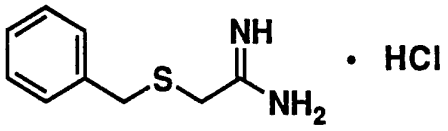
Example 12 : Isobutylcarbamidine hydrochloride; J. Gen. Chem. 14, 280-291 (1944)



Prepared as in example 1 from isovalerylnitrile to afford the title compound as a white solid. 1H-NMR(D2O) 0.9 (d, 6H), 2.05 (m, 1H), 2.25 (d, 2H); Mass Spectra, M+H=101; Elemental analysis Calcd. for C5H13N2Cl2 + 1/10 N1H4Cl1:

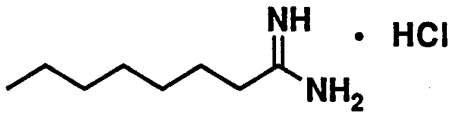
C, 42.30; H, 9.51; N,20.72. Found C, 42.45, H, 9.47, N, 20.69.

Example 13 : Benzylthioacetamidine hydrochloride; Fr. 1,429,279



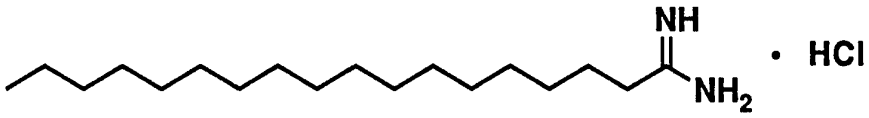
Prepared as in example 1 from benzylthioacetonitrile to afford the title compound as an off-white solid. 1H-NMR(D2O) 3.35 (s, 2H), 3.73 (s, 2H), 7.22 (m, 5H); Mass Spectra, M+H=181.

Example 14 : Heptlycarbamidine hydrochloride; Chem. Abst. 41:5468i



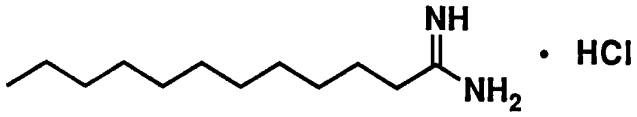
Prepared as in example 1 from heptylcyanide to afford the title compound as a white solid. Mass Spectra, M+H=143.

Example 15 : Heptadecylcarbamidine hydrochloride; J. Chem. Soc. 738-742 (1947)



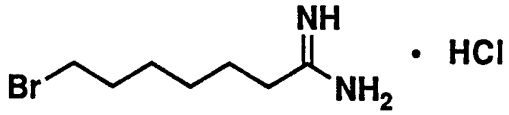
Prepared as in example 1 from heptadecylcyanide to afford the title compound as a white solid. Mass Spectra, M+H=283.

Example 16 : Undecylcarbamidine hydrochloride; Chem. Abst. 51:12808f

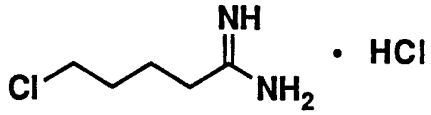


Prepared as in example 1 from undecylcyanide to afford the title compound as a white solid. Mass Spectra, M+H=199.

Example 17 : 1-Bromo-6-carbamidylhexane hydrochloride Prepared as in example 1 from 1-bromo-6-cyanohexane to afford the title compound as a white solid. Mass Spectra, M+H=207.

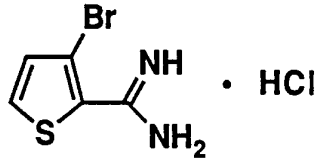


Example 18 : X-10191 1-Chloro-4-carbamidylbutane hydrochloride



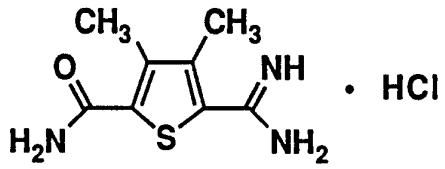
Prepared as in example 1 from 1-chloro-4-cyanobutane to afford the title compound as a white solid. Mass Spectra, M+H=135.

Example 19 : 2-Carbamidyl-3-bromothiophene hydrochloride



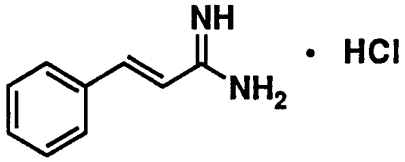
Prepared as in example 1 from 2-cyano-3-bromothiophene to afford the title compound as a white solid. Mass Spectra, M+H=205.

Example 20 : 2-Amidyl, 3,4-dimethyl, 5-carbamidylthiophene hydrochloride



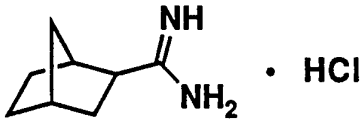
Prepared as in example 1 from 2-amidyl, 3,4-dimethyl, 5-cyanothiophene to afford the title compound as a white solid. Mass Spectra, M+H=198.

Example 21 : Styrylcarbamidine hydrochloride; J. Am. Chem. Soc. 78, 1434-7 (1956)



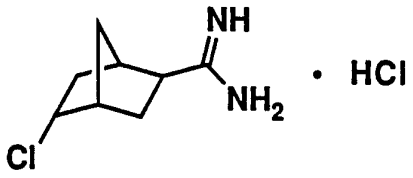
Prepared as in example 1 from cinnamonitrile to afford the title compound as a white solid. Mass Spectra, M+H=147.

Example 22 : 2-carbamidylnorbornane hydrochloride



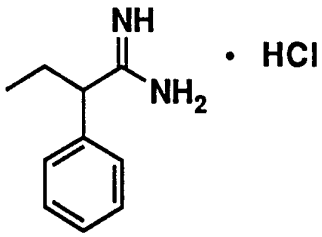
Prepared as in example 1 from 2-norbornanecarbonitrile to afford the title compound as a white solid. Mass Spectra, M+H=139.

Example 23 : 2-Chloro-5-carbamidylnorbornane hydrochloride



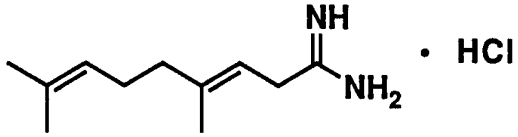
Prepared as in example 1 from 2-Chloro-5-cyanonorbornane to afford the title compound as a white solid. Mass Spectra, M+H=173.

Example 24 : 2-Phenylbutyramidine hydrochloride; Compt. Rend. 246, 2905-6 (1958)



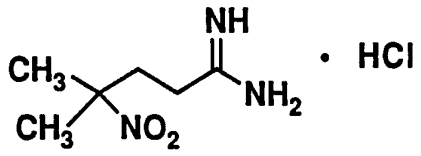
Prepared as in example 1 from 2-phenylbutyronitrile to afford the title compound as a white solid. Mass Spectra, M+H=163.

Example 25 : 2,6-Dimethyl, 7-carbamidylhepta-2-6-diene hydrochloride



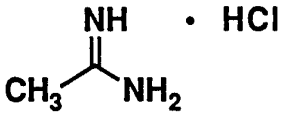
Prepared as in example 1 from 2,6-Dimethyl, 7-cyanohepta-2-6-diene to afford the title compound as a white solid. Mass Spectra, M+H=167.

Example 26 : 2-Methyl, 2-nitro, 4-carbamidylbutane hydrochloride

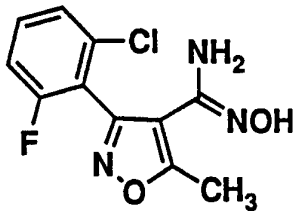


Prepared as in example 1 from 2-Methyl, 2-nitro, 4-cyanobutane to afford the title compound as a white solid. Mass Spectra, M+H=160.

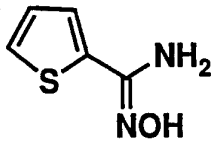
Example 27 : Acetamidine; Aldrich



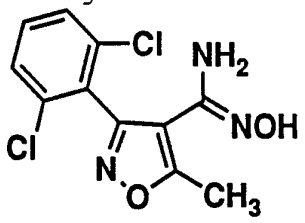
Example 28 : 3-(2-chloro,6-fluorophenyl), 4-carbamidoxime, 5-methylisoxazole; Maybridge Chemical Co. Ltd.



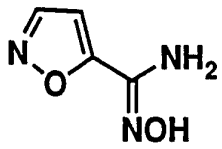
Example 29 : 2-carboxamidoximylthiophene; Boll. sci. fac. chim. ind. Bologna 15(3), 57-62 (1957).



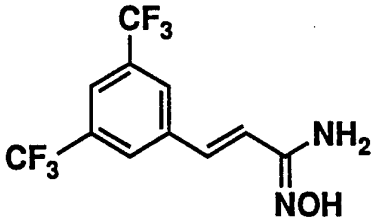
Example 30 : 3-(2,6-dichlorophenyl), 4-carbamidoxime, 5-methylisoxazole; Maybridge Chemical Co. Ltd.



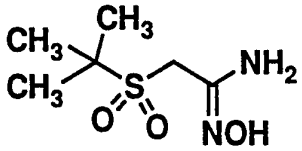
Example 31 : 5-carboxamidoximylisoxazole; Maybridge Chemical Co. Ltd.



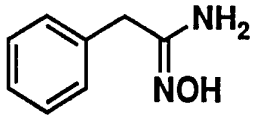
Example 32 : 3,5-bistrifluoromethyl-styrylcarboxamidoxime; Maybridge Chemical Co. Ltd.



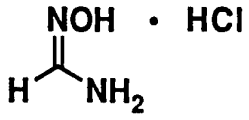
Example 33 : t-Butylsulfonylacetamidoxime; Maybridge Chemical Co. Ltd.



Example 34 : Phenylacetamidoxime; Aldrich



Example 35 Formamidoxime; Aldrich



Biological Data

The activity of the above listed compounds as NO synthase inhibitors has been determined in the following assays:

Citrulline Assay for Nitric Oxide Synthase

Nitric oxide synthase activity was measured by monitoring the conversion of [3H]-arginine to [3H]-citrulline. Mouse inducible nitric oxide synthase (miNOS) was prepared from an extract of LPS-treated RAW 264.7 cells and partially purified by DEAE-Sepharose chromatography. Rat brain constitutive nitric oxide synthase (rnNOS) was preparedfrom an extract of rat cerebellum and partially purified by DEAE-Sepharose chromatography. Enzyme and inhibitors were incubated at 37°C for 15 minutes in a reaction volume of 100 µL with the following components added to start the reaction: 50 mM Tris (pH 7.6), 1 mg/ml bovine serum albumin, 1 mM DTT, 2 mM CaCl2, 10 µM FAD, 10 µM tetrahydrobiopterin, 30 µM L-arginine containing L-[2,3-3H]-arginine at 300 cpm/pmole and 1 mM NADPH. For constitutive NOS, 50 nM calmodulin was also added. The reaction was terminated by

addition of cold stop buffer containing 10 mM EGTA, 100 mM HEPES, pH 5.5 and 1 mM citrulline. [3H]-Citrulline was separated by chromatography on Dowex 50W X-8 cation exchange resin and radioactivity determined with a liquid scintillation counter.

Raw Cell Nitrite Assay

RAW 264.7 cells are plated to confluency on a 96-well tissue culture plate grown overnight (17h) in the presence of LPS to induce NOS. A row of 3-6 wells were left untreated and served as controls for subtraction of nonspecific background. The media was removed from each well and the cells are washed twice with Krebs-Ringers-Hepes (25mM, pH 7.4) with 2 mg/ml glucose. The cells are then placed on ice and incubated with 50 µL of buffer containing L-arginine (30 µM) +/- inhibitors for 1h. The assay is initiated by warming the plate to 37°C in a water bath for 1h. Production of nitrite by intracellular iNOS is linear with time. To terminate the cellular assay, the plate of cells is placed on ice and the nitrite-containing buffer removed and analyzed for nitrite using a previously published fluorescent determination for nitrite. All values are the average of triplicate wells and are compared to a background-subtracted induced set of cells (100% value).

TABLE I

Compound	iNOS IC_{50} [μM]	cNOS	Raw Cell IC_{50} [μM]
Example 1	39% @10 μM		
Example 2	2.3	10	46
Example 3	27	31	
Example 4	2.0	7.3	
Example 5	16	44	
Example 6	39% @10 μM		
Example 7	35% @10 μM		
Example 8	1.0	1.3	7.0
Example 9	3.0	3.0	158
Example 10	33% @10 μM		
Example 11	2.0	7.0	60

Compound	iNOS IC_{50} [μM]	cNOS	Raw Cell IC_{50} [μM]
Example 12	35% @10 μM*	26% @10 μM	
Example 13	43% @100 μM*	35% @100 μM**	
Example 14	0% @100 μM*	0% @100 μM**	
Example 15	0% @100 μM*	51% @100 μM**	
Example 16	2% @100 μM*	65% @100 μM**	
Example 17	10% @100 μM*	0% @100 μM**	
Example 18	74% @10 μM*	45% @10 μM**	
Example 19	30% @100 μM*	18% @100 μM**	
Example 20	0% @100 μM*	0.4% @100 μM**	
Example 21	18% @100 μM*	80% @100 μM**	
Example 22	7% @100 μM*	1% @100 μM**	
Example 23	57% @100 μM*	22% @100 μM**	
Example 24	0% @100 μM*	0% @100 μM**	
Example 25	1% @100 μM*	0% @100 μM**	
Example 26	10% @100 μM*	2% @100 μM**	

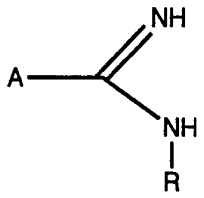
* hiNOS Data

** hecNOS Data

From the foregoing description, one skilled in the art can easily ascertain the essential characteristics of this invention, and without departing from the spirit and scope thereof, can make various changes and modifications of the invention to adapt it to various usages and conditions.

WHAT IS CLAIMED IS:

1. A method of inhibiting nitric oxide synthesis in a subject in need of such inhibition by administering a therapeutically effective amount of a compound having the formula:



(I)

and salts, and pharmaceutically acceptable ester and prodrugs thereof, wherein:

A is hydrogen, lower alkyl , lower alkenyl , lower alkynyl group, alkylthioalkyl group, alkyloxyalkyl group, alkylsulfonylalkyl group, cycloalkyl group, bicycloalkyl group, cycloalkenyl group, cycloalkylalkyl group,, phenylalkyl group, phenylalkenyl group, biphenylalkyl group, heterocyclic group, biaryl group, or aryl wherein each said radical may optionally be substituted by one or more of the following substituents such as alkyl, alkoxy, hydroxy, halogen, nitro, cyano, haloalkyl, carboxylic, carboxamide, amino, alkylamino or dialkylamino; and

R is H, OH or lower alkyl group.

2. The method of inhibiting nitric oxide synthesis as recited in Claim 1 wherein;

A is hydrogen, lower alkyl of 1 to about 10 carbon atoms, lower alkenyl group of 2 to about 6 carbon atoms, lower alkynyl group of 2 to about 6 carbon atoms, alkylthioalkyl group of 2 to about 6 carbon atoms, alkyloxyalkyl of 2 to about 6 carbon atoms,

alkylsulfonylalkyl group of 2 to about 6 carbon atoms, cycloalkyl group of 3 to about 8 carbon atoms, bicycloalkyl group of 6 to about 10 carbon atoms, cycloalkenyl group of 3 to about 8 carbon atoms, cycloalkylalkyl group of 4 to about 10 carbon atoms, phenylalkyl group, phenylalkenyl group, biphenylalkyl group, heterocyclic group, or aryl substituted heterocyclic groups and which each group may be optionally be substituted by one or more of the following substituents: lower alkyl of 1 to about 4 carbon atoms, alkoxy of 1 to about 4 carbon atoms, hydroxy, halogen, nitro, cyano, haloalkyl, carboxyl, carboxamide, amino, monoalkylamino or dialkylamino; and

R is H, OH or lower alkyl group of 1 to about 6 carbon atoms.

3. The method of inhibiting nitric oxide synthesis as recited in Claim 1 wherein:

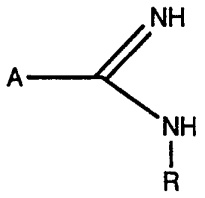
A is lower alkyl of 2 to about 10 carbon atoms, lower alkenyl of 2 to 6 carbon atoms, cycloalkyl group of 3 to 8 carbon atoms, bicycloalkyl group of 7 carbon atoms, alkylthioalkyl group of 2 to about 6 carbon atoms, alkyloxyalkyl of 2 to about 6 carbon atoms, alkylsulfonylalkyl group of 2 to about 6 carbon atoms , heterocyclic group, or aryl substituted heterocyclic group and which each may be optionally be substituted by one or more of the following lower alkyl, alkoxy, haloalkyl or halogen, nitro; and

R is H, OH or lower alkyl group of 1 to about 6 carbon atoms.

4. The method of inhibiting nitric oxide synthesis as recited in Claim 1 wherein the compound is selected from the group consisting of:
cyclopropylcarbamidine; isopropylcarbamidine;

1-propylcarbamidine; ethylcarbamidine; 2-methoxyacetamidine; 3-amidinopyridine; 2-amidinothiophene; 2-amidinofuran; 2-chloroacetamidine, 2-methylmercaptoacetamide, Isobutylcarbamidine hydrochloride, Benzylthioacetamidine hydrochloride, Heptadecylcarbamidine hydrochloride, Undecylcarbamidine hydrochloride, 1-Bromo-6-carbamidylhexane hydrochloride, 1-Chloro-4-carbamidylbutane hydrochloride, 2-Carbamidyl-3-bromothiophene hydrochloride, 2-Amidyl, 3,4-dimethyl, 5-carbamidylthiophene hydrochloride, Styrylcarbamidine hydrochloride, 2-carbamidylnorbornane hydrochloride, 2-Chloro-5-carbamidylnorbornane hydrochloride, 2-Phenylbutyramidine hydrochloride, 2,6-Dimethyl, 7-carbamidylhepta-2-6-diene hydrochloride, 2-Methyl, 2-nitro, 4-carbamidylbutane hydrochloride, Acetamidine, 3-(2-chloro,6-fluorophenyl), 4-carbamidoxime, 5-methylisoxazole, 2-carboxamidoximylthiophene, 3-(2,6-dichlorophenyl), 4-carbamidoxime, 5-methylisoxazole, 5-carboxamidoximylisoxazole, 3,5-bistrifluoromethylstyrylcarboxamidoxime, t-Butylsulfonylacetamidoxime, Phenylacetamidoxime, and Formamidoxime.

5. A method of selectively inhibiting nitric oxide synthesis produced by inducible NO synthase over nitric oxide produced by the constitutive forms of NO synthase in a subject in need of such selective inhibition by administering a therapeutically effective amount of a compound having the formula:



(I)

and salts, and pharmaceutically acceptable ester and

prodrugs thereof, wherein:

A is hydrogen, lower alkyl , lower alkenyl , lower alkynyl group, alkylthioalkyl group, alkyloxyalkyl group, alkylsulfonylalkyl group, cycloalkyl group, bicycloalkyl group, cycloalkenyl group, cycloalkylalkyl group,, phenylalkyl group, phenylalkenyl group, biphenylalkyl group, heterocyclic group, biaryl group, or aryl wherein each said radical may optionally be substituted by one or more of the following substituents such as alkyl, alkoxy, hydroxy, halogen, nitro, cyano, haloalkyl, carboxylic, carboxamide, amino, alkylamino or dialkylamino; and

R is H, OH or lower alkyl group.

6. The method as recited in Claim 5 wherein;

A is hydrogen, lower alkyl of 1 to about 10 carbon atoms, lower alkenyl group of 2 to about 6 carbon atoms, lower alkynyl group of 2 to about 6 carbon atoms, alkylthioalkyl group of 2 to about 6 carbon atoms, alkyloxyalkyl of 2 to about 6 carbon atoms, alkylsulfonylalkyl group of 2 to about 6 carbon atoms, cycloalkyl group of 3 to about 8 carbon atoms, bicycloalkyl group of 6 to about 10 carbon atoms, cycloalkenyl group of 3 to about 8 carbon atoms, cycloalkylalkyl group of 4 to about 10 carbon atoms, phenylalkyl group, phenylalkenyl group, biphenylalkyl group, heterocyclic group, or aryl substituted heterocyclic groups and which each group may be optionally be substituted by one or more of the following substituents: lower alkyl of 1 to about 4 carbon atoms, alkoxy of 1 to about 4 carbon atoms, hydroxy, halogen, nitro, cyano, haloalkyl, carboxyl, carboxamide, amino, monoalkylamino or dialkylamino; and

R is H, OH or lower alkyl group of 1 to about 6 carbon atoms.

7. The method as recited in Claim 5 wherein:

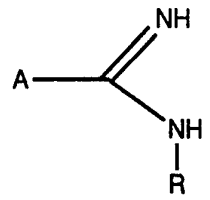
A is lower alkyl of 2 to about 10 carbon atoms, lower alkenyl of 2 to 6 carbon atoms, cycloalkyl group of 3 to 8 carbon atoms, bicycloalkyl group of 7 carbon atoms, alkylthioalkyl group of 2 to about 6 carbon atoms, alkyloxyalkyl of 2 to about 6 carbon atoms, alkylsulfonylalkyl group of 2 to about 6 carbon atoms , heterocyclic group, or aryl substituted heterocyclic group and which each may be optionally be substituted by one or more of the following lower alkyl, alkoxy, haloalkyl or halogen, nitro; and

R is H, OH or lower alkyl group of 1 to about 6 carbon atoms.

8. The method of inhibiting nitric oxide synthesis as recited in Claim 5 wherein the compound is selected from the group consisting of:
cyclopropylcarbamidine; isopropylcarbamidine; 1-propylcarbamidine; ethylcarbamidine; 2-methoxyacetamidine; 3-amidinopyridine; 2-amidinothiophene; 2-amidinofuran; 2-chloroacetamidine 2-Methylmercaptoacetamide, Isobutylcarbamidine hydrochloride, Benzylthioacetamidine hydrochloride, Heptadecylcarbamidine hydrochloride, Undecylcarbamidine hydrochloride, 1-Bromo-6-carbamidylhexane hydrochloride, 1-Chloro-4-carbamidylbutane hydrochloride, 2-Carbamidyl-3-bromothiophene hydrochloride, 2-Amidyl, 3,4-dimethyl, 5-carbamidylthiophene hydrochloride, Styrylcarbamidine hydrochloride, 2-carbamidylnorbornane hydrochloride, 2-Chloro-5-carbamidylnorbornane hydrochloride, 2-Phenylbutyramidine hydrochloride, 2,6-Dimethyl, 7-

carbamidylhepta-2-6-diene hydrochloride, 2-Methyl, 2-nitro, 4-carbamidylbutane hydrochloride, Acetamidine, 3-(2-chloro,6-fluorophenyl), 4-carbamidoxime, 5-methylisoxazole, 2-carboxamidoximylthiophene, 3-(2,6-dichlorophenyl), 4-carbamidoxime, 5-methylisoxazole, 5-carboxamidoximylisoxazole, 3,5-bistrifluoromethylstyrylcarboxamidoxime, t-Butylsulfonylacetamidoxime, Phenylacetamidoxime, and Formamidoxime.

9. A method of lowering nitric oxide levels in a subject in need of such by administering a therapeutically effective amount of a compound having the formula:



(I)

and salts, and pharmaceutically acceptable ester and prodrugs thereof, wherein:

A is hydrogen, lower alkyl , lower alkenyl , lower alkynyl group, alkylthioalkyl group, alkyloxyalkyl group, alkylsulfonylalkyl group, cycloalkyl group, bicycloalkyl group, cycloalkenyl group, cycloalkylalkyl group,, phenylalkyl group, phenylalkenyl group, biphenylalkyl group, heterocyclic group, biaryl group, or aryl wherein each said radical may optionally be substituted by one or more of the following substituents such as alkyl, alkoxy, hydroxy, halogen, nitro, cyano, haloalkyl, carboxylic, carboxamide, amino, alkylamino or dialkylamino; and

R is H, OH or lower alkyl group.

10. The method as recited in Claim 9 wherein;

A is hydrogen, lower alkyl of 1 to about 10 carbon atoms, lower alkenyl group of 2 to about 6 carbon atoms, lower alkynyl group of 2 to about 6 carbon atoms, alkylthioalkyl group of 2 to about 6 carbon atoms, alkyloxyalkyl of 2 to about 6 carbon atoms, alkylsulfonylalkyl group of 2 to about 6 carbon atoms, cycloalkyl group of 3 to about 8 carbon atoms, bicycloalkyl group of 6 to about 10 carbon atoms, cycloalkenyl group of 3 to about 8 carbon atoms, cycloalkylalkyl group of 4 to about 10 carbon atoms, phenylalkyl group, phenylalkenyl group, biphenylalkyl group, heterocyclic group, or aryl substituted heterocyclic groups and which each group may be optionally be substituted by one or more of the following substituents: lower alkyl of 1 to about 4 carbon atoms, alkoxy of 1 to about 4 carbon atoms, hydroxy, halogen, nitro, cyano, haloalkyl, carboxyl, carboxamide, amino, monoalkylamino or dialkylamino; and

R is H, OH or lower alkyl group of 1 to about 6 carbon atoms.

11. The method as recited in Claim 9 wherein;

A is lower alkyl of 2 to about 10 carbon atoms, lower alkenyl of 2 to 6 carbon atoms, cycloalkyl group of 3 to 8 carbon atoms, bicycloalkyl group of 7 carbon atoms, alkylthioalkyl group of 2 to about 6 carbon atoms, alkyloxyalkyl of 2 to about 6 carbon atoms, alkylsulfonylalkyl group of 2 to about 6 carbon atoms , heterocyclic group, or aryl substituted heterocyclic group and which each may be optionally be substituted by one or more of the following lower alkyl, alkoxy, haloalkyl or halogen, nitro; and

R is H, OH or lower alkyl group of 1 to about 6 carbon atoms.

12. The method as recited in Claim 9 wherein the compound is selected from the group consisting of: cyclopropylcarbamidine; isopropylcarbamidine; 1-propylcarbamidine; ethylcarbamidine; 2-methoxyacetamidine; 3-amidinopyridine; 2-amidinothiophene; 2-amidinofuran; 2-chloroacetamidine 2-Methylmercaptoacetamide, Isobutylcarbamidine hydrochloride, Benzylthioacetamidine hydrochloride, Heptadecylcarbamidine hydrochloride, Undecylcarbamidine hydrochloride, 1-Bromo-6-carbamidylhexane hydrochloride, 1-Chloro-4-carbamidylbutane hydrochloride, 2-Carbamidyl-3-bromothiophene hydrochloride, 2-Amidyl, 3,4-dimethyl, 5-carbamidylthiophene hydrochloride, Styrylcarbamidine hydrochloride, 2-carbamidylnorbornane hydrochloride, 2-Chloro-5-carbamidylnorbornane hydrochloride, 2-Phenylbutyramidine hydrochloride, 2,6-Dimethyl, 7-carbamidylhepta-2-6-diene hydrochloride, 2-Methyl, 2-nitro, 4-carbamidylbutane hydrochloride, Acetamidine, 3-(2-chloro,6-fluorophenyl), 4-carbamidoxime, 5-methylisoxazole, 2-carboxamidoximylthiophene, 3-(2,6-dichlorophenyl), 4-carbamidoxime, 5-methylisoxazole, 5-carboxamidoximylisoxazole, 3,5-bistrifluoromethylstyrylcarboxamidoxime, t-Butylsulfonylacetamidoxime, Phenylacetamidoxime, and Formamidoxime

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US 94/11724

A. CLASSIFICATION OF SUBJECT MATTER
IPC 6 A61K31/00 A61K31/155

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 6 A61K C07D C07C

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO,A,93 03714 (THE UPJOHN COMPANY) 4 March 1993 see the whole document ---	1-12
X,P	WO,A,94 02135 (H.L.ELFORD ET AL.) 3 February 1994 see the whole document ---	1-12
X	US,A,3 978 202 (PALLOS ET AL.) 31 August 1976 see the whole document ---	1-12
X	US,A,3 978 219 (PALLOS ET AL.) 31 August 1976 see the whole document ---	1-12
	-/--	

[X] Further documents are listed in the continuation of box C.

[X] Patent family members are listed in annex.

* Special categories of cited documents :

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

Date of the actual completion of the international search	Date of mailing of the international search report
1 February 1995	10.02.95

Name and mailing address of the ISA	Authorized officer
European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax (+31-70) 340-3016	Theuns, H

Form PCT/ISA/210 (second sheet) (July 1992)

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P,X	DATABASE WPI Derwent Publications Ltd., London, GB; AN 94-363554 & JP,A,6 287 180 (YAMANOUCHI PHARM CO LTD) 11 October 1994 see abstract	1-12
X	'The Merck Index' 1989 , MERCK & CO. INC. , RAHWAY, N.J., USA see Monograph 37	1-12
X	WO,A,92 14453 (J.N.CAMPBELL) 3 September 1992 see claims 11,12	1-12
X	US,A,4 634 783 (FUJII ET AL.) 6 January 1987 see the whole document	1-12
P,X	EP,A,0 568 289 (EISAI CO., LTD.) 3 November 1993 see the whole document	1-12
X	EP,A,0 518 819 (CIBA-GEIGY AG) 16 December 1992 see the whole document	1-12
P,X	EP,A,0 601 977 (CIBA-GEIGY AG) 15 June 1994 see the whole document	1-12
P,X	WO,A,94 11341 (CIBA-GEIGY AG) 26 May 1994 see the whole document	1-12
P,X	WO,A,94 16719 (SMITHKLINE BEECHAM PLC) 4 August 1994 see the whole document	1-12
X	US,A,5 246 965 (MAIN) 21 September 1993 see the whole document	1-12

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US94/11724

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. [X] Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
REMARK: Although claims 1-12 are directed to a method of treatment of the human/animal body the search has been based on the alleged effects of the compound/composition.

2. [X] Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
For further information please see annex.

3. [ ] Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. [ ] As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.

2. [ ] As all searchable claims could be searches without effort justifying an additional fee, this Authority did not invite payment of any additional fee.

3. [ ] As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:

4. [ ] No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

[ ] The additional search fees were accompanied by the applicant's protest.

[ ] No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/210

CONTINUATION OF BOX I.2:

In view of the large number of compounds which are defined by formula (I) and which have only a (substituted) amidino group in common, the search was limited to the characterising part of the compounds (Art. 6 PCT; Guideline B-II, 7, last sentence, and B-III, 3.7).

The expression "heterocyclic group" is not a clear and limited description of a group. By the use of this expression a complete search would involve a major part of the chemistry-related IPC documentation. Such a search is economically not feasible.

The definition of the therapeutic usefulness by the expression "inhibiting nitric oxide synthesis in a subject in need of such inhibition" is not a proper definition of the intended therapeutic use, because it is not fully known which conditions fulfil this requirement, and which conditions do not fulfil this requirement. This has as an effect that in order to judge whether a known compound of formula (I) having a pharmaceutical utility would fulfil the requirements as expressed in the claims in each case tests should be performed in order to establish whether the treatment of each disease in question would benefit from inhibition of nitric oxide synthesis.

It is clear that in this situation a complete search is virtually impossible.

The definition "aryl substituted heterocyclic group" for A in claim 2 and further claims is not comprised in the definition for A in formula (I) as expressed in claim 1.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No.
PCT/US 94/11724

Patent document cited in search report	Publication date	Patent family member(s)		Publication date
WO-A-9303714	04-03-93	AU-A-	2407592	16-03-93
		CA-A-	2113817	04-02-93
		EP-A-	0600973	15-06-94
		JP-T-	6510760	01-12-94
WO-A-9402135	03-02-94	US-A-	5350770	27-09-94
US-A-3978202	31-08-76	NONE		
US-A-3978219	31-08-76	NONE		
WO-A-9214453	03-09-92	CA-A-	2104873	03-09-92
		EP-A-	0573581	15-12-93
		JP-T-	6507392	25-08-94
US-A-4634783	06-01-87	JP-C-	1700995	14-10-92
		JP-B-	3068859	30-10-91
		JP-A-	59139357	10-08-84
		DE-A,C	3402628	02-08-84
		FR-A,B	2540118	03-08-84
		GB-A,B	2134901	22-08-84
EP-A-0568289	03-11-93	US-A-	5340833	23-08-94
		JP-A-	6049058	22-02-94
EP-A-0518819	16-12-92	AU-B-	653603	06-10-94
		AU-A-	1807392	17-12-92
		JP-A-	5239009	17-09-93
		NZ-A-	243079	25-11-94
EP-A-0601977	15-06-94	AU-B-	5218093	23-06-94
		CA-A-	2110838	10-06-94
		FI-A-	935452	10-06-94
		HU-A-	65778	28-07-94
		JP-A-	6263710	20-09-94
		NO-A-	934483	10-06-94
WO-A-9411341	26-05-94	AU-B-	5599494	08-06-94
WO-A-9416719	04-08-94	AU-B-	5863694	15-08-94

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No.
PCT/US 94/11724

Patent document cited in search report	Publication date	Patent family member(s)		Publication date
US-A-5246965	21-09-93	AU-A-	1807292	17-12-92
		EP-A-	0518818	16-12-92
		HU-A-	61977	29-03-93
		JP-A-	5239008	17-09-93