

Dictionary
of
Organic
Compounds

SIXTH EDITION

VOLUME FOUR

F—Mer

Dictionary of Organic Compounds

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F-0-00001-M-0-00454



CHAPMAN & HALL

Electronic Publishing Division

London · Glasgow · Weinheim · New York · Tokyo · Melbourne · Madras

Published by Chapman & Hall, 2-6 Boundary Row, London SE1 8HN, UK

Chapman & Hall, 2-6 Boundary Row, London SE1 8HN, UK

Blackie Academic & Professional, Wester Cleddens Road, Bishopbriggs,
Glasgow G64 2NZ, UK

Chapman & Hall, Pappelallee 3, 69469 Weinheim, Germany

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Tokyo 102, Japan

Chapman & Hall Australia, 102 Dodds Street, South Melbourne, Victoria 3205, Australia

Chapman & Hall India, R. Seshadri, 32 Second Main Road, CIT East, Madras 600 035, India

First edition in three volumes published by

Eyre & Spottiswoode (Publishers) Ltd:

Volume I published 1934

Volume II published 1936

Volume III published 1937

Second edition of Volume I published 1943

Volumes II and III reprinted with supplement 1944

Volumes I, II and III reprinted 1947

Third edition in four volumes published 1953

Fourth edition in five volumes published 1965

Fifth edition in seven volumes published 1982 by Chapman and Hall

This Sixth edition in nine volumes published 1996

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Typeset and printed in Great Britain at the University Press, Cambridge

ISBN 0 412 54090 8 (Nine volume set)

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A catalogue record for this book is available from the British Library

Library of Congress Catalog Card Number: 94-69994

0957381

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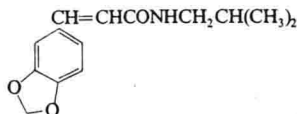
The specific information in this publication on the hazardous and toxic properties of certain substances is included to alert the reader to possible dangers associated with the use of those compounds. The absence of such information should not however be taken as an indication of safety in use or misuse.

F

Fagaramide

F-0-00001

3-(1,3-Benzodioxol-5-yl)-N-(2-methylpropyl)-2-propenamide, 9CI. N-Isobutyl-3,4-methylenedioxybenzamide



$C_{14}H_{17}NO_3$ M 247.2

(E)-form [495-86-3]

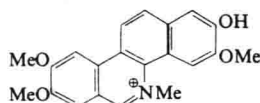
Alkaloid from the bark of *Zanthoxylum macrophyllum* and *Anthocleista vogelii*, the root of *Fagara tessmannii*, the root bark of *F. xanthoxyloides* and the wood of *Piper novae-hollandiae* (Rutaceae, Piperaceae). Plates (EtOAc). Mp 119.5° (softens at 105°).

Thoms, H. et al, Ber., 1911, **44**, 3717 (isol, struct, synth)
Goodson, J.A., Biochem. J., 1921, **15**, 123 (isol, struct)
Loder, J.W. et al, Aust. J. Chem., 1969, **22**, 1531 (isol, uv, ms)
Okorie, D.A., Phytochemistry, 1976, **15**, 1799 (isol, ir, pmr, ms)
Nagao, Y. et al, Tet. Lett., 1980, **21**, 841 (synth)

Fagaronine

F-0-00002

2-Hydroxy-3,8,9-trimethoxy-5-methylbenzo[c]phenanthridinium(1+), 9CI [52259-65-1]



$C_{21}H_{20}NO_4^+$ M 350.3 (ion)

Alkaloid from the roots of *Fagara xanthoxyloides* (Rutaceae). Active as a tumour inhibitor; very active antileukaemic agent. Inhibitor of RNA reverse transcriptase activity in oncogenic viruses; shows anti-HIV activity. An intercalating agent. Bactericidal.

► DI9861000.

Chloride:

$C_{21}H_{20}ClNO_4$ M 385.8 Yellow needles (EtOAc/MeOH). Mp 202° (198-200°), Mp 260-261° (double Mp).

Messmer, W.M. et al, J. Pharm. Sci., 1972, **61**, 1858 (ir, uv, ms, pmr, isol struct)

Gillespie, J.P. et al, J.O.C., 1974, **39**, 3239 (synth)

Tin-Wa, M. et al, J. Pharm. Sci., 1974, **63**, 1476 (struct)

Sethi, V.S. et al, Biochem. Biophys. Res. Commun., 1975, **63**, 1070 (pharmacol)

Phillips, S.D. et al, J. Het. Chem., 1981, **18**, 223 (rev)

Ishii, H. et al, Chem. Pharm. Bull., 1983, **31**, 2963 (synth)

Hanaoka, M. et al, Chem. Pharm. Bull., 1987, **35**, 2348 (synth, uv, ir, pmr, ms)

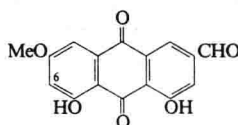
Ishii, H. et al, J.C.S. Perkin I, 1987, 671 (synth, ir, pmr)

Šmidrkal, J., Coll. Czech. Chem. Comm., 1988, **53**, 3184 (synth, uv, pmr)
Tan, G.T., J. Nat. Prod., 1991, **54**, 143 (anti-HIV activity)

Fallacinal

F-0-00003

9,10-Dihydro-4,5-dihydroxy-7-methoxy-9,10-dioxo-2-anthracenecarboxaldehyde, 9CI. 4,5-Dihydroxy-7-methoxyanthraquinone-2-carboxaldehyde. 3-Formyl-1,8-dihydroxy-6-methoxyanthraquinone. Phallacinal [4517-57-1]



$C_{16}H_{10}O_6$ M 298.2

Pigment of lichens e.g. *Anthracotheceum ochraceoflavum*, *Caloplaca* spp., *Cetraria cucullata*, *Fulgensia fulgens* and *Xanthoria* spp. Orange-yellow needles (C_6H_6 or Me_2CO). Mp 259.5-260.5° (251°).

6-Chloro: [41680-24-4]. **Chlorofallacinal**. 6-Chloro-4,5-dihydroxy-7-methoxyanthraquinone-2-carboxaldehyde. 2-Chloro-6-formyl-1,8-dihydroxy-3-methoxyanthraquinone

$C_{16}H_9ClO_6$ M 332.6 Isol. from the thalli an unidentified C. sp. Posn. of Cl suggested on biogenetic grounds.

Murakami, T., Chem. Pharm. Bull., 1956, **4**, 298.

Keogh, M.F. et al, Phytochemistry, 1972, **11**, 1495.

Nakano, H. et al, Phytochemistry, 1972, **11**, 3505 (Chlorofallacinal)

Gonzalez, G.A. et al, An. Quim., 1973, **69**, 805; CA, **79**, 104996 (ir, ms, nmr)

Yosioka, I. et al, Chem. Pharm. Bull., 1973, **21**, 1547.

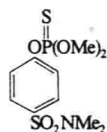
Renner, B. et al, Naturwissenschaften, 1978, **65**, 439.

Hirose, Y. et al, Chem. Pharm. Bull., 1982, **30**, 4186 (synth)

Famphur

F-0-00004

O-[4-(Dimethylamino)sulfonyl]phenyl O,O-dimethyl phosphorothioate, 9CI. O,O-Dimethyl phosphorothioate, O-ester with p-hydroxy-N,N-dimethylbenzenesulfonamide, 8CI. *Varbex*. *Famophos*. CL 38023. ENT 25644. *Warbex*. *Bo-Ana*. *Warbexol* [52-85-7]



$C_{10}H_{16}NO_4P_2S_2$ M 325.3

Insecticide and acaricide, cholinesterase inhibitor. Cryst. (toluene/cyclohexane). Spar. sol. H_2O , aliphatic hydrocarbons. Mp 52.5-53.5°.

► Nausea and headache result from occup. exposure. Skin irritant. LD₅₀ (rat, orl) 28 mg/kg. LD₅₀ (rbt, skn) 1460 mg/kg. TF7650000.

Turner, E.C. et al, J. Am. Vet. Med. Assoc., 1962, **141**, 360 (pharmacol)

O'Brien, R.D. et al, J. Agric. Food Chem., 1965, **13**, 366 (metab, tox)

U.S. Pat., 3 179 560, (1965); CA, **63**, 1738 (synth)

Gatterdam, P.E. et al, J. Agric. Food Chem., 1967, **15**, 845 (metab)

Black, W.D. et al, Toxicol. Appl. Pharmacol., 1979, **50**, 167 (tox)

Ding, X.D. et al, J. Agric. Food Chem., 1984, **32**, 622 (hplc)

Dangerous Prop. Ind. Mater. Rep., 1987, **7**, 91 (rev, tox)

Pesticide Manual, 9th edn., 1991, No. 5995.

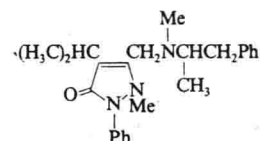
Lewis, R.J., *Sax's Dangerous Properties of Industrial Materials*, 8th edn., Van Nostrand Reinhold, 1992, FAB600.

Handbook of Pesticide Toxicology, (eds. Hayes, W.J. et al), Academic Press, 1991, 1039.

Famprofazone, BAN, INN

F-0-00005

4-Isopropyl-2-methyl-3-[[methyl(α-methylphenethyl)amino]methyl]-1-phenyl-3-pyrazolin-5-one, 8CI. 5-[N,α-Dimethyl(phenethylamino)methyl]-4-isopropylnorantipyrene [22881-35-2]



$C_{24}H_{31}N_3O$ M 377.5

Analgesic, antipyretic. Powder (EtOH). Mp 132-133°.

► UQ9451000.

Swiss Pat., 275 620, (1951); CA, **47**, 1745f.

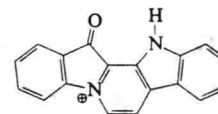
Martindale, The Extra Pharmacopoeia, 28th/29th edn., Pharmaceutical Press, London, 1982/1989, 2645.

Mrongovius, R. et al, Eur. J. Med. Chem. (Chim. Ther.), 1984, **19**, 161 (synth, pharmacol, metab)

Fascaplysin

F-0-00006

12,13-Dihydro-13-oxopyrido[1,2-a:3,4-b']diindol-5-ium(1+), 9CI



$C_{18}H_{11}N_2O^+$ M 271.2 (ion)

Pigment from the marine sponge

Fascaplysinopsis sp. Exhibits antimicrobial and cytotoxic props.

Chloride: [114719-57-2].

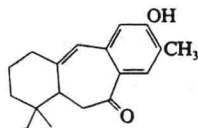
$C_{18}H_{11}ClN_2O$ M 306.7 Red cryst. (MeOH or $CHCl_3$). Mp 232-235° dec.

Roll, D.M. et al, J.O.C., 1988, **53**, 3276 (isol, uv, ir, pmr, cmr, ms, cryst struct)

Faveline **F-0-00007**

1,2,3,4,11,11a-Hexahydro-7-hydroxy-1,1,8-trimethyl-10H-dibenzo[a,d]cyclohepten-10-one, 9CI

[136624-32-3]



$C_{18}H_{22}O_2$ M 270.3

Isol. from the bark of *Cnidioscolus phyllacanthus*. Cytotoxic. Cryst. ($CHCl_3$ /hexane). Mp 192–194°. $[\alpha]_D^{25}$ –303.9 (c, 0.73 in $CHCl_3$).

Me ether: [136624-31-2].

$C_{19}H_{24}O_2$ M 284.3 Constit. of *C. phyllacanthus*. Cytotoxic. Cryst. (hexane). Mp 135–136°. $[\alpha]_D^{25}$ –344.2 (c, 0.91 in $CHCl_3$).

Deoxo: [136624-33-4]. **Deoxofaveline**

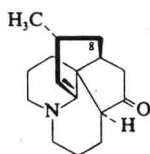
$C_{18}H_{20}O$ M 256.3 Constit. of *C. phyllacanthus*. Cytotoxic. Cryst. (hexane). Mp 149–151°. $[\alpha]_D^{25}$ +242.1 (c, 0.6 in $CHCl_3$).

Endo, Y. et al, *Tet. Lett.*, 1991, 32, 3083 (isol, pmr, cmr)

Ghosh, A.K. et al, *J.C.S. Perkin 1*, 1994, 327 (synth)

Fawcettidine **F-0-00008**

[14912-31-3]



$C_{16}H_{23}NO$ M 245.3

Alkaloid from *Lycopodium fawcettii*, *L. alopecuroides* and *L. phlegmaria* (Lycopodiaceae). Liq. Bp_{0.1} 125°.

Picrate: Mp 222–223°.

Methiodide: Mp 224–226°.

8S-Hydroxy: [62023-85-2].

Anhydroaposeratinine

$C_{16}H_{23}NO_2$ M 261.3 Alkaloid from *L. verticillatum* (Lycopodiaceae). Mp 151°. $[\alpha]_D^{25}$ +211 ($CHCl_3$).

8S-Hydroxy, Ac: Mp 144°.

Burnell, R.H., *J.C.S.*, 1959, 3091 (isol)

Ishii, H. et al, *Tet. Lett.*, 1966, 6215 (struct, ir)

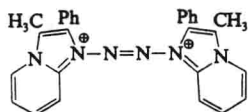
Ishii, H. et al, *Chem. Pharm. Bull.*, 1970, 18, 1880 (struct)

Nyembo, L. et al, *Bull. Soc. Chim. Belg.*, 1976, 85, 595; *CA*, 86, 90108u

(Anhydroaposeratinine)

Fazadinium(2+) **F-0-00009**

1,1'-Azobis[3-methyl-2-phenylimidazo[1,2-a]pyridinium], 9CI. Phenazinium [36653-54-0]



$C_{28}H_{24}N_8^{2+}$ M 444.5 (ion)

Dibromide: [49564-56-9]. **Fazadinium bromide**, BAN, INN. **Fazadon**. AH 8165. **Fazadan**

$C_{28}H_{24}Br_2N_8$ M 604.3 Neuromuscular blocking agent. Yellow cryst. + 2H₂O (H₂O). Mp 215–219° (softens at 196°).

LD₅₀ (rat, ivn) 0.91 mg/kg. NJ5830100.

Glover, E.E. et al, *J.C.S.(C)*, 1971, 3280 (synth)

Bolger, L. et al, *Nature (London)*, 1972, 238, 354 (pharmacol)

Playle, A.C., *Drugs of Today (Barcelona)*, 1977, 13, 95 (rev, pharmacol)

Bell, J.A. et al, *Br. J. Pharmacol.*, 1979, 66, 433P (metab)

Kienlen, J. et al, *Ann. Anesthesiol. Fr.*, 1981, 22, 631 (rev, pharmacol)

d'Hollander, A. et al, *Eur. J. Clin. Pharmacol.*, 1983, 24, 407 (pharmacol)

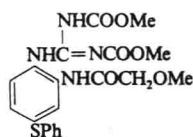
Martindale, *The Extra Pharmacopoeia*, 30th edn., Pharmaceutical Press, London, 1993, 1204.

Febantel, BAN, USAN, INN **F-0-00010**

[[2-[(Methoxyacetyl)amino]-4-(phenylthio)phenyl]carbonimidoyl]biscarbamic acid dimethyl ester, 9CI. **Amatron**. Bayverm.

Rintal. Bay Vh 5757

[58306-30-2]



$C_{26}H_{22}N_4O_6S$ M 446.4

Veterinary anthelmintic. Mp 129–130°.

LD₅₀ (rat, ori) 10605 mg/kg. FB3955000.

Ger. Pat., 2 423 679, (1975); *CA*, 84, 73949k (synth, pharmacol)

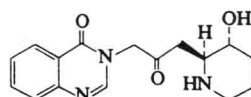
Wollweber, H. et al, *Arzneim.-Forsch.*, 1978, 28, 2193; 1984, 34, 531 (pharmacol, synth, props)

Martindale, *The Extra Pharmacopoeia*, 30th edn., Pharmaceutical Press, London, 1993, 43.

Lewis, R.J., *Sax's Dangerous Properties of Industrial Materials*, 8th edn., Van Nostrand Reinhold, 1992, BKN250.

Febrifugine† **F-0-00011**

3-[3-(3-Hydroxy-2-piperidinyl)-2-oxopropyl]-4(3H)-quinazolinone, 9CI. **Dichroine B**. β -Dichroine. γ -Dichroine



Absolute configuration

$C_{16}H_{19}N_3O_3$ M 301.3

(+)-form [24159-07-7]

Alkaloid from the roots and leaves of the Chinese drug Ch'ang Shan (*Dichroa febrifuga*) and from *Hydrangea* spp. (Hydrangeaceae). Shows high degree of antimalarial activity (ca. 100 times as active as Quinine) but use limited by toxicity. Also shows antipyretic and emetic activities; an effective coccidiostat. Various synthetic analogues show similar activities and much lower toxicity. Antineoplastic agent. Needles (EtOH). Mp 139–140° or 154–156° (dimorph). $[\alpha]_D^{25}$ +6 (c, 0.5 in $CHCl_3$); +28 (c, 0.5 in EtOH).

Hydrochloride (1:2): Mp 220–222° dec.

Bis(benzenesulfonyl): Mp 148–148.5°.

Semicarbazone: Mp 187–188° dec.

Oxime: Mp 224–225° dec.

N-Nitroso: Mp 171.5–172.5°.

(±)-form [39037-90-6]

Synthetic. Mp 178.5–180.5°.

Hydrobromide: Cryst. (EtOH). Mp 229–231°.

Koepfli, J.B. et al, *J.A.C.S.*, 1947, 69, 1837;

1949, 71, 1048 (isol, uv)

Kuehl, F.A. et al, *J.A.C.S.*, 1948, 70, 2091 (isol)

Abbondi, F. et al, *J.O.C.*, 1952, 17, 14 (isol)

Baker, B.R. et al, *J.O.C.*, 1955, 20, 136 (synth)

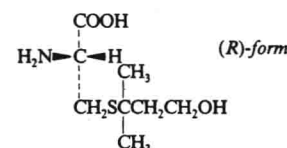
Barringer, D.F. et al, *J.O.C.*, 1973, 38, 1933,

1937 (pmr, abs config)

Johns, S., *Alkaloids (N.Y.)*, 1986, 29, 129 (rev, pharmacol)

Feniline **F-0-00012**

S-(3-Hydroxy-1,1-dimethylpropyl)cysteine, 9CI. 3-[(3-Hydroxy-1,1-dimethylpropyl)thio]alanine, 8CI



$C_8H_{17}NO_3S$ M 207.2

(R)-form [471-09-0]

L-form

Occurs in the urine of cats. Amorph. ppt. (Me_2CO aq.). Mp 185° (177°). $[\alpha]_D^{25}$ –12.5 (c, 2 in H₂O), $[\alpha]_D^{25}$ +9.0 (c, 2 in 1M HCl).

(±)-form

Cryst. (EtOH aq.). Mp 181° dec.

N-2,4-Dinitrophenyl: Cryst. (EtOH aq.). Mp 120–122°.

Schoeberl, A. et al, *Chem. Ber.*, 1968, 101, 373 (synth)

Felypressin, BAN, INN, USAN **F-0-00013**

2-L-Phenylalanine-8-L-lysinevasopressin, 9CI. **Collupressine**. **Octapressin**. **Octopressin**. **Phelypressin**. PLV 2

[56-59-7]

Cys-Phe-Phe-Gln-Asn-Cys-Pro-Lys-Gly-NH₂

$C_{46}H_{65}N_{13}O_{11}S_2$ M 1040.2

Synthetic deriv. of vasopressin.

Vasodilator. No phys. props. reported.

Component of numerous preparations.

Boissonnas, R.A. et al, *Helv. Chim. Acta*, 1960, 43, 190 (synth)

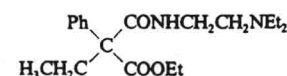
Meienhofer, J. et al, *J.A.C.S.*, 1960, 82, 6336 (synth)

Schmidt, L. et al, *Arzneim.-Forsch.*, 1962, 12, 1019 (pharmacol)

Martindale, *The Extra Pharmacopoeia*, 30th edn., Pharmaceutical Press, London, 1993, 953.

Fenalamide, USAN, INN **F-0-00014**

α -[[2-(Diethylamino)ethylamino]carbonyl]- α -ethylbenzeneacetic acid ethyl ester, 9CI. Ethyl N-[2-(diethylamino)ethyl]-2-ethyl-2-phenylmalonamate, 8CI. **Phemamide**. **Spasmamide**. Sch 5706. SH 30858. **Femamide** [4551-59-1]



$C_{19}H_{29}N_2O_3$ M 334.4

LD₅₀ (rat, ori) 610 mg/kg. ON9110800.

(±)-form

Smooth muscle relaxant. Oil. Bp₃ 182-188°.

Hydrochloride: [15734-93-7].

Mp 74-80°.

Hydrobromide: [5957-23-3].

Mp 71-74°.

U.K. Pat., 870 887, (1959); CA, 56, 550c (synth)

Carminati, G.M., Arch. Int. Pharmacodyn.

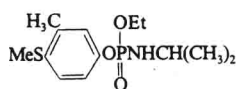
Ther., 1965, 153, 312 (pharmacol)

Martindale, The Extra Pharmacopoeia, 30th edn., Pharmaceutical Press, London, 1993, 1369.

Lewis, R.J., Sax's Dangerous Properties of Industrial Materials, 8th edn., Van Nostrand Reinhold, 1992, FAJ150.

Fenamiphos, BSI**F-0-00015**

Ethyl 3-methyl-4-(methylthio)phenyl 1-methylethylphosphoramidate, 9CI. Ethyl 4-methylthio-m-tolyl isopropylphosphoramidate, 8CI. Nemacur. Phenamiphos [22224-92-6]



C₁₃H₂₂N₃O₃PS M 303.3

Systemic agricultural nematocide.

Cholinesterase inhibitor. Solid. Spar. sol. H₂O. Mp 49°.

► LD₅₀ (rat, ori) 8 mg/kg. LD₅₀ (rbt, skn) 178 mg/kg. ACGIH TLV 0.1 mg m⁻³ (Sk). TB3675000.

S-Oxide: [31972-43-7].

C₁₃H₂₂N₃O₄PS M 319.3 n_D²⁰ 1.5274.

S,S-Dioxide: [31972-44-8].

C₁₃H₂₂N₃O₅PS M 335.3 Mp 77°.

Ger. Pat., 1 934 001, (1971); CA, 74, 141266f (oxides)

Waggoner, T.B. et al, J. Agric. Food Chem., 1972, 20, 157 (metab)

Waggoner, T.B. et al, Residue Rev., 1974, 53, 79 (rev, pharmacol)

Ripley, B.D. et al, J. Assoc. Off. Anal. Chem., 1983, 66, 1084 (glc)

Bowman, B.T. et al, J. Environ. Sci. Health, Part B, 1983, 18, 221 (props)

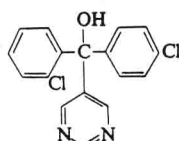
Pesticide Manual, 9th edn., 1991, No. 6010.

Lewis, R.J., Sax's Dangerous Properties of Industrial Materials, 8th edn., Van Nostrand Reinhold, 1992, FAK000.

Fenarimol, BSI, ISO, ANSI**F-0-00016**

α-(2-Chlorophenyl)-α-(4-chlorophenyl)-5-pyrimidinemethanol, 9CI. 2,4'-Dichloro-α-(5-pyrimidinyl)benzhydrol alcohol. Bloc. Rimidin. Rubigan

[60168-88-9]



C₁₇H₁₂Cl₂N₂O M 331.2

► UV9279400.

(±)-form

Agricultural and horticultural fungicide. Cryst. Mp 117-119°.

► LD₅₀ (rat, ori) 2500 mg/kg. Exp. reprod. effects.

U.K. Pat., 1 218 623, (1967); CA, 72, 100745b.

Dây, E.W. et al, Anal. Methods Pestic. Plant Growth Regul., 1984, 13, 173 (dein)

Sisler, H.D. et al, Pestic. Sci., 1984, 15, 167 (rev, activity)

Taylor, H.M. et al, J. Med. Chem., 1987, 30, 1359 (synth)

Albinati, A. et al, Acta Cryst. C, 1988, 44, 1782 (cryst struct)

Pesticide Manual, 9th edn., 1991, No. 6030.

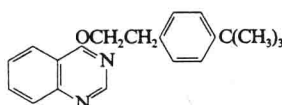
Gardyan, C. et al, Z. Lebensm.-Unters.-Forsch., 1991, 192, 40 (hplc)

Agrochemicals Handbook, 3rd edn., Royal Society of Chemistry, 1992, A193.

Lewis, R.J., Sax's Dangerous Properties of Industrial Materials, 8th edn., Van Nostrand Reinhold, 1992, FAK100.

Fenazaquin, BSI**F-0-00017**

4-[2-[4-(1,1-Dimethylethyl)phenyl]ethoxy]quinazoline, 9CI. 4-tert-Butylphenethyl 4-quinazolinyl ether. Magister. EL-436 [120928-09-8]



C₂₀H₂₂N₂O M 306.4

Acaricide. Cryst. Mp 70-71°.

► LD₅₀ (rat, ori) 50-500 mg/kg. VA1382000.

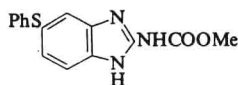
Eur. Pat., 326 329, (1989); CA, 112, 35882x (synth, activity)

Eur. Pat., 380 264, (1990); CA, 114, 6539b (synth)

Pesticide Manual, 9th edn., 1991, No. 6055.

Fenbendazole, BAN, USAN, INN**F-0-00018**

Methyl 5-phenylthio-1H-benzimidazol-2-yl carbamate. Axilur. Panacur. Pancacur. Safe-Guard. Worm-A-Rest. Hoe 881 [43210-67-9]



C₁₅H₁₃N₃O₂S M 299.3

Veterinary anthelmintic. Powder. Mp 233° dec.

► Mutagenic props. DD6520500.

Baeder, C. et al, Experientia, 1974, 30, 753.

Ger. Pat., 2 332 486, (1974); CA, 83, 43324q.

Averkin, E.A. et al, J. Med. Chem., 1975, 18, 1164 (synth, pharmacol)

Duewel, D., Pestic. Sci., 1977, 8, 550 (props)

Martindale, The Extra Pharmacopoeia, 30th edn., Pharmaceutical Press, London, 1993, 43.

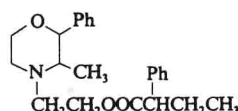
Lewis, R.J., Sax's Dangerous Properties of Industrial Materials, 8th edn., Van Nostrand Reinhold, 1992, FAL100.

Fenbutrazate, INN**F-0-00019**

α-Ethylbenzeneacetic acid 2-(3-methyl-2-phenyl-4-morpholinyl)ethyl ester, 9CI. 2-Phenylbutyric acid 2-(3-methyl-2-phenylmorpholino)ethyl ester, 8CI.

Phenbutrazate, BAN

[4378-36-3]



C₂₃H₂₉N₃O₃ M 367.4

Anorexic drug. Central stimulant. Honey-coloured, viscous oil. Sol. MeOH. Bp_{0.05} 235-240°.

► ET5979000.

Hydrochloride: [6474-85-7]. R 381

Insol. Et₂O, C₆H₆, sol. EtOH, Me₂CO. Mp 154°.

► LD₅₀ (mus, ori) 3200 mg/kg. ET5980000.

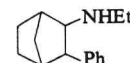
Hengen, O., Arzneim.-Forsch., 1955, 5, 526; 1957, 7, 524 (pharmacol)

U.S. Pat., 3 018 222, (1962); CA, 58, 6840e (synth)

Martindale, The Extra Pharmacopoeia, 30th edn., Pharmaceutical Press, London, 1993, 1230.

Fencamfamin, BAN, INN**F-0-00020**

N-Ethyl-3-phenylbicyclo[2.2.1]heptan-2-amine, 9CI. N-Ethyl-3-phenyl-2-norbornanamine, 8CI. 2-Ethylamino-3-phenylbicyclo[2.2.1]heptane. Euwitol. Glucoenergan. W 1206 [1209-98-9]



C₁₅H₂₁N M 215.3

Used as CNS stimulant and appetite suppressant. Liq. Bp_{0.1} 128-130°.

► RB6650000.

Hydrochloride: [2240-14-4]. Norcamphane. H 610

Principal ingredient of Reactivan. Cryst. (Me₂CO). Mp 192°. Reactivan also contains vitamins B₁, B₆, B₁₂, C.

► LD₅₀ (rat, ori) 83 mg/kg. RB6825000.

Hotovy, R. et al, Arzneim.-Forsch., 1961, 11, 20, 24 (pharmacol)

Ger. Pat., 1 110 159, (1961); CA, 56, 2352 (synth)

Ger. Pat., 1 219 478, (1966); CA, 65, 12136 (synth)

Valzelli, L. et al, Psychopharmacologia, 1971, 20, 91 (pharmacol)

Martindale, The Extra Pharmacopoeia, 30th edn., Pharmaceutical Press, London, 1993, 1226.

Lewis, R.J., Sax's Dangerous Properties of Industrial Materials, 8th edn., Van Nostrand Reinhold, 1992, EOM000.

Fencarbamide, INN**F-0-00021**

Phencarbamide, USAN. S-[2-(Diethylamino)ethyl] diphenylcarbamothioate, 9CI. S-[2-(Diethylamino)ethyl] diphenylthiocarbamate, 8CI. Diphencarbamide. Escorpal. Rispassulf. Bayer 1355. Wh 3363 [3735-90-8]

Ph₂NCOSCH₂CH₂NEt₂

C₁₉H₂₄N₂OS M 328.4

Antispasmodic and anticholinergic drug used during parturition. Cryst. Sol. MeOH, Et₂O, CHCl₃, petrol; prac. insol. H₂O.

► LD₅₀ (rat, ori) 370 mg/kg. FA4439000.

Hydrochloride: [58-13-9].

Cryst. (MeCOEt). Sol. H₂O. Mp 180-182°.

► LD₅₀ (rat, ori) 410 mg/kg. FA4440000.

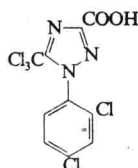
Anon, Angew. Chem., Int. Ed., 1964, 3, 68 (rev)

Wirth, W. et al, Arch. Int. Pharmacodyn. Ther., 1964, 151, 515 (pharmacol)

Risse, K.H. et al, CA, 1964, 60, 7939 (synth)

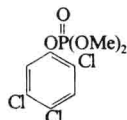
Madan, B.R. *et al*, *Arch. Int. Pharmacodyn. Ther.*, 1970, **185**, 53 (pharmacol)
 Martindale, *The Extra Pharmacopoeia*, 30th edn., Pharmaceutical Press, London, 1993, 430.
 Lewis, R.J., *Sax's Dangerous Properties of Industrial Materials*, 8th edn., Van Nostrand Reinhold, 1992, PDC850, PDC875.

Fenchlorazole, BSI **F-0-00022**
 1-(2,4-Dichlorophenyl)-5-(trichloromethyl)-1H-1,2,4-triazole-3-carboxylic acid, 9CI
 [103112-36-3]



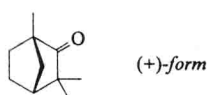
$C_{10}H_4Cl_5N_3O_2$ M 375.4
 Mp 108-112°. Inactive compared to ethyl ester.
 ▶ LD₅₀ (rat, ori) > 5000 mg/kg. XZ4195000.
 Et ester: [103112-35-2]. Fenchlorazole-ethyl. HOE 070 542
 $C_{12}H_8Cl_5N_3O_2$ M 403.4 Herbicide safener.
 Ger. Pat., 3 525 205, (1986); CA, **105**, 37496d (synth, activity)
 Bieringer, H. *et al*, *Proc. Br. Crop Prot. Conf. Weeds*, 1989, 77, 495 (activity)
Pesticide Manual, 9th edn., 1991, No. 6065.

Fenchlorphos-oxon **F-0-00023**
 Dimethyl 2,4,5-trichlorophenylphosphite, 9CI.
 Ronnel oxon
 [3983-45-7]



$C_8H_8Cl_3O_4P$ M 305.4
 Insecticide. d₄²⁵ 1.44. Bp_{0.4} 151-154°. n_D²⁵ 1.5335.
 ▶ TC5425000.
 U.S. Pat., 2 599 515, (1952); CA, **46**, 8322a (synth)
 Baughmann, R.G. *et al*, *J. Agric. Food Chem.*, 1978, **26**, 403 (cryst struct)
 Sanchez-Baeza, F. *et al*, *Tet. Lett.*, 1990, **31**, 3359 (synth, pmr, cmr)
 Durand, G. *et al*, *Biol. Mass. Spectrom.*, 1991, **20**, 3 (ms)

Fenchone **F-0-00024**
 1,3,3-Trimethylbicyclo[2.2.1]heptan-2-one, 9CI.
 Fenchol (obsol.)
 [1195-79-5]

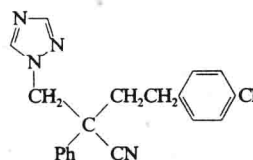


$C_{10}H_{16}O$ M 152.2
 Fp 5-6°.
 ▶ Skin irritant. LD₅₀ (rat, ori) 4400 mg/kg. Fl. p. 65°. RB7875000.
 (+)-form [7787-20-4]
 Widespread in plants, e.g. found in fennel

oil (*Foeniculum vulgare*), *Blumea lacera*, *Prunella vulgaris*. Can be used as H acceptor in dehydrogenations. Oil with camphoraceous odour. Prac. insol. H₂O, v. sol. EtOH. Mp 3-5°, Fp 5-6°. Bp 193.5°.
 [α]_D²⁰ +63.

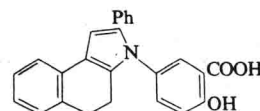
E-Oxime: [25072-90-6].
 $C_{10}H_{17}NO$ M 167.2 Mp 132°.
 Z-Oxime: [25072-89-3].
 $C_{10}H_{17}NO$ M 167.2 Mp 164-165°.
 [α]_D²⁰ +65.2 (c. 5.8 in MeOH).
 (-)-form [4695-62-9]
 Isol. from various essential oils e.g. *Thuja occidentalis* and *Artemisia frigida*. [α]_D²⁰ -67.
 ▶ RB7875200.
 (±)-form [18492-37-0]
 Bp 192-194°.
 Oxime: [2157-47-3].
 Mp 158-159°.
 [74163-80-7]
 Aldrich Library of FT-IR Spectra: Vapor Phase, 3, 527C (ir)
 Aldrich Library of ¹³C and ¹H FT NMR Spectra, 1, 678B (nmr)
 Aldrich Library of FT-IR Spectra, 1st edn., 1, 442B (ir)
 Komppa, G. *et al*, *Ber.*, 1935, **68**, 2001 (synth)
 Bays, D.E. *et al*, *J.C.S.(B)*, 1966, 885 (cd)
 Thomas, A.F. *et al*, *Helv. Chim. Acta*, 1967, **50**, 826 (ms)
 Baker, K.M. *et al*, *Tetrahedron*, 1968, **24**, 1663 (pmr)
 Lipman, E. *et al*, *Org. Magn. Reson.*, 1970, **2**, 581 (cmr)
 Boyle, P.H. *et al*, *J.C.S.(C)*, 1971, 2136 (synth)
 Opdyke, D.L.J., *Food Cosmet. Toxicol.*, 1976, **14**, 769 (rev, tox)
 Fieser and Fieser's *Reagents for Organic Synthesis*, Wiley, 1980, 8, 228.
 Buchbauer, G. *et al*, *Annalen*, 1981, 2093 (synth)
 Wijekoon, W.M.D. *et al*, *J. Phys. Chem.*, 1983, **87**, 3034 (pmr, cmr, uv, cd)
 Lewis, R.J., *Sax's Dangerous Properties of Industrial Materials*, 8th edn., Van Nostrand Reinhold, 1992, TLW250.

Fencobuconazole, BSI **F-0-00025**
 α-[2-(4-Chlorophenyl)ethyl]-α-phenyl-1H-1,2,4-triazole-1-propanenitrile, 9CI. 4-(4-Chlorophenyl)-2-phenyl-2-(1H-1,2,4-triazol-1-ylmethyl)butyronitrile. RH 7592
 [114369-43-6]



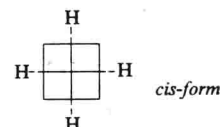
$C_{19}H_{17}ClN_4$ M 336.8
 ▶ XZ5257500.
 (±)-form [119611-00-6]
 Fungicide. Steroid demethylation inhibitor. Cryst. Mp 124-126°.
 ▶ LD₅₀ (rat, ori) > 2000 mg/kg. XZ5258000.
 [119611-00-6]
 Ger. Pat., 3 721 786, (1988); CA, **108**, 200219p (synth, activity)
Pesticide Manual, 9th edn., 1991, 157.
 Jeffers, S.N., *Plant Dis.*, 1991, **75**, 502 (activity)

Fendosal, BAN, INN, USAN **F-0-00026**
 5-(4,5-Dihydro-2-phenyl-3H-benz[e]indol-3-yl)-2-hydroxybenzoic acid, 9CI. Alnovin. HP 129.
 P 710129
 [53597-27-6]



$C_{25}H_{19}NO_3$ M 381.4
 Analgesic, antiinflammatory and antipyretic agent. Cryst. (AcOH). Mp 223-225° dec.
 ▶ LD₅₀ (rat, ori) 450 mg/kg. DG8576200.
 Anderson, V.B. *et al*, *J. Med. Chem.*, 1976, **19**, 318 (synth, pharmacol)
 Lassman, H.B. *et al*, *Agents Actions*, 1978, **8**, 209 (pharmacol)
 Warrander, A. *et al*, *Drug Metab. Dispos.*, 1981, **9**, 161 (metab)
 Martindale, *The Extra Pharmacopoeia*, 30th edn., Pharmaceutical Press, London, 1993, 14.
 Lewis, R.J., *Sax's Dangerous Properties of Industrial Materials*, 8th edn., Van Nostrand Reinhold, 1992, FA0100.

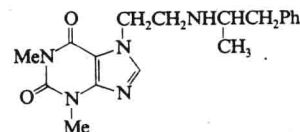
[4.4.4]Fenestrane **F-0-00027**
 Tetracyclo[3.3.1.0^{3,9}.0^{7,9}]nonane, 9CI.
 Tetracyclo[5.1.1.0^{3,8}.0^{5,8}]nonane. Windowpane
 [40220-30-2]



C_9H_{12} M 120.1
 A molecule of considerable interest from point of view of obt. planar tetracoordinate carbon but may be too strained for its synth. to be achieved.

Georgian, V. *et al*, *Tet. Lett.*, 1972, 4315.
 Wiberg, K.B. *et al*, *J.A.C.S.*, 1976, **98**, 1212.
 Minkin, V.I. *et al*, *Zh. Org. Khim.*, 1978, **14**, 3.
 Hoeve, W.T. *et al*, *J.O.C.*, 1980, **45**, 2925, 2980.
 Würthein, E.-U. *et al*, *Tet. Lett.*, 1981, 843.
 Krohn, K., *Org. Synth. Highlights*, 1991, 371 (rev)

Fenetylline, INN **F-0-00028**
 3,7-Dihydro-1,3-dimethyl-7-[2-(1-methyl-2-phenylethyl)aminoethyl]-1H-purine-2,6-dione, 9CI. 7-[2-[(α-Methylphenethylamino)]ethyl]theophylline. Amphetaminetheophylline. Amfetyline. Fenethylline, BAN. Biocapton
 [3736-08-1]

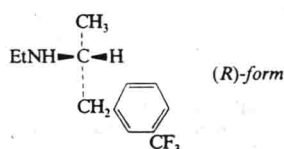


$C_{18}H_{23}N_5O_2$ M 341.4
 ▶ XH6110000.
 (±)-form
 Central stimulant, antidepressant. Metabolites in man incl. amphetamine.
 Hydrochloride: [1892-80-4]. Fenethylline hydrochloride, USAN. Captagon. H 814
 Mp 227-229°, Mp 237-239°.
 ▶ LD₅₀ (rat, ori) 100 mg/kg. XH6125000.

- Wagemann, U. *et al*, *Arzneim.-Forsch.*, 1959, **9**, 311 (*pharmacol*)
Ger. Pat., 1 123 329, (1962); *CA*, **57**, 5933 (*synth*)
 Ellison, T. *et al*, *Eur. J. Pharmacol.*, 1974, **13**, 123 (*metab*)
 Gagauzov, I. *et al*, *CA*, 1984, **101**, 130491 (*synth*)
 Nickel, B. *et al*, *Drug Alcohol Depend.*, 1986, **235** (*pharmacol, metab*)
 Negwer, M., *Organic-Chemical Drugs and their Synonyms*, 6th edn., Akademie-Verlag, Berlin, 1987, 4919 (*synonyms*)
 Yoshimura, H. *et al*, *Xenobiotica*, 1988, **18**, 929 (*metab*)
 Martindale, *The Extra Pharmacopoeia*, 30th edn., Pharmaceutical Press, London, 1993, 1226.
 Lewis, R.J., *Sax's Dangerous Properties of Industrial Materials*, 8th edn., Van Nostrand Reinhold, 1992, CBF825.

Fenfluramine, BAN F-0-00029

N-Ethyl- α -methyl-3-(trifluoromethyl)benzeneethanamine, 9CI. N-Ethyl- α -methyl-m-(trifluoromethyl)phenethylamine, 8CI. 2-Ethylamino-1-(3-(trifluoromethyl)phenyl)propane. Many other names
 [458-24-2]



$C_{12}H_{16}F_3N$ M 231.2

▶ SH6820000.

(R)-form [37577-24-5] **Levofenfluramine**

$[\alpha]_D^{25}$ -9.5 (c, 8.0 in EtOH).

▶ SH6824000.

(S)-form [3239-44-9] **Dexfenfluramine, INN.**

Adifax. S5614

Appetite suppressant. $[\alpha]_D^{25}$ +9.5 (c, 8.0 in EtOH).

▶ SH6822000.

Hydrochloride: [3239-45-0].

Cryst. (EtOAc). Mp 160-161°.

(±)-form [5220-89-3]

Anorexic agent. Bp₁₂ 108-112°.

▶ SH6821000.

Hydrochloride: [404-82-0]. **Fenfluramine**

hydrochloride, *USAN*. *Ponderax*. *Pondimin*. Anorexic. Mp 166°.

▶ Common adverse effects are drowsiness, diarrhoea, dizziness and lethargy.

Psychotomimetic agent if abused. LD₅₀ (rat, ori) 69 mg/kg. SH6825000.

Belg. Pat., 609 630, (1964); *CA*, **60**, 1647 (*synth*)

U.S. Pat., 3 198 834, (1965); *CA*, **63**, 11425 (*synth*)

Beckett, A.H. *et al*, *Tetrahedron*, 1968, **24**, 1283 (*abs config*)

Beckett, A. *et al*, *J. Chromatogr.*, 1972, **69**, 285 (*glc*)

Morgan, C.D. *et al*, *J. Pharmacol. Exp. Ther.*, 1972, **180**, 127 (*metab*)

Smith, H.E. *et al*, *J.O.C.*, 1975, **40**, 1562 (*cd, conformn*)

Caccia, S. *et al*, *Arch. Int. Pharmacodyn. Ther.*, 1982, **258**, 15 (*metab*)

Caccia, S. *et al*, *Eur. J. Clin. Pharmacol.*, 1985, **29**, 221 (*pharmacol*)

Negwer, M., *Organic-Chemical Drugs and their Synonyms*, 6th Ed., Akademie-Verlag, Berlin, 1987, 2410.

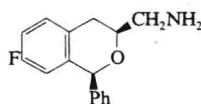
McTavish, D. *et al*, *Drugs*, 1992, **43**, 713 (*rev*)

Martindale, *The Extra Pharmacopoeia*, 30th edn., Pharmaceutical Press, London, 1993, 1226.

Lewis, R.J., *Sax's Dangerous Properties of Industrial Materials*, 8th edn., Van Nostrand Reinhold, 1992, ENJ000, PDM250.

Fenimorex, BAN, USAN, INN F-0-00030

7-Fluoro-3,4-dihydro-1-phenyl-1H-2-benzopyran-3-methanamine, 9CI. R 800
 [34887-52-0]



$C_{16}H_{16}FNO$ M 257.3

▶ DJ2548200.

(IRS,3RS)-form

cis-form

Anorexic agent.

Hydrochloride: Cryst. (EtOH/Et₂O). Mp 227.5-228°.

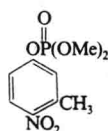
U.S. Pat., 3 743 659, (1973); *CA*, **79**, 78614d (*synth*)

U.S. Pat., 3 851 062, (1974); *CA*, **82**, 133170v (*pharmacol*)

Fenitrooxon F-0-00031

Dimethyl 3-methyl-4-nitrophenyl phosphate, 9CI. 8CI. Dimethyl 4-nitro-m-tolyl phosphate. Sumioxon

[2255-17-6]



$C_9H_{12}NO_6P$ M 261.1

Liq. Product of photodegradation and metab. of Fenitrothion, F-0-00032. Bp_{0.07} 134-136°. n_D^{20} 1.5165.

▶ LD₅₀ (rat, ori) 24 mg/kg, LD₅₀ (rat, skn) 1300 mg/kg. TC5260000.

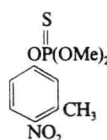
Miyamoto, J. *et al*, *Agric. Biol. Chem.*, 1963, **27**, 669 (*synth*)

Ohkawa, H. *et al*, *Agric. Biol. Chem.*, 1974, **38**, 2247 (*tlc*)

Lewis, R.J., *Sax's Dangerous Properties of Industrial Materials*, 8th edn., Van Nostrand Reinhold, 1992, PHD750.

Fenitrothion, BAN, BSI F-0-00032

O,O-Dimethyl O-(3-methyl-4-nitrophenyl) phosphorothioate, 9CI. O,O-Dimethyl O-4-nitro-m-tolyl phosphorothioate, 8CI. O,O-Dimethyl O-(3-methyl-4-nitrophenyl) thiophosphate. Accothion. Cyfen. Folithion. Metathion. Methylnitrothios. Phenitrothion. Sumithion. Arbogal
 [122-14-5]



$C_9H_{12}NO_4PS$ M 277.2

Contact insecticide. Cholinesterase inhibitor.

Yellow oil. Insol. H₂O, spar. sol. petrol. d_4^{25} 1.323. Bp_{0.1} 140-145°, Bp_{0.05} 118°. n_D^{25}

1.5528. Stable to acidic but not basic hydrol. S-Me isomer is formed at high temp. Component of Nuvantop and Zeproxx.

▶ Symptoms from occupational exposure incl. malaise, headache, anorexia. LD₅₀ (rat, ori) 250 mg/kg; LD₅₀ (rat, skn) 750 mg/kg. TG0350000.

Schrader, G., *Angew. Chem.*, 1961, **73**, 331 (*synth, props*)

Nishizawa, Y. *et al*, *Residue Rev.*, 1970, **60** (*bibl, rev*)

Stan, H.-J. *et al*, *Fresenius' Z. Anal. Chem.*, 1977, **287**, 271 (*glc, ms*)

Volpé, G. *et al*, *Chromatographia*, 1981, **14**, 333 (*hplc*)

Stan, H.-J. *et al*, *Biomed. Mass Spectrom.*, 1982, **9**, 483 (*ms*)

Agrochemicals Handbook, The Royal Society of Chemistry, 1983, A199.

Wang, Y., *CA*, 1986, **104**, 129985 (*synth*)

Pesticide Manual, 9th edn., 1991, No. 6120.

Martindale, *The Extra Pharmacopoeia*, 30th edn., Pharmaceutical Press, London, 1993, 1127.

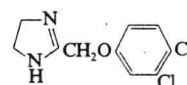
Lewis, R.J., *Sax's Dangerous Properties of Industrial Materials*, 8th edn., Van Nostrand Reinhold, 1992, DSQ000.

Handbook of Pesticide Toxicology, (Eds. Hayes, W.J. *et al*), Academic Press, 1991, 1020.

Fenmetozole, INN F-0-00033

2-[(3,4-Dichlorophenoxy)methyl]-4,5-dihydro-1H-imidazole, 9CI. 2-[(3,4-Dichlorophenoxy)methyl]-2-imidazoline, 8CI

[41473-09-0]



$C_{10}H_{10}Cl_2N_2O$ M 245.1

Antidepressant, narcotic antagonist.

Hydrochloride: [23712-05-2]. **Fenmetozole hydrochloride**, *USAN*. *DH 524*
 Mp 244-245°.

▶ LD₅₀ (rat, ori) 100 mg/kg. NJ2625000.

U.S. Pat., 3 449 355, (1969); *CA*, **71**, 81365p (*synth, pharmacol*)

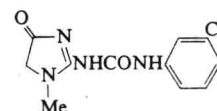
Reitz, R. *et al*, *Proc. Soc. Exp. Biol. Med.*, 1973, **144**, 391 (*props*)

Martindale, *The Extra Pharmacopoeia*, 30th edn., Pharmaceutical Press, London, 1993, 254.

Lewis, R.J., *Sax's Dangerous Properties of Industrial Materials*, 8th edn., Van Nostrand Reinhold, 1992, DGA800.

Fenobam, USAN, INN F-0-00034

N-(3-Chlorophenyl)-N'-(4,5-dihydro-1-methyl-4-oxo-1H-imidazol-2-yl)urea, 9CI. McN 3377
 [57653-26-6]



$C_{11}H_{11}ClN_4O_2$ M 266.6

Tranquilliser, muscle relaxant. Cryst. (Me₂CO/Et₂O). Mp 180-180.5°.

U.S. Pat., 3 983 135, (1976); *CA*, **86**, 55441 (*synth, pharmacol*)

Martindale, *The Extra Pharmacopoeia*, 28th/29th edn., Pharmaceutical Press, London, 1982/1989, 12740.
 Pellizer, G. et al, *J. Pharm. Sci.*, 1983, 72, 189 (pmr, cmr)

Fenobucarb, BSI F-0-00035

2-(1-Methylpropyl)phenyl methylcarbamate.
Osbac. Bassa. Baycarb. 2-sec-Butylphenyl N-methylcarbamate. BPMC, JMAF
 [3766-81-2]



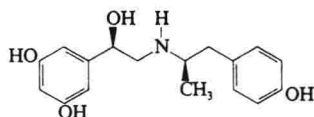
$C_{12}H_{17}NO_2$ M 207.2
 Insecticide against rice hoppers. Mp 32°.

► LD₅₀ (rat, orl) 410 mg/kg. FB5425000.
 [75863-93-3]

Fraser, J. et al, *J. Sci. Food Agric.*, Suppl. 8, 1968 (synth, use)
 Grou, E. et al, *J. Chromatogr.*, 1983, 260, 502 (hplc)
Pesticide Manual, 9th edn., 1991, No. 6130.
 Kulkarni, G.H. et al, *Tetrahedron*, 1991, 47, 1249 (synth, pmr)
Agrochemicals Handbook, 3rd edn., Royal Society of Chemistry, 1992, A958.
 Lewis, R.J., *Sax's Dangerous Properties of Industrial Materials*, 8th edn., Van Nostrand Reinhold, 1992, MOV000.

Fenoterol, BAN, USAN F-0-00036

5-[1-Hydroxy-2-[[2-(4-hydroxyphenyl)-1-methylethyl]amino]ethyl]-1,3-benzenediol, 9CI.
 3,5-Dihydroxy-α-[[p-hydroxy-α-methylphenethyl]amino]methylbenzyl alcohol, 8CI. *Airum. Dosberotec. Partusisten. Respilac. Segamol*
 [13392-18-2]



$C_{17}H_{21}NO_4$ M 303.3

► DO1490000.

(R,R)-form

Bronchodilator.

► DO1490000.

Hydrochloride: [1944-10-1].

Mp 183°.

Hydrobromide: [1944-12-3]. *Berotec. Th 1165a*
 Cryst. (MeOH/Et₂O). Mp 222-223°.

► Human reprod. effects reported. LD₅₀ (rat, orl) 1600 mg/kg. Exp. reprod. effects. DO1500000.

U.S. Pat., 3 135 797, (1964); CA, 61, 3022 (synth)

Beale, J.P., *Cryst. Struct. Commun.*, 1972, 1, 67, 223 (cryst struct)

Beale, J.P. et al, *J. Pharm. Pharmacol.*, 1972, 24, 277 (cryst struct)

Ger. Pat., 2 413 102, (1975); CA, 84, 30653s (synth)

Heel, R.C. et al, *Drugs*, 1978, 15, 3 (rev)

Kojima, S. et al, *Arzneim.-Forsch.*, 1980, 30, 959 (metab)

Kelly, H.W., *Clin. Pharm.*, 1985, 4, 393 (rev)

Svedmyr, N., *Pharmacotherapy (Carlisle, Mass.)*, 1985, 5, 109 (rev)

Martindale, *The Extra Pharmacopoeia*, 30th edn., Pharmaceutical Press, London, 1993, 1245.

Lewis, R.J., *Sax's Dangerous Properties of Industrial Materials*, 8th edn., Van Nostrand Reinhold, 1992, FAQ100.

Fenothiocarb, BSI, ISO, JMAF F-0-00037

S-(4-Phenoxybutyl) dimethylcarbamothioate, 9CI. *Phenothiocarb. Panocon. KCO-3001*
 [62850-32-2]



$C_{13}H_{19}NO_2S$ M 253.3

Acaricide. Cryst. Mp 40-41°. Bp_{0.02} 155°.

► LD₅₀ (rat, orl) 1150 mg/kg. FD3825000.

Ger. Pat., 2 636 620, (1977); CA, 87, 1196b

(synth, activity)

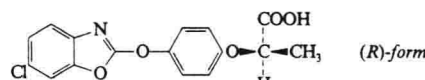
Ogawa, H., *Jpn. Pestic. Inf.*, 1985, 46, 11 (props)

Unai, T. et al, *Appl. Entomol. Zool.*, 1986, 21, 283 (metab)

Pesticide Manual, 10th edn., 1994, No. 298.

Fenoxaprop, BSI, ISO, ANSI F-0-00038

2-[4-[(6-Chloro-2-benzoxazolyl)oxy]phenoxy]propanoic acid, 9CI
 [73519-55-8]



$C_{16}H_{12}ClNO_5$ M 333.7

(R)-form [113158-40-0]

Fenoxaprop-P

► UA2452300.

Et ester: [71283-80-2]. *Fenoxaprop-P-ethyl. Hoe 046360*

$C_{18}H_{16}ClNO_5$ M 361.7 Post-emergent herbicide. More active isomer. Phys. and chem. props. reported to be the same as racemate.

► LD₅₀ (rat, orl) 3040 mg/kg.

(±)-form [95617-09-7]

Herbicide. Fatty acid synth. inhibitor.

► LD₅₀ (rat, orl) 2357 mg/kg. UA2452500.

Et ester: [82110-72-3].

Herbicide. Solid. Mp 84-85°.

[66441-23-4, 113776-20-8, 113776-21-9, 116573-18-3]

Eur. Pat., 62 905, (1982); CA, 98, 107282u (synth, activity)

Wink, O. et al, *Pestic. Sci.*, 1988, 22, 31 (activity)

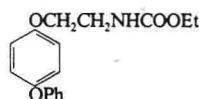
Kobek, K. et al, *Z. Naturforsch., C*, 1988, 43, 47 (activity)

Lefsrud, C. et al, *Pestic. Biochem. Physiol.*, 1989, 34, 218 (activity, metab)

Pesticide Manual, 9th edn., 1991, No. 6160.

Fenoxycarb, BSI, ISO, ANSI F-0-00039

Ethyl [2-(4-phenoxyphenoxy)ethyl]carbamate, 9CI. *Logic. Torus. Ro 13-5223*
 [72490-01-8]



$C_{17}H_{19}NO_4$ M 301.3

Insecticide. Shows strong juvenile hormone activity. Disrupts development of pests. Cryst. Mp 49-54°.

► LD₅₀ (rat, orl) >10000 mg/kg. FD0423000.

[79127-80-3]

Eur. Pat., 4 334, (1979); CA, 92, 58473x (synth, activity)

Dorn, S. et al, *Z. Pflanzenkrankh.*

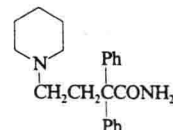
Pflanzenschutz., 1981, 88, 269 (activity)

Haenni, R.P. et al, *Anal. Methods Pestic. Plant Growth Regul.*, 1988, 16, 21 (props, tox activity)

Pesticide Manual, 9th edn., 1991, No. 6170.

Fenpipramide, BAN, INN F-0-00040

α,α-Diphenyl-1-piperidinebutanamide, 9CI. 2,2-Diphenyl-4-piperidinobutyramide. *Hoe 9980. R 14. U 0229*
 [77-01-0]



$C_{21}H_{26}N_2O$ M 322.4

Antispasmodic. Leaflets (EtOH). Mp 188°. Constit. of Efosin.

► LD₅₀ (mus, ivn) 30 mg/kg. TM5073300.

Hydrochloride: [14007-53-5].

Deliquescent solid. Mp 220°.

Methobromide: [125-60-0]. 1-(4-Amino-4-oxo-3,3-diphenylbutyl)-1-methylpiperidinium bromide, 9CI. *Fenpiverinium bromide, INN. Fenpipramide methylbromide. Resantin. Hoe 12494*

$C_{22}H_{28}BrN_2O$ M 417.3 Dimorphic cryst. (2-propanol/EtOAc). Mp 177.5-178.5°, Mp 216-216.5°.

► LD₅₀ (mus, orl) 800 mg/kg. TN4694150.

Walton, E. et al, *J.C.S.*, 1949, 648 (synth, pharmacol)

Janssen, P.A.J. et al, *J.A.C.S.*, 1955, 77, 4423 (synth, uv)

Moffett, R.B. et al, *J.A.C.S.*, 1957, 79, 4451 (synth)

Janssen, P.A.J. et al, *J. Med. Chem.*, 1959, 1, 187 (pharmacol, props)

Martindale, *The Extra Pharmacopoeia*, 30th edn., Pharmaceutical Press, London, 1993, 1370.

Lewis, R.J., *Sax's Dangerous Properties of Industrial Materials*, 8th edn., Van Nostrand Reinhold, 1992, RDA375.

Fenpiprane, BAN, INN F-0-00041

1-(3,3-Diphenylpropyl)piperidine, 8CI. 1,1-Diphenyl-3-piperidinopropane
 [3540-95-2]



$C_{20}H_{25}N$ M 279.4

Spasmolytic and antiallergic drug. Cryst. Mp 41-42.5°. Bp₈ 210-220°, Bp₁ 183-184°.

Hydrochloride: [3329-14-4]. *Hoe 10116*

Cryst. Mp 216-217°. Constit. of Aspason and Efosin.

► LD₅₀ (mus, ivn) 25 mg/kg. TM7861000.

Bockmühl, M. et al, *Annalen*, 1948, 561, 52 (synth)

Ruddy, A.W. et al, *J.A.C.S.*, 1950, 72, 718 (synth)

White, A.C. et al, *Br. J. Pharmacol.*, 1951, 6, 560 (pharmacol)

Klosa, J., *J. Prakt. Chem.*, 1966, 34, 312 (synth)

Maisey, R.F. et al, *J. Med. Chem.*, 1971, 14, 161 (pharmacol)

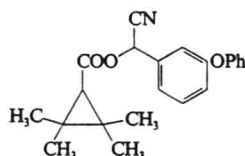
Martindale, *The Extra Pharmacopoeia*, 28th/29th edn., Pharmaceutical Press, London, 1982/1989, 12743.

Bohnen, K. et al, *Br. Crop Prot. Conf. - Pests Dis., Proc.*, 1986, 27 (props)
Pesticide Manual, 9th edn., 1991, No. 6210.

Konno, T. et al, *Br. Crop Prot. Conf. - Pests Dis., Proc.*, 1990, 71.
Pesticide Manual, 10th edn., 1994, No. 306.

Fenpropathrin, BSI, ISO F-0-00042

Cyano(3-phenoxyphenyl)methyl 2,2,3,3-tetramethylcyclopropanecarboxylate, 9CI. α -Cyano-3-phenoxybenzyl 2,2,3,3-tetramethylcyclopropanecarboxylate. Meothrin. Danitol. Rody. Herald [39515-41-8]



$C_{22}H_{23}NO_3$ M 349.4
Pyrethroid insecticide.

- Occup. exposure causes transient facial tingling. LD₅₀ (rat, ori) 18 mg/kg. LD₅₀ (rat, skn) 870 mg/kg. GZ2090000.

(±)-form

Mp 49-50°. Mixt. of diastereoisomers. (1R)-Form also known.

[64257-84-7, 67890-38-4, 67890-41-9]

Janes, N.F. et al, *J.C.S. Perkin 1*, 1977, 1878

(cmr)

Elliott, M. et al, *Pestic. Sci.*, 1978, 9, 105

(activity)

Japan. Pat., 79 130 536, (1979); CA, 92, 71076q

(synth)

Grueter, H. et al, *Pestic. Sci.*, 1980, 11, 148

(synth)

Matsuo, T. et al, *Pestic. Sci.*, 1980, 11, 202 (abs

config, activity)

Fujita, Y., *Jpn. Pestic. Inf.*, 1981, 38, 21

(activity, use, tox)

Roberts, T.R., *Prog. Pestic. Biochem.*, 1981, 1,

115 (rev, metab)

Horiba, M. et al, *Agric. Biol. Chem.*, 1982, 46,

3041 (abs config)

Mikami, N. et al, *J. Agric. Food Chem.*, 1985,

33, 980 (metab)

Pesticide Manual, 9th edn., 1991, No. 6200.

Lewis, R.J., *Sax's Dangerous Properties of*

Industrial Materials, 8th Ed., Van Nostrand-

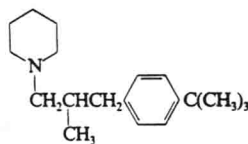
Reinhold, 1992, DAB825.

Handbook of Pesticide Toxicology, (Eds. Hayes,

W.J. et al), Academic Press, 1991, 596.

Fenpropidin, BSI, ISO F-0-00043

1-[3-[4-(1,1-Dimethylethyl)phenyl]-2-methylpropyl]piperidine, 9CI. 1-[3-(4-tert-Butylphenyl)-2-methylpropyl]piperidine. Patrol. Tern [67306-00-7]



$C_{19}H_{31}N$ M 273.4

- LD₅₀ (rat, ori) 1447 mg/kg. TM7292000.

(±)-form

Fungicide. Steroid redn. inhibitor. Pale yellow liq. Bp_{0.004} 100°. pK_a 10.5.

[6739-82-8]

Ger. Pat., 2 727 482, (1979); CA, 90, 152004a

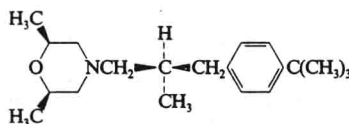
(synth, activity)

Baloch, R.I. et al, *Phytochemistry*, 1984, 23,

2219 (activity)

Fenpropimorph, BSI, ISO F-0-00044

4-[3-(4-tert-Butylphenyl)-2-methylpropyl]-2,6-dimethylmorpholine. Corbel [67306-03-0]



(R)-form

$C_{20}H_{33}NO$ M 303.4

Agricultural fungicide used against powdery mildews.

- QE2060000.

(R)-form [67564-91-4]

Shows mod. activity against wheat mildew and is prac. inactive against brown rust disease.

- QE1940000.

(S)-form

Very active against both wheat mildew and brown rust disease.

(±)-form

Oil. Bp_{0.05} 120°. pK_a 7.02 (basic).

- LD₅₀ (rat, ori) 3000 mg/kg.

[67564-92-5, 67564-93-6, 76492-89-2, 76492-90-5, 76492-92-7, 127738-75-4, 128141-83-3, 130462-20-3]

Bohnen, K. et al, *Meded. Fac. Landbouwwet., Rijksuniv. Gent*, 1979, 44, 487, 499 (rev, props, tox, use)

Himmele, W. et al, *Angew. Chem., Int. Ed.*,

1980, 19, 184.

Ger. Pat., 2 907 614, (1980); CA, 94, 65703s

(synth, resoln)

Polak, A., *Ann. N.Y. Acad. Sci.*, 1988, 544, 221

(rev)

Jensen, J.S. et al, *Acta Cryst. C*, 1990, 46, 779

(cryst struct)

Pesticide Manual, 9th edn., 1991, No. 6220.

Basarab, G.S. et al, *ACS Symp. Ser.*, 1992, 504,

414 (synth, activity)

Agrochemicals Handbook, 3rd edn., Royal

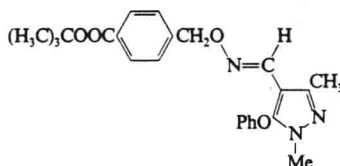
Society of Chemistry, 1992, A201.

Sandholm Jensen, J. et al, *Pestic. Sci.*, 1992, 36,

309 (conformn)

Fenpyroximate, BSI F-0-00045

1,1-Dimethylethyl 4-[[[(1,3-dimethyl-5-phenoxymethyl-4-yl)methylene]amino]oxy]methyl]benzoate, 9CI. Danitron. Ortus. NN 1850 [134098-61-6]



$C_{24}H_{27}N_3O_4$ M 421.4

Strictly, the name Fenpyroximate applies

only to the coml. available (E)-form.

Acaricide. Cryst. (MeOH). Mp 101-102°.

- LD₅₀ (rat, ori) 245 mg/kg. DG9081200.

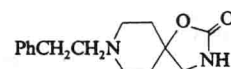
[111812-58-9]

Eur. Pat., 234 045, (1987); CA, 108, 21882c

(synth)

Fenspiride, INN F-0-00046

8-(2-Phenylethyl)-1-oxa-3,8-diazaspiro[4.5] decan-2-one, 9CI. Many other names [5053-06-5]



$C_{15}H_{20}N_2O_2$ M 260.3

Bronchodilator, antiadrenergic drug. Mp 156°.

- RO0373000.

Hydrochloride: [5053-08-7]. *Fenspiride*

hydrochloride, USAN. Decaspiride.

Espiran. *Respiride*

Mp 232-233° dec.

- LD₅₀ (rat, ori) 437 mg/kg. RO0375000.

LeDouarec, J.C. et al, *Arzneim.-Forsch.*, 1969,

19, 1263 (pharmacol)

Ranier, G. et al, *Eur. J. Med. Chem. (Chim.*

Ther.), 1969, 4, 185 (synth, pharmacol)

Beiler, J.M. et al, *Arch. Int. Pharmacodyn.*

Ther., 1971, 193, 111; 1973, 201, 90

(pharmacol, metab)

Hajdukovic, G. et al, *C. R. Hebd. Seances Acad.*

Sci. Ser. C, 1971, 272, 330 (pmr, conformn)

Duhault, J. et al, *Arzneim.-Forsch.*, 1972, 22,

1947 (pharmacol)

Sardi, B. et al, *Boll. Chim. Farm.*, 1978, 117,

343 (tox)

Evrard, Y. et al, *Sem. Hop.*, 1986, 62, 1375 (rev,

pharmacol)

Negwer, M., *Organic-Chemical Drugs and their*

Synonyms, 6th edn., Akademie-Verlag, Berlin,

1987, 3638 (synonyms)

Martindale, The Extra Pharmacopoeia, 30th

edn., Pharmaceutical Press, London, 1993,

1370.

Lewis, R.J., *Sax's Dangerous Properties of*

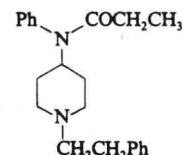
Industrial Materials, 8th edn., Van Nostrand

Reinhold, 1992, DAI200.

Fentanyl, BAN

F-0-00047

N-Phenyl-N-[1-(2-phenylethyl)-4-piperidinyl]propanamide, 9CI. N-(1-Phenethyl-4-piperidyl)propionanilide, 8CI. Phentanyl. Sublimaze. Many other names [437-38-7]



$C_{22}H_{28}N_2O$ M 336.4

Narcotic analgesic. Cryst. Mp 83-84°.

Component of Thalamonal.

- Adverse effects incl. somnolence and respiratory depression when used therapeutically. LD₅₀ (mus, ipr) 76 mg/kg. LD₅₀ (rat, ori) 18 mg/kg. UE5550000.

Citrate (1:1): [990-73-8]. *Fentanyl citrate*,

USAN. *Hypnorm*

Cryst. Slightly sol. H₂O. Mp 149-151°.

Component of Innovar.

- LD₅₀ (rat, ori) 18 mg/kg. UE5600000.

Gardocki, J.F. et al, *Toxicol. Appl. Pharmacol.*,

1964, 6, 48 (pharmacol)

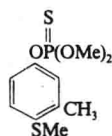
U.S. Pat., 3 141 823, (1964); CA, 61, 10689

(synth)

- Jonczyk, A. *et al*, *Przem. Chem.*, 1978, **57**, 131 (synth)
- Zee, S.-H. *et al*, *J. Chin. Chem. Soc. (Taipei)*, 1980, **27**, 147; *CA*, **94**, 174821 (synth)
- Peeters, O.M. *et al*, *J. Cryst. Mol. Struct.*, 1980, **27**, 147 (cryst struct)
- Andrews, C.J.H. *et al*, *Clin. Anaesthesiol.*, 1983, **1**, 97 (rev, pharmacol)
- Mather, L.E., *Clin. Pharmacokinet.*, 1983, **8**, 422 (rev)
- Casy, A.F. *et al*, *Eur. J. Med. Chem. (Chim. Ther.)*, 1983, **18**, 56 (cmr)
- Goromaru, T. *et al*, *Chem. Pharm. Bull.*, 1985, **33**, 3922 (metab)
- Tollenaere, J.P. *et al*, *Prog. Drug Res.*, 1986, **30**, 91 (rev, conformn)
- Negwer, M., *Organic-Chemical Drugs and their Synonyms*, 6th edn., Akademie-Verlag, Berlin, 1987, 6613 (synonyms)
- Martindale, *The Extra Pharmacopoeia*, 30th edn., Pharmaceutical Press, London, 1993, 1076.
- Lewis, R.J., *Sax's Dangerous Properties of Industrial Materials*, 8th edn., Van Nostrand Reinhold, 1992, PDW500, PDW750.

Fenthion, BSI, BAN F-0-00048

O,O-Dimethyl O-[(3-methyl-4-methylthio)phenyl] phosphorothioate, 9CI. O,O-Dimethyl-O-(4-methylthio-m-tolyl) phosphorothioate, 8CI. 4-Methylmercapto-3-methylphenyl dimethyl thiophosphate. Mercaptophos. Baytex. Lebaycid. Queletox. Spotton. Talodex. Bayer 29493. ENT 25540. S 1752 [55-38-9]



$C_{10}H_{15}O_3PS_2$ M 278.3

Insecticide with low mammalian toxicity. Used against mosquito larvae in tropical fresh waters. Cholinesterase inhibitor. Liq. with slight garlic odour. V. spar. sol. H_2O , d_4^{20} 1.25. $Bp_{0.01}$ 87°. n_D^{20} 1.5698. Thermally stable to 210°, alkali-resistant to pH 9. Metab. via sulfoxide and sulfone.

- Component of Tiguvon.
- LD_{50} (rat, ori) 180 mg/kg; LD_{50} (rat, skn) 330 mg/kg. Can cause delayed neurotoxic symptoms and retinal changes. Occup. exposure can cause numbness, tingling of extremities and muscle weakness. ACGIH TLV: long-term 0.2 mg m^{-3} (sk). TF9625000.

S-Oxide: [3761-41-9]. O,O-Dimethyl O-[(3-methyl-4-methylsulfinyl)phenyl] phosphorothioate. Fenthion sulfoxide $C_{10}H_{15}O_4PS_2$ M 294.3 Metab. of Fenthion. Stereochem. not detd.

► TF9400000.

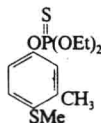
S,S-Dioxide: [3761-42-0]. O,O-Dimethyl O-[(3-methyl-4-methylsulfonyl)phenyl] phosphorothioate. Fenthion sulfone $C_{10}H_{15}O_5PS_2$ M 310.3 Metab. of Fenthion.

- TF9420000.
- Ger. Pat.*, 1 116 656, (1961); *CA*, **56**, 14170 (synth, props)
- Ibrahim, F.B. *et al*, *J. Agric. Food Chem.*, 1966, **14**, 369 (uv)
- Keith, L.H. *et al*, *J. Assoc. Off. Anal. Chem.*, 1968, **51**, 1063 (pmr)
- Getz, M.E. *et al*, *J. Assoc. Off. Anal. Chem.*, 1968, **51**, 1101 (tlc)

- Ross, R.T. *et al*, *Anal. Chim. Acta*, 1970, **52**, 139 (P nmr)
- Stan, H.-J. *et al*, *Fresenius' Z. Anal. Chem.*, 1977, **287**, 271; *Biomed. Mass Spectrom.*, 1982, **9**, 483 (glc, ms)
- Rainsford, K.D., *Pestic. Biochem. Physiol.*, 1978, **8**, 302 (tox, metab)
- Agrochemicals Handbook*, The Royal Society of Chemistry, 1983, A203.
- Ripley, B.D., *J. Assoc. Off. Anal. Chem.*, 1983, **66**, 1084 (derivs, glc)
- Stan, H.-J. *et al*, *J. Chromatogr.*, 1983, **279**, 173 (glc)
- Pesticide Manual*, 9th edn., 1991, No. 6280.
- Ballantyne, B. *et al*, *Clinical and Experimental Toxicology of Organophosphates and Carbamates*, Butterworth-Heinemann, Oxford, 1992.
- Martindale, *The Extra Pharmacopoeia*, 30th edn., Pharmaceutical Press, London, 1993, 1127.
- Lewis, R.J., *Sax's Dangerous Properties of Industrial Materials*, 8th edn., Van Nostrand Reinhold, 1992, DSS400, DSS800, FAQ900.
- Handbook of Pesticide Toxicology*, (Eds. Hayes, W.J. *et al*), Academic Press, 1991, 1023.
- Chemical Hazards of the Workplace*, (Eds. Proctor, N.H. *et al*), 3rd edn., VNR, 1991, 301.

Fenthion-ethyl F-0-00049

O,O-Diethyl O-[(3-methyl-4-(methylthio)phenyl]phosphorothioate, 9CI. O,O-Diethyl O-[(4-(methylthio)-m-tolyl]phosphorothioate, 8CI [1716-09-2]



$C_{12}H_{19}O_3PS_2$ M 306.3

Insecticide. $Bp_{0.01}$ 104°.

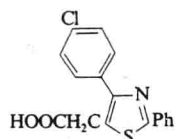
- TF4100000.
- 4-S-Oxide: [6587-44-6]. O,O-Diethyl O-[(3-methyl-4-(methylsulfinyl)phenyl] phosphorothioate, 9CI

$C_{12}H_{19}O_4PS_2$ M 322.3 d_4^{20} 1.23. n_D^{20} 1.5471.

- [24505-67-7]
- Ger. Pat.*, 1 116 656, (1961); *CA*, **56**, 14170i (synth, activity)
- Neely, W.B. *et al*, *J. Agric. Food Chem.*, 1970, **18**, 45 (activity)
- Lewis, R.J., *Sax's Dangerous Properties of Industrial Materials*, 8th Ed., Van Nostrand-Reinhold, 1992, LIN400.

Fentiazac, BAN, INN, JAN F-0-00050

4-(4-Chlorophenyl)-2-phenyl-5-thiazoleacetic acid, 9CI. 8CI. Donorest. Flogene. Norvedan. Many other names [18046-21-4]



$C_{17}H_{12}ClNO_2S$ M 329.8

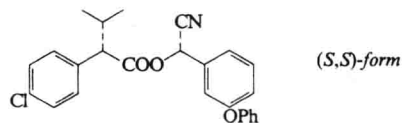
Antipyretic, antiinflammatory drug. Needles (C_6H_6). Mp 161-162°.

- LD_{50} (rat, ori) 409 mg/kg. Exp. reprod. and teratogenic effects. XJ1890000.

- Brown, K. *et al*, *J. Med. Chem.*, 1974, **17**, 1177 (synth, pharmacol)
- Destro, R., *Acta Cryst. B*, 1978, **34**, 959 (cryst struct)
- Marmo, E., *Curr. Med. Res. Opin.*, 1979, **6**, 53 (rev, pharmacol, tox)
- Anon., *Drugs of Today (Barcelona)*, 1979, **15**, 323 (rev)
- Fumero, S. *et al*, *Arzneim.-Forsch.*, 1980, **30**, 1253 (metab)
- Franklin, R.A. *et al*, *Xenobiotica*, 1984, **14**, 955 (metab)
- Negwer, M., *Organic-Chemical Drugs and their Synonyms*, 6th edn., Akademie-Verlag, Berlin, 1987, 4231 (synonyms)
- Martindale, *The Extra Pharmacopoeia*, 30th edn., Pharmaceutical Press, London, 1993, 15.
- Lewis, R.J., *Sax's Dangerous Properties of Industrial Materials*, 8th edn., Van Nostrand Reinhold, 1992, CK1750.

Fenvalerate, BAN, BSI, ISO F-0-00051

Cyano(3-phenoxyphenyl)methyl 4-chloro- α -(1-methylethyl)benzeneacetate, 9CI. α -Cyano-3-phenoxybenzyl 2-(4-chlorophenyl)-3-methylbutyrate. Belmark. Pydrin. Sumicidin [51630-58-1]



$C_{25}H_{22}ClNO_3$ M 419.9

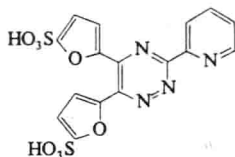
Agricultural, public health and animal husbandry insecticide. Viscous yellow liq. d_4^{25} 1.175. Bp_{37} 300°.

- LD_{50} (rat, ori) 451 mg/kg. CY1576350.
- (S,S)-form [66230-04-4] **Esfenvalerate, BSI, ISO.** Sumi-alpha. Asana. Fenvalerate α Insecticide. Has greater insecticidal activity than racemate. Cryst. d_4^{26} 1.26. Mp 59-60°. $[\alpha]_D^{25}$ -15.0 (c, 2.0 in MeOH).
- CY1576367.
- [66267-77-4, 67614-32-8, 67614-33-9, 67890-39-5, 67890-40-8, 73367-17-6, 73367-18-7]
- U.K. Pat.*, 1 439 615, (1972); *CA*, **80**, 120746w.
- Hattori, J., *Jpn. Pestic. Inf.*, 1977, **33**, 13 (rev, props)
- Aketa, K. *et al*, *Agric. Biol. Chem.*, 1978, **42**, 895 (synth, pmr, activity)
- Ger. Pat.*, 2 737 297, (1978); *CA*, **88**, 152268y (esfenvalerate)
- Yoshioka, H., *Rev. Plant. Prot. Res.*, 1978, **11**, 39 (rev, synth, tox)
- Horiba, M. *et al*, *Agric. Biol. Chem.*, 1982, **46**, 3041 (abs config, bibl)
- Papadopolou-Mourkidou, E., *Residue Rev.*, 1983, **89**, 179 (rev, detn)
- Wheeler, T.N. *et al*, *J. Agric. Food Chem.*, 1984, **32**, 1125 (synth, activity)
- U.S. Pat.*, 4 432 908, (1984); *CA*, **100**, 209428y (cryst struct)
- Oo'uchi, H., *Jpn. Pestic. Inf.*, 1985, **46**, 21 (esfenvalerate)
- Mikami, N. *et al*, *Pestic. Sci.*, 1985, **16**, 46 (metab)
- Miyamoto, J. *et al*, *ACS Symp. Ser.*, 1986, **299**, 268 (rev, tox)
- Lidgard, R.O. *et al*, *Biomed. Environ. Mass Spectrom.*, 1986, **13**, 677 (ms)
- Papadopolou-Mourkidou, E., *Anal. Methods Pestic. Plant Growth Regul.*, 1988, **16**, 31 (hplc)
- IARC Monog.*, 1991, **53**, 309 (rev, tox)
- Pesticide Manual*, 9th edn., 1991, No. 5630, 6340.

Agrochemicals Handbook, 3rd edn., Royal Society of Chemistry, 1992, A990, A207.
Lewis, R.J., *Sax's Dangerous Properties of Industrial Materials*, 8th edn., Van Nostrand Reinhold, 1992, FAR100.

Ferene **F-0-00052**

5,5'-[3-(2-Pyridinyl)-1,2,4-triazine-5,6-diyl]bis-2-furansulfonic acid, 9CI
[90691-98-8]



$C_{16}H_{10}N_4O_8S_2$ M 450.4

Di-Na salt: [79551-14-7]. *Ferene S*
Used as 0.18mM aq. soln. for photometric detn. of Fe(II) (λ_{max} 593 nm, ϵ 34500).
Yellow cryst. + $1H_2O$ (EtOH).

[90358-66-0]

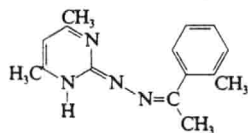
Aldrich Library of ^{13}C and 1H FT NMR Spectra, 3, 414C (nmr)

Higgins, T., *CA*, 1981, **95**, 183278.

Hennessy, D.J. *et al.* *Can. J. Chem.*, 1984, **62**, 721 (synth, detn, Fe)

Ferimzone, BSI **F-0-00053**

4,6-Dimethyl-2(1H)-pyrimidinone [1-(2-methylphenyl)ethylidene]hydrazone, 9CI



(Z)-form

$C_{15}H_{18}N_4$ M 254.3

(E)-form [77359-18-3]

Needles (hexane). Mp 110-111°.

(Z)-form [89269-64-7]

TF 164

Fungicide. Prisms (hexane). Mp 178-180°.

► LD₅₀ (rat, orl) 725 mg/kg. UW7430000.

Konishi, K. *et al.* *Agric. Biol. Chem.*, 1986, **50**, 2427 (synth, ir, uv, pmr)

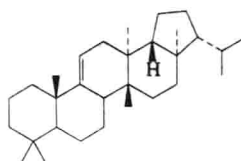
Okuno, T. *et al.* *Phytochemistry*, 1989, **79**, 827 (activity)

Pesticide Manual, 10th edn., 1994, No. 312.

9(11)-Fernene **F-0-00054**

Davallene. Fernene

[1615-99-2]



$C_{30}H_{50}$ M 410.7

Constit. of *Dryopteris crenata*, *D.*

crassirhizoma and of *Adiantum pedatum*.

Fairly widely distributed in ferns. Cryst. Mp 175-177°. [α]_D -26.4 (CHCl₃).

21 α H-form [29428-21-5]

21-Epiferen-9(11)-ene

Present in the epicuticular layer of ferns. Mp 151-153.5°. [α]_D -15.3.

Ageta, H. *et al.* *Tet. Lett.*, 1963, 1447; 1966,

6069 (isol, struct, pmr)

Ghisalberti, E.L. *et al.* *Phytochemistry*, 1970, **9**, 1817 (biosynth)

Barton, D.H.R. *et al.* *J.C.S.(C)*, 1971, 110 (biosynth)

Wollenweber, E. *et al.* *Z. Naturforsch., C*, 1981, **36**, 896 (isol)

Sengupta, P. *et al.* *Indian J. Chem., Sect. B*, 1983, **22**, 882 (cryst struct)

Murakami, T. *et al.* *Prog. Chem. Org. Nat. Prod.*, 1988, **54**, 1 (occur)

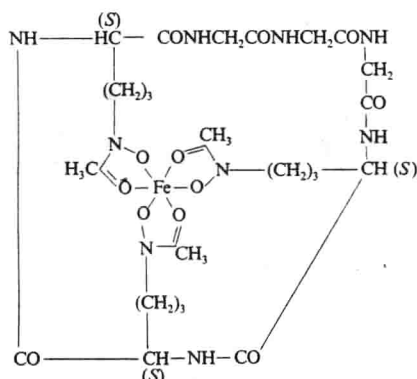
Masuda, K. *et al.* *Chem. Pharm. Bull.*, 1989, **37**, 1673 (pmr, cmr)

Wilkins, A.L. *et al.* *Chem. Pharm. Bull.*, 1989, **37**, 1673 (cmr)

Ageta, H. *et al.* *Chem. Pharm. Bull.*, 1994, **42**, 39 (pmr, cmr)

Ferrichrome, 9CI

[15630-64-5]



$C_{27}H_{42}FeN_9O_{12}$ M 740.5

Metab. of *Ustilago sphaerogena* grown under special conditions. Growth promotor. Yellow needles (MeOH). Sol. H₂O, MeOH, spar. sol. Me₂CO, CHCl₃. Mp 248-251° dec. [α]_D²⁰ +304 (c, 0.425 in H₂O).

[34787-28-5]

Neilands, J.B., *J.A.C.S.*, 1952, **74**, 4846 (isol)

Emery, T. *et al.* *J.A.C.S.*, 1960, **82**, 3658; 1961, **83**, 1626 (struct)

Llinas, M. *et al.* *J. Mol. Biol.*, 1972, **68**, 265 (pmr, conformn)

Isowa, Y. *et al.* *Bull. Chem. Soc. Jpn.*, 1974, **47**, 215 (ir, synth)

Llinas, M. *et al.* *J. Mol. Biol.*, 1976, **104**, 853 (cmr, struct)

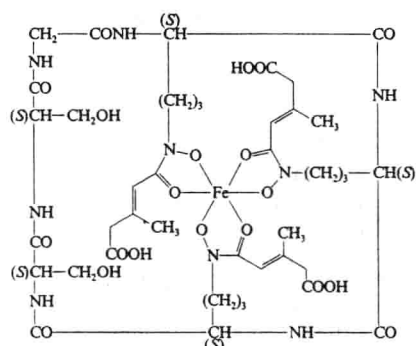
Naegli, H.-U. *et al.* *Helv. Chim. Acta*, 1978, **61**, 2088 (synth, ir, uv)

Van der Helm, D. *et al.* *J.A.C.S.*, 1980, **102**, 4224 (cryst struct)

Dell, A. *et al.* *Biomed. Mass Spectrom.*, 1982, **9**, 158 (ms)

Ferrichrome A

[15258-80-7]



$C_{41}H_{58}FeN_9O_{20}$ M 1052.8

Metab. product of *Ustilago sphaerogena*.

Iron transport compound, growth factor.

Hygroscopic, amber cryst. + 4H₂O (H₂O).

Sol. EtOH; insol. Me₂CO, CHCl₃.

[25312-33-8]

Garibaldi, J.A. *et al.* *J.A.C.S.*, 1955, **77**, 2429 (isol, ir)

Emery, T. *et al.* *J.A.C.S.*, 1960, **82**, 3658; 1961, **83**, 1626 (struct)

Keller-Schierlein, W., *Helv. Chim. Acta*, 1963, 1920 (ir, struct)

Zalkin, A. *et al.* *J.A.C.S.*, 1966, **88**, 1810 (cryst struct, abs config)

Llinas, M. *et al.* *J. Mol. Biol.*, 1972, **68**, 265 (pmr, conformn)

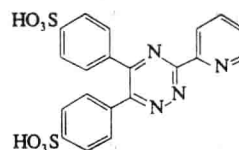
Van der Helm, D. *et al.* *Acta Cryst. B*, 1981, **37**, 323 (cryst struct)

Dell, A. *et al.* *Biomed. Mass Spectrom.*, 1982, **9**, 158 (ms)

Ferrozine

F-0-00057

4,4'-[3-(2-Pyridinyl)-1,2,4-triazine-5,6-diyl]bisbenzenesulfonic acid, 9CI. 3-(2-Pyridinyl)-5,6-bis(4-sulfophenyl)-1,2,4-triazine
[32796-55-7]



$C_{20}H_{14}N_4O_6S_2$ M 470.4

Used for photometric detn. of Fe(II) (λ_{max} 560 nm, ϵ 28000), Co, Cr and indirectly As, Mo, SO₂.

Di-Na salt: [28048-33-1].

Cryst. (H₂O). Mp > 350°.

► DB7345000.

Stookey, L.L., *Anal. Chem.*, 1970, **42**, 779 (synth, detn, Fe)

Attari, A. *et al.* *Anal. Chem.*, 1972, **44**, 151 (detn, SO₂)

Kundra, S.K. *et al.* *Anal. Chem.*, 1974, **46**, 1605 (detn, Co)

Kellen, G.J. *et al.* *Anal. Chem.*, 1976, **48**, 1538 (detn, As)

Simonzadeh, N. *et al.* *Talanta*, 1979, **26**, 935 (detn, Mo)

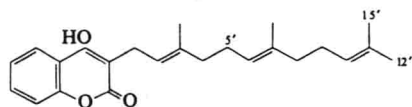
Thompson, J.C. *et al.* *Anal. Chem.*, 1984, **56**, 755 (detn, Fe)

Onishi, H., *Photometric Determination of Traces of Metals, Part IIa: Individual Metals.*

Aluminium to Lithium, John Wiley, New York, 4th Ed., 1986, 730.

Ferulenol **F-0-00058**

4-Hydroxy-3-[(3,7,11-trimethyl-2,6,10-dodecatrienyl)-2H-1-benzopyran-2-one, 9Cl. 3-(1-Farnesyl)-4-hydroxycoumarin [6805-34-1]



$C_{24}H_{30}O_3$ M 366.4

Constit. of the latex of *Ferula communis*.

Shows haemorrhagic action. Mp 64-65°.

12'-Hydroxy: [106909-08-4]. (E)- ω -

Hydroxyferulenol

$C_{24}H_{30}O_4$ M 382.4 Constit. of *F. communis*. Shows haemorrhagic action. Yellow oil.

15'-Hydroxy: [110107-13-6]. (Z)- ω -

Hydroxyferulenol

$C_{24}H_{30}O_4$ M 382.4 Constit. of *F. communis* and *F. communis* var. *genuina*. Yellow oil.

12'-Acetoxy: [119234-02-5]. (E)- ω -

Acetoxyferulenol

$C_{26}H_{32}O_5$ M 424.5 Constit. of *F. communis*. Oil.

15'-Acetoxy: [119259-89-1]. (Z)- ω -

Acetoxyferulenol

$C_{26}H_{32}O_5$ M 424.5 Constit. of *F. communis*. Oil.

12'-Oxo: [106894-38-6]. (E)- ω -**Oxoferulenol**

$C_{24}H_{28}O_4$ M 380.4 Constit. of *F. communis*. Oil.

5'-Hydroxy: ϵ -**Hydroxyferulenol**

$C_{24}H_{30}O_4$ M 382.4 Isol. from *F. communis* var. *genuina*.

Carboni, S. et al, *Tet. Lett.*, 1964, 2783 (struct, pmr)

Valle, M.G. et al, *Phytochemistry*, 1987, 26, 253 (isol)

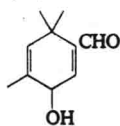
Lamnaouer, D. et al, *Phytochemistry*, 1987, 26, 1613; 1991, 30, 2383 (isol, pmr, cmr, deriv)

Appendino, G. et al, *Phytochemistry*, 1988, 27, 3619 (deriv)

Ferulol **F-0-00059**

For a fuller version of this Entry see *Dictionary of Natural Products*, F-00145

3-Hydroxy-4,6,6-trimethyl-1,4-cyclohexadiene-1-carboxaldehyde [55568-77-9]



$C_{10}H_{14}O_2$ M 166.2

Derivs. are known with both positive and negative specific rotations and may belong to opposite enantiomeric series. Constit. of the roots of *Selinum carvifolium*. Derivs. of Ferulol reported from *Astrantia*, *Cenolophium*, *Eryngium*, *Ferula*, *Hacquetia*, *Hladnikia*, *Ligusticum*, *Peucedanum*, *Selinum* and *Silaum* spp. Oil. $[\alpha]_D^{20} +15$ (c, 1 in $CHCl_3$).

Angeloyl: [28333-43-9].

$C_{15}H_{20}O_3$ M 248.3 Constit. of *Ferula hispanica*. Oil. Bp_{0.1} 130°. $[\alpha]_D^{20} -52$ (c, 4.5 in Et_2O).

O-(3-Methyl-2-butenoyl): [33530-15-3].

$C_{15}H_{20}O_3$ M 248.3 Constit. of *L. seguieri*. Oil. $[\alpha]_D^{20} +15$ (c, 1 in $CHCl_3$).

[41628-55-1]

Bohlmann, F. et al, *Chem. Ber.*, 1969, 102, 2211; 1971, 104, 1957; 1974, 107, 1769.

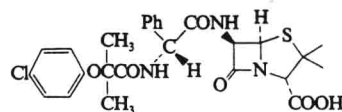
Lemmich, J. et al, *Acta Chem. Scand.*, 1971, 25, 995.

Fibracillin, INN**F-0-00060**

α -(p-Chlorophenoxy)isobutyrylamicillin.

Alongapen. Fibrapen. Mitibron. AL 70-35

[51154-48-4]



$C_{26}H_{28}ClN_3O_6S$ M 546.0

Semisynthetic. Antibacterial agent.

[51154-50-8]

Ger. Pat., 2 207 723, (1972); CA, 77, 152167 (synth)

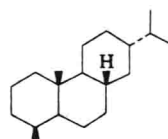
Ger. Pat., 2 327 558, (1974); CA, 80, 120916 (synth)

Martindale, *The Extra Pharmacopoeia*, 30th edn., Pharmaceutical Press, London, 1993, 166.

Fichtelite**F-0-00061**

18-Norabietane

[2221-95-6]



$C_{19}H_{34}$ M 262.4

Constit. of *Pinus pumila*; found in peat and other fossil substances of plant origin.

Cryst. (MeOH). Mp 46.5°. Bp₇₁₉ 355°, Bp₄₃ 235-236°.

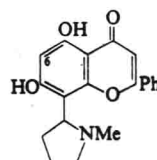
Ruzicka, L. et al, *Helv. Chim. Acta*, 1923, 6, 692 (bibl)

Burgstahler, A.W. et al, *J.O.C.*, 1969, 34, 1562 (bibl)

Taber, D.F. et al, *J.A.C.S.*, 1980, 102, 5085 (synth)

Ficine**F-0-00062**

[2520-36-7]



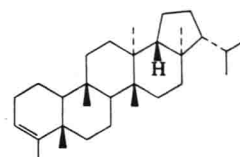
$C_{20}H_{19}NO_4$ M 337.3

Alkaloid from *Ficus pantoniana* (Moraceae). Mp 235°.

Johns, S.R. et al, *Tet. Lett.*, 1965, 1987 (w, ir, pmr, ms, struct)

3-Filicene**F-0-00063****Filicene**

[2472-29-9]



$C_{30}H_{50}$ M 410.7

Constit. of *Adiantum monochlamys* and *A. pedatum* and other ferns. Cryst. Mp 242-243° (228.5-229.5°). $[\alpha]_D +50$ ($CHCl_3$).

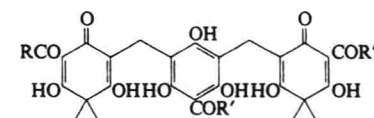
3 α ,4 α -Epoxide: [23983-65-5]. 3 α ,4 α -Epoxyfilicane. **Adiantoxide**. Isol. from *Adiantum capillus-veneris*. Cryst. Mp 229-231°. $[\alpha]_D^{20} +46.8$.

Ageta, H. et al, *Tet. Lett.*, 1964, 3413; 1966, 6069 (Filicene)

Berti, G. et al, *Tetrahedron*, 1969, 25, 2939

(Adiantoxide) Shiojima, K. et al, *Chem. Pharm. Bull.*, 1993, 41, 262 (Filicene, pmr, cmr)

Ageta, H. et al, *Chem. Pharm. Bull.*, 1994, 42, 39 (pmr, cmr)

Filixic acids**F-0-00064**

Filixic acid

ABA, R = R'' = CH_3 , R' = $CH_2CH_2CH_3$

ABP, R = CH_3 , R' = $CH_2CH_2CH_3$,

R'' = CH_2CH_3

ABB, R = CH_3 , R' = R'' = $CH_2CH_2CH_3$

PBP, R = R'' = CH_2CH_3 , R' = $CH_2CH_2CH_3$

PBB, R = CH_2CH_3 , R' = R'' = $CH_2CH_2CH_3$

BBB, R = R' = R'' = $CH_2CH_2CH_3$

The alphabetical suffixes refer to the acyl substituents (A = acetyl, P = propionyl, B = butyryl). Isol. from ferns *Dryopteris* and *Aspidium* spp.

Filixic acid ABA [38226-84-5]

$C_{32}H_{56}O_{12}$ M 612.6

From *D. dickinsii*. Yellow needles (Me₂CO). Mp 163-166°.

Filixic acid ABP [57765-54-5]

$C_{33}H_{58}O_{12}$ M 626.6

Filixic acid ABB [37318-24-4]

$C_{34}H_{60}O_{12}$ M 640.6

Filixic acid PBP [51005-85-7]

$C_{34}H_{60}O_{12}$ M 640.6

Light yellow needles (Me₂CO). Mp 192-194°. MF incorr. given as C_{30} by Hisada et al.

Filixic acid PBB [49582-09-4]

$C_{35}H_{62}O_{12}$ M 654.7

Mp 184-186°.

Filixic acid BBB [4482-83-1]

Filixic acid. Filicin. Trisalsapidin BBB

$C_{36}H_{64}O_{12}$ M 668.7

Cryst. (Me₂CO), light yellow tablets (EtOAc). Mp 184-185° (168-170°).

► LK4250000.

Penttilä, A. et al, *Acta Chem. Scand.*, 1963, 17, 191 (isol, struct)

Hisada, S. et al, *Phytochemistry*, 1972, 11, 1850, 2881; 1973, 12, 1493 (isol)