

# International Nonproprietary Names for Pharmaceutical Substances

In accordance with article 3 of the Procedure for the Selection of Recommended International Nonproprietary Names for Pharmaceutical Substances,<sup>1</sup> notice is hereby given that the following names are under consideration by the World Health Organization as Proposed International Nonproprietary Names.

Comments on, or formal objections to, the proposed names may be forwarded by any person to the Pharmaceuticals unit of the World Health Organization within four months of the date of their publication in the *WHO Chronicle*, e.g. for List 45 Prop. INN not later than 30 September 1981.

*The inclusion of a name in the lists of proposed international nonproprietary names does not imply any recommendation for the use of the substance in medicine or pharmacy.*

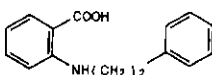
## Proposed International Nonproprietary Names (Prop. INN): List 45<sup>2</sup>

Proposed International  
Nonproprietary Name (Latin, English)

Chemical Name or Description, Molecular and Graphic Formulae  
Chemical Abstracts Service (CAS) registry number

acidum enfenamicum  
enfenamic acid

N-phenethylanthranilic acid  
C<sub>15</sub>H<sub>15</sub>NO<sub>2</sub> 23049-93-6



Comprehensive information on the INN programme can be found in WHO Technical Report Series, No. 581, 1975 (*Nonproprietary Names for Pharmaceutical Substances* Twentieth Report of the WHO Expert Committee), ISBN 92 4 120581 4 (price: Sw. fr. 6.-); an account of this publication will be found in page 16 of this Supplement (Annex 2). All names from Lists 1-37 of Proposed International Nonproprietary Names, together with a molecular formula index, will be found in: *International Nonproprietary Names for Pharmaceutical Substances. Cumulative list No. 5, 1977*, World Health Organization, Geneva, 1977 (ISBN 92 4 056011 4) (price: Sw. fr. 48.-). This publication consists, in the main, of a computer printout which groups together all the proposed and recommended international nonproprietary names (INN)—in Latin, English, French, Russian, and Spanish—published up to March 1977. The printout also indicates in which of the 37 individual lists of proposed names and 16 lists of recommended names, each INN was originally published, and gives references to national nonproprietary names, pharmacopoeia monographs, and other sources. In addition, the list contains molecular formulae and Chemical Abstracts Service registry numbers. For easy reference, national nonproprietary names that differ from INN, molecular formulae, and Chemical Abstracts Service registry numbers are indexed in a series of annexes. A final annex describes the procedure for selecting recommended INN and outlines the general principles to be followed in devising these names. All the textual material published in this volume appears in both English and French.

These publications may be obtained, direct or through booksellers, from the sales agents listed on the back cover of the *WHO Chronicle*. Orders from countries where sales agents have not yet been appointed may be addressed to: World Health Organization, Distribution and Sales Service, 1211 Geneva 27, Switzerland.

<sup>1</sup> See Annex 1, p. 15.

<sup>2</sup> Other lists of proposed international nonproprietary names can be found in *Chron. Wld Hlth Org.*, 1953, 7, 299; 1954, 8, 216, 313; 1956, 10, 28, 1957, 11, 231; 1958, 12, 102; *WHO Chronicle*, 1959, 13, 105, 152; 1960, 14, 168, 244; 1961, 15, 314; 1962, 16, 385; 1963, 17, 389; 1964, 18, 433; 1965, 19, 446; 1966, 20, 216; 1967, 21, 70, 478, 1968, 22, 112, 407; 1969, 23, 183, 418, 1970, 24,

119, 413; 1971, 25, 123, 415; 1972, 26, 121, 414; 1973, 27, 120, 330; 1974, 28, 133, supplements to *WHO Chronicle*, 1974, Vol. 28, No. 9, 1975, Vol. 29, No. 3, No. 9, 1976, Vol. 30, No. 3, No. 9; 1977, Vol. 31, No. 3, No. 9, 1978, Vol. 32, No. 3, No. 9; 1979, Vol. 33, No. 3, No. 9; 1980, Vol. 34, No. 3, No. 9.

Lists or recommended international nonproprietary names were published

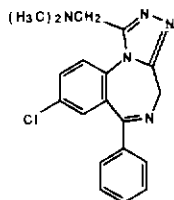
in *Chron. Wld Hlth Org.*, 1955, 9, 185; *WHO Chronicle*, 1959, 13, 106, 463; 1962, 16, 101; 1965, 19, 165, 206, 249; 1966, 20, 421, 1967, 21, 538; 1968, 22, 463; 1969, 23, 490; 1970, 24, 526; 1971, 25, 476; 1972, 26, 476; 1973, 27, 453; supplements to *WHO Chronicle*, 1974, Vol. 28, No. 10; 1975, Vol. 29, No. 10, 1976, Vol. 30, No. 10; 1977, Vol. 31, No. 10; 1978, Vol. 32, No. 10; 1979, Vol. 33, No. 10, 1980, Vol. 34, No. 10.

Proposed International  
Nonproprietary Name (Latin, English)

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Chemical Abstracts Service (CAS) registry number

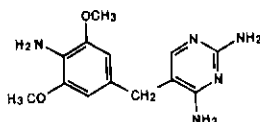
adinazolamum  
adinazolam

8-chloro-1-[(dimethylamino)methyl]-6-phenyl-4H-s-triazolo[4,3-a][1,4]benzo-  
diazepine  
 $C_{19}H_{18}ClN_5$  37115-32-5



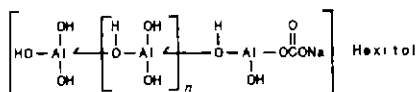
aditerenum  
aditeren

2,4-diamino-5-(4-amino-3,5-dimethoxybenzyl)pyrimidine  
 $C_{13}H_{17}N_5O_2$  56066-19-4



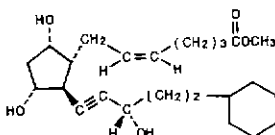
alexitolum natricum  
alexitol sodium

sodium polyhydroxyaluminium monocarbonate hexitol complex  
where  $n = 0$  or an integer, controlled by the preparative conditions  
66813-51-2



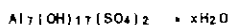
alfaprostolum  
alfaprostol

methyl (Z)-7-[(1R,2S,3R,5S)-2-[(3S)-5-cyclohexyl-3-hydroxy-1-pentynyl]-3,5-  
dihydroxycyclopentyl]-5-heptenoate  
 $C_{24}H_{38}O_5$  74176-31-1



alusulfum  
alusulf

heptaaluminum heptadecahydroxide bis(sulfate) hydrate  
 $Al_7H_{17}O_{25}S_2 \cdot xH_2O$  61115-28-4

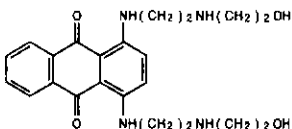


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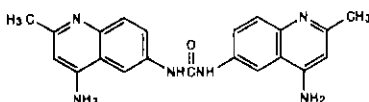
ametantronum  
ametantrone

1,4-bis[[2-[(2-hydroxyethyl)amino]ethyl]amino]anthraquinone  
 $C_{22}H_{28}N_4O_4$  64862-96-0



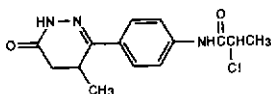
aminoquinuridum  
aminoquinuride

1,3-bis(4-amino-2-methyl-6-quinolyl)urea  
 $C_{21}H_{20}N_6O$  3811-56-1



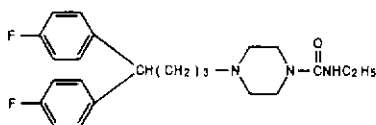
amipizonum  
amipizone

2-chloro-4'-(1,4,5,6-tetrahydro-4-methyl-6-oxo-3-pyridazinyl)propionanilide  
 $C_{14}H_{16}ClN_3O_2$  69635-63-8



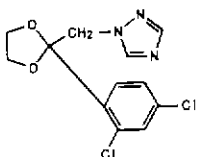
amperozidum  
amperozide

4-[4-bis(p-fluorophenyl)butyl]-N-ethyl-1-piperazinecarboxamide  
 $C_{23}H_{29}F_2N_3O$  75558-90-6



azonazolum  
azonazole

1-[[2-(2,4-dichlorophenyl)-1,3-dioxolan-2-yl]methyl]-1H-1,2,4-triazole  
 $C_{12}H_{11}Cl_2N_3O_2$  60207-31-0

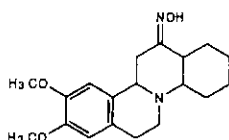


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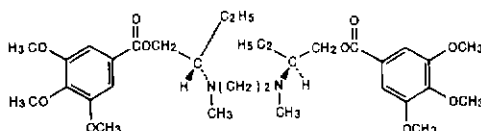
benolizimum  
benolizime

1,2,3,4,4a,6,7,11b,12,13a-decahydro-9,10-dimethoxy-13H-dibenzo[a,f]quinolizin-  
13-one oxime  
 $C_{19}H_{26}N_2O_3$  61864-30-0



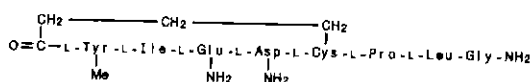
butobendinum  
butobendine

(+)-(S,S)-ethylenebis[(methylimino)(2-ethylethylene)] bis(3,4,5-trimethoxyben-  
zoate)  
 $C_{32}H_{46}N_2O_{10}$  55769-65-8



carbetocinum  
carbetocin

1-butyric acid-2-[3-(p-methoxyphenyl)-L-alanine]oxytocin  
 $C_{45}H_{69}N_{11}O_{12}S$  37025-55-1

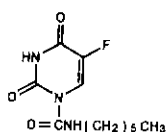


carmellosum  
carmellose

polycarboxymethyl ether of cellulose  
9000-11-7

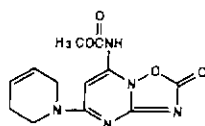
carmofurum  
carmofur

5-fluoro-N-hexyl-3,4-dihydro-2,4-dioxo-1(2H)-pyrimidinecarboxamide  
 $C_{11}H_{16}FN_3O_3$  61422-45-5



carprazidilum  
carprazidil

methyl 5-{3,6-dihydro-1(2H)-pyridyl}-2-oxo-2H-[1,2,4]oxadiazolo[2,3-a]pyrimi-  
dine-7-carbamate  
 $C_{12}H_{13}N_5O_4$  68020-77-9

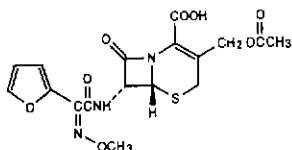


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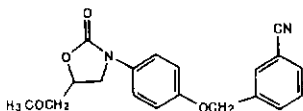
cefuracetimum  
cefuracetime

(6*R*,7*R*)-7-[2-(2-furyl)glyoxylamido]-3-(hydroxymethyl)-8-oxo-5-thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid 7<sup>2</sup>-(*Z*)-(O-methyloxime), acetate (ester)  
C<sub>17</sub>H<sub>17</sub>N<sub>3</sub>O<sub>6</sub>S 39685-31-9



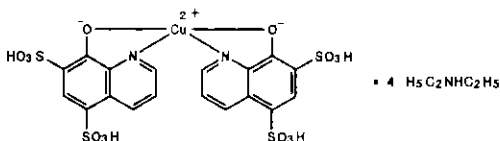
cimoxatonum  
cimoxatone

$\alpha$ -[*p*-[5-(methoxymethyl)-2-oxo-3-oxazolidinyl]phenoxy]-*m*-tolunitrile  
C<sub>19</sub>H<sub>18</sub>N<sub>2</sub>O<sub>4</sub> 73815-11-9



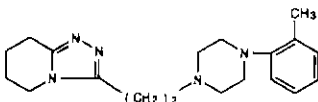
cuproxolinum  
cuproxoline

bis(dihydrogen 8-hydroxy-5,7-quinolinedisulfonato)copper, compound with diethylamine (1 : 4)  
C<sub>18</sub>H<sub>12</sub>CuN<sub>2</sub>O<sub>14</sub>S<sub>4</sub> · 4C<sub>4</sub>H<sub>11</sub>N or C<sub>34</sub>H<sub>56</sub>CuN<sub>6</sub>O<sub>14</sub>S<sub>4</sub> 13007-93-7



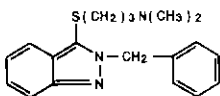
dapiprazolum  
dapiprazole

5,6,7,8-tetrahydro-3-[2-(4-*o*-tolyl-1-piperazinyl)ethyl]-*s*-triazolo[4,3-*a*]pyridine  
C<sub>19</sub>H<sub>27</sub>N<sub>5</sub> 72822-12-9



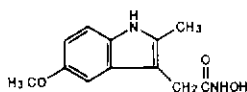
dazidaminum  
dazidamine

2-benzyl-3-[[3-(dimethylamino)propyl]thio]-2*H*-indazole  
C<sub>19</sub>H<sub>23</sub>N<sub>3</sub>S 75522-73-5



deboxametum  
deboxamet

5-methoxy-2-methylindole-3-acetohydroxamic acid  
C<sub>12</sub>H<sub>14</sub>N<sub>2</sub>O<sub>3</sub> 34024-41-4



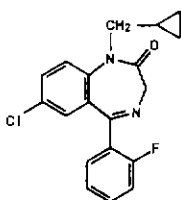
$$\text{H}_3\text{C}(\text{CH}_2)_9\text{OCH}_2\underset{\text{OH}}{\text{CH}}\text{CH}_2\text{NH}_2$$

The chemical structure shows a benzothiazepine core. A chlorine atom is attached to the benzene ring at the 2-position. A sulfonamide group (-SO<sub>2</sub>NH-) is attached to the 4-position of the tetrahydrobenzothiazepine ring. The nitrogen of the sulfonamide group is further substituted with a 2,4-dichlorophenyl ring.

CN1CCCC1[C@H]2C[C@@H](C)[C@H]3C=C[C@H](C=C3)C2=OC1CCNCC1CC(=O)NC2CCCCC2

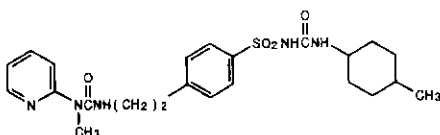
flutoprazepamum  
flutoprazepam

7-chloro-1-(cyclopropylmethyl)-5-(*o*-fluorophenyl)-1,3-dihydro-2*H*-1,4-benzodiazepin-2-one  
 $C_{19}H_{15}ClFN_2O$  25967-29-7



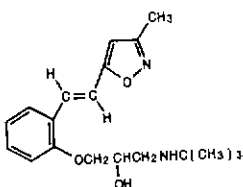
glisamuridum  
glisamuride

1-methyl-3-[*p*-[[3-(4-methylcyclohexyl)ureido]sulfonyl]phenethyl]-1-(2-pyridyl)-urea  
 $C_{23}H_{31}N_5O_4S$  52430-65-6



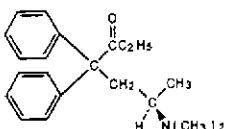
isoxaprololum  
isoxaprolol

(±)-(E)-1-(*tert*-butylamino)-3-[*o*-[2-(3-methyl-5-isoxazolyl)vinyl]phenoxy]-2-propanol  
 $C_{19}H_{28}N_2O_3$  75949-60-9



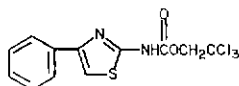
levomethadonum  
levomethadone

(-)-6-(dimethylamino)-4,4-diphenyl-3-heptanone  
 $C_{21}H_{27}NO$  125-58-6



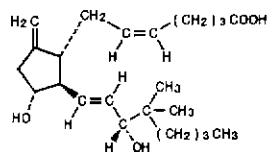
lotifazolum  
lotifazole

2,2,2-trichloroethyl 4-phenyl-2-thiazolecarbamate  
 $C_{12}H_9Cl_3N_2O_2S$  71119-10-3



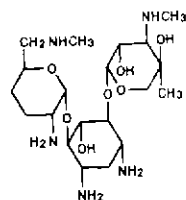
meteneprostum  
meteneprost

(Z)-7-[(1*R*,2*R*,3*R*)-3-hydroxy-2-[(*E*)-(3*R*)-3-hydroxy-4,4-dimethyl-1-octenyl]-5-methylenecyclopentyl]-5-heptenoic acid  
 $C_{23}H_{38}O_4$  61263-35-2



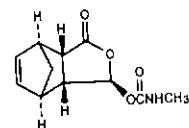
micronomicinum  
micronomicin

*O*-2-amino-2,3,4,6-tetradecoxy-6-(methylamino)- $\alpha$ -D-erythro-hexopyranosyl-(1  $\rightarrow$  4)-*O*-[3-deoxy-4-*C*-methyl-3-(methylamino)- $\beta$ -L-arabinopyranosyl-(1  $\rightarrow$  6)]-2-deoxy-D-streptamine  
 $C_{26}H_{41}N_5O_7$  52093-21-7



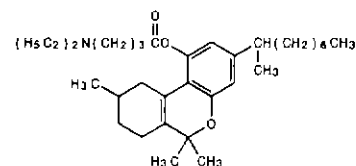
moxadolum  
moxadolen

methylcarbamic acid, ester with (3*R*\*,3*aR*\*,4*S*\*,7*R*\*,7*aS*\*)-3*a*,4,7,7*a*-tetrahydro-3-hydroxy-4,7-methanoisobenzofuran-1(3*H*)-one  
 $C_{11}H_{13}NO_4$  75992-53-9



naboctatum  
naboctate

7,8,9,10-tetrahydro-6,6,9-trimethyl-3-(1-methyloctyl)-6*H*-dibenzo[*b,d*]pyran-1-yl 4-(diethylamino)butyrate  
 $C_{33}H_{53}NO_3$  74912-19-9



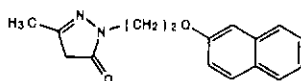


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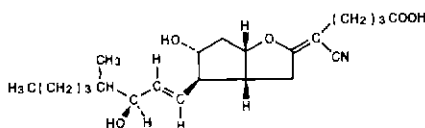
nafazatromum  
nafazatrom

3-methyl-1-[2-(2-naphthoxy)ethyl]-2-pyrazolin-5-one  
 $C_{16}H_{16}N_2O_2$  59040-30-1



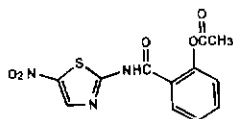
nileprost  
nileprost

(*E*)-(3*aR*,4*R*,5*R*,6*aS*)- $\delta$ -cyano-3,3*a*,4,5,6,6*a*-hexahydro-5-hydroxy-4-[(*E*)-(3*S*,4*RS*)-3-hydroxy-4-methyl-1-octenyl]-2*H*-cyclopenta[*b*]furan- $\Delta^{2,\delta}$ -valeric acid  
 $C_{22}H_{33}NO_5$  71097-83-1



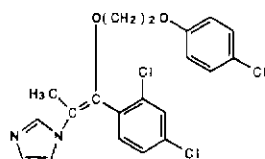
nitazoxanidum  
nitazoxanide

*N*-(5-nitro-2-thiazolyl)salicylamide acetate (ester)  
 $C_{12}H_9N_3O_5S$  55981-09-4



omoconazolum  
omoconazole

(*E*)-1-[2,4-dichloro- $\beta$ -[2-(*p*-chlorophenoxy)ethoxy]- $\alpha$ -methylstyryl]imidazole  
 $C_{20}H_{17}Cl_3N_2O_2$  74512-12-2

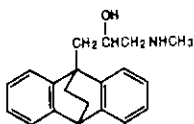


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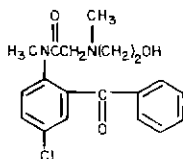
oxaprotilinum  
oxaprotiline

$\alpha$ -[(methylamino)methyl]-9,10-ethanoanthracene-9(10H)-ethanol  
 $C_{20}H_{23}NO$  56433-44-4



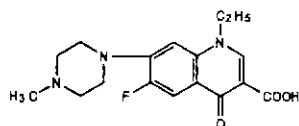
oxazafonum  
oxazafone

2'-benzoyl-4'-chloro-2-[(2-hydroxyethyl)methylamino]-N-methylacetanilide  
 $C_{19}H_{21}ClN_2O_3$  70541-17-2



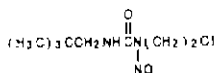
pefloxacinum  
pefloxacin

1-ethyl-6-fluoro-1,4-dihydro-7-(4-methyl-1-piperazinyl)-4-oxo-3-quinolinecarboxylic acid  
 $C_{17}H_{20}FN_3O_3$  70458-92-3



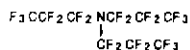
pentamustinum  
pentamustine

1-(2-chloroethyl)-3-neopentyl-1-nitrosourea  
 $C_8H_{15}ClN_3O_2$  73105-03-0



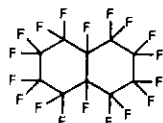
perfluaminum  
perfluamine

heneicosafluorotripropylamine  
 $C_9F_{21}N$  338-83-0



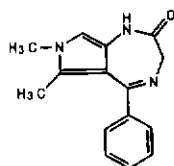
perflunafenum  
perflunafene

octadecafluorodecahydronaphthalene  
 $C_{10}F_{18}$  306-94-5



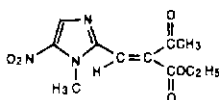
premazepamum  
premazepam

3,7-dihydro-6,7-dimethyl-5-phenylpyrrolo[3,4-*e*]-1,4-diazepin-2(1*H*)-one  
 $C_{15}H_{15}N_3O$  57435-86-6



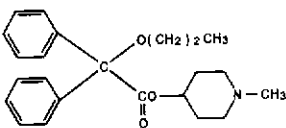
propenidazolium  
propenidazole

ethyl *trans*- $\alpha$ -acetyl-1-methyl-5-nitroimidazole-2-acrylate  
 $C_{17}H_{13}N_3O_5$  76448-31-2



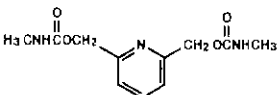
propiverinum  
propiverine

1-methyl-4-piperidyl diphenylpropoxyacetate  
 $C_{23}H_{29}NO_3$  60569-19-9



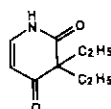
pyricarbatum  
pyricarbate

2,6-pyridinediyl dimethylene bis(methylcarbamate)  
 $C_{11}H_{15}N_3O_4$  1882-26-4



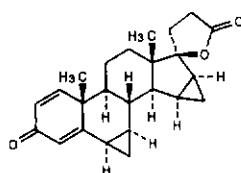
pyrithyldionum  
pyrithyldione

3,3-diethyl-2,4(1*H*,3*H*)-pyridinedione  
 $C_9H_{13}NO_2$  77-04-3



spirorenonum  
spirorenone

(6*R*,7*R*,8*R*,9*S*,10*R*,13*S*,14*R*,15*S*,16*S*,17*S*)-3',4',6,7,8,9,11,12,13,14,15,16,20,21-tetradecahydro-10,13-dimethylspiro[17*H*-dicyclopropa[6,7:15,16]cyclopenta[*a*]phenanthrene-17,2'(5'*H*)-furan]-3(10*H*),5'-dione or 17-hydroxy-6β,7β:15β,16β-dimethylene-3-oxo-17α-pregna-1,4-diene-21-carboxylic acid, γ-lactone  
 $C_{24}H_{28}O_3$  74220-07-8

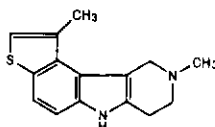


thymostimulinum  
thymostimulin

polypeptide immunostimulant factor extracted from thymus of mammalian species. The source of the product should be indicated, e.g. thymostimulin (calf)

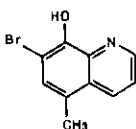
tienocarinum  
tienocarbine

7,8,9,10-tetrahydro-1,9-dimethyl-6*H*-pyrido[4,3-*b*]thieno[3,2-*e*]indole  
 $C_{15}H_{16}N_2S$  75458-65-0



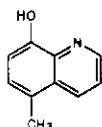
tilbroquinolum  
tilbroquinol

7-bromo-5-methyl-8-quinolinol  
 $C_{10}H_8BrNO$  7175-09-9



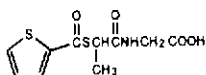
tiliquinolum  
tiliquinol

5-methyl-8-quinolinol  
 $C_{10}H_9NO$  5541-67-3



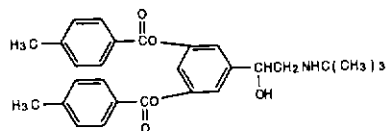
tiofacicum  
tiofacic

*N*-(2-mercaptopropionyl)glycine 2-thiophenecarboxylate (ester)  
C<sub>10</sub>H<sub>11</sub>NO<sub>4</sub>S<sub>2</sub> 72324-18-6



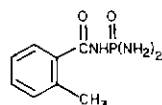
tobuterolum  
tobuterol

(±)-5-[2-(*tert*-butylamino)-1-hydroxyethyl]-*m*-phenylene di-*p*-toluate  
C<sub>28</sub>H<sub>31</sub>NO<sub>5</sub> 75626-99-2



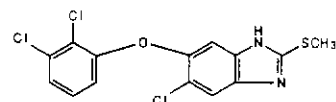
tolfamidum  
tolfamide

*N*-(diaminophosphinyl)-*o*-toluamide  
C<sub>8</sub>H<sub>12</sub>N<sub>3</sub>O<sub>2</sub>P 70788-29-3



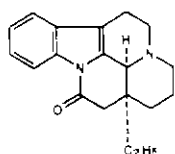
triclabendazolum  
triclabendazole

5-chloro-6-(2,3-dichlorophenoxy)-2-(methylthio)benzimidazole  
C<sub>14</sub>H<sub>9</sub>Cl<sub>3</sub>N<sub>2</sub>OS 68786-66-3



vinburninum  
vinburnine

3α,16α-eburnamonine  
C<sub>19</sub>H<sub>22</sub>N<sub>2</sub>O 4880-88-0



# AMENDMENTS TO PREVIOUS LISTS

Cumulative List No. 5, 1977

## International Nonproprietary Names (INN) for Pharmaceutical Substances:

p. 116	<i>delete</i>	<i>insert</i>
	levarterenolum	norepinephrinum
	levarterenol	norepinephrine

Supplement to Vol. 29, No. 3

## International Nonproprietary Names (Prop. INN): List 33

p. 2	<i>delete</i>	<i>insert</i>
	acidum etodolicum	etodolacum
	etodolic acid	etodolac

Supplement to Vol 33, No. 9

## International Nonproprietary Names (Prop. INN): List 42

p. 17	<i>delete</i>	<i>insert</i>
	sulerginum	disulerginum
	sulergine	disulergine

Supplement to Vol. 34, No. 3

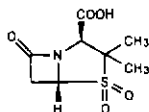
## International Nonproprietary Names (Prop. INN): List 43

p. 17	<i>delete</i>	<i>insert</i>
	triaconazolum	terconazolum
	triaconazole	terconazole

Supplement to Vol. 34, No. 9

## International Nonproprietary Names (Prop. INN): List 44

p. 3	<i>delete</i>	<i>insert</i>
	alisactidum	alsactidum
	alisactide	alsactide
p. 10	enilconazolum	<i>replace CAS registry number by: 35554-44-0</i>
	enilconazole	
p. 21	sulbactamum	<i>insert the following graphic formula:</i>
	sulbactam	



p. 25	<i>delete</i>	<i>insert</i>
	vintenatum	vincatenatum
	vintenate	vincatenate
p. 27	aclatonii napadisilas	<i>cancel amendment and retain graphic formula in List 43 prop INN</i>
	aclatonium napadisilate	

## Annex 1

### PROCEDURE FOR THE SELECTION OF RECOMMENDED INTERNATIONAL NONPROPRIETARY NAMES FOR PHARMACEUTICAL SUBSTANCES \*

The following procedure shall be followed by the World Health Organization in the selection of recommended international nonproprietary names for pharmaceutical substances, in accordance with the World Health Assembly resolution WHA3.11:

1. Proposals for recommended international nonproprietary names shall be submitted to the World Health Organization on the form provided therefor.

2. Such proposals shall be submitted to the Director-General of the World Health Organization to the members of the Expert Advisory Panel on the International Pharmacopoeia and Pharmaceutical Preparations designated for this purpose, for consideration in accordance with the "General principles for guidance in devising International Nonproprietary Names" appended to this procedure. The name used by the person discovering or first developing and marketing a pharmaceutical substance shall be accepted, unless there are compelling reasons to the contrary.

3. Subsequent to the examination provided for in article 2, the Director-General of the World Health Organization shall give notice that a proposed international nonproprietary name is being considered.

A. Such notice shall be given by publication in the *Chronicle of the World Health Organization*<sup>1</sup> and by letter to Member States and to national pharmacopoeia commissions or other bodies designated by Member States.

(i) Notice may also be sent to specific persons known to be concerned with a name under consideration.

B. Such notice shall:

- (i) set forth the name under consideration;
- (ii) identify the person who submitted a proposal for naming the substance, if so requested by such person;
- (iii) identify the substance for which a name is being considered;
- (iv) set forth the time within which comments and objections will be received and the person and place to whom they should be directed;
- (v) state the authority under which the World Health Organization is acting and refer to these rules of procedure.

C. In forwarding the notice, the Director-General of the World Health Organization shall request that Member States take such steps as are necessary to prevent the acquisition of proprietary rights in the proposed name during the period it is under consideration by the World Health Organization.

4. Comments on the proposed name may be forwarded by any person to the World Health Organization within four months of the date of publication, under article 3, of the name in the *Chronicle of the World Health Organization*.<sup>1</sup>

5. A formal objection to a proposed name may be filed by any interested person within four months of the date of publication, under article 3, of the name in the *Chronicle of the World Health Organization*.<sup>1</sup>

A. Such objection shall:

- (i) identify the person objecting;
- (ii) state his interest in the name;
- (iii) set forth the reasons for his objection to the name proposed.

6. Where there is a formal objection under article 5, the World Health Organization may either reconsider the proposed name or use its good offices to attempt to obtain withdrawal of the objection. Without prejudice to the consideration by the World Health Organization of a substitute name or names, a name shall not be selected by the World Health Organization as a recommended international nonproprietary name while there exists a formal objection thereto filed under article 5 which has not been withdrawn.

7. Where no objection has been filed under article 5, or all objections previously filed have been withdrawn, the Director-General of the World Health Organization shall give notice in accordance with subsection A of article 3 that the name has been selected by the World Health Organization as a recommended international nonproprietary name.

8. In forwarding a recommended international nonproprietary name to Member States under article 7, the Director-General of the World Health Organization shall:

A. request that it be recognized as the nonproprietary name for the substance; and

B. request that Member States take such steps as are necessary to prevent the acquisition of proprietary rights in the name, including prohibiting registration of the name as a trade-mark or trade-name.

\* Text adopted by the Executive Board of WHO in resolution EB15.R7 (Off. Rec. Wild Hlth Org., 1955, 60, 3) and amended by the Board in resolution EB43.R9 (Off. Rec. Wild Hlth Org., 1969, 173, 10).

<sup>1</sup> The title of this publication was changed to *WHO Chronicle* in January 1959.

### GENERAL PRINCIPLES FOR GUIDANCE IN DEVISING INTERNATIONAL NONPROPRIETARY NAMES FOR PHARMACEUTICAL SUBSTANCES

1. International Nonproprietary Names (INN) should be distinctive in sound and spelling. They should not be inconveniently long and should not be liable to confusion with names in common use.

2. The INN for a substance belonging to a group of pharmacologically related substances should, where appropriate, show this relationship. Names that are likely to convey to a patient an anatomical, physiological,

pathological or therapeutic suggestion should be avoided.

*These primary principles are to be implemented by using the following secondary principles*

3. In devising the INN of the first substance in a new pharmacological group, consideration should be given to the possibility of devising suitable INN for related substances, belonging to the new group.

4. In devising INN for acids, one-word names are preferred; their salts should be named without modifying the acid name, e.g. "oxacillin" and "oxacillin sodium", "ibufenac" and "ibufenac sodium".

5. INN for substances which are used as salts should in general apply to the active base or the active acid. Names for different salts or esters of the same active substance should differ

only in respect of the name of the inactive acid or the inactive base.

For quaternary ammonium substances, the cation and anion should be named appropriately as separate components of a quaternary substance and not in the amine-salt style.

6. The use of an isolated letter or number should be avoided; hyphenated construction is also undesirable.

7. To facilitate the translation and pronunciation of INN, "f" should be

used instead of "ph", "t" instead of "th", "e" instead of "ae" or "oe", and "i" instead of "y"; the use of the letters "h" and "k" should be avoided.

8. Provided that the names suggested are in accordance with these principles, names proposed by the person discovering or first developing and marketing a pharmaceutical preparation, or names already officially in use in any country, should receive preferential consideration.

9. Group relationship in INN (see Guiding Principle 2) should if possible be shown by using a stem from the following list. The stem should only be used for substances of the appropriate group. Where a stem is shown without any hyphens it may be used anywhere in the name.

Subsidiary group relationships should be shown by devising INN which show similarities to and are analogous with a previously named substance.

Latin	English	French
-actidum	-actide	-actide
andr	andr	andr
-arolum	-arol	-arol
-azepamum	-azepam	-azépam
bol	bol	bol
-buzonium	-buzone	-buzone
-cainum	-caine	-caine
cef-	cef-	céf-
-cillinum	-cillin	-cilline
cort	cort	cort
-cyclinum	-cycline	-cycline
estr	estr	estr
-fibratum	-fibrate	-fibrate
-forminum	-formin	-formine
gest	gest	gest
gli-	gli-	gli-
io-	io-	io-
-ium	-ium	-ium
-metacinum	-metacin	-métacine
-mycinum	-mycin	-mycine
-nidazolum	-nidazole	-nidazole
-ololum	-olol	-olol
-onidum	-onide	-onide
-orexum	-orex	-orex
-praminum	-pramine	-pramine
-profenum	-profen	-profène
prost	prost	prost
-relinum	-relin	-réline
sulfa-	sulfa-	sulfa-
-terolum	-terol	-térol
-tizidum	-tizide	-tizide
-verinum	-verine	-vérine

synthetic polypeptides with a corticotrophin-like action  
steroids, androgens  
anticoagulants of the dicoumarol group  
substances of the diazepam group  
steroids, anabolic  
anti-inflammatory analgesics of the phenylbutazone group  
local anaesthetics  
antibiotics, derivatives of cephalosporanic acid  
antibiotics, derivatives of 6-aminopenicillanic acid  
corticosteroids, except those of the prednisolone group  
antibiotics of the tetracycline group  
estrogenic substances  
substances of the clofibrate group  
hypoglycemics of the phenformin group  
steroids, progestogens  
sulfonamide hypoglycemics  
iodine-containing contrast media  
quaternary ammonium compounds  
anti-inflammatory substances of the indometacin group  
antibiotics, produced by *Streptomyces* strains  
antiprotozoal substances of the metronidazole group  
 $\beta$ -adrenergic blocking agents of the propranolol group  
steroids for topical use, containing an acetal group  
anorexigenic agents, phenethylamine derivatives  
substances of the imipramine group  
anti-inflammatory substances of the ibuprofen group  
prostaglandins  
hypophyseal hormone release-stimulating peptides  
sulfonamides, anti-infective  
bronchodilators, phenethylamine derivatives  
diuretics of the chlorothiazide group  
spasmolytics with a papaverine-like action

## Annex 2

### NONPROPRIETARY NAMES FOR PHARMACEUTICAL SUBSTANCES: TWENTIETH REPORT OF THE WHO EXPERT COMMITTEE

In its twentieth report<sup>1</sup> the WHO Expert Committee on Nonproprietary Names for Pharmaceutical Substances reviewed the general principles for devising, and the procedures for selecting, international nonproprietary names (INN) in the light of developments in pharmaceutical compounds in recent years. The most significant recent change has been the extension to the naming of synthetic chemical substances of the practice previously used for substances originating in or derived from

natural products. This practice involves employing a characteristic "stem" indicative of a common property of the members of a group. The reasons for, and the implications of, the change are fully discussed. Also reported is the intention to change the practice with regard to the nomenclature of individual members of polymeric series.

Other sections of the report concern instructions to be followed by bodies making application for international nonproprietary names, the

availability of computer-printed cumulative lists of international nonproprietary names, information supplied by WHO Member States concerning their official use of national or international names for pharmaceutical products, and proposals relative to the withdrawal of international nonproprietary names allocated to substances that are no longer in use.

The official texts relating to the procedures for selecting, and general guidance for devising, international nonproprietary names are reproduced



in two annexes to the report. Other annexes give examples of international nonproprietary names that incorporate selected stems, the most frequently used initial groups of letters in international nonproprietary

names, a historical review of the programme of selecting international nonproprietary names, some useful literature references, and a model of the form to be used in all applications for international nonproprietary names.

<sup>1</sup> WHO Technical Report Series, No. 581, 1975 (*Nonproprietary Names for Pharmaceutical Substances*. Twentieth Report of the WHO Expert Committee), ISBN 92 4 120581 4. Price. Sw fr 8.-