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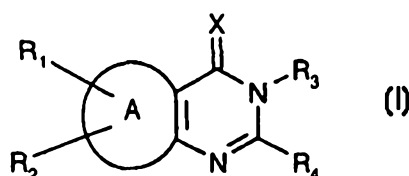
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(54) Title: NOVEL PYRIMIDIN-4-ONE AND PYRIMIDIN-4-THIONE AS FUNGICIDE

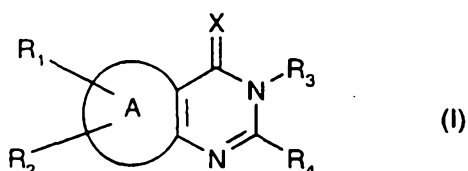


(57) Abstract

Novel pyrimidin-4-one and pyrimidin-4-thione derivatives of formula (I) wherein A is phenyl, thienyl (including all 3 isomers), thiazolyl, pyridyl or pyridazinyl; X is oxygen or sulfur; R₁ is hydrogen, halogen or trimethylsilyl; R₂ is hydrogen, halogen or trimethylsilyl; at least one of R₁ and R₂ is not hydrogen; R₃ is C₁-C₈alkyl, C₁-C₈alkenyl, C₁-C₈alkinyl which are unsubstituted or mono to tri-substituted by C₃-C₆cycloalkyl, halogen, C₁-C₆alkoxy or C₁-C₆haloalkoxy; O-C₁-C₆alkyl, O-C₂-C₆alkenyl, O-C₂-C₆alkynyl, which are unsubstituted or mono to tri-substituted by C₃-C₆cycloalkyl, halogen or C₁-C₆alkoxy; N-C₁-C₆alkyl; or N=CHC₁-C₆alkyl; R₄ is C₁-C₈alkyl, C₁-C₈alkenyl, C₁-C₈alkinyl which are unsubstituted or mono to tri-substituted by C₃-C₆cycloalkyl, halogen, cyano, C₁-C₆alkoxy or C₁-C₆haloalkoxy; C₁-C₄alkoxy-C₁-C₄alkoxy; C₁-C₄alkoxy-C₁-C₄alkylthio; nitro; -CO-C₁-C₆alkyl; C₃-C₆cycloalkyl; or phenyl, which is unsubstituted or mono to tri-substituted by halogen, C₁-C₆alkyl, C₁-C₆haloalkyl, C₁-C₆alkoxy, C₁-C₆haloalkoxy, cyano, nitro, amino, mono-C₁-C₆alkylamino, di-C₁-C₆alkylamino, C₁-C₆alkylthio, phenyl or phenoxy and in which the phenyl part is unsubstituted or mono to tri-substituted by halogen, C₁-C₆alkyl, C₁-C₆haloalkyl, C₁-C₆alkoxy or C₁-C₆haloalkoxy. The novel compounds have plant-protective properties and are suitable for protecting plants against infestation by phytopathogenic microorganisms, in particular fungi.

NOVEL PYRIMIDIN-4-ONE and PYRIMIDIN-4-THIONE AS FUNGICIDE

The present invention relates to novel pyrimidin-4-one and pyrimidin-4-thione derivatives of formula I, which have pesticidal activity, in particular fungicidal activity,



wherein

A is phenyl, thienyl (including all 3 isomers), thiazolyl, pyridyl or pyridazinyl;

X is oxygen or sulfur;

R₁ is hydrogen, halogen or trimethylsilyl;

R₂ is hydrogen, halogen or trimethylsilyl; at least one of R₁ and R₂ is not hydrogen;

R₃ is C₁-C₈alkyl, C₁-C₈alkenyl, C₁-C₈alkynyl which are unsubstituted or mono to tri-substituted by C₃-C₆cycloalkyl, halogen, C₁-C₆alkoxy or C₁-C₆haloalkoxy; O-C₁-C₆alkyl, O-C₂-C₆alkenyl, O-C₂-C₆alkynyl, which are unsubstituted or mono to tri-substituted by C₃-C₆cycloalkyl, halogen or C₁-C₆alkoxy; N-C₁-C₆alkyl; or N=CHC₁-C₆alkyl;

R₄ is C₁-C₈alkyl, C₁-C₈alkenyl, C₁-C₈alkynyl which are unsubstituted or mono to tri-substituted by C₃-C₆cycloalkyl, halogen, cyano, C₁-C₆alkoxy or C₁-C₆haloalkoxy; C₁-C₄alkoxy-C₁-C₄alkoxy; C₁-C₄alkoxy-C₁-C₄alkylthio; nitro; -CO-C₁-C₆alkyl; C₃-C₆cycloalkyl; or phenyl, which is unsubstituted or mono to tri-substituted by halogen, C₁-C₆alkyl, C₁-C₆haloalkyl, C₁-C₆alkoxy, C₁-C₆haloalkoxy, cyano, nitro, amino, mono-C₁-C₆alkylamino, di-C₁-C₆alkylamino, C₁-C₆alkylthio, phenyl or phenoxy and in which the phenyl part is unsubstituted or mono to tri-substituted by halogen, C₁-C₆alkyl, C₁-C₆haloalkyl, C₁-C₆alkoxy or C₁-C₆haloalkoxy.

The invention also relates to the preparation of these compounds, to agrochemical compositions comprising as active ingredient at least one of these compounds, as well as to the use of the active ingredients or compositions for pest control, in particular as fungicides, in agriculture and horticulture.

The compounds I and, optionally, their tautomers may be obtained in the form of their salts. Because the compounds I have at least one basic center they can, for example, form acid addition salts. Said acid addition salts are, for example, formed with mineral acids, typically

sulfuric acid, a phosphoric acid or a hydrogen halide, with organic carboxylic acids, typically acetic acid, oxalic acid, malonic acid, maleic acid, fumaric acid or phthalic acid, with hydroxycarboxylic acids, typically ascorbic acid, lactic acid, malic acid, tartaric acid or citric acid, or with benzoic acid, or with organic sulfonic acids, typically methanesulfonic acid or p-toluenesulfonic acid.

Together with at least one acidic group, the compounds of formula I can also form salts with bases. Suitable salts with bases are, for example, metal salts, typically alkali metal salts or alkaline earth metal salts, e.g. sodium salts, potassium salts or magnesium salts, or salts with ammonia or an organic amine, e.g. morpholine, piperidine, pyrrolidine, a mono-, di- or trialkylamine, typically ethylamine, diethylamine, triethylamine or dimethylpropylamine, or a mono-, di- or trihydroxyalkylamine, typically mono-, di- or triethanolamine. Where appropriate, the formation of corresponding internal salts is also possible. Within the scope of this invention, agrochemical acceptable salts are preferred.

Where asymmetrical carbon atoms are present in the compounds of formula I, these compounds are in optically active form. Owing to the presence of double bonds, the compounds can be obtained in the [E] and/or [Z] form. Atropisomerism can also occur. The invention relates to the pure isomers, such as enantiomers and diastereomers, as well as to all possible mixtures of isomers, e.g. mixtures of diastereomers, racemates or mixtures of racemates.

The general terms used hereinabove and hereinbelow have the following meanings, unless otherwise defined:

Alkyl groups on their own or as structural element of other groups such as alkoxy are, in accordance with the number of carbon atoms, straight-chain or branched and will typically be methyl, ethyl, n-propyl, isopropyl, n-butyl, sec-butyl, isobutyl, tert-butyl, n-amyl, tert-amyl, 1-hexyl, 3-hexyl, 1-heptyl or 1-octyl.

Alkenyl will be understood as meaning straight-chain or branched alkenyl such as allyl, methallyl, 1-methylvinyl, but-2-en-1-yl, 1-pentenyl, 1-hexenyl, 1-heptenyl or 1-octenyl. Preferred alkenyl radicals contain 3 to 4 carbon atoms in the chain.

Alkynyl can likewise, in accordance with the number of carbon atoms, be straight-chain or branched and is typically propargyl, but-1-yn-1-yl, but-1-yn-3-yl, 1-pentynyl, 1-hexynyl, 1-heptynyl or 1-octynyl. The preferred meaning is propargyl.

Halogen and halo substituents will be understood generally as meaning fluorine, chlorine, bromine or iodine. Fluorine, chlorine or bromine are preferred meanings.

Haloalkyl can contain identical or different halogen atoms, typically fluoromethyl, difluoromethyl, difluorochloromethyl, trifluoromethyl, chloromethyl, dichloromethyl, trichloromethyl, 2,2,2-trifluoroethyl, 2-fluoroethyl, 2-chloroethyl, 2,2,2-trichloroethyl, 3,3,3-trifluoropropyl.

Cycloalkyl is, depending on the ring size, cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl or cyclooctyl.

Preferred compounds are those of formula I, wherein A is thienyl, including all 3 isomers (subgroup A).

Within the scope of said subgroup A, those compounds of formula I are preferred wherein R_1 is hydrogen, fluorine, chlorine, bromine or iodine;

R_2 is hydrogen, fluorine, chlorine, bromine or iodine; at least one of R_1 and R_2 is not hydrogen;

R_3 is C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl which are unsubstituted or mono to tri-substituted by C_3 - C_6 cycloalkyl, halogen or C_1 - C_4 alkoxy; O- C_1 - C_6 alkyl, O- C_2 - C_6 alkenyl, O- C_2 - C_6 alkynyl, which are unsubstituted or mono to tri-substituted by C_3 - C_6 cycloalkyl, halogen or C_1 - C_4 alkoxy; N- C_1 - C_6 alkyl; or N=CHC $_1$ - C_6 alkyl;

R_4 is C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl which are unsubstituted or mono to tri-substituted by C_3 - C_6 cycloalkyl, halogen or C_1 - C_4 alkoxy; or phenyl which is unsubstituted or mono to tri-substituted by fluorine, chlorine, bromine, C_1 - C_4 alkyl, C_1 - C_4 haloalkyl, C_1 - C_4 alkoxy, C_1 - C_4 haloalkoxy, phenyl or phenoxy and in which the phenyl part is unsubstituted or mono to tri-substituted by fluorine, chlorine, bromine, C_1 - C_4 alkyl, C_1 - C_4 haloalkyl, C_1 - C_4 alkoxy or C_1 - C_4 haloalkoxy (subgroup B).

A special group within the scope of subgroup B is that of the compounds of formula I, wherein

R₃ is C₁-C₆alkyl, C₂-C₆alkenyl, C₂-C₆alkynyl which are unsubstituted or mono to tri-substituted by C₃-C₄cycloalkyl, fluorine, chlorine, bromine or C₁-C₄alkoxy; O-C₁-C₆alkyl; O-C₂-C₆alkenyl; O-C₂-C₆alkynyl; N-C₁-C₆alkyl; or N=CHC₁-C₆alkyl;

R₄ is C₁-C₆alkyl, C₂-C₆alkenyl, C₂-C₆alkynyl which are unsubstituted or mono to tri-substituted by C₃-C₄cycloalkyl, fluorine, chlorine, bromine or C₁-C₄alkoxy; or phenyl which is unsubstituted or mono to tri-substituted by fluorine, chlorine, bromine, C₁-C₄alkyl, C₁-C₄haloalkyl, C₁-C₄alkoxy, C₁-C₄haloalkoxy, phenyl or phenoxy and in which the phenyl part is unsubstituted or mono to tri-substituted by fluorine, chlorine or bromine (subgroup C).

A preferred group within the scope of subgroup C is that of the compounds of the formula I, wherein

A is thienyl[2.3-d],

X is oxygen,

R₁ is hydrogen, chlorine or bromine;

R₂ is hydrogen, chlorine or bromine; at least one of R₁ and R₂ is not hydrogen;

R₃ is C₃-C₅alkyl or O-C₁-C₄alkyl;

R₄ is C₂-C₅alkyl or phenyl which is unsubstituted or mono to tri-substituted by fluorine, chlorine, bromine, C₁-C₄alkyl or phenoxy and in which the phenoxy is unsubstituted or mono to tri-substituted by fluorine, chlorine or bromine (subgroup D1).

Another preferred group within the scope of subgroup C is that of compounds of the formula I, wherein

A is thienyl[2.3-d],

X is sulfur,

R₁ is hydrogen, chlorine or bromine;

R₂ is hydrogen, chlorine or bromine; at least one of R₁ and R₂ is not hydrogen;

R₃ is C₃-C₅alkyl or O-C₁-C₄alkyl;

R₄ is C₂-C₅alkyl or phenyl which is unsubstituted or mono to tri-substituted by fluorine, chlorine, bromine, C₁-C₄alkyl or phenoxy and in which the phenoxy is unsubstituted or mono to tri-substituted by fluorine, chlorine or bromine (subgroup D2).

Another preferred group within the scope of subgroup C is that of the compounds of the formula I, wherein

A is thienyl[3.2-d],

X is oxygen,

R₁ is hydrogen, chlorine or bromine;

R₂ is hydrogen, chlorine or bromine; at least one of R₁ and R₂ is not hydrogen;

R₃ is C₃-C₅alkyl or O-C₁-C₄alkyl;

R₄ is C₂-C₅alkyl or phenyl which is unsubstituted or mono to tri-substituted by fluorine, chlorine, bromine, C₁-C₄alkyl or phenoxy and in which the phenoxy is unsubstituted or mono to tri-substituted by fluorine, chlorine or bromine (subgroup E).

Another preferred group of compounds are those of formula I, wherein

A is pyridyl (subgroup F).

Within the scope of said group F, those compounds of formula I are preferred wherein

X is oxygen;

R₁ is hydrogen, fluorine, chlorine, bromine or iodine;

R₂ is hydrogen, fluorine, chlorine, bromine or iodine; at least one of R₁ and R₂ is not hydrogen;

R₃ is C₁-C₆alkyl, C₂-C₆alkenyl, C₂-C₆alkynyl which are unsubstituted or mono to tri-substituted by C₃-C₆cycloalkyl, halogen or C₁-C₄alkoxy; O-C₁-C₆alkyl, O-C₂-C₆alkenyl, O-C₂-C₆alkynyl, which are unsubstituted or mono to tri-substituted by C₃-C₆cycloalkyl, halogen or C₁-C₄alkoxy;

R₄ is C₁-C₆alkyl, C₂-C₆alkenyl, C₂-C₆alkynyl which are unsubstituted or mono to tri-substituted by C₃-C₆cycloalkyl, halogen or C₁-C₄alkoxy; or phenyl which is unsubstituted or mono to tri-substituted by fluorine, chlorine, bromine, C₁-C₄alkyl, C₁-C₄haloalkyl, C₁-C₄alkoxy, C₁-C₄haloalkoxy, phenyl or phenoxy and in which the phenyl part is unsubstituted or mono to tri-substituted by fluorine, chlorine, bromine, C₁-C₄alkyl, C₁-C₄haloalkyl, C₁-C₄alkoxy or C₁-C₄haloalkoxy (subgroup G).

Other preferred group of compounds are those of formula I, wherein

A is phenyl (subgroup H).

Within the scope of said group H, those compounds of formula I are preferred wherein

X is oxygen;

R₁ is hydrogen, fluorine, chlorine, bromine or iodine;

R₂ is hydrogen, fluorine, chlorine, bromine or iodine; at least one of R₁ and R₂ is not hydrogen;

R₃ is C₁-C₆alkyl, C₂-C₆alkenyl, C₂-C₆alkynyl which are unsubstituted or mono to tri-substituted by C₃-C₆cycloalkyl, halogen or C₁-C₄alkoxy; O-C₁-C₆alkyl, O-C₂-C₆alkenyl, O-C₂-C₆alkynyl, which are unsubstituted or mono to tri-substituted by C₃-C₆cycloalkyl, halogen or C₁-C₄alkoxy; N-C₁-C₆alkyl; or N=CHC₁-C₆alkyl;

R₄ is C₁-C₆alkyl, C₂-C₆alkenyl, C₂-C₆alkynyl which are unsubstituted or mono to tri-substituted by C₃-C₆cycloalkyl, halogen or C₁-C₄alkoxy; or phenyl which is unsubstituted or mono to tri-substituted by fluorine, chlorine, bromine, C₁-C₄alkyl, C₁-C₄haloalkyl, C₁-C₄alkoxy, C₁-C₄haloalkoxy, phenyl or phenoxy and in which the phenyl part is unsubstituted or mono to tri-substituted by fluorine, chlorine, bromine, C₁-C₄alkyl, C₁-C₄haloalkyl, C₁-C₄alkoxy or C₁-C₄haloalkoxy (subgroup J1).

Another preferred group within the scope of subgroup H is that of compounds of the formula I, wherein

X is sulfur;

R₁ is hydrogen, fluorine, chlorine or bromine;

R₂ is hydrogen, fluorine, chlorine or bromine; at least one of R₁ and R₂ is not hydrogen;

R₃ is C₁-C₆alkyl, C₂-C₆alkenyl, C₂-C₆alkynyl which are unsubstituted or mono to tri-substituted by C₃-C₆cycloalkyl, halogen or C₁-C₄alkoxy; O-C₁-C₆alkyl, O-C₂-C₆alkenyl, O-C₂-C₆alkynyl, which are unsubstituted or mono to tri-substituted by C₃-C₆cycloalkyl, halogen or C₁-C₄alkoxy;

R₄ is C₁-C₆alkyl, C₂-C₆alkenyl, C₂-C₆alkynyl which are unsubstituted or mono to tri-substituted by C₃-C₆cycloalkyl, halogen or C₁-C₄alkoxy; or phenyl which is unsubstituted or mono to tri-substituted by fluorine, chlorine, bromine, C₁-C₄alkyl, C₁-C₄haloalkyl, C₁-C₄alkoxy, C₁-C₄haloalkoxy, phenyl or phenoxy and in which the phenyl part is unsubstituted or mono to tri-substituted by fluorine, chlorine, bromine, C₁-C₄alkyl, C₁-C₄haloalkyl, C₁-C₄alkoxy or C₁-C₄haloalkoxy (subgroup J2).

Other preferred groups of compounds are those of formula I, wherein

A is thiazolyl (subgroup K).

Within the scope of said group K, those compounds of formula I are preferred wherein

X is oxygen;

R₁ is hydrogen, fluorine, chlorine, bromine or iodine;

R₂ is hydrogen, fluorine, chlorine, bromine or iodine; at least one of R₁ and R₂ is not hydrogen;

R₃ is C₁-C₆alkyl, C₂-C₆alkenyl, C₂-C₆alkinyl which are unsubstituted or mono to tri-substituted by C₃-C₆cycloalkyl, halogen or C₁-C₄alkoxy; O-C₁-C₆alkyl, O-C₂-C₆alkenyl, O-C₂-C₆alkynyl, which are unsubstituted or mono to tri-substituted by C₃-C₆cycloalkyl, halogen or C₁-C₄alkoxy;

R₄ is C₁-C₆alkyl, C₂-C₆alkenyl, C₂-C₆alkinyl which are unsubstituted or mono to tri-substituted by C₃-C₆cycloalkyl, halogen or C₁-C₄alkoxy; or phenyl which is unsubstituted or mono to tri-substituted by fluorine, chlorine, bromine, C₁-C₄alkyl, C₁-C₄haloalkyl, C₁-C₄alkoxy, C₁-C₄haloalkoxy, phenyl or phenoxy and in which the phenyl part is unsubstituted or mono to tri-substituted by fluorine, chlorine, bromine, C₁-C₄alkyl, C₁-C₄haloalkyl, C₁-C₄alkoxy or C₁-C₄haloalkoxy (subgroup L).

Another preferred group of compounds are those of formula I, wherein

A is pyridazinyl (subgroup M).

Within the scope of said group M, those compounds of formula I are preferred wherein

X is oxygen;

R₁ is hydrogen, fluorine, chlorine, bromine or iodine;

R₂ is hydrogen, fluorine, chlorine, bromine or iodine; at least one of R₁ and R₂ is not hydrogen;

R₃ is C₁-C₆alkyl, C₂-C₆alkenyl, C₂-C₆alkinyl which are unsubstituted or mono to tri-substituted by C₃-C₆cycloalkyl, halogen or C₁-C₄alkoxy; O-C₁-C₆alkyl, O-C₂-C₆alkenyl, O-C₂-C₆alkynyl, which are unsubstituted or mono to tri-substituted by C₃-C₆cycloalkyl, halogen or C₁-C₄alkoxy;

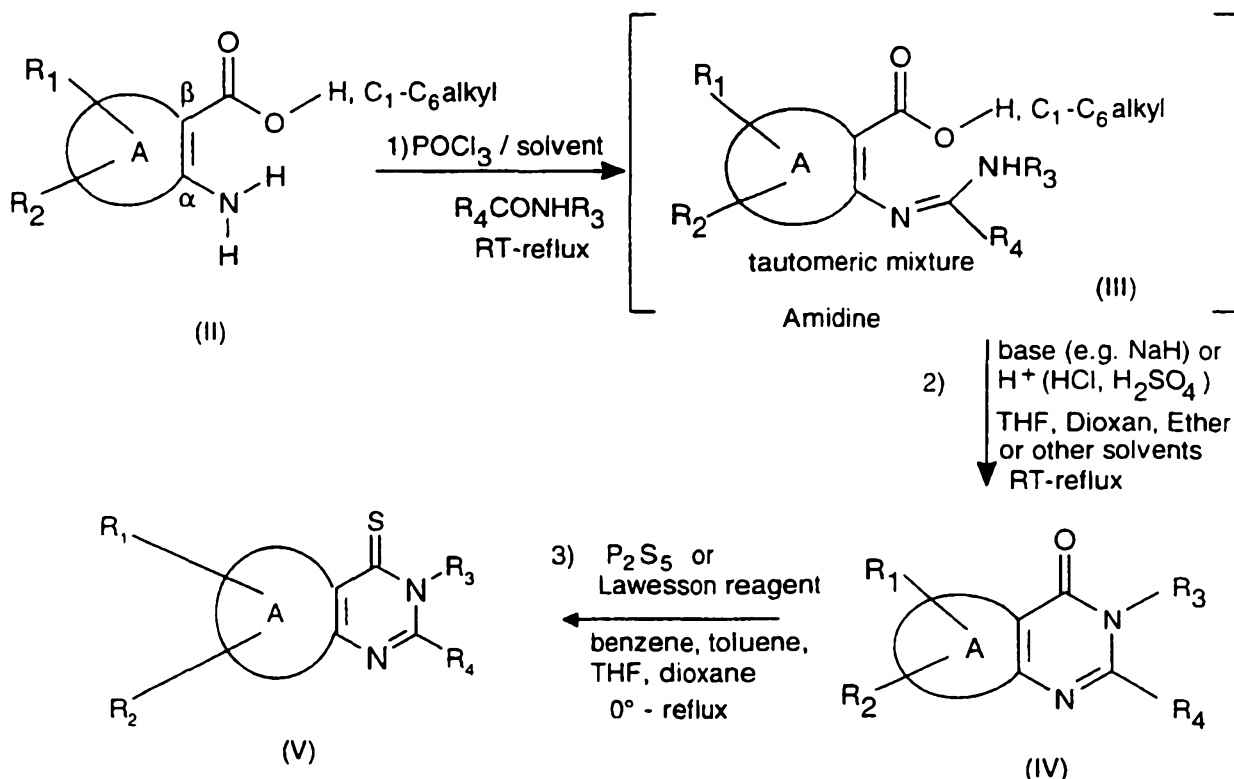
R₄ is C₁-C₆alkyl, C₂-C₆alkenyl, C₂-C₆alkinyl which are unsubstituted or mono to tri-substituted by C₃-C₆cycloalkyl, halogen or C₁-C₄alkoxy; or phenyl which is unsubstituted or mono to tri-substituted by fluorine, chlorine, bromine, C₁-C₄alkyl, C₁-C₄haloalkyl, C₁-C₄alkoxy,

C₁-C₄haloalkoxy, phenyl or phenoxy and in which the phenyl part is unsubstituted or mono to tri-substituted by fluorine, chlorine, bromine, C₁-C₄alkyl, C₁-C₄haloalkyl, C₁-C₄alkoxy or

C₁-C₄haloalkoxy (subgroup N).

The compounds of formula I can be prepared as follows :

Scheme 1



The compounds of formula I are preferably prepared starting from α -amino- β -carboalkoxy-heterocycles or α -amino- β -carbocyclic acid heterocycles, some of which, where Het = thienyl, are commercially available (2 isomers). The methyl thiophene-2-amino-3-carboxylate can be prepared, for example, in accordance with Acta Pharm. Suecica 1968, Vol. 5, p.563, according to S.Gronowitz et al. Other heterocycles can be prepared according to instructions in the literature. The synthesis of, for example, ethyl 5-aminothiazole-4-carboxylate and ethyl 5-amino-2-methylthiazole-4-carboxylate is described by Golankiewicz et al. in Tetrahedron 1985, 41, 5989. The reaction of the α -amino- β -carboalkoxyheterocycles or α -amino- β -carbocyclic acid heterocycles with amides (R_4CONHR_3) (step 1 in scheme 1) is conveniently carried out in the presence of $POCl_3$, $SOCl_2$ or SO_2Cl_2 , in solvents, such as $ClCH_2CH_2Cl$, $CHCl_3$, CH_2Cl_2 , benzene, toluene, hexane, cyclohexane or others in the temperature range from RT to reflux temperature. The resulting amidines (III) either cyclise spontaneously to the pyrimidin-4-ones or were converted into the cyclised products by

treatment with bases such as *t*-Butyl-O-K, NaH, KH, *n*-BuLi, NaOH, Na₂CO₃ or others in solvents such as THF, dioxane, hexane, toluene, DMSO, DMF, dimethylacetamid or others at temperatures between 20°C and reflux-temperature.

The replacement of the 4-one group with sulfur to the 4-thione group (step 3 in scheme 1) is carried out by reaction with P₂S₅ or Lawesson-reagent in tetrahydrofurane, dioxane or toluene as solvent in the temperature range of RT to reflux temperature.

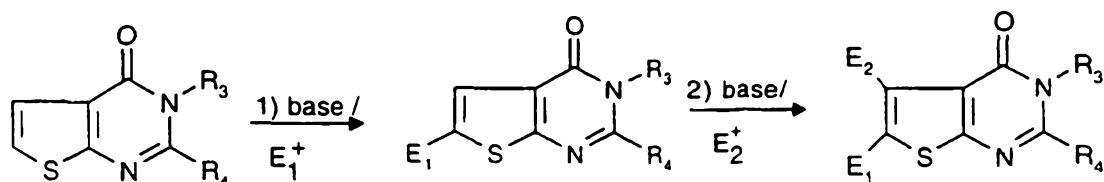
The above synthesis route is the first disclosure of how to prepare 3H-thieno[2.3-d]-pyrimidin-4-one derivatives within the structural pattern of formula I herein.

Methods for the preparation of compounds of the general formula I wherein R₁ = R₂ = hydrogen are described in Chemical Scripta 1981, 18, 135, Synthesis 1977, 180, Chem.Pharm.Bull. 1989, 37,2122 and DE-OS-2411273.

The invention also relates to the intermediates of the formula III, IV and V, and especially to those wherein A represents thienyl[2.3-d].

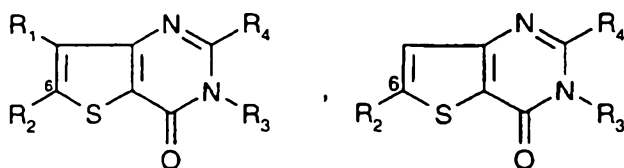
The introduction of further substituents into the 5-ring of the thienopyrimidin-4-ones may also conveniently be carried out using metallorganic methodology. Thieno[3.2-d]-pyrimidin-4-ones and thieno[2.3-d]pyrimidin-4-ones, for example, can be deprotonised selectively in 6-position. Particularly suitable bases for this purpose are lithium diisopropylamide (LDA), lithium cyclohexylisopropylamide (LICA) or secondary butyl lithium/TMEDA. A great number of the radicals R₁ or R₂ indicated above can be introduced by reacting the resulting anions with electrophiles (step 1 in scheme 2), typically Br₂, NBS, F₂, ICl, Cl₂, F⁺ reagents, trimethylsilyl chloride.

Scheme 2 : Synthesis of special heterocycles



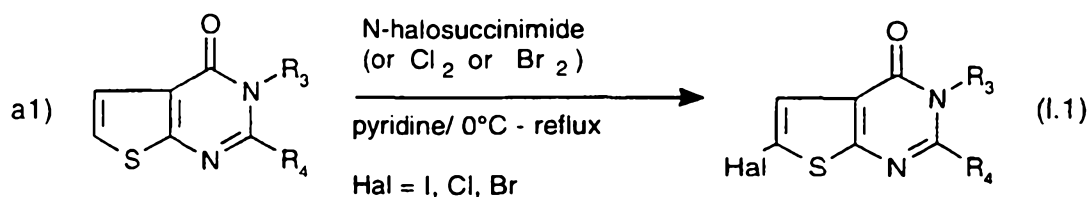
E_{1,2}⁺ = NBS (N-Bromsuccinimide), NCS (N-Chlorsuccinimide), I₂, Cl₂, Br₂, FCl, F⁺ reagents, TMS and similar Si reagents.

The following compounds can likewise be prepared in general accordance with the methods described in scheme 2:



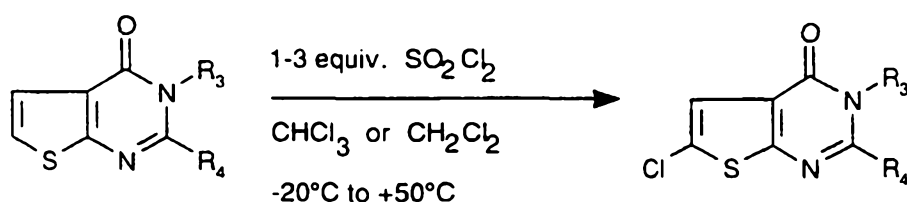
Scheme 3 : Synthesis of special thienopyrimidin-4-ones (special methods for the introduction of halogen into the thiophene ring)

a) Thieno[2.3-d]pyrimidin-4-ones :



1-3 molar equivalents of N-bromosuccinimide or N-chlorosuccinimide (or Cl_2 gas or Br_2) are used for halogenation. The solvent used is, for example, pyridine in the temperature range from 0°C to reflux. The reaction time is 1 to 24 hours.

a2) "Pure" chlorinating method :



The described reactions are carried out in per se known manner, e.g. in the presence or absence of a suitable solvent or diluent or of a mixture thereof, if appropriate with cooling, at room temperature or with heating, e.g. in the temperature range from about -20°C to the boiling temperature of the reaction medium, preferably in the range from about -20°C to about $+150^\circ\text{C}$ and, if required, in a closed vessel, under pressure, in an inert gas atmosphere and/or under anhydrous conditions.

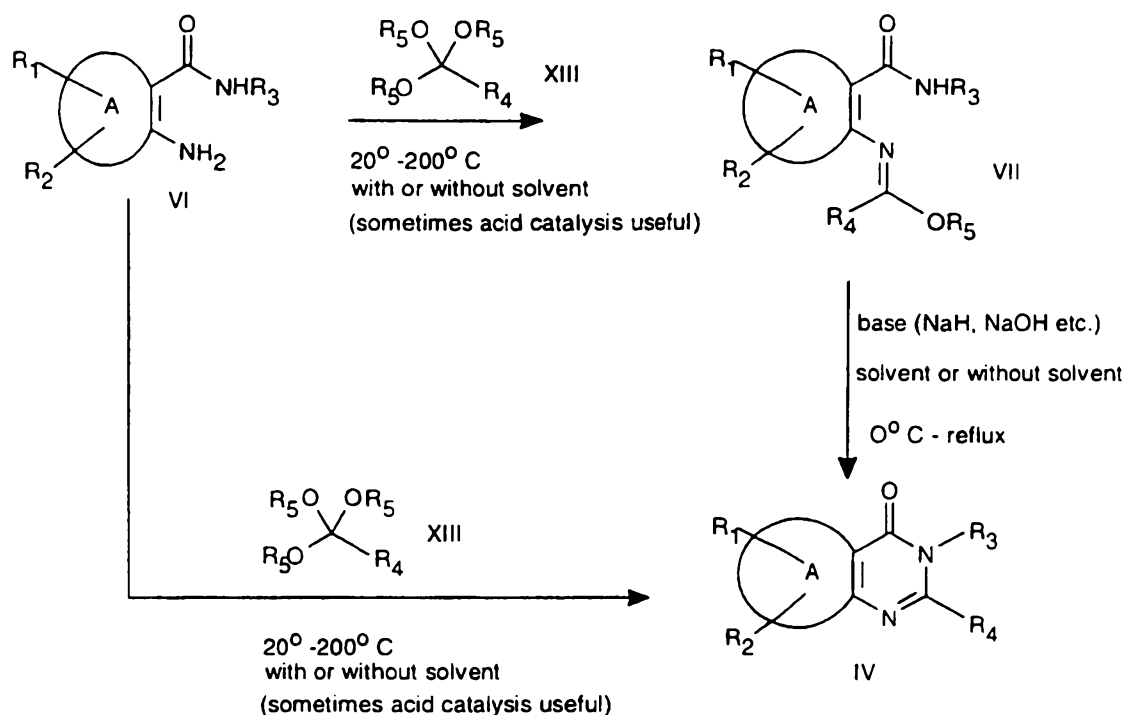
Illustrative examples of such solvents or diluents are: aromatic, aliphatic and alicyclic hydrocarbons and halogenated hydrocarbons, typically benzene, toluene, xylene, chlorobenzene, bromobenzene, petroleum ether, hexane, cyclohexane, dichloromethane, trichloromethane,

dichloroethane or trichloroethane; ethers, typically diethyl ether, tert-butylmethyl ether, tetrahydrofuran or dioxane; ketones, typically acetone or methyl ethyl ketone; alcohols, typically methanol, ethanol, propanol, butanol, ethylene glycol or glycerol; esters, typically ethyl acetate or butyl acetate; amides, typically N,N-dimethylformamide, N,N-dimethylacetamide, N-methylpyrrolidone or hexamethylphosphoric acid triamide; nitriles, typically acetonitrile; and sulfoxides, typically dimethylsulfoxide. Bases used in excess, such as triethylamine, pyridine, N-methylmorpholine or N,N-diethylaniline, can also be used as solvents or diluents.

Suitable bases are, for example, alkali metal hydroxide or alkaline earth metal hydroxide, alkali metal hydride or alkaline earth metal hydride, alkali metal amide or alkaline earth metal amide, alkali metal alkanolate or alkaline earth metal alkanolate, alkali metal carbonate or alkaline earth metal carbonate, alkali metal dialkylamide or alkaline earth metal dialkylamide, or alkali metal alkylsilylamide or alkaline earth metal alkylsilylamide, alkylamines, alkylenediamines, optionally N-alkylated, optionally unsaturated cycloalkylamines, basic heterocycles, ammonium hydroxides and carbocyclic amines. Examples meriting mention are sodium hydroxide, sodium hydride, sodium amide, sodium methanolate, sodium carbonate, potassium tert-butanolate, potassium carbonate, lithium diisopropylamide, potassium bis(trimethylsilyl)amide, calcium hydride, triethylamine, triethylenediamine, cyclohexylamine, N-cyclohexyl-N,N-dimethylamine, N,N-diethylaniline, pyridine, 4-(N,N-dimethylamino)pyridine, N-methylmorpholine, benzyltrimethylammonium hydroxide, and 1,8-diazabicyclo[5.4.0]undec-5-ene (DBU).

The compounds of the formula I can also be prepared as follows

Scheme 4:



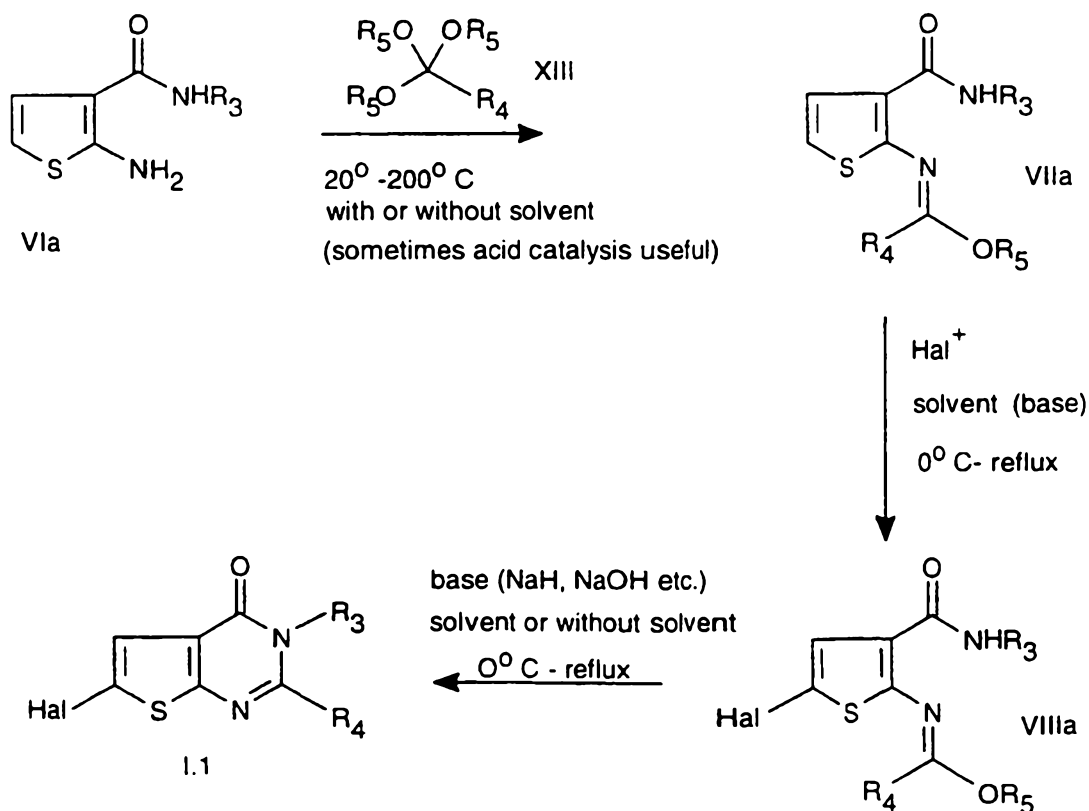
$R_5 = \text{C}_1\text{-C}_6\text{-alkyl}$

The amino carboxylic acidamide of formula VI reacts with the orthoester of formula XIII in the presence or absence of a suitable solvent or diluent, if required in the presence of an acid catalyst at room temperature or with heating, e.g. in the temperature range from about 20 to 200°C . Illustrative examples of solvents or diluents are ethers like tert.butylmethylether, tetrahydrofuran, dimethylether; amides like N,N-dimethylformamide or N-methyl-pyrrolidone; sulfoxides, typically dimethylsulfoxid and alcohols like methanol, ethanol, propanol, butanol, ethylene glycol or glycerol. As catalyst can be used hydrogen halides, methanesulfonic acid, trifluoromethyl acetic acid, p-toluenesulfonic acid and others in the absence of water. Commonly used bases are sodium hydroxid, potassium hydroxid, sodium hydrogencarbonate, sodium carbonate, sodium hydride, potassium hydride, potassium carbonate and others.

Especially for the thienopyrimidinone, scheme 5 describes the reaction of the aminothiophene-carboxylic-acid amide VIa with the orthoester XIII in the presence or absence of a solvent, if required in the presence of an acid catalyst in the temperature range from 20 to 200°C. The resulting intermediate VIIa is then halogenated in the presence of a solvent at temperatures from 20°C to reflux. The halogenated intermediate VIIIa is then cyclised in the presence of a base, in the presence or absence of a suitable solvent at temperatures from 20°C to reflux. Halogenation reagents are typically N-Bromsuccinimide, N-Chlorosuccinimide, N-iodosuccinimide, Chlorgas, Br₂, thionylchloride and others. Preferably solvents used for the halogenation are tert.-butylmethylether, tetrahydrofurane, chloroform, methylenechloride, pyridine and quinoline.

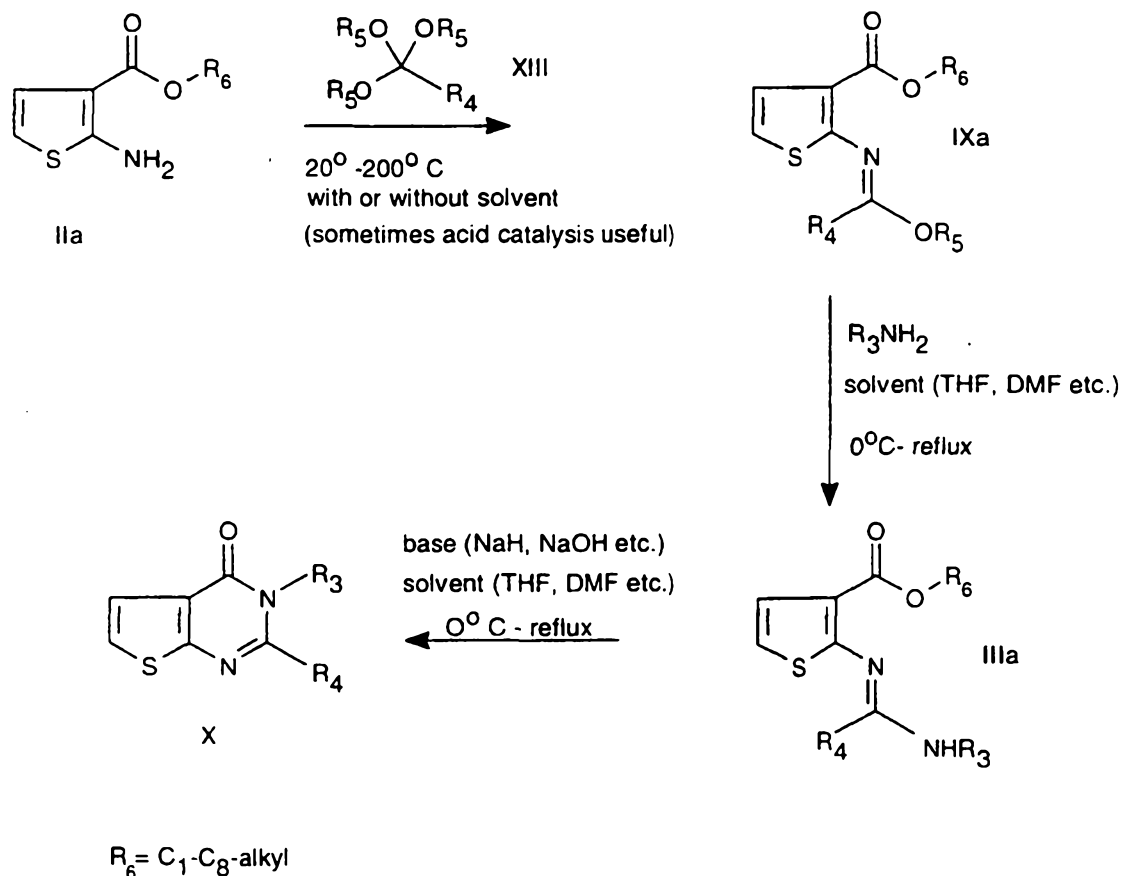
Scheme 5:

(Thienopyrimidinones)

Hal⁺ = NCS, NBS, NIS, Cl₂, Br₂, SO₂Cl₂solvents for halogenation: THF, TBME, CHCl₃, CH₂Cl₂, pyridine, quinoline and others

Another alternative is described in scheme 6, in which the amino-carboalkoxy-thiophene of formula IIa reacts with the orthoester XIII to the intermediate IXa, than the compound IXa is transformed to the amidine IIIa and cyclised to the thienopyrimidinone X. The halogenation of X to obtain I.1 is as described in Scheme 3.

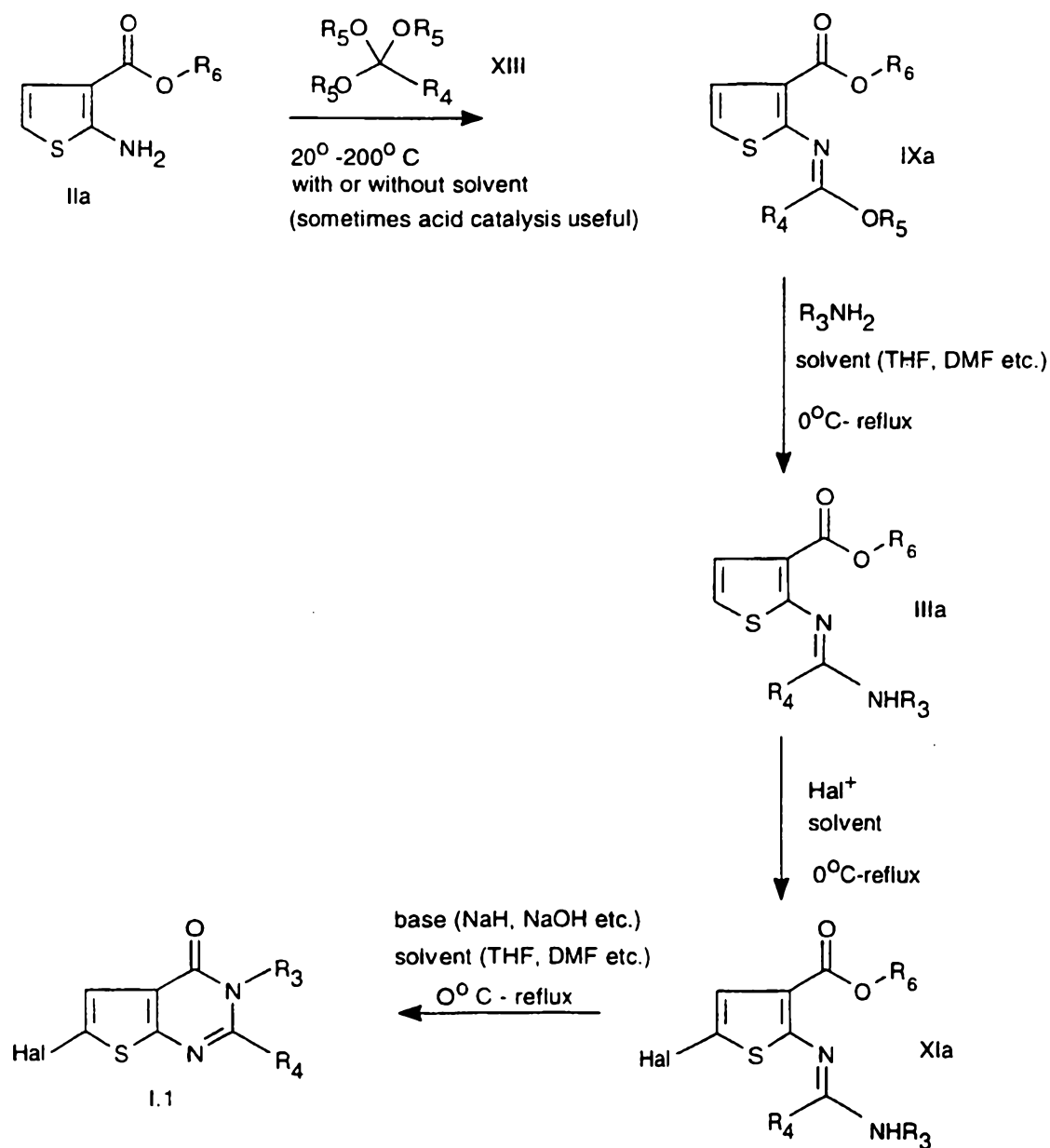
Scheme 6:



The reaction conditions from IIa to IXa is as described in the schemes 4 or 5, as well as the cyclisation from IIIa to X is as described above. The reaction from IXa to IIIa requires as solvent for example tetrahydrofuran, N,N-dimethylformamide or others at a temperature range from 0°C to reflux.

The scheme 7 describes the reaction of the compound IIa with the orthoester XIII to obtain the intermediate IXa, which is converted to the intermediate IIIa and than halogenated to the thiophene XIa. Cyclisation of XIa gives the compound I.1.

Scheme 7

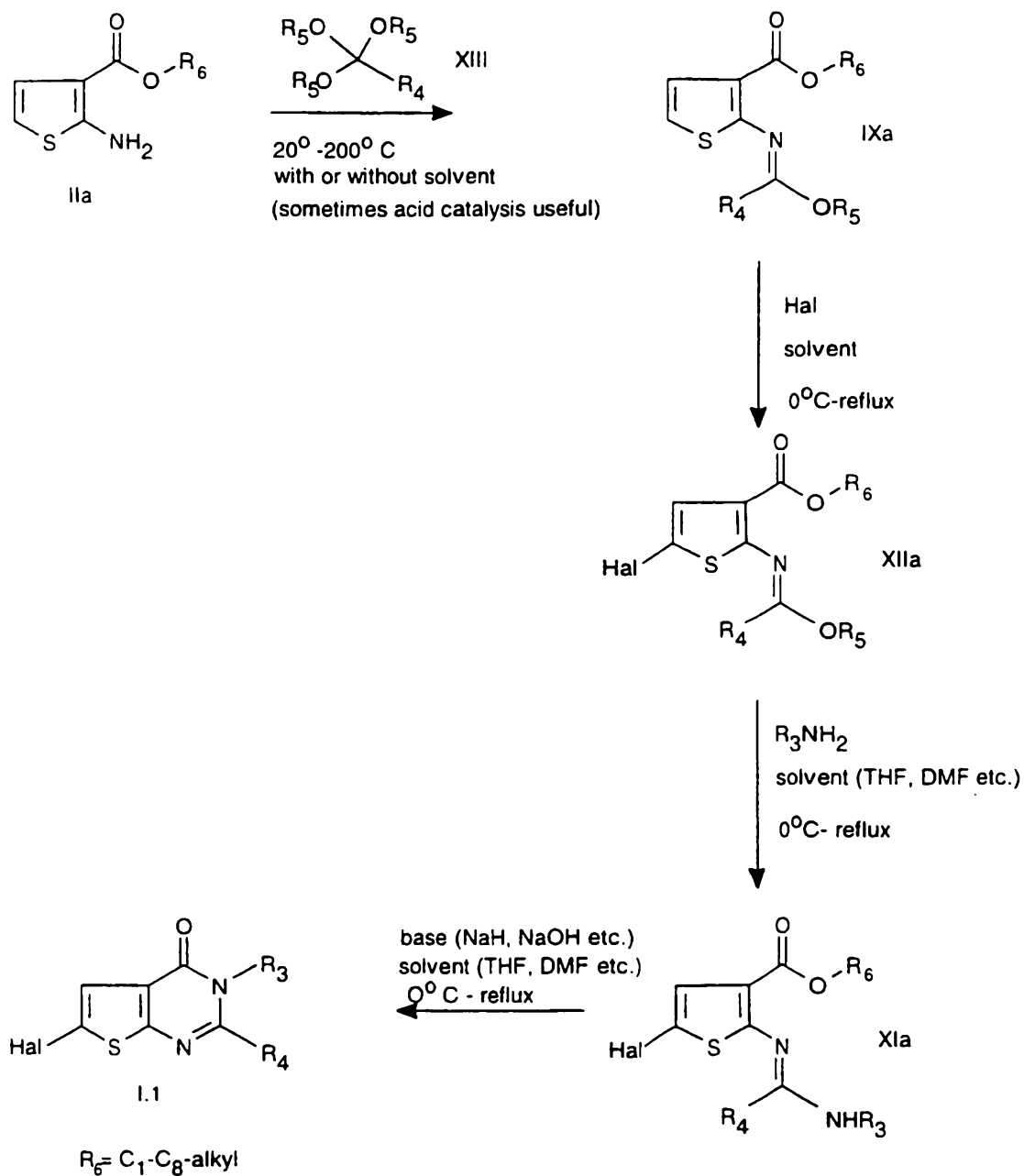


$\text{R}_6 = \text{C}_1\text{-C}_8\text{-alkyl}$

The reactions conditions described in process 4 are as described analogously in the schemes 4, 5 or 6.

Scheme 8 describes the alternative route comprising the reaction of the compound IIa with the orthoester XIII to the intermediate IXa, which is halogenated to the intermediate XIa and then converted to the thiophene XIa. Cyclisation of XIa gives the end product I.1. The reactions conditions are as described in the schemes 4, 5, 6 and 7.

Scheme 8:



Quinazolinone derivatives having fungicidal properties are known from WO-94/26722 or EP-A-276825 and thienopyrimidinones are known from WO-97/02262.

Surprisingly, it has now been found that the novel compounds of formula I have, for practical purposes, a very advantageous spectrum of activities for protecting plants against diseases that are caused by fungi as well as by bacteria and viruses.

The compounds of formula I can be used in the agricultural sector and related fields as active ingredients for controlling plant pests. The novel compounds are distinguished by excellent activity at low rates of application, by being well tolerated by plants and by being environmentally safe. They have very useful curative, preventive and systemic properties and are used for protecting numerous cultivated plants. The compounds of formula I can be used to inhibit or destroy the pests that occur on plants or parts of plants (fruit, blossoms, leaves, stems, tubers, roots) of different crops of useful plants, while at the same time protecting also those parts of the plants that grow later e.g. from phytopathogenic micro-organisms.

It is also possible to use compounds of formula I as dressing agents for the treatment of plant propagation material, in particular of seeds (fruit, tubers, grains) and plant cuttings (e.g. rice), for the protection against fungal infections as well as against phytopathogenic fungi occurring in the soil.

The compounds I are, for example, effective against the phytopathogenic fungi of the following classes: Fungi imperfecti (e.g. Botrytis, Pyricularia, Helminthosporium, Fusarium, Septoria, Cercospora and Alternaria) and Basidiomycetes (e.g. Rhizoctonia, Hemileia, Puccinia). Additionally, they are also effective against the Ascomycetes classes (e.g. Venturia and Erysiphe, Podosphaera, Monilinia, Uncinula) and of the Oomycetes classes (e.g. Phytophthora, Pythium, Plasmopara). Furthermore, the novel compounds of formula I are effective against phytopathogenic bacteria and viruses (e.g. against Xanthomonas spp, Pseudomonas spp, Erwinia amylovora as well as against the tobacco mosaic virus).

Within the scope of this invention, target crops to be protected typically comprise the following species of plants: cereal (wheat, barley, rye, oat, rice, maize, sorghum and related species); beet (sugar beet and fodder beet); pomes, drupes and soft fruit (apples, pears, plums, peaches, almonds, cherries, strawberries, raspberries and blackberries); leguminous plants (beans, lentils, peas, soybeans); oil plants (rape, mustard, poppy, olives, sunflowers, coconut, castor oil plants, cocoa beans, groundnuts); cucumber plants (pumpkins, cucumbers, melons); fiber plants (cotton, flax, hemp, jute); citrus fruit (oranges, lemons, grapefruit, mandarins); vegetables (spinach, lettuce, asparagus, cabbages, carrots, onions, tomatoes, potatoes, paprika); lauraceae (avocado, cinnamomum, camphor) or

plants such as tobacco, nuts, coffee, eggplants, sugar cane, tea, pepper, vines, hops, bananas and natural rubber plants, as well as ornamentals.

The compounds of formula I are normally used in the form of compositions and can be applied to the crop area or plant to be treated, simultaneously or in succession with further compounds. These further compounds can be e.g. fertilizers or micronutrient donors or other preparations which influence the growth of plants. They can also be selective herbicides as well as insecticides, fungicides, bactericides, nematocides, molluscicides or mixtures of several of these preparations, if desired together with further carriers, surfactants or application promoting adjuvants customarily employed in the art of formulation.

The compounds of formula I can be mixed with other fungicides, resulting in some cases in unexpected synergistic activities.

Mixing components which are particularly preferred are azoles such as azaconazole, bitertanol, propiconazole, difenoconazole, diniconazole, cyproconazole, epoxiconazole, fluquinconazole, flusilazole, flutriafol, hexaconazole, imazalil, imibenconazole, ipconazole, tebuconazole, tetraconazole, fenbuconazole, metconazole, myclobutanil, perfurazoate, penconazole, bromuconazole, pyrifenoxy, prochloraz, triadimefon, triadimenol, triflumizole or triticonazole; pyrimidinyl carbinols such as ancymidol, fenarimol or nuarimol; 2-amino-pyrimidine such as bupirimate, dimethirimol or ethirimol; morpholines such as dodemorph, fenpropidin, fenpropimorph, spiroxamin or tridemorph; anilinopyrimidines such as cyprodinil, pyrimethanil or mepanipyrim; pyrroles such as fenpiclonil or fludioxonil; phenylamides such as benalaxyl, furalaxyl, metalaxyl, R-metalaxyl, ofurace or oxadixyl; benzimidazoles such as benomyl, carbendazim, debacarb, fuberidazole or thiabendazole; dicarboximides such as chlozolinate, dichlozoline, iprodione, myclozoline, procymidone or vinclozolin; carboxamides such as carboxin, fenfuram, flutolanil, mepronil, oxycarboxin or thifluzamide; guanidines such as guazatine, dodine or iminoctadine; strobilurines such as azoxystrobin, kresoxim-methyl, metominostrobin, SSF-129 or 2-[α [(α -methyl-3-trifluoromethyl-benzyl)imino]-oxy]-o-tolyl]-glyoxylic acid-methylester-O-methyloxime; dithiocarbamates such as ferbam, mancozeb, maneb, metiram, propineb, thiram, zineb or ziram; N-halomethylthio-dicarboximides such as captafol, captan, dichlofluanid, fluoromide, folpet or tolyfluanid; copper compounds such as Bordeaux mixture, copper hydroxide, copper oxychloride, copper sulfate, cuprous oxide, mancopper or oxine-copper; nitrophenol derivatives such as dinocap or nitrothal-isopropyl; organo phosphorous derivatives such as edifenphos,

iprobenphos, isoprothiolane, phosdiphen, pyrazophos or toclofos-methyl; and other compounds of diverse structures such as acibenzolar-S-methyl, anilazine, blastocidin-S, chinomethionat, chloroneb, chlorothalonil, cymoxanil, dichlone, diclomezine, dicloran, diethofencarb, dimethomorph, dithianon, etridiazole, famoxadone, fentin, ferimzone, fluazinam, flusulfamide, fenhexamid, fosetyl-aluminium, hymexazol, kasugamycin, methasulfocarb, pencycuron, phthalide, polyoxins, probenazole, propamocarb, pyroquilon, quinoxyfen, quintozene, sulfur, triazoxide, tricyclazole, triforine or validamycin.

Preferred compound for mixing with the above-mentioned mixing components is compound no. 3.30.

Another preferred compound for mixing with the above-mentioned mixing components is compound no. 3.31.

Another preferred compound for mixing with the above-mentioned mixing components is compound no. 3.58.

Another preferred compound for mixing with the above-mentioned mixing components is compound no. 3.59.

Suitable carriers and adjuvants can be solid or liquid and are substances useful in formulation technology, e.g. natural or regenerated mineral substances, solvents, dispersants, wetting agents, tackifiers, thickeners, binders or fertilizers.

A preferred method of applying a compound of formula I, or an agrochemical composition which contains at least one of said compounds, is foliar application. The frequency of application and the rate of application will depend on the risk of infestation by the corresponding pathogen. However, the compounds of formula I can also penetrate the plant through the roots via the soil (systemic action) by drenching the locus of the plant with a liquid formulation, or by applying the compounds in solid form to the soil, e.g. in granular form (soil application). In crops of water rice such granulates can be applied to the flooded rice field. The compounds of formula I may also be applied to seeds (coating) by impregnating the seeds or tubers either with a liquid formulation of the fungicide or coating them with a solid formulation.

The compounds of formula I are used in unmodified form or, preferably, together with the adjuvants conventionally employed in the art of formulation. To this end they are conveniently formulated in known manner to emulsifiable concentrates, coatable pastes, directly sprayable or dilutable solutions, dilute emulsions, wettable powders, soluble powders, dusts, granulates, and also encapsulations e.g. in polymeric substances. As with the type of the compositions, the methods of application, such as spraying, atomizing, dusting, scattering, coating or pouring, are chosen in accordance with the intended objectives and the prevailing circumstances.

Advantageous rates of application are normally from 5 g to 2 kg of active ingredient (a.i.) per hectare (ha), preferably from 10 g to 1 kg a.i./ha, most preferably from 20 g to 600 g a.i./ha. When used as seed drenching agent, convenient dosages are from 10 mg to 1 g of active substance per kg of seeds.

The formulation, i.e. the compositions containing the compound of formula I and, if desired, a solid or liquid adjuvant, are prepared in known manner, typically by intimately mixing and/or grinding the compound with extenders, e.g. solvents, solid carriers and, optionally, surface active compounds (surfactants).

Suitable carriers and adjuvants may be solid or liquid and correspond to the substances ordinarily employed in formulation technology, such as, e.g. natural or regenerated mineral substances, solvents, dispersants, wetting agents, tackifiers, thickeners binding agents or fertilizers. Such carriers are for example described in WO 97/33890.

Further surfactants customarily employed in the art of formulation are known to the expert or can be found in the relevant literature.

The agrochemical formulations will usually contain from 0.1 to 99 % by weight, preferably from 0.1 to 95 % by weight, of the compound of formula I, 99.9 to 1 % by weight, preferably 99.8 to 5 % by weight, of a solid or liquid adjuvant, and from 0 to 25 % by weight, preferably from 0.1 to 25 % by weight, of a surfactant.

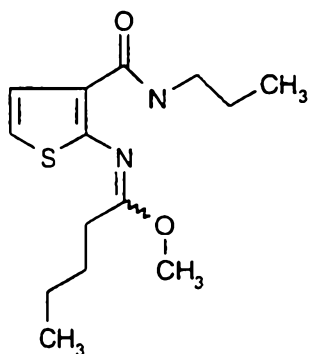
Whereas it is preferred to formulate commercial products as concentrates, the end user will normally use dilute formulations.

The compositions may also contain further adjuvants such as stabilizers, antifoams, viscosity regulators, binders or tackifiers as well as fertilizers, micronutrient donors or other formulations for obtaining special effects.

The following non-limitative Examples illustrate the above-described invention in more detail. Temperatures are given in degrees Celsius. The following abbreviations are used: Et = ethyl; i-propyl = isopropyl; Me = methyl; m.p.= melting point. "NMR" means nuclear magnetic resonance spectrum. MS = mass spectrum. "%" is percent by weight, unless corresponding concentrations are indicated in other units.

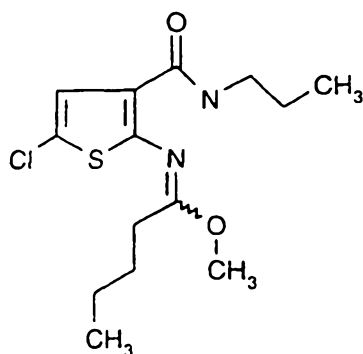
Preparation examples:

Example P-1 : 2-(1-n-butyl-1-methoxymethyleneamino)thiophene-3-carboxylic-acid propylamide



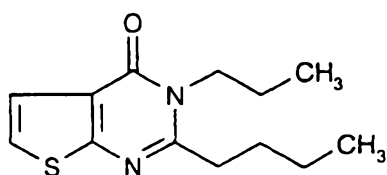
In a distillation apparatus, a mixture of 1.84 g of 2-aminothiophene-3-carboxylic-acid propylamide and 2.43 g of trimethyl orthovalerate is heated for 2 hours at 130°C. Methanol, which arises during the reaction is directly distilled out of the reaction flask. After cooling, the crude product is purified by column chromatography (eluant: hexane/ethylacetate = 1:2). Yield : 1.9 g pure 2-(1-n-butyl-1-methoxymethyleneamino)thiophene-3-carboxylic-acid propylamide; m.p. 68-70°C.

Example P-2 : 5-Chloro-2-(1-n-butyl-1-methoxymethyleneamino)thiophene-3-carboxylic-acid propylamide



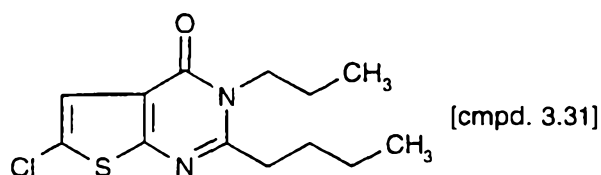
In a sulfonation flask 0.85 g 2-(1-n-butyl-1-methoxymethyleneamino)thiophene-3-carboxylic acid propylamide are added with stirring to 10 ml absolute pyridine. The internal temperature is then raised to 60°C and 0.5 g of N-chlorosuccinimide (NCS) are added in two portions. After stirring for 1 hour at 60°C, the pyridine is removed in a water jet vacuum. The residue is taken up in ethylacetate and the organic phase is washed twice with water. After drying of the organic phase, the solvent is removed in a water jet vacuum and the raw material purified by column chromatography over silica gel (eluant : hexane/ethylacetate = 3:1). Yield : 0.6 g 5-Chloro-2-(1-n-butyl-1-methoxymethyleneamino)thiophene-3-carboxylic acid propylamide in the form of brownish crystals; m.p. 110-112°C.

Example P-3 : 2-n-Butyl-3-n-propyl-3H-thieno[2,3-d]pyrimidin-4-one



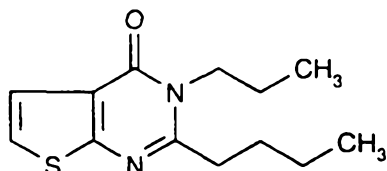
In a sulfonation flask, 0.85 g of 2-(1-n-butyl-1-methoxymethyleneamino)-thiophene-3-carboxylic acid propylamide is dissolved in 20 ml of absolute THF and 0.15 g of a ca. 55% NaH dispersion is added in small portions. The mixture is stirred for 15 minutes at room temperature and 1 hour at reflux temperature. Then the solvent is removed in a water jet vacuum and the residue taken up in ethylacetate. The organic phase is washed twice with water and after drying of the organic phase with sodium sulfate, the solvent is removed in a water jet vacuum. The resulting crude product (yield: 0.8g of 2-n-butyl-3-n-propyl-3H-thieno[2,3-d]pyrimidine-4-one in the form of a brown liquid) can be used without further purification for the halogenation step.

Example P-4 : 2-n-butyl-6-chloro-3-n-propyl-3H-thieno[2,3-d]pyrimidine-4-one



In a sulfonation flask, 0.36g of 5-Chloro-2-(1-n-butyl-1-methoxymethylene-amino)thiophene-3-carboxylic-acid propylamide is dissolved in 20 ml of absolute THF and 0.085g of a ca. 55% NaH dispersion is added in one portion. The mixture is stirred for 15 minutes at room temperature and 1 hour at reflux temperature. Then the solvent is removed in a water jet vacuum and the residue taken up in ethylacetate. The organic phase is washed twice with water and after drying of the organic phase with sodium sulfate, the solvent is removed in a water jet vacuum. The resulting crude product is purified by column chromatography over silica gel (eluant: hexane/ethylacetate = 5:1). Yield: 0.2 g 2-n-butyl-6-chloro-3-n-propyl-3H-thieno[2,3-d]pyrimidin-4-one in the form of a slightly yellowish powder; m.p. 67-69°C.

Example P-3a : 2-n-butyl-3-n-propyl-3H-thieno[2.3-d]pyrimidin-4-one

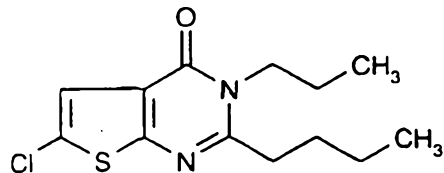


In a sulfonation flask, 11.0 g (70 mmol) of 2-amino-3-carbomethoxythiophen and 10.9 g (76 mmol) valeric acid propylamide are added to 60 ml 1,2-dichloroethane. Under stirring and at room temperature 7 ml of phosphoroychloride is slowly added dropwise. After 3 hours at reflux temperature the mixture is poored into ice water and adjusted light alkali with sodium hydrogencarbonat. The resulting mixture is then extracted three times with methylene-chloride and the separated organic phase dried over sodium sulfate. The solvent is then removed in a water-jet vacuum.

In a sulfonation flask, the crude product is added to 100 ml of absolute tetrahydrofurane and under stirring 4.36 g (0.1 mol) NaH in 50 ml abs. THF is carefully added. After stirring for 2 hours at reflux temperature, the solvent is removed in a water-jet vacuum and the residue is taken up in ethyl acetate/water. The water-phase is extracted with additionally ethyl acetate. The organic phase is dried over sodium sulfate and the solvent removed in a water-jet vacuum. The crude product is purified by column chromatography over silica gel

(eluant : TBME/hexane = 1:2). 12.0 g of 2-n-butyl-3-n-propyl-3H-thieno[2.3-d]pyrimidin-4-one are obtained in the form of a yellow powder having a melting point of 70-72°C.

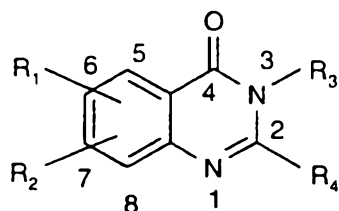
Example P-4a 2-n-butyl-6-chloro-3-n-propyl-3H-thieno-[2.3-d]pyrimidin-4-one



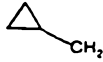
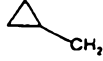






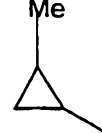
[cmpd. no. 3.31]





In a sulfonation flask, 2.0 g (8 mmol) of 2-n-butyl-3-propyl-3H-thieno[2.3-d]pyrimidin-4-one are added, with stirring, to 15 ml of absolute pyridine. The internal temperature is then raised to 80°C and then 1.87 g (14 mmol) of N-chlorosuccinimide (NCS) are added in smallish portions. After stirring for 3 hours at 90°C 1.0 g of NCS is added and the mixture is stirred another 3 hours at 90°C. The pyridine is removed in a water-jet vacuum and the crude product so obtained is purified by column chromatography over silica gel (eluant: n-hexane/tert.butylmethylether = 3:1), giving 0.9 g of 2-n-butyl-6-chloro-3-propyl-3H-thieno-[2.3-d]pyrimidin-4-one in the form of a beige powder having a melting point of 67-69°C.

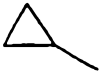
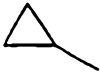
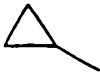
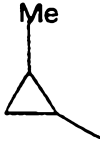
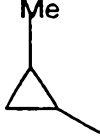
Table 1: A = Phenyl













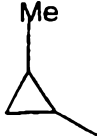




compd. No.	R ₁	R ₂	R ₃	R ₄	phys. data m.p. °C
1.1	6-Br	H	Me	Me	
1.2	6-Cl	H	Me	Et	
1.3	6-Br	H	Me	n-Propyl	
1.4	6-Cl	H	Me	n-Propyl	
1.5	H	7-Cl	Me	n-Propyl	
1.6	6-Br	H	Me	n-Butyl	
1.7	6-Cl	H	Me	n-Butyl	
1.8	H	7-Cl	Me	n-Butyl	
1.9	6-Br	H	Me	i-Butyl	
1.10	6-Cl	H	Me	i-Butyl	
1.11	6-Br	H	Me	n-Pentyl	
1.12	6-Br	H	Me		
1.13	6-Cl	H	Me		
1.14	6-Br	H	Et	Me	
1.15	6-Cl	H	Et	Et	
1.16	6-Br	H	Et	n-Propyl	
1.17	6-Cl	H	Et	n-Propyl	
1.18	H	7-Cl	Et	n-Propyl	
1.19	6-Br	H	Et	n-Butyl	
1.20	6-Cl	H	Et	n-Butyl	
1.21	H	7-Cl	Et	n-Butyl	
1.22	6-Br	H	Et	i-Butyl	

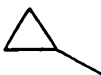
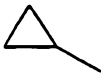
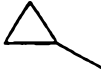
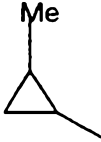
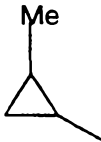
1.23	6-Cl	H	Et	i-Butyl	
1.24	6-Br	H	Et	n-Pentyl	
1.25	6-Br	H	Et		
1.26	6-Cl	H	Et		
1.27	6-Br	H	n-Propyl	Me	
1.28	6-Cl	H	n-Propyl	Et	
1.29	6-Br	H	n-Propyl	n-Propyl	
1.30	6-Cl	H	n-Propyl	n-Propyl	
1.31	H	7-Cl	n-Propyl	n-Propyl	
1.32	H	7-I	n-Propyl	n-Propyl	
1.33	6-Br	H	n-Propyl		
1.34	6-Cl	H	n-Propyl		
1.35	H	7-Cl	n-Propyl		
1.36	6-Br	H	n-Propyl	n-Butyl	130-135
1.37	6-Cl	H	n-Propyl	n-Butyl	
1.38	H	7-Cl	n-Propyl	n-Butyl	
1.39	H	7-I	n-Propyl	n-Butyl	
1.40	6-Br	H	n-Propyl	i-Butyl	
1.41	6-Cl	H	n-Propyl	i-Butyl	
1.42	6-Br	H	n-Propyl		
1.43	6-Cl	H	n-Propyl		
1.44	H	7-Cl	n-Propyl		
1.45	6-Br	H	n-Propyl	Me 	




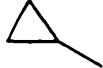
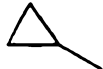
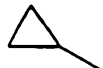
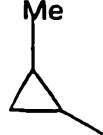
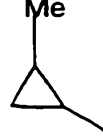
1.46	6-Cl	H	n-Propyl	Me 
1.47	6-Br	H	n-Propyl	Cyclobutyl
1.48	6-Br	H	n-Propyl	n-Pentyl
1.49	6-Cl	H	n-Propyl	n-Pentyl
1.50	H	7-Cl	n-Propyl	n-Pentyl
1.51	6-Br	H	n-Propyl	Cyclopentyl
1.52	6-Br	H	n-Propyl	n-Hexyl
1.53	6-Br	H	n-Propyl	Cyclohexyl
1.54	6-Br	H	n-Propyl	Phenyl
1.55	6-Br	H	n-Propyl	4-Chloro-phenyl
1.56	6-Cl	H	n-Propyl	4-Chloro-phenyl
1.57	6-Br	H	n-Propyl	4-Phenoxy-phenyl
1.58	6-Br	H	n-Butyl	Me
1.59	6-Cl	H	n-Butyl	Et
1.60	6-Br	H	n-Butyl	n-Propyl
1.61	6-Cl	H	n-Butyl	n-Propyl
1.62	H	7-Cl	n-Butyl	n-Propyl
1.63	H	7-I	n-Butyl	n-Propyl
1.64	6-Br	H	n-Butyl	
1.65	6-Cl	H	n-Butyl	
1.66	H	7-Cl	n-Butyl	
1.67	6-Br	H	n-Butyl	n-Butyl
1.68	6-Cl	H	n-Butyl	n-Butyl
1.69	H	7-Cl	n-Butyl	n-Butyl
1.70	H	7-I	n-Butyl	n-Butyl
1.71	6-Br	H	n-Butyl	i-Butyl



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1.73	6-Br	H	n-Butyl	
1.74	6-Cl	H	n-Butyl	
1.75	H	7-Cl	n-Butyl	
1.76	6-Br	H	n-Butyl	Me 
1.77	6-Cl	H	n-Butyl	Me 
1.78	6-Br	H	n-Butyl	Cyclobutyl
1.79	6-Br	H	n-Butyl	n-Pentyl
1.80	6-Cl	H	n-Butyl	n-Pentyl
1.81	H	7-Cl	n-Butyl	n-Pentyl
1.82	6-Br	H	n-Butyl	Cyclopentyl
1.83	6-Br	H	n-Butyl	n-Hexyl
1.84	6-Br	H	n-Butyl	Cyclohexyl
1.85	6-Br	H	n-Butyl	Phenyl
1.86	6-Br	H	n-Butyl	4-Chloro-phenyl
1.87	6-Cl	H	n-Butyl	4-Chloro-phenyl
1.88	6-Br	H	n-Butyl	4-Phenoxy-phenyl
1.89	6-Br	H	i-Butyl	n-Propyl
1.90	6-Cl	H	i-Butyl	n-Propyl
1.91	6-Br	H	i-Butyl	n-Butyl
1.92	6-Cl	H	i-Butyl	n-Butyl

1.93	6-Br	H		n-Propyl
1.94	6-Cl	H		n-Propyl
1.95	6-Br	H		n-Propyl
1.96	6-Br	H		n-Butyl
1.97	6-Br	H	n-Pentyl	Me
1.98	6-Cl	H	n-Pentyl	Et
1.99	6-Br	H	n-Pentyl	n-Propyl
1.100	6-Cl	H	n-Pentyl	n-Propyl
1.101	H	7-Cl	n-Pentyl	n-Propyl
1.102	H	7-I	n-Pentyl	n-Propyl
1.103	6-Br	H	n-Pentyl	
1.104	6-Cl	H	n-Pentyl	
1.105	H	7-Cl	n-Pentyl	
1.106	6-Br	H	n-Pentyl	n-Butyl
1.107	6-Cl	H	n-Pentyl	n-Butyl
1.108	H	7-Cl	n-Pentyl	n-Butyl
1.109	H	7-I	n-Pentyl	n-Butyl
1.110	6-Br	H	n-Pentyl	i-Butyl
1.111	6-Cl	H	n-Pentyl	i-Butyl
1.112	6-Br	H	n-Pentyl	
1.113	6-Cl	H	n-Pentyl	
1.114	H	7-Cl	n-Pentyl	

1.115	6-Br	H	n-Pentyl	Me 
1.116	6-Cl	H	n-Pentyl	Me 
1.117	6-Br	H	n-Pentyl	Cyclobutyl
1.118	6-Br	H	n-Pentyl	n-Pentyl
1.119	6-Cl	H	n-Pentyl	n-Pentyl
1.120	H	7-Cl	n-Pentyl	n-Pentyl
1.121	6-Br	H	n-Pentyl	Cyclopentyl
1.122	6-Br	H	n-Pentyl	n-Hexyl
1.123	6-Br	H	n-Pentyl	Cyclohexyl
1.124	6-Br	H	n-Pentyl	Phenyl
1.125	6-Br	H	n-Pentyl	4-Chloro-phenyl
1.126	6-Cl	H	n-Pentyl	4-Chloro-phenyl
1.127	6-Br	H	n-Pentyl	4-Phenoxy-phenyl
1.128	6-Br	H	OEt	Me
1.129	6-Cl	H	OEt	Et
1.130	6-Br	H	OEt	n-Propyl
1.131	6-Cl	H	OEt	n-Propyl
1.132	H	7-Cl	OEt	n-Propyl
1.133	H	7-I	OEt	n-Propyl
1.134	6-Br	H	OEt	
1.135	6-Cl	H	OEt	
1.136	H	7-Cl	OEt	
1.137	6-Br	H	OEt	n-Butyl

1.138	6-Cl	H	OEt	n-Butyl
1.139	H	7-Cl	OEt	n-Butyl
1.140	H	7-I	OEt	n-Butyl
1.141	6-Br	H	OEt	i-Butyl
1.142	6-Cl	H	OEt	i-Butyl
1.143	6-Br	H	OEt	
1.144	6-Cl	H	OEt	
1.145	H	7-Cl	OEt	
1.146	6-Br	H	OEt	Me 
1.147	6-Cl	H	OEt	Me 
1.148	6-Br	H	OEt	Cyclobutyl
1.149	6-Br	H	OEt	n-Pentyl
1.150	6-Cl	H	OEt	n-Pentyl
1.151	H	7-Cl	OEt	n-Pentyl
1.152	6-Br	H	OEt	Cyclopentyl
1.153	6-Br	H	OEt	n-Hexyl
1.154	6-Br	H	OEt	Cyclohexyl
1.155	6-Br	H	OEt	Phenyl
1.156	6-Br	H	OEt	4-Chloro-phenyl
1.157	6-Cl	H	OEt	4-Chloro-phenyl
1.158	6-Br	H	OEt	4-Phenoxy-phenyl
1.159	6-Br	H	O-n-Propyl	Me

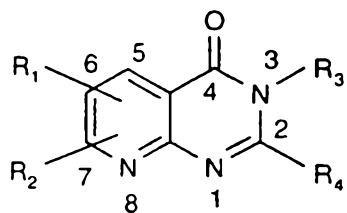
1.160	6-Cl	H	O-n-Propyl	Et
1.161	6-Br	H	O-n-Propyl	n-Propyl
1.162	6-Cl	H	O-n-Propyl	n-Propyl
1.163	H	7-Cl	O-n-Propyl	n-Propyl
1.164	H	7-I	O-n-Propyl	n-Propyl
1.165	6-Br	H	O-n-Propyl	
1.166	6-Cl	H	O-n-Propyl	
1.167	H	7-Cl	O-n-Propyl	
1.168	6-Br	H	O-n-Propyl	n-Butyl
1.169	6-Cl	H	O-n-Propyl	n-Butyl
1.170	H	7-Cl	O-n-Propyl	n-Butyl
1.171	H	7-I	O-n-Propyl	n-Butyl
1.172	6-Br	H	O-n-Propyl	i-Butyl
1.173	6-Cl	H	O-n-Propyl	i-Butyl
1.174	6-Br	H	O-n-Propyl	
1.175	6-Cl	H	O-n-Propyl	
1.176	H	7-Cl	O-n-Propyl	
1.177	6-Br	H	O-n-Propyl	Me 
1.178	6-Cl	H	O-n-Propyl	Me 
1.179	6-Br	H	O-n-Propyl	Cyclobutyl
1.180	6-Br	H	O-n-Propyl	n-Pentyl
1.181	6-Cl	H	O-n-Propyl	n-Pentyl
1.182	H	7-Cl	O-n-Propyl	n-Pentyl





1.183	6-Br	H	O-n-Propyl	Cyclopentyl
1.184	6-Br	H	O-n-Prpyl	n-Hexyl
1.185	6-Br	H	O-n-Prpyl	Cyclohexyl
1.186	6-Br	H	O-n-Prpyl	Phenyl
1.187	6-Br	H	O-n-Prpyl	4-Chloro-phenyl
1.188	6-Cl	H	O-n-Prpyl	4-Chloro-phenyl
1.189	6-Br	H	O-n-Prpyl	4-Phenoxy-phenyl
1.190	6-Br	H	Et	CH ₂ OMe
1.191	6-Cl	H	Et	CH ₂ OMe
1.192	6-Br	H	n-Prpyl	CH ₂ OMe
1.193	6-Cl	H	n-Prpyl	CH ₂ OMe
1.194	H	7-Cl	n-Prpyl	CH ₂ OMe
1.195	6-Br	H	n-Butyl	CH ₂ OMe
1.196	6-Cl	H	n-Butyl	CH ₂ OMe
1.197	6-Br	H		CH ₂ OMe
1.198	6-Br	H	n-Pentyl	CH ₂ OMe
1.199	6-Br	H	Et	CH ₂ OEt
1.200	6-Cl	H	Et	CH ₂ OEt
1.201	6-Br	H	n-Propyl	CH ₂ OEt
1.202	6-Cl	H	n-Propyl	CH ₂ OEt
1.203	H	7-Cl	n-Propyl	CH ₂ OEt
1.204	6-Br	H	n-Butyl	CH ₂ OEt
1.205	6-Cl	H	n-Butyl	CH ₂ OEt
1.206	6-Br	H		CH ₂ OEt
1.207	6-Br	H	n-Pentyl	CH ₂ OEt
1.208	6-Br	H	n-Propyl	CH ₂ CN
1.209	6-Br	H	n-Butyl	CH ₂ CN
1.210	6-Br	H	n-Propyl	t-Butyl
1.211	6-Br	H	n-Propyl	t-Butyl



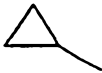
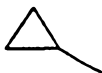
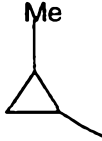
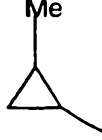
- 34 -




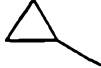
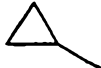

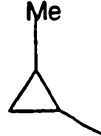
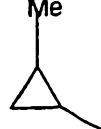
1.212	6-Cl	H	n-Propyl	CF ₃
1.213	6-Br	H	n-Propyl	CF ₃
1.214	6-Br	H	n-Butyl	CF ₃







Tabelle 2: A = Pyridyl



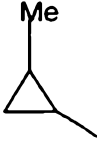
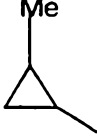






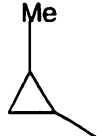
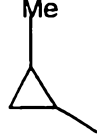
cmpd. No.	R ₁	R ₂	R ₃	R ₄	phys. data m.p.°C
2.1	6-Br	H	Me	Me	
2.2	6-Cl	H	Me	Et	
2.3	6-Br	H	Me	n-Propyl	
2.4	6-Cl	H	Me	n-Propyl	
2.5	6-Br	H	Me	n-Butyl	
2.6	6-Cl	H	Me	n-Butyl	
2.7	6-Br	H	Me	i-Butyl	
2.8	6-Cl	H	Me	i-Butyl	
2.9	6-Br	H	Me	n-Pentyl	
2.10	6-Br	H	Me		
2.11	6-Cl	H	Me		
2.12	6-Br	H	Et	Me	
2.13	6-Cl	H	Et	Et	
2.14	6-Br	H	Et	n-Propyl	
2.15	6-Cl	H	Et	n-Propyl	
2.16	6-Br	H	Et	n-Butyl	
2.17	6-Cl	H	Et	n-Butyl	
2.18	6-Br	H	Et	i-Butyl	
2.19	6-Cl	H	Et	i-Butyl	
2.20	6-Br	H	Et	n-Pentyl	
2.21	6-Br	H	Et		
2.22	6-Cl	H	Et		
2.23	6-Br	H	n-Propyl	Me	





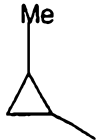
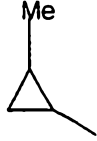
2.24	6-Cl	H	n-Propyl	Et	
2.25	6-Br	H	n-Propyl	n-Propyl	
2.26	6-Cl	H	n-Propyl	n-Propyl	
2.27	7-I	H	n-Propyl	n-Propyl	
2.28	6-Br	H	n-Propyl		
2.29	6-Cl	H	n-Propyl		
2.30	6-Br	H	n-Propyl	n-Butyl	Oil, ¹ H-NMR
2.31	6-Cl	H	n-Propyl	n-Butyl	
2.32	6-I	H	n-Propyl	n-Butyl	
2.33	6-Br	H	n-Propyl	i-Butyl	
2.34	6-Cl	H	n-Propyl	i-Butyl	
2.35	6-Br	H	n-Propyl		
2.36	6-Cl	H	n-Propyl		
2.37	6-Br	H	n-Propyl	Me 	
2.38	6-Cl	H	n-Propyl	Me 	
2.39	6-Br	H	n-Propyl	Cyclobutyl	
2.40	6-Br	H	n-Propyl	n-Pentyl	
2.41	6-Cl	H	n-Propyl	n-Pentyl	
2.42	6-Br	H	n-Propyl	Cyclopentyl	
2.43	6-Br	H	n-Propyl	n-Hexyl	
2.44	6-Br	H	n-Propyl	Cyclohexyl	
2.45	6-Br	H	n-Propyl	Phenyl	
2.46	6-Br	H	n-Propyl	4-Chloro-phenyl	

2.47	6-Cl	H	n-Propyl	4-Chloro-phenyl
2.48	6-Br	H	n-Propyl	4-Phenoxy-phenyl
2.49	6-Br	H	n-Butyl	Me
2.50	6-Cl	H	n-Butyl	Et
2.51	6-Br	H	n-Butyl	n-Propyl
2.52	6-Cl	H	n-Butyl	n-Propyl
2.53	H	7-Cl	n-Butyl	n-Propyl
2.54	H	7-I	n-Butyl	n-Propyl
2.55	6-Br	H	n-Butyl	
2.56	6-Cl	H	n-Butyl	
2.57	H	7-Cl	n-Butyl	
2.58	6-Br	H	n-Butyl	n-Butyl
2.59	6-Cl	H	n-Butyl	n-Butyl
2.60	6-I	H	n-Butyl	n-Butyl
2.61	6-Br	H	n-Butyl	i-Butyl
2.62	6-Cl	H	n-Butyl	i-Butyl
2.63	6-Br	H	n-Butyl	
2.64	6-Cl	H	n-Butyl	
2.65	6-I	H	n-Butyl	
2.66	6-Br	H	n-Butyl	Me 
2.67	6-Cl	H	n-Butyl	Me 

2.68	6-Br	H	n-Butyl	Cyclobutyl
2.69	6-Br	H	n-Butyl	n-Pentyl
2.70	6-Cl	H	n-Butyl	n-Pentyl
2.71	6-Br	H	n-Butyl	Cyclopentyl
2.72	6-Br	H	n-Butyl	n-Hexyl
2.73	6-Br	H	n-Butyl	Cyclohexyl
2.74	6-Br	H	n-Butyl	Phenyl
2.75	6-Br	H	n-Butyl	4-Chloro-phenyl
2.76	6-Cl	H	n-Butyl	4-Chloro-phenyl
2.77	6-Br	H	n-Butyl	4-Phenoxy-phenyl
2.78	6-Br	H	i-Butyl	n-Propyl
2.79	6-Cl	H	i-Butyl	n-Prpyl
2.80	6-Br	H	i-Butyl	n-Butyl
2.81	6-Cl	H	i-Butyl	n-Butyl
2.82	6-Br	H		n-Propyl
2.83	6-Cl	H		n-Propyl
2.84	6-Br	H		n-Propyl
2.85	6-Br	H		n-Butyl
2.86	6-Br	H	n-Pentyl	Me
2.87	6-Cl	H	n-Pentyl	Et
2.88	6-Br	H	n-Pentyl	n-Propyl
2.89	6-Cl	H	n-Pentyl	n-Propyl
2.90	6-Br	H	n-Pentyl	
2.91	6-Cl	H	n-Pentyl	
2.92	6-Br	H	n-Pentyl	n-Butyl

2.93	6-Cl	H	n-Pentyl	n-Butyl
2.94	6-I	H	n-Pentyl	n-Butyl
2.95	6-Br	H	n-Pentyl	i-Butyl
2.96	6-Cl	H	n-Pentyl	i-Butyl
2.97	6-Br	H	n-Pentyl	
2.98	6-Cl	H	n-Pentyl	
2.99	6-Br	H	n-Pentyl	Me 
2.100	6-Cl	H	n-Pentyl	Me 
2.101	6-Br	H	n-Pentyl	Cyclobutyl
2.102	6-Br	H	n-Pentyl	n-Pentyl
2.103	6-Cl	H	n-Pentyl	n-Pentyl
2.104	6-Br	H	n-Pentyl	Cyclopentyl
2.105	6-Br	H	n-Pentyl	n-Hexyl
2.106	6-Br	H	n-Pentyl	Cyclohexyl
2.107	6-Br	H	n-Pentyl	Phenyl
2.108	6-Br	H	n-Pentyl	4-Chloro-phenyl
2.109	6-Cl	H	n-Pentyl	4-Chloro-phenyl
2.110	6-Br	H	n-Pentyl	4-Phenoxy-phenyl
2.111	6-Br	H	OEt	Me
2.112	6-Cl	H	OEt	Et
2.113	6-Br	H	OEt	n-Propyl
2.114	6-Cl	H	OEt	n-Propyl

2.115	6-Br	H	OEt	
2.116	6-Cl	H	OEt	
2.117	6-Br	H	OEt	n-Butyl
2.118	6-Cl	H	OEt	n-Butyl
2.119	6-I	H	OEt	n-Butyl
2.120	6-Br	H	OEt	i-Butyl
2.121	6-Cl	H	OEt	i-Butyl
2.122	6-Br	H	OEt	
2.123	6-Cl	H	OEt	
2.124	6-Br	H	OEt	Me 
2.125	6-Cl	H	OEt	Me 
2.126	6-Br	H	OEt	Cyclobutyl
2.127	6-Br	H	OEt	n-Pentyl
2.128	6-Cl	H	OEt	n-Pentyl
2.129	6-Br	H	OEt	Cyclopentyl
2.130	6-Br	H	OEt	n-Hexyl
2.131	6-Br	H	OEt	Cyclohexyl
2.132	6-Br	H	OEt	Phenyl
2.133	6-Br	H	OEt	4-Chloro-phenyl
2.134	6-Cl	H	OEt	4-Chloro-phenyl
2.135	6-Br	H	OEt	4-Phenoxy-phenyl
2.136	6-Br	H	O-n-Propyl	Me

2.137	6-Cl	H	O-n-Propyl	Et
2.138	6-Br	H	O-n-Propyl	n-Propyl
2.139	6-Cl	H	O-n-Propyl	n-Propyl
2.140	6-Br	H	O-n-Propyl	
2.141	6-Cl	H	O-n-Propyl	
2.142	6-Br	H	O-n-Propyl	n-Butyl
2.143	6-Cl	H	O-n-Propyl	n-Butyl
2.144	6-Br	H	O-n-Propyl	i-Butyl
2.145	6-Cl	H	O-n-Propyl	i-Butyl
2.146	6-Br	H	O-n-Propyl	
2.147	6-Cl	H	O-n-Propyl	
2.148	6-Br	H	O-n-Propyl	Me 
2.149	6-Cl	H	O-n-Propyl	Me 
2.150	6-Br	H	O-n-Propyl	Cyclobutyl
2.151	6-Br	H	O-n-Propyl	n-Pentyl
2.152	6-Cl	H	O-n-Propyl	n-Pentyl
2.153	6-Br	H	O-n-Propyl	Cyclopentyl
2.154	6-Br	H	O-n-Propyl	n-Hexyl
2.155	6-Br	H	O-n-Propyl	Cyclohexyl
2.156	6-Br	H	O-n-Propyl	Phenyl
2.157	6-Br	H	O-n-Propyl	4-Chloro-phenyl
2.158	6-Cl	H	O-n-Propyl	4-Chloro-phenyl



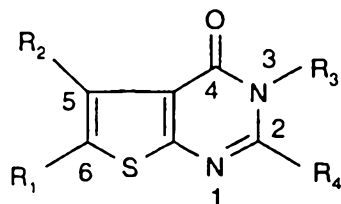
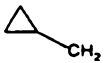
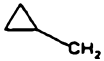
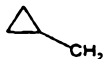



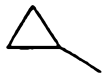
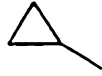
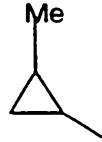
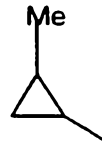



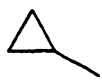
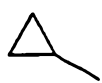


2.159	6-Br	H	O-n-Propyl	4-Phenoxy-phenyl
2.160	6-Br	H	Et	CH ₂ OMe
2.161	6-Cl	H	Et	CH ₂ OMe
2.162	6-Br	H	n-Propyl	CH ₂ OMe
2.163	6-Cl	H	n-Propyl	CH ₂ OMe
2.164	6-Br	H	n-Butyl	CH ₂ OMe
2.165	6-Cl	H	n-Butyl	CH ₂ OMe
2.166	6-Br	H		CH ₂ OMe
2.167	6-Br	H	n-Pentyl	CH ₂ OMe
2.168	6-Br	H	Et	CH ₂ OEt
2.169	6-Cl	H	Et	CH ₂ OEt
2.170	6-Br	H	n-Propyl	CH ₂ OEt
2.171	6-Cl	H	n-Propyl	CH ₂ OEt
2.172	6-Br	H	n-Butyl	CH ₂ OEt
2.173	6-Cl	H	n-Butyl	CH ₂ OEt
2.174	6-Br	H		CH ₂ OEt
2.175	6-Br	H	n-Pentyl	CH ₂ OEt
2.176	6-Br	H	n-Prpyl	CH ₂ CN
2.177	6-Br	H	n-Butyl	CH ₂ CN
2.178	6-Br	H	n-Propyl	t-Butyl
2.179	6-Br	H	n-Propyl	t-Butyl
2.180	6-Br	H	n-Propyl	CF ₃
2.181	6-Br	H	n-Butyl	CF ₃

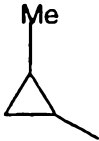




Tabelle 3: A = Thienyl[2.3-d]





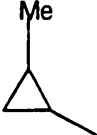
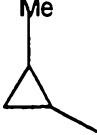






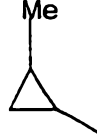

compd. No.	R ₁	R ₂	R ₃	R ₄	phys. data m.p. °C
3.1	Cl	H	Me	Me	139-141
3.2	Cl	H	Me	Et	
3.3	Br	H	Me	n-Propyl	
3.4	Cl	H	Me	n-Propyl	
3.5	Br	H	Me	n-Butyl	
3.6	Cl	H	Me	n-Butyl	63-65
3.7	Br	H	Me	i-Butyl	
3.8	Cl	H	Me	i-Butyl	87-89
3.9	Br	H	Me	n-Pentyl	
3.10	Br	H	Me		
3.11	Cl	H	Me		
3.12	Br	H	Et	Me	
3.13	Cl	H	Et	Et	
3.14	Br	H	Et	n-Propyl	
3.15	Cl	H	Et	n-Propyl	80-82
3.16	Br	H	Et	n-Butyl	
3.17	Cl	H	Et	n-Butyl	Oil, ¹ H-NMR
3.18	Br	H	Et	i-Butyl	
3.19	Cl	H	Et	i-Butyl	
3.20	Br	H	Et	n-Pentyl	
3.21	Br	H	Et		



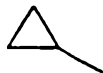

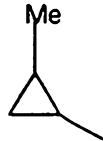
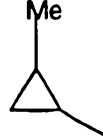
3.22	Cl	H	Et		
3.23	Br	H	n-Propyl	Me	
3.24	Cl	H	n-Propyl	Et	
3.25	Br	H	n-Propyl	n-Propyl	
3.26	Cl	H	n-Propyl	n-Propyl	74-76
3.27	I	H	n-Propyl	n-Propyl	
3.28	Br	H	n-Propyl		
3.29	Cl	H	n-Propyl		
3.30	Br	H	n-Propyl	n-Butyl	63-66
3.31	Cl	H	n-Propyl	n-Butyl	67-69
3.32	I	H	n-Propyl	n-Butyl	
3.33	Br	H	n-Propyl	i-Butyl	
3.34	Cl	H	n-Propyl	i-Butyl	Oil, ¹ H-NMR
3.35	Br	H	n-Propyl		
3.36	Cl	H	n-Propyl		Oil, ¹ H-NMR
3.37	Br	H	n-Propyl	Me 	
3.38	Cl	H	n-Propyl	Me 	
3.39	Br	H	n-Propyl	Cyclobutyl	
3.40	Br	H	n-Propyl	n-Pentyl	
3.41	Cl	H	n-Propyl	n-Pentyl	
3.42	Br	H	n-Propyl	Cyclopentyl	
3.43	Br	H	n-Propyl	n-Hexyl	
3.44	Br	H	n-Propyl	Cyclohexyl	
3.45	Br	H	n-Propyl	Phenyl	

3.46	Br	H	n-Propyl	4-Chloro-phenyl	
3.47	Cl	H	n-Propyl	4-Chloro-phenyl	126-128
3.48	Br	H	n-Propyl	4-Phenoxy-phenyl	
3.49	Br	H	n-Butyl	Me	
3.50	Cl	H	n-Butyl	Et	
3.51	Br	H	n-Butyl	n-Propyl	
3.52	Cl	H	n-Butyl	n-Propyl	Oil
3.53	I	H	n-Butyl	n-Propyl	
3.54	I	H	n-Butyl	n-Propyl	
3.55	Br	H	n-Butyl		
3.56	Cl	H	n-Butyl		54-56
3.57	I	H	n-Butyl		
3.58	Br	H	n-Butyl	n-Butyl	Oil
3.59	Cl	H	n-Butyl	n-Butyl	57-58
3.60	I	H	n-Butyl	n-Butyl	
3.61	Br	H	n-Butyl	i-Butyl	
3.62	Cl	H	n-Butyl	i-Butyl	
3.63	Br	H	n-Butyl		
3.64	Cl	H	n-Butyl		
3.65	I	H	n-Butyl		
3.66	Br	H	n-Butyl	Me 	

3.67	Cl	H	n-Butyl		
3.68	Br	H	n-Butyl	Cyclobutyl	
3.69	Br	H	n-Butyl	n-Pentyl	
3.70	Cl	H	n-Butyl	n-Pentyl	
3.71	Br	H	n-Butyl	Cyclopentyl	
3.72	Br	H	n-Butyl	n-Hexyl	
3.73	Br	H	n-Butyl	Cyclohexyl	
3.74	Cl	H	n-Butyl	Phenyl	
3.75	Br	H	n-Butyl	4-Chloro-phenyl	
3.76	Cl	H	n-Butyl	4-Chloro-phenyl	
3.77	Br	H	n-Butyl	4-Phenoxy-phenyl	
3.78	Br	H	i-Butyl	n-Propyl	
3.79	Cl	H	i-Butyl	n-Propyl	Oil, ¹ H-NMR
3.80	Br	H	i-Butyl	n-Butyl	
3.81	Cl	H	i-Butyl	n-Butyl	
3.82	Br	H		n-Propyl	
3.83	Cl	H		n-Propyl	
3.84	Cl	H		n-Propyl	
3.85	Br	H		n-Butyl	
3.86	Br	H	n-Pentyl	Me	
3.87	Cl	H	n-Pentyl	Et	
3.88	Br	H	n-Pentyl	n-Propyl	
3.89	Cl	H	n-Pentyl	n-Propyl	

3.90	Br	H	n-Pentyl	
3.91	Cl	H	n-Pentyl	
3.92	Br	H	n-Pentyl	n-Butyl
3.93	Cl	H	n-Pentyl	n-Butyl
3.94	I	H	n-Pentyl	n-Butyl
3.95	Br	H	n-Pentyl	i-Butyl
3.96	Cl	H	n-Pentyl	i-Butyl
3.97	Br	H	n-Pentyl	
3.98	Cl	H	n-Pentyl	
3.99	Br	H	n-Pentyl	
3.100	Cl	H	n-Pentyl	
3.101	Cl	H	n-Pentyl	Cyclobutyl
3.102	Br	H	n-Pentyl	n-Pentyl
3.103	Cl	H	n-Pentyl	n-Pentyl
3.104	Cl	H	n-Pentyl	Cyclopentyl
3.105	Br	H	n-Pentyl	n-Hexyl
3.106	Cl	H	n-Pentyl	Cyclohexyl
3.107	Br	H	n-Pentyl	Phenyl
3.108	Br	H	n-Pentyl	4-Chloro-phenyl
3.109	Cl	H	n-Pentyl	4-Chloro-phenyl
3.110	Br	H	n-Pentyl	4-Phenoxy-phenyl
3.111	Br	H	OEt	Me

3.112	Cl	H	OEt	Et	
3.113	Br	H	OEt	n-Propyl	
3.114	Cl	H	OEt	n-Propyl	
3.115	Br	H	OEt		
3.116	Cl	H	OEt		
3.117	Br	H	OEt	n-Butyl	75-77
3.118	Cl	H	OEt	n-Butyl	66-69
3.119	I	H	OEt	n-Butyl	
3.120	Br	H	OEt	i-Butyl	
3.121	Cl	H	OEt	i-Butyl	
3.122	Br	H	OEt		
3.123	Cl	H	OEt		
3.124	Br	H	OEt	Me 	
3.125	Cl	H	OEt	Me 	
3.126	Br	H	OEt	Cyclobutyl	
3.127	Br	H	OEt	n-Pentyl	
3.128	Cl	H	OEt	n-Pentyl	
3.129	Br	H	OEt	Cyclopentyl	
3.130	Br	H	OEt	n-Hexyl	
3.131	Br	H	OEt	Cyclohexyl	
3.132	Br	H	OEt	Phenyl	
3.133	Br	H	OEt	4-Chloro-phenyl	
3.134	Cl	H	OEt	4-Chloro-phenyl	

3.135	Cl	H	OEt	4-Phenoxy-phenyl
3.136	Br	H	O-n-Propyl	Me
3.137	Cl	H	O-n-Propyl	Et
3.138	Br	H	O-n-Propyl	n-Propyl
3.139	Cl	H	O-n-Propyl	n-Propyl
3.140	Br	H	O-n-Propyl	
3.141	Cl	H	O-n-Propyl	
3.142	Br	H	O-n-Propyl	n-Butyl
3.143	Cl	H	O-n-Propyl	n-Butyl
3.144	Br	H	O-n-Propyl	i-Butyl
3.145	Cl	H	O-n-Propyl	i-Butyl
3.146	Br	H	O-n-Propyl	
3.147	Cl	H	O-n-Propyl	
3.148	Br	H	O-n-Propyl	Me 
3.149	Cl	H	O-n-Propyl	Me 
3.150	Br	H	O-n-Propyl	Cyclobutyl
3.151	Br	H	O-n-Propyl	n-Pentyl
3.152	Cl	H	O-n-Propyl	n-Pentyl
3.153	Br	H	O-n-Propyl	Cyclopentyl
3.154	Cl	H	O-n-Propyl	n-Hexyl
3.155	Br	H	O-n-Propyl	Cyclohexyl
3.156	Cl	H	O-n-Propyl	Phenyl
3.157	Br	H	O-n-Propyl	4-Chloro-phenyl



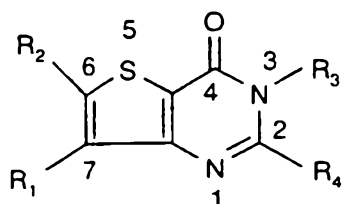









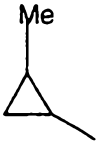



3.158	Cl	H	O-n-Propyl	4-Chloro-phenyl	
3.159	Br	H	O-n-Propyl	4-Phenoxy-phenyl	
3.160	Br	H	Et	CH ₂ OMe	
3.161	Cl	H	Et	CH ₂ OMe	
3.162	Br	H	n-Propyl	CH ₂ OMe	
3.163	Cl	H	n-Propyl	CH ₂ OMe	Oil, ¹ H-NMR
3.164	Br	H	n-Butyl	CH ₂ OMe	
3.165	Cl	H	n-Butyl	CH ₂ OMe	
3.166	Br	H		CH ₂ OMe	
3.167	Br	H	n-Pentyl	CH ₂ OMe	
3.168	Br	H	Et	CH ₂ OEt	
3.169	Cl	H	Et	CH ₂ OEt	
3.170	Br	H	n-Propyl	CH ₂ OEt	
3.171	Cl	H	n-Propyl	CH ₂ OEt	40-41
3.172	Br	H	n-Butyl	CH ₂ OEt	
3.173	Cl	H	n-Butyl	CH ₂ OEt	
3.174	Br	H		CH ₂ OEt	
3.175	Br	H	n-Pentyl	CH ₂ OEt	
3.176	Br	H	n-Propyl	CH ₂ CN	
3.177	Cl	H	n-Butyl	CH ₂ CN	
3.178	Br	H	n-Propyl	t-Butyl	
3.179	Cl	H	n-Propyl	t-Butyl	
3.180	Br	H	n-Propyl	CF ₃	
3.181	Cl	H	n-Butyl	CF ₃	
3.182	Cl	H	n-Pentyl	CF ₃	
3.183	Cl	Cl	n-Propyl	n-Propyl	
3.184	Cl	Cl	n-Propyl	n-Butyl	
3.185	Br	Br	n-Propyl	n-Butyl	
3.186	Br	Br	n-Butyl	n-Butyl	






Tabelle 4: A=Thienyl[3.2-d]













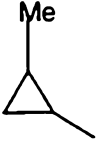
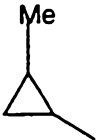



Cmpd. No.	R ₁	R ₂	R ₃	R ₄	phys. data
4.1	Br	H	Me	Me	
4.2	H	Cl	Me	Et	
4.3	Br	H	Me	n-Propyl	
4.4	H	Cl	Me	n-Propyl	
4.5	H	Cl	Me	n-Propyl	
4.6	Br	H	Me	n-Butyl	
4.7	H	Cl	Me	n-Butyl	
4.8	H	Cl	Me	n-Butyl	
4.9	Br	H	Me	i-Butyl	
4.10	H	Cl	Me	i-Butyl	
4.11	Br	H	Me	n-Pentyl	
4.12	Br	H	Me		
4.13	H	Cl	Me		
4.14	Br	H	Et	Me	
4.15	H	Cl	Et	Et	
4.16	Br	H	Et	n-Propyl	
4.17	H	Cl	Et	n-Propyl	
4.18	H	Cl	Et	n-Propyl	
4.19	Br	H	Et	n-Butyl	
4.20	H	Cl	Et	n-Butyl	
4.21	H	Cl	Et	n-Butyl	
4.22	Br	H	Et	i-Butyl	

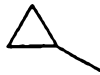
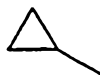
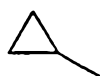

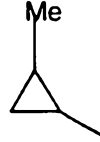
4.23	H	Cl	Et	i-Butyl	
4.24	Br	H	Et	n-Pentyl	
4.25	Br	H	Et		
4.26	H	Cl	Et		
4.27	Br	H	n-Propyl	Me	
4.28	H	Cl	n-Propyl	Et	
4.29	Br	H	n-Propyl	n-Propyl	
4.30	H	Cl	n-Propyl	n-Propyl	
4.31	H	Cl	n-Propyl	n-Propyl	
4.32	H	I	n-Propyl	n-Propyl	
4.33	Br	H	n-Propyl		
4.34	H	Cl	n-Propyl		
4.35	H	Cl	n-Propyl		
4.36	Br	H	n-Propyl	n-Butyl	120-121
4.37	H	Cl	n-Propyl	n-Butyl	
4.38	H	Cl	n-Propyl	n-Butyl	
4.39	H	I	n-Propyl	n-Butyl	
4.40	Br	H	n-Propyl	i-Butyl	
4.41	H	Cl	n-Propyl	i-Butyl	
4.42	Br	H	n-Propyl		
4.43	H	Cl	n-Propyl		
4.44	H	Cl	n-Propyl		
4.45	H	Cl	n-Propyl	Me 	









4.46	H	Cl	n-Propyl	
4.47	Br	H	n-Propyl	Cyclobutyl
4.48	Br	H	n-Propyl	n-Pentyl
4.49	H	Cl	n-Propyl	n-Pentyl
4.50	H	Cl	n-Propyl	n-Pentyl
4.51	Br	H	n-Propyl	Cyclopentyl
4.52	Br	H	n-Propyl	n-Hexyl
4.53	Br	H	n-Propyl	Cyclohexyl
4.54	Br	H	n-Propyl	Phenyl
4.55	Br	H	n-Propyl	4-Chloro-phenyl
4.55	H	Cl	n-Propyl	4-Chloro-phenyl
4.56	Br	H	n-Propyl	4-Phenoxy-phenyl
4.57	Br	H	n-Butyl	Me
4.58	H	Cl	n-Butyl	Et
4.59	Br	H	n-Butyl	n-Propyl
4.60	H	Cl	n-Butyl	n-Propyl
4.61	H	Cl	n-Butyl	n-Propyl
4.62	H	I	n-Butyl	n-Propyl
4.62	Br	H	n-Butyl	
4.63	H	Cl	n-Butyl	
4.64	H	Cl	n-Butyl	
4.65	Br	H	n-Butyl	n-Butyl
4.66	H	Cl	n-Butyl	n-Butyl
4.67	H	Cl	n-Butyl	n-Butyl
4.68	H	I	n-Butyl	n-Butyl
4.69	Br	H	n-Butyl	i-Butyl



4.70	H	Cl	n-Butyl	i-Butyl
4.71	Br	H	n-Butyl	
4.72	H	Cl	n-Butyl	
4.73	H	Cl	n-Butyl	
4.74	Br	H	n-Butyl	
4.75	H	Cl	n-Butyl	
4.76	Br	H	n-Butyl	Cyclobutyl
4.77	Br	H	n-Butyl	n-Pentyl
4.78	H	Cl	n-Butyl	n-Pentyl
4.79	H	Cl	n-Butyl	n-Pentyl
4.80	Br	H	n-Butyl	Cyclopentyl
4.81	Br	H	n-Butyl	n-Hexyl
4.82	Br	H	n-Butyl	Cyclohexyl
4.83	Br	H	n-Butyl	Phenyl
4.84	Br	H	n-Butyl	4-Chloro-phenyl
4.85	H	Cl	n-Butyl	4-Chloro-phenyl
4.86	Br	H	n-Butyl	4-Phenoxy-phenyl
4.87	Br	H	i-Butyl	n-Propyl
4.88	H	Cl	i-Butyl	n-Propyl
4.89	Br	H	i-Butyl	n-Butyl
4.90	H	Cl	i-Butyl	n-Butyl

4.91	Br	H		n-Propyl
4.92	H	Cl		n-Propyl
4.93	Br	H		n-Propyl
4.94	Br	H		n-Butyl
4.95	Br	H	n-Pentyl	Me
4.96	H	Cl	n-Pentyl	Et
4.97	Br	H	n-Pentyl	n-Propyl
4.98	Cl	H	n-Pentyl	n-Propyl
4.99	H	Cl	n-Pentyl	n-Propyl
4.100	H	I	n-Pentyl	n-Propyl
4.101	Br	H	n-Pentyl	
4.102	H	Cl	n-Pentyl	
4.103	H	Cl	n-Pentyl	
4.104	Br	H	n-Pentyl	n-Butyl
4.105	H	Cl	n-Pentyl	n-Butyl
4.106	H	Cl	n-Pentyl	n-Butyl
4.107	H	I	n-Pentyl	n-Butyl
4.108	Br	H	n-Pentyl	i-Butyl
4.109	H	Cl	n-Pentyl	i-Butyl
4.110	Br	H	n-Pentyl	
4.111	H	Cl	n-Pentyl	
4.112	H	Cl	n-Pentyl	

4.113	Br	H	n-Pentyl	
4.114	H	Cl	n-Pentyl	
4.115	Br	H	n-Pentyl	Cyclobutyl
4.116	Br	H	n-Pentyl	n-Pentyl
4.117	Cl	Cl	n-Pentyl	n-Pentyl
4.118	H	Cl	n-Pentyl	n-Pentyl
4.119	Br	H	n-Pentyl	Cyclopentyl
4.120	Br	H	n-Pentyl	n-Hexyl
4.121	Br	H	n-Pentyl	Cyclohexyl
4.122	Br	H	n-Pentyl	Phenyl
4.123	Br	H	n-Pentyl	4-Chloro-phenyl
4.124	H	Cl	n-Pentyl	4-Chloro-phenyl
4.125	Br	H	n-Pentyl	4-Phenoxy-phenyl
4.126	Br	H	OEt	Me
4.127	Cl	H	OEt	Et
4.128	Br	H	OEt	n-Propyl
4.129	H	Cl	OEt	n-Propyl
4.130	H	Cl	OEt	n-Propyl
4.131	H	I	OEt	n-Propyl
4.132	Br	H	OEt	
4.133	H	Cl	OEt	
4.134	H	Cl	OEt	
4.135	Br	H	OEt	n-Butyl

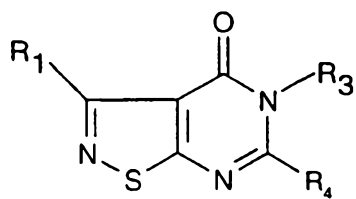
4.136	H	Cl	OEt	n-Butyl
4.137	H	Cl	OEt	n-Butyl
4.138	H	I	OEt	n-Butyl
4.139	Br	H	OEt	i-Butyl
4.140	H	Cl	OEt	i-Butyl
4.141	Br	H	OEt	
4.142	H	Cl	OEt	
4.143	H	Cl	OEt	
4.144	Br	H	OEt	
4.145	H	Cl	OEt	
4.146	Br	H	OEt	Cyclobutyl
4.147	Br	H	OEt	n-Pentyl
4.148	H	Cl	OEt	n-Pentyl
4.149	H	Cl	OEt	n-Pentyl
4.150	Br	H	OEt	Cyclopentyl
4.151	Br	H	OEt	n-Hexyl
4.152	Br	H	OEt	Cyclohexyl
4.153	Br	H	OEt	Phenyl
4.154	Br	H	OEt	4-Chloro-phenyl
4.155		Cl	OEt	4-Chloro-phenyl
4.156	Br	H	OEt	4-Phenoxy-phenyl
4.157	Br	H	O-n-Propyl	Me

4.158	H	Cl	O-n-Propyl	Et
4.159	Br	H	O-n-Propyl	n-Propyl
4.160	H	Cl	O-n-Propyl	n-Propyl
4.161	H	Cl	O-n-Propyl	n-Propyl
4.162	H	I	O-n-Propyl	n-Propyl
4.163	Br	H	O-n-Propyl	
4.164	H	Cl	O-n-Propyl	
4.165	H	Cl	O-n-Propyl	
4.166	Br	H	O-n-Propyl	n-Butyl
4.167	H	Cl	O-n-Propyl	n-Butyl
4.168	H	Cl	O-n-Propyl	n-Butyl
4.169	H	II	O-n-Propyl	n-Butyl
4.170	Br	H	O-n-Propyl	i-Butyl
4.171	H	Cl	O-n-Propyl	i-Butyl
4.172	Br	H	O-n-Propyl	
4.173	H	Cl	O-n-Propyl	
4.174	H	Cl	O-n-Propyl	
4.175	Br	H	O-n-Propyl	Me 
4.176	H	Cl	O-n-Propyl	Me 
4.177	Br	H	O-n-Propyl	Cyclobutyl
4.178	Br	H	O-n-Propyl	n-Pentyl
4.179	H	Cl	O-n-Propyl	n-Pentyl
4.180	H	Cl	O-n-Propyl	n-Pentyl

4.181	Br	H	O-n-Propyl	Cyclopentyl
4.182	Br	H	O-n-Propyl	n-Hexyl
4.183	Br	H	O-n-Propyl	Cyclohexyl
4.184	Br	H	O-n-Propyl	Phenyl
4.185	Br	H	O-n-Propyl	4-Chloro-phenyl
4.186	H	Cl	O-n-Propyl	4-Chloro-phenyl
4.187	Br	H	O-n-Propyl	4-Phenoxy-phenyl
4.188	Br	H	Et	CH ₂ OMe
4.189	H	Cl	Et	CH ₂ OMe
4.190	Br	H	n-Propyl	CH ₂ OMe
4.191	H	Cl	n-Propyl	CH ₂ OMe
4.192	H	Cl	n-Propyl	CH ₂ OMe
4.193	Br	H	n-Butyl	CH ₂ OMe
4.194	H	Cl	n-Butyl	CH ₂ OMe
4.195	Br	H		CH ₂ OMe
4.195	Br	H	n-Pentyl	CH ₂ OMe
4.196	Br	H	Et	CH ₂ OEt
4.197	H	Cl	Et	CH ₂ OEt
4.198	Br	H	n-Propyl	CH ₂ OEt
4.199	H	Cl	n-Propyl	CH ₂ OEt
4.200	H	Cl	n-Propyl	CH ₂ OEt
4.201	Br	H	n-Butyl	CH ₂ OEt
4.202	H	Cl	n-Butyl	CH ₂ OEt
4.203	Br	H		CH ₂ OEt
4.204	Br	H	n-Pentyl	CH ₂ OEt
4.205	Br	H	n-Propyl	CH ₂ CN
4.206	Br	H	n-Butyl	CH ₂ CN
4.207	Br	H	n-Propyl	t-Butyl
4.208	Br	H	n-Propyl	t-Butyl

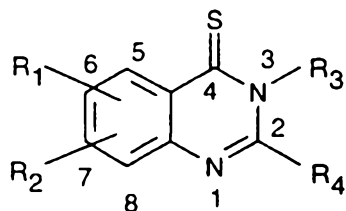
4.209	H	Cl	n-Propyl	CF ₃
4.210	Br	H	n-Propyl	CF ₃
4.211	H	Cl	n-Butyl	CF ₃
4.212	Br	H	n-Butyl	CF ₃
4.213	Cl	Cl	n-Propyl	n-Propyl
4.214	Cl	Cl	n-Propyl	n-Butyl
4.215	Br	Br	n-Propyl	n-Butyl
4.216	Br	Br	n-Butyl	n-Butyl



Tabelle 5: A= Thiazolyl












Cmpd. No.	R ₁	R ₃	R ₄	phys. data
5.1	H	Et	n-Propyl	
5.2	H	n-Propyl	n-Propyl	
5.3	H	n-Propyl	n-Butyl	
5.4	H	n-Butyl	n-Butyl	
5.5	Me	n-Propyl	n-Propyl	
5.6	Me	n-Propyl	n-Butyl	
5.7	Me	n-Butyl	n-Butyl	
5.8	H	n-Propyl	Phenyl	

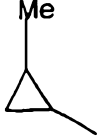



Table 6: A = Phenyl




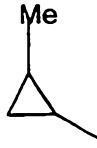













compd. No.	R ₁	R ₂	R ₃	R ₄	phys. data m.p. °C
6.1	6-Br	H	Me	Me	
6.2	6-Cl	H	Me	Et	
6.3	6-Br	H	Me	n-Propyl	
6.4	6-Cl	H	Me	n-Propyl	
6.5	H	7-Cl	Me	n-Propyl	
6.6	6-Br	H	Me	n-Butyl	
6.7	6-Cl	H	Me	n-Butyl	
6.8	H	7-Cl	Me	n-Butyl	
6.9	6-Br	H	Me	i-Butyl	
6.10	6-Cl	H	Me	i-Butyl	
6.11	6-Br	H	Me	n-Pentyl	
6.12	6-Br	H	Me		
6.13	6-Cl	H	Me		
6.14	6-Br	H	Et	Me	
6.15	6-Cl	H	Et	Et	
6.16	6-Br	H	Et	n-Propyl	
6.17	6-Cl	H	Et	n-Propyl	
6.18	H	7-Cl	Et	n-Propyl	
6.19	6-Br	H	Et	n-Butyl	
6.20	6-Cl	H	Et	n-Butyl	
6.21	H	7-Cl	Et	n-Butyl	
6.22	6-Br	H	Et	i-Butyl	

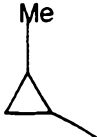
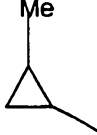



6.23	6-Cl	H	Et	i-Butyl
6.24	6-Br	H	Et	n-Pentyl
6.25	6-Br	H	Et	
6.26	6-Cl	H	Et	
6.27	6-Br	H	n-Propyl	Me
6.28	6-Cl	H	n-Propyl	Et
6.29	6-Br	H	n-Propyl	n-Propyl
6.30	6-Cl	H	n-Propyl	n-Propyl
6.31	H	7-Cl	n-Propyl	n-Propyl
6.32	H	7-I	n-Propyl	n-Propyl
6.33	6-Br	H	n-Propyl	
6.34	6-Cl	H	n-Propyl	
6.35	H	7-Cl	n-Propyl	
6.36	6-Br	H	n-Propyl	n-Butyl
6.37	6-Cl	H	n-Propyl	n-Butyl
6.38	H	7-Cl	n-Propyl	n-Butyl
6.39	H	7-I	n-Propyl	n-Butyl
6.40	6-Br	H	n-Propyl	i-Butyl
6.41	6-Cl	H	n-Propyl	i-Butyl
6.42	6-Br	H	n-Propyl	
6.43	6-Cl	H	n-Propyl	
6.44	H	7-Cl	n-Propyl	
6.45	6-Br	H	n-Propyl	Me 




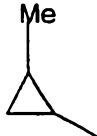
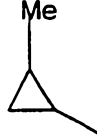
Oil, ¹H-NMR








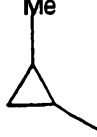
6.46	6-Cl	H	n-Propyl	
6.47	6-Br	H	n-Propyl	Cyclobutyl
6.48	6-Br	H	n-Propyl	n-Pentyl
6.49	6-Cl	H	n-Propyl	n-Pentyl
6.50	H	7-Cl	n-Propyl	n-Pentyl
6.51	6-Br	H	n-Propyl	Cyclopentyl
6.52	6-Br	H	n-Propyl	n-Hexyl
6.53	6-Br	H	n-Propyl	Cyclohexyl
6.54	6-Br	H	n-Propyl	Phenyl
6.55	6-Br	H	n-Propyl	4-Chloro-phenyl
6.56	6-Cl	H	n-Propyl	4-Chloro-phenyl
6.57	6-Br	H	n-Propyl	4-Phenoxy-phenyl
6.58	6-Br	H	n-Butyl	Me
6.59	6-Cl	H	n-Butyl	Et
6.60	6-Br	H	n-Butyl	n-Propyl
6.61	6-Cl	H	n-Butyl	n-Propyl
6.62	H	7-Cl	n-Butyl	n-Propyl
6.63	H	7-I	n-Butyl	n-Propyl
6.64	6-Br	H	n-Butyl	
6.65	6-Cl	H	n-Butyl	
6.66	H	7-Cl	n-Butyl	
6.67	6-Br	H	n-Butyl	n-Butyl
6.68	6-Cl	H	n-Butyl	n-Butyl
6.69	H	7-Cl	n-Butyl	n-Butyl
6.70	H	7-I	n-Butyl	n-Butyl
6.71	6-Br	H	n-Butyl	i-Butyl

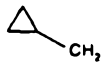
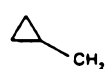
6.72	6-Cl	H	n-Butyl	i-Butyl
6.73	6-Br	H	n-Butyl	
6.74	6-Cl	H	n-Butyl	
6.75	H	7-Cl	n-Butyl	
6.76	6-Br	H	n-Butyl	Me 
6.77	6-Cl	H	n-Butyl	Me 
6.78	6-Br	H	n-Butyl	Cyclobutyl
6.79	6-Br	H	n-Butyl	n-Pentyl
6.80	6-Cl	H	n-Butyl	n-Pentyl
6.81	H	7-Cl	n-Butyl	n-Pentyl
6.82	6-Br	H	n-Butyl	Cyclopentyl
6.83	6-Br	H	n-Butyl	n-Hexyl
6.84	6-Br	H	n-Butyl	Cyclohexyl
6.85	6-Br	H	n-Butyl	Phenyl
6.86	6-Br	H	n-Butyl	4-Chloro-phenyl
6.87	6-Cl	H	n-Butyl	4-Chloro-phenyl
6.88	6-Br	H	n-Butyl	4-Phenoxy-phenyl
6.89	6-Br	H	i-Butyl	n-Propyl
6.90	6-Cl	H	i-Butyl	n-Propyl
6.91	6-Br	H	i-Butyl	n-Butyl
6.92	6-Cl	H	i-Butyl	n-Butyl

6.93	6-Br	H		n-Propyl
6.94	6-Cl	H		n-Propyl
6.95	6-Br	H		n-Propyl
6.96	6-Br	H		n-Butyl
6.97	6-Br	H	n-Pentyl	Me
6.98	6-Cl	H	n-Pentyl	Et
6.99	6-Br	H	n-Pentyl	n-Propyl
6.100	6-Cl	H	n-Pentyl	n-Propyl
6.101	H	7-Cl	n-Pentyl	n-Propyl
6.102	H	7-I	n-Pentyl	n-Propyl
6.103	6-Br	H	n-Pentyl	
6.104	6-Cl	H	n-Pentyl	
6.105	H	7-Cl	n-Pentyl	
6.106	6-Br	H	n-Pentyl	n-Butyl
6.107	6-Cl	H	n-Pentyl	n-Butyl
6.108	H	7-Cl	n-Pentyl	n-Butyl
6.109	H	7-I	n-Pentyl	n-Butyl
6.110	6-Br	H	n-Pentyl	i-Butyl
6.111	6-Cl	H	n-Pentyl	i-Butyl
6.112	6-Br	H	n-Pentyl	
6.113	6-Cl	H	n-Pentyl	
6.114	H	7-Cl	n-Pentyl	

6.115	6-Br	H	n-Pentyl	Me 
6.116	6-Cl	H	n-Pentyl	Me 
6.117	6-Br	H	n-Pentyl	Cyclobutyl
6.118	6-Br	H	n-Pentyl	n-Pentyl
6.119	6-Cl	H	n-Pentyl	n-Pentyl
6.120	H	7-Cl	n-Pentyl	n-Pentyl
6.121	6-Br	H	n-Pentyl	Cyclopentyl
6.122	6-Br	H	n-Pentyl	n-Hexyl
6.123	6-Br	H	n-Pentyl	Cyclohexyl
6.124	6-Br	H	n-Pentyl	Phenyl
6.125	6-Br	H	n-Pentyl	4-Chloro-phenyl
6.126	6-Cl	H	n-Pentyl	4-Chloro-phenyl
6.127	6-Br	H	n-Pentyl	4-Phenoxy-phenyl
6.128	6-Br	H	OEt	Me
6.129	6-Cl	H	OEt	Et
6.130	6-Br	H	OEt	n-Propyl
6.131	6-Cl	H	OEt	n-Propyl
6.132	H	7-Cl	OEt	n-Propyl
6.133	H	7-I	OEt	n-Propyl
6.134	6-Br	H	OEt	
6.135	6-Cl	H	OEt	
6.136	H	7-Cl	OEt	
6.137	6-Br	H	OEt	n-Butyl

6.138	6-Cl	H	OEt	n-Butyl
6.139	H	7-Cl	OEt	n-Butyl
6.140	H	7-I	OEt	n-Butyl
6.141	6-Br	H	OEt	i-Butyl
6.142	6-Cl	H	OEt	i-Butyl
6.143	6-Br	H	OEt	
6.144	6-Cl	H	OEt	
6.145	H	7-Cl	OEt	
6.146	6-Br	H	OEt	
6.147	6-Cl	H	OEt	
6.148	6-Br	H	OEt	Cyclobutyl
6.149	6-Br	H	OEt	n-Pentyl
6.150	6-Cl	H	OEt	n-Pentyl
6.151	H	7-Cl	OEt	n-Pentyl
6.152	6-Br	H	OEt	Cyclopentyl
6.153	6-Br	H	OEt	n-Hexyl
6.154	6-Br	H	OEt	Cyclohexyl
6.155	6-Br	H	OEt	Phenyl
6.156	6-Br	H	OEt	4-Chloro-phenyl
6.157	6-Cl	H	OEt	4-Chloro-phenyl
6.158	6-Br	H	OEt	4-Phenoxy-phenyl
6.159	6-Br	H	O-n-Propyl	Me

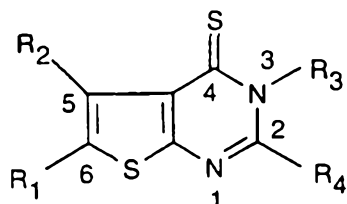
6.160	6-Cl	H	O-n-Propyl	Et
6.161	6-Br	H	O-n-Propyl	n-Propyl
6.162	6-Cl	H	O-n-Propyl	n-Propyl
6.163	H	7-Cl	O-n-Propyl	n-Propyl
6.164	H	7-I	O-n-Propyl	n-Propyl
6.165	6-Br	H	O-n-Propyl	
6.166	6-Cl	H	O-n-Propyl	
6.167	H	7-Cl	O-n-Propyl	
6.168	6-Br	H	O-n-Propyl	n-Butyl
6.169	6-Cl	H	O-n-Propyl	n-Butyl
6.170	H	7-Cl	O-n-Propyl	n-Butyl
6.171	H	7-I	O-n-Propyl	n-Butyl
6.172	6-Br	H	O-n-Propyl	i-Butyl
6.173	6-Cl	H	O-n-Propyl	i-Butyl
6.174	6-Br	H	O-n-Propyl	
6.175	6-Cl	H	O-n-Propyl	
6.176	H	7-Cl	O-n-Propyl	
6.177	6-Br	H	O-n-Propyl	Me 
6.178	6-Cl	H	O-n-Propyl	Me 
6.179	6-Br	H	O-n-Propyl	Cyclobutyl
6.180	6-Br	H	O-n-Propyl	n-Pentyl
6.181	6-Cl	H	O-n-Propyl	n-Pentyl
6.182	H	7-Cl	O-n-Propyl	n-Pentyl




6.183	6-Br	H	O-n-Propyl	Cyclopentyl
6.184	6-Br	H	O-n-Prpyl	n-Hexyl
6.185	6-Br	H	O-n-Prpyl	Cyclohexyl
6.186	6-Br	H	O-n-Prpyl	Phenyl
6.187	6-Br	H	O-n-Prpyl	4-Chloro- phenyl
6.188	6-Cl	H	O-n-Prpyl	4-Chloro- phenyl
6.189	6-Br	H	O-n-Prpyl	4-Phenoxy- phenyl
6.190	6-Br	H	Et	CH ₂ OMe
6.191	6-Cl	H	Et	CH ₂ OMe
6.192	6-Br	H	n-Prpyl	CH ₂ OMe
6.193	6-Cl	H	n-Prpyl	CH ₂ OMe
6.194	H	7-Cl	n-Prpyl	CH ₂ OMe
6.195	6-Br	H	n-Butyl	CH ₂ OMe
6.196	6-Cl	H	n-Butyl	CH ₂ OMe
6.197	6-Br	H		CH ₂ OMe
6.198	6-Br	H	n-Pentyl	CH ₂ OMe
6.199	6-Br	H	Et	CH ₂ OEt
6.200	6-Cl	H	Et	CH ₂ OEt
6.201	6-Br	H	n-Propyl	CH ₂ OEt
6.202	6-Cl	H	n-Propyl	CH ₂ OEt
6.203	H	7-Cl	n-Propyl	CH ₂ OEt
6.204	6-Br	H	n-Butyl	CH ₂ OEt
6.205	6-Cl	H	n-Butyl	CH ₂ OEt
6.206	6-Br	H		CH ₂ OEt
6.207	6-Br	H	n-Pentyl	CH ₂ OEt
6.208	6-Br	H	n-Propyl	CH ₂ CN
6.209	6-Br	H	n-Butyl	CH ₂ CN
6.210	6-Br	H	n-Propyl	t-Butyl
6.211	6-Br	H	n-Propyl	t-Butyl

- 70 -

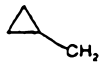


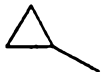
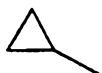

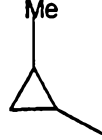
6.212	6-Cl	H	n-Propyl	CF ₃
6.213	6-Br	H	n-Propyl	CF ₃
6.214	6-Br	H	n-Butyl	CF ₃








Tabelle 7: A = Thienyl[2.3-d]













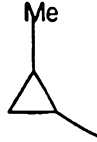
cmpd. No.	R ₁	R ₂	R ₃	R ₄	phys. data m.p.°C
7.1	Br	H	Me	Me	
7.2	Cl	H	Me	Et	
7.3	Br	H	Me	n-Propyl	
7.4	Cl	H	Me	n-Propyl	
7.5	Br	H	Me	n-Butyl	
7.6	Cl	H	Me	n-Butyl	113-114
7.7	Br	H	Me	i-Butyl	
7.8	Cl	H	Me	i-Butyl	
7.9	Br	H	Me	n-Pentyl	
7.10	Br	H	Me		
7.11	Cl	H	Me		
7.12	Br	H	Et	Me	
7.13	Cl	H	Et	Et	
7.14	Br	H	Et	n-Propyl	
7.15	Cl	H	Et	n-Propyl	
7.16	Br	H	Et	n-Butyl	
7.17	Cl	H	Et	n-Butyl	
7.18	Br	H	Et	i-Butyl	
7.19	Cl	H	Et	i-Butyl	
7.20	Br	H	Et	n-Pentyl	
7.21	Br	H	Et		





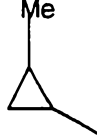

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

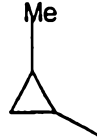
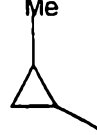
7.22	Cl	H	Et		
7.23	Br	H	n-Propyl	Me	
7.24	Cl	H	n-Propyl	Et	
7.25	Br	H	n-Propyl	n-Propyl	
7.26	Cl	H	n-Propyl	n-Propyl	
7.27	I	H	n-Propyl	n-Propyl	
7.28	Br	H	n-Propyl		
7.29	Cl	H	n-Propyl		
7.30	Br	H	n-Propyl	n-Butyl	
7.31	Cl	H	n-Propyl	n-Butyl	Oil, ¹ H-NMR
7.32	I	H	n-Propyl	n-Butyl	
7.33	Br	H	n-Propyl	i-Butyl	
7.34	Cl	H	n-Propyl	i-Butyl	57-60
7.35	Br	H	n-Propyl		
7.36	Cl	H	n-Propyl		
7.37	Br	H	n-Propyl	Me 	
7.38	Cl	H	n-Propyl	Me 	
7.39	Br	H	n-Propyl	Cyclobutyl	
7.40	Br	H	n-Propyl	n-Pentyl	
7.41	Cl	H	n-Propyl	n-Pentyl	
7.42	Br	H	n-Propyl	Cyclopentyl	
7.43	Br	H	n-Propyl	n-Hexyl	
7.44	Br	H	n-Propyl	Cyclohexyl	
7.45	Br	H	n-Propyl	Phenyl	

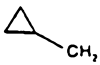
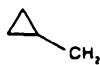
7.46	Br	H	n-Propyl	4-Chloro-phenyl
7.47	Cl	H	n-Propyl	4-Chloro-phenyl
7.48	Br	H	n-Propyl	4-Phenoxy-phenyl
7.49	Br	H	n-Butyl	Me
7.50	Cl	H	n-Butyl	Et
7.51	Br	H	n-Butyl	n-Propyl
7.52	Cl	H	n-Butyl	n-Propyl
7.53	I	H	n-Butyl	n-Propyl
7.54	I	H	n-Butyl	n-Propyl
7.55	Br	H	n-Butyl	
7.56	Cl	H	n-Butyl	
7.57	I	H	n-Butyl	
7.58	Br	H	n-Butyl	n-Butyl
7.59	Cl	H	n-Butyl	n-Butyl
7.60	I	H	n-Butyl	n-Butyl
7.61	Br	H	n-Butyl	i-Butyl
7.62	Cl	H	n-Butyl	i-Butyl
7.63	Br	H	n-Butyl	
7.64	Cl	H	n-Butyl	
7.65	I	H	n-Butyl	
7.66	Br	H	n-Butyl	Me 

7.67	Cl	H	n-Butyl	
7.68	Br	H	n-Butyl	Cyclobutyl
7.69	Br	H	n-Butyl	n-Pentyl
7.70	Cl	H	n-Butyl	n-Pentyl
7.71	Br	H	n-Butyl	Cyclopentyl
7.72	Br	H	n-Butyl	n-Hexyl
7.73	Br	H	n-Butyl	Cyclohexyl
7.74	Cl	H	n-Butyl	Phenyl
7.75	Br	H	n-Butyl	4-Chloro-phenyl
7.76	Cl	H	n-Butyl	4-Chloro-phenyl
7.77	Br	H	n-Butyl	4-Phenoxy-phenyl
7.78	Br	H	i-Butyl	n-Propyl
7.79	Cl	H	i-Butyl	n-Propyl
7.80	Br	H	i-Butyl	n-Butyl
7.81	Cl	H	i-Butyl	n-Butyl
7.82	Br	H		n-Propyl
7.83	Cl	H		n-Propyl
7.84	Cl	H		n-Propyl
7.85	Br	H		n-Butyl
7.86	Br	H	n-Pentyl	Me
7.87	Cl	H	n-Pentyl	Et
7.88	Br	H	n-Pentyl	n-Propyl
7.89	Cl	H	n-Pentyl	n-Propyl

7.90	Br	H	n-Pentyl	
7.91	Cl	H	n-Pentyl	
7.92	Br	H	n-Pentyl	n-Butyl
7.93	Cl	H	n-Pentyl	n-Butyl
7.94	I	H	n-Pentyl	n-Butyl
7.95	Br	H	n-Pentyl	i-Butyl
7.96	Cl	H	n-Pentyl	i-Butyl
7.97	Br	H	n-Pentyl	
7.98	Cl	H	n-Pentyl	
7.99	Br	H	n-Pentyl	Me 
7.100	Cl	H	n-Pentyl	Me 
7.101	Cl	H	n-Pentyl	Cyclobutyl
7.102	Br	H	n-Pentyl	n-Pentyl
7.103	Cl	H	n-Pentyl	n-Pentyl
7.104	Cl	H	n-Pentyl	Cyclopentyl
7.105	Br	H	n-Pentyl	n-Hexyl
7.106	Cl	H	n-Pentyl	Cyclohexyl
7.107	Br	H	n-Pentyl	Phenyl
7.108	Br	H	n-Pentyl	4-Chloro-phenyl
7.109	Cl	H	n-Pentyl	4-Chloro-phenyl
7.110	Br	H	n-Pentyl	4-Phenoxy-phenyl
7.111	Br	H	OEt	Me

7.112	Cl	H	OEt	Et
7.113	Br	H	OEt	n-Propyl
7.114	Cl	H	OEt	n-Propyl
7.115	Br	H	OEt	
7.116	Cl	H	OEt	
7.117	Br	H	OEt	n-Butyl
7.118	Cl	H	OEt	n-Butyl
7.119	I	H	OEt	n-Butyl
7.120	Br	H	OEt	i-Butyl
7.121	Cl	H	OEt	i-Butyl
7.122	Br	H	OEt	
7.123	Cl	H	OEt	
7.124	Br	H	OEt	Me 
7.125	Cl	H	OEt	Me 
7.126	Br	H	OEt	Cyclobutyl
7.127	Br	H	OEt	n-Pentyl
7.128	Cl	H	OEt	n-Pentyl
7.129	Br	H	OEt	Cyclopentyl
7.130	Br	H	OEt	n-Hexyl
7.131	Br	H	OEt	Cyclohexyl
7.132	Br	H	OEt	Phenyl
7.133	Br	H	OEt	4-Chloro- phenyl
7.134	Cl	H	OEt	4-Chloro- phenyl

7.135	Cl	H	OEt	4-Phenoxy-phenyl
7.136	Br	H	O-n-Propyl	Me
7.137	Cl	H	O-n-Propyl	Et
7.138	Br	H	O-n-Propyl	n-Propyl
7.139	Cl	H	O-n-Propyl	n-Propyl
7.140	Br	H	O-n-Propyl	
7.141	Cl	H	O-n-Propyl	
7.142	Br	H	O-n-Propyl	n-Butyl
7.143	Cl	H	O-n-Propyl	n-Butyl
7.144	Br	H	O-n-Propyl	i-Butyl
7.145	Cl	H	O-n-Propyl	i-Butyl
7.146	Br	H	O-n-Propyl	
7.147	Cl	H	O-n-Propyl	
7.148	Br	H	O-n-Propyl	Me 
7.149	Cl	H	O-n-Propyl	Me 
7.150	Br	H	O-n-Propyl	Cyclobutyl
7.151	Br	H	O-n-Propyl	n-Pentyl
7.152	Cl	H	O-n-Propyl	n-Pentyl
7.153	Br	H	O-n-Propyl	Cyclopentyl
7.154	Cl	H	O-n-Propyl	n-Hexyl
7.155	Br	H	O-n-Propyl	Cyclohexyl
7.156	Cl	H	O-n-Propyl	Phenyl
7.157	Br	H	O-n-Propyl	4-Chloro-phenyl

7.158	Cl	H	O-n-Propyl	4-Chloro-phenyl
7.159	Br	H	O-n-Propyl	4-Phenoxy-phenyl
7.160	Br	H	Et	CH ₂ OMe
7.161	Cl	H	Et	CH ₂ OMe
7.162	Br	H	n-Propyl	CH ₂ OMe
7.163	Cl	H	n-Propyl	CH ₂ OMe
7.164	Br	H	n-Butyl	CH ₂ OMe
7.165	Cl	H	n-Butyl	CH ₂ OMe
7.166	Br	H		CH ₂ OMe
7.167	Br	H	n-Pentyl	CH ₂ OMe
7.168	Br	H	Et	CH ₂ OEt
7.169	Cl	H	Et	CH ₂ OEt
7.170	Br	H	n-Propyl	CH ₂ OEt
7.171	Cl	H	n-Propyl	CH ₂ OEt
7.172	Br	H	n-Butyl	CH ₂ OEt
7.173	Cl	H	n-Butyl	CH ₂ OEt
7.174	Br	H		CH ₂ OEt
7.175	Br	H	n-Pentyl	CH ₂ OEt
7.176	Br	H	n-Propyl	CH ₂ CN
7.177	Cl	H	n-Butyl	CH ₂ CN
7.178	Br	H	n-Propyl	t-Butyl
7.179	Cl	H	n-Propyl	t-Butyl
7.180	Br	H	n-Propyl	CF ₃
7.181	Cl	H	n-Butyl	CF ₃
7.182	Cl	H	n-Pentyl	CF ₃
7.183	Cl	Cl	n-Propyl	n-Propyl
7.184	Cl	Cl	n-Propyl	n-Butyl
7.185	Br	Br	n-Propyl	n-Butyl
7.186	Br	Br	n-Butyl	n-Butyl

Examples for specific formulations-combination are as disclosed e.g. in WO 97/33890, e.g. for wettable powders, emulsifiable concentrates, dusts, extruder granules, coated granules, solutions and suspension concentrates.

Biological Examples: Fungicidal actions

Example B-1: Action against *Colletotrichum lagenarium* on cucumbers

After a growth period of 2 weeks, cucumber plants are sprayed with an aqueous spray mixture (concentration 0.002%) prepared from a wettable powder formulation of the test compound and infected 2 days later with a spore suspension (1.5×10^5 spores/ml) of the fungus and incubated for 36 hours at 23°C and high humidity. Incubation is then continued at normal humidity and c. 22°C. Evaluation of the fungal infestation is made 8 days after infection.

The compounds of the Tables 1-7 show good to excellent activity, preferably the compounds 1.36, 2.30, 3.1, 3.6, 3.8, 3.15, 3.17, 3.26, 3.30, 3.31, 3.47, 3.52, 3.56, 3.58, 3.59, 3.79, 3.117, 3.118, 3.163, 3.171, 4.36, 6.29, 7.6, 7.31 and 7.34.

Example B-2: Residual-protective action against *Venturia inaequalis* on apples

Apple cuttings with fresh shoots 10 to 20 cm long are sprayed to drip point with a spray mixture (0.02% a.i.) prepared from a wettable powder formulation of the test compound. The plants are infected 24 hours later with a conidia suspension of the fungus. The plants are then incubated for 5 days at 90 to 100 % relative humidity and stood in a greenhouse for a further 10 days at 20 to 24°C. Evaluation of the fungal infestation is made 12 days after infection.

Compounds of Tables 1-7 show good activity, preferably the compounds 1.36, 2.30, 3.1, 3.6, 3.8, 3.15, 3.17, 3.26, 3.30, 3.31, 3.34, 3.36, 3.47, 3.52, 3.56, 3.58, 3.59, 3.79, 3.117, 3.118, 3.163, 3.171, 4.36, 6.29, 7.6, 7.31 and 7.34.

Example B-3: Action against *Erysiphe graminis* on barley

a) Residual-protective action

Barley plants about 8 cm in height are sprayed to drip point with a spray mixture (0.02% a.i.) prepared from a wettable powder formulation of the test compound, and the treated plants are dusted with conidia of the fungus 3 to 4 hours later. The infected plants are stood in a greenhouse at 22°C. Evaluation of the fungal infection is made 12 days after infection.

b) Systemic action

Barley plants about 8 cm in height are drenched with an aqueous spray mixture (0.002% a.i., based on the volume of the soil) prepared from a wettable powder formulation of the test compound. Care is taken that the spray mixture does not come into contact with the growing parts of the plants. The treated plants are dusted 48 hours later with conidia of the fungus. The infected plants are then stood in a greenhouse at 22°C. Evaluation of the fungal infestation is made 12 days after infection.

Compared with the control plants, infestation of the plants treated with compounds of formula I from Tables 1-7, for example the compounds 1.36, 2.30, 3.1, 3.6, 3.8, 3.15, 3.17, 3.26, 3.30, 3.31, 3.34, 3.36, 3.47, 3.52, 3.56, 3.58, 3.59, 3.79, 3.117, 3.118, 3.163, 3.171, 4.36, 6.29, 7.6, 7.31 and 7.34 is 20 % or less.

Example B-4: Action against Podosphaera leucotricha on apple shoots

Apple cuttings with fresh shoots about 15cm long are sprayed with a spray mixture (0.06% a.i.). The plants are infected 24 hours later with a conidia suspension of the fungus and stood in a climatic chamber at 70% relative humidity and 20°C. Evaluation of the fungal infestation is made 12 days after infection.

Compounds of Tables 1-7 show good activity. The following compounds exhibit especially strong efficacy : 1.36, 2.30, 3.1, 3.6, 3.8, 3.15, 3.17, 3.26, 3.30, 3.31, 3.34, 3.36, 3.47, 3.52, 3.56, 3.58, 3.59, 3.79, 3.117, 3.118, 3.163, 3.171, 4.36, 6.29, 7.6, 7.31 and 7.34 (0-5% infestation).

Example B-5: Action against Plasmopara viticola on vines

a) Residual-preventive action: Vine cuttings of the Chasselas variety are raised in a greenhouse. At the 10-leaf stage, 3 plants are sprayed with a spray mixture (200 ppm a.i.). After the spray coating has dried, the plants are infected uniformly on the underside of the leaves with a spore suspension of the fungus. The plants are then kept in a humidity chamber for 8 days, after which time marked symptoms of disease are observed on the control plants. The number and size of the infected areas on the untreated plants act as an indicator of the efficacy of the tested compounds.

b) Curative action: Vine cuttings of the Chasselas variety are raised in a greenhouse and sprayed at the 10-leaf stage on the underside of the leaves with a spore suspension of *Plasmopara viticola*. After 24 hours in the humidity chamber, the plants are sprayed with a

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spray mixture (200 ppm a.i.). The plants are then kept for another 7 days in the humidity chamber. After this time the control plants exhibit symptoms of the disease. The number and size of the infected areas on the untreated plants act as an indicator of the efficacy of the tested compounds.

Compounds of Tables 1-7 show good efficacy, preferably the compounds 1.36, 2.30, 3.1, 3.6, 3.8, 3.15, 3.17, 3.26, 3.30, 3.31, 3.34, 3.36, 3.47, 3.52, 3.56, 3.58, 3.59, 3.79, 3.117, 3.118, 3.163, 3.171, 4.36, 6.29, 7.6, 7.31 and 7.34.

Example B-6: Action against *Uncinula necator* on vines

5 week old vine cuttings are sprayed with a spray mixture (200 ppm a.i.) prepared from a wettable powder formulation of the test compound. The plants are infected 24 hours later by conidia from strongly infested vine leaves that are shaken off over the test plants. The plants are then incubated at 26°C and 60% relative humidity. The evaluation of the fungal infestation is made ca. 14 days after infection.

Compared with the control plants, infestation of the plants treated with compounds of formula I from the Tables 1-7, for example the compounds 1.36, 2.30, 3.1, 3.6, 3.8, 3.15, 3.17, 3.26, 3.30, 3.31, 3.34, 3.36, 3.47, 3.52, 3.56, 3.58, 3.59, 3.79, 3.117, 3.118, 3.163, 3.171, 4.36, 6.29, 7.6, 7.31 and 7.34 is 20 % or less.

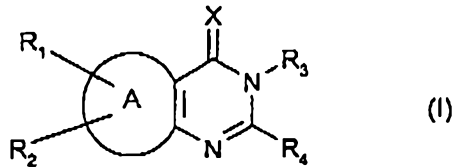
The reference to any prior art in this specification is not, and should not be taken as, an acknowledgment or any form of suggestion that that prior art forms part of the common general knowledge in Australia.

Throughout this specification and the claims which follow, unless the context requires otherwise, the word "comprise", and variations such as "comprises" and "comprising", will be understood to imply the inclusion of a stated integer or step or group of integers or steps but not the exclusion of any other integer or step or group of integers or steps.



THE CLAIMS DEFINING THE INVENTION ARE AS FOLLOWS:

1. A compound of formula I



wherein

A is thienyl (including all 3 isomers), thiazolyl, pyridyl or pyridazinyl;

X is oxygen or sulfur;

R₁ is hydrogen, halogen or trimethylsilyl;

R₂ is hydrogen, halogen or trimethylsilyl; at least one of R₁ and R₂ is not hydrogen;

R₃ is C₁-C₈alkyl, C₁-C₈alkenyl, C₁-C₈alkinyl which are unsubstituted or mono to tri-substituted by C₃-C₆cycloalkyl, halogen, C₁-C₆alkoxy or C₁-C₆haloalkoxy; O-C₁-C₆alkyl, O-C₂-C₆alkenyl, O-C₂-C₆alkynyl, which are unsubstituted or mono to tri-substituted by C₃-C₆cycloalkyl, halogen or C₁-C₆alkoxy; N-C₁-C₆alkyl; or N=CHC₁-C₆alkyl;

R₄ is C₁-C₈alkyl, C₁-C₈alkenyl, C₁-C₈alkinyl which are unsubstituted or mono to tri-substituted by C₃-C₆cycloalkyl, halogen, cyano, C₁-C₆alkoxy or C₁-C₆haloalkoxy; nitro; -CO-C₁-C₆alkyl; C₃-C₆cycloalkyl; or phenyl, which is unsubstituted or mono to tri-substituted by halogen, C₁-C₆alkyl, C₁-C₆haloalkyl, C₁-C₆alkoxy, C₁-C₆haloalkoxy, cyano, nitro, amino, mono-C₁-C₆alkylamino, di-C₁-C₆alkyl-amino, C₁-C₆alkylthio, phenyl or phenoxy and in which the phenyl part is unsubstituted or mono to tri-substituted by halogen, C₁-C₆alkyl, C₁-C₆haloalkyl, C₁-C₆alkoxy or C₁-C₆haloalkoxy.

2. A compound of formula I according to claim 1, wherein

A is thienyl, including all 3 isomers.

3. A compounds of formula I according to claim 2 wherein

R₁ is hydrogen, fluorine, chlorine, bromine or iodine;

R₂ is hydrogen, fluorine, chlorine, bromine or iodine; at least one of R₁ and R₂ is not hydrogen;



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R_3 is C_1-C_6 alkyl, C_2-C_6 alkenyl, C_2-C_6 alkynyl which are unsubstituted or mono to tri-substituted by C_3-C_6 cycloalkyl, halogen or C_1-C_4 alkoxy; $O-C_1-C_6$ alkyl, $O-C_2-C_6$ alkenyl, $O-C_2-C_6$ alkynyl, which are unsubstituted or mono to tri-substituted by C_3-C_6 cycloalkyl, halogen or C_1-C_4 alkoxy; $N-C_1-C_6$ alkyl; or $N=CHC_1-C_6$ alkyl;

R_4 is C_1-C_6 alkyl, C_2-C_6 alkenyl, C_2-C_6 alkynyl which are unsubstituted or mono to tri-substituted by C_3-C_6 cycloalkyl, halogen or C_1-C_4 alkoxy; or phenyl which is unsubstituted or mono to tri-substituted by fluorine, chlorine, bromine, C_1-C_4 alkyl, C_1-C_4 haloalkyl, C_1-C_4 alkoxy, C_1-C_4 haloalkoxy, phenyl or phenoxy and in which the phenyl part is unsubstituted or mono to tri-substituted by fluorine, chlorine, bromine, C_1-C_4 alkyl, C_1-C_4 haloalkyl, C_1-C_4 alkoxy or C_1-C_4 haloalkoxy.

4. A compound of formula I according to claim 3, wherein

R_3 is C_1-C_6 alkyl, C_2-C_6 alkenyl, C_2-C_6 alkynyl which are unsubstituted or mono to tri-substituted by C_3-C_4 cycloalkyl, fluorine, chlorine, bromine or C_1-C_4 alkoxy; $O-C_1-C_6$ alkyl; $O-C_2-C_6$ alkenyl; $O-C_2-C_6$ alkynyl; $N-C_1-C_6$ alkyl; or $N=CHC_1-C_6$ alkyl;

R_4 is C_1-C_6 alkyl, C_2-C_6 alkenyl, C_2-C_6 alkynyl which are unsubstituted or mono to tri-substituted by C_3-C_4 cycloalkyl, fluorine, chlorine, bromine or C_1-C_4 alkoxy; or phenyl which is unsubstituted or mono to tri-substituted by fluorine, chlorine, bromine, C_1-C_4 alkyl, C_1-C_4 haloalkyl, C_1-C_4 alkoxy, C_1-C_4 haloalkoxy, phenyl or phenoxy and in which the phenyl part is unsubstituted or mono to tri-substituted by fluorine, chlorine or bromine.

5. A compound of the formula I according to claim 4, wherein

A is thienyl[2.3-d],

X is oxygen,

R_1 is hydrogen, chlorine or bromine;

R_2 is hydrogen, chlorine or bromine; at least one of R_1 and R_2 is not hydrogen;

R_3 is C_3-C_5 alkyl or $O-C_1-C_4$ alkyl;

R_4 is C_2-C_5 alkyl or phenyl which is unsubstituted or mono to tri-substituted by fluorine, chlorine, bromine, C_1-C_4 alkyl or phenoxy and in which phenoxy is unsubstituted or mono to tri-substituted by fluorine, chlorine or bromine.

6. A compound of the formula I according to claim 4, wherein



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A is thienyl[2.3-d],

X is sulfur,

R₁ is hydrogen, chlorine or bromine;

R₂ is hydrogen, chlorine or bromine; at least one of R₁ and R₂ is not hydrogen;

R₃ is C₃-C₅alkyl or O-C₁-C₄alkyl;

R₄ is C₂-C₅alkyl or phenyl which is unsubstituted or mono to tri-substituted by fluorine, chlorine, bromine, C₁-C₄alkyl or phenoxy and in which phenoxy is unsubstituted or mono to tri-substituted by fluorine, chlorine or bromine.

7. A compound of the formula I according to claim 4, wherein

A is thienyl[3.2-d],

X is oxygen,

R₁ is hydrogen, chlorine or bromine;

R₂ is hydrogen, chlorine or bromine; at least one of R₁ and R₂ is not hydrogen;

R₃ is C₃-C₅alkyl or O-C₁-C₄alkyl;

R₄ is C₂-C₅alkyl or phenyl which is unsubstituted or mono to tri-substituted by fluorine, chlorine, bromine, C₁-C₄alkyl or phenoxy and in which phenoxy is unsubstituted or mono to tri-substituted by fluorine, chlorine or bromine.

8. A compound of formula I according to claim 1, wherein

A is pyridyl.

9. A compound of formula I according to claim 1, wherein

A is thiazolyl.

10. A compound of formula I according to claim 1, wherein

A is pyridazinyl.

11. A composition for controlling and preventing pests, wherein the active ingredient is a compound as claimed in claim 1 together with a suitable carrier.



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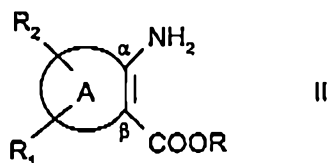
12. Use of a compound of formula I according to claim 1 for protecting plants against infestation by phytopathogenic microorganisms.

13. A method of controlling or preventing infestation of cultivated plants by phytopathogenic microorganisms by application of a compound of formula I as claimed in claim 1 to plants, to parts thereof or to the locus thereof.

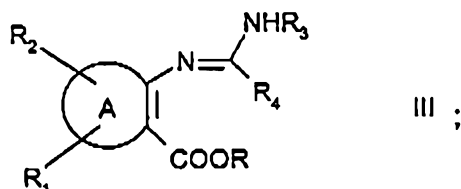
14. A method according to claim 13, wherein the phytopathogenic microorganism is a fungal organism.

15. A method for the preparation of a compound of formula I according to claim 1, which comprises

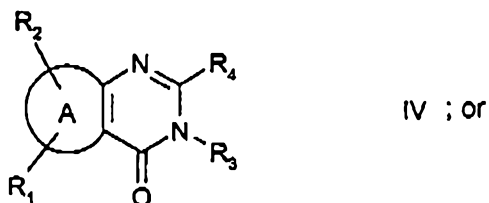
a) converting an α -amino- β -carboalkoxyheterocycle of formula II, wherein R_1 and R_2 have the meanings stated for formula I and R is hydrogen, C_1 - C_8 alkyl,



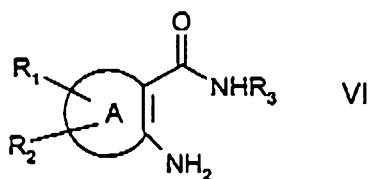
with $POCl_3$ in the presence of a solvent and R_4CONHR_3 into an amidine of formula III, wherein R_3 and R_4 have the meanings stated for formula I



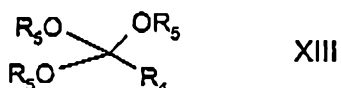
b) and treating the amidine, in the presence of a solvent and if necessary in the presence of a base, and obtaining, with ring closure, the pyrimidin-4-one derivative of formula IV



c) reacting an amino carboxylic acidamide of formula VI

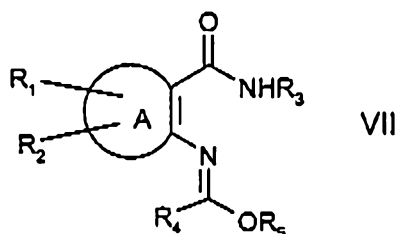


wherein A, R₁, R₂ and R₃ have the meanings stated for formula I with an orthoester of formula XIII



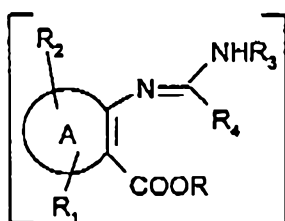
wherein R₄ has the meaning stated for formula I and R₅ is C₁-C₂alkyl, in the presence or absence of a solvent, in the presence or absence of an acid catalyst at 20-200°C, and obtaining the pyrimidin-4-one derivative of formula IV; and

d) if the intermediate VII is formed



treating the compound VII in the presence of a solvent and if necessary in the presence of a base, and obtaining with ring closure the pyrimidin-4-one derivative of formula IV.

16. A compound of formula III



III

wherein

A is thienyl[2.3-d] or thienyl[3.2-d];



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R₁ is hydrogen, halogen or trimethylsilyl;

R₂ is hydrogen, halogen or trimethylsilyl; and at least one of R₁ and R₂ is not hydrogen;

R₃ is C₁-C₈alkyl, C₁-C₈alkenyl, C₁-C₈alkynyl which are unsubstituted or mono to tri-substituted by C₃-C₈cycloalkyl, halogen, C₁-C₆alkoxy or C₁-C₆haloalkoxy; O-C₁-C₈alkyl, O-C₂-C₆alkenyl, O-C₂-C₆alkynyl, which are unsubstituted or mono to tri-substituted by C₃-C₆cycloalkyl, halogen or C₁-C₆alkoxy; N-C₁-C₆alkyl; or N=CHC₁-C₆alkyl;

R₄ is C₁-C₈alkyl, C₁-C₈alkenyl, C₁-C₈alkynyl which are unsubstituted or mono to tri-substituted by C₃-C₆cycloalkyl, halogen, cyano, C₁-C₆alkoxy or C₁-C₆haloalkoxy; C₁-C₄alkoxy-C₁-C₄alkoxy; C₁-C₄alkoxy-C₁-C₄alkylthio; nitro; -CO-C₁-C₆alkyl; C₃-C₈cycloalkyl; or phenyl, which is unsubstituted or mono to tri-substituted by halogen, C₁-C₆alkyl, C₁-C₆haloalkyl; C₁-C₆alkoxy, C₁-C₆haloalkoxy, cyano, nitro, amino, mono-C₁-C₆alkylamino, di-C₁-C₆alkylamino, C₁-C₆alkylthio, phenyl or phenoxy and in which the phenyl part is unsubstituted or mono to tri-substituted by halogen, C₁-C₆alkyl, C₁-C₆haloalkyl, C₁-C₆alkoxy or C₁-C₆haloalkoxy; and

R is hydrogen or C₁-C₆alkyl.

17. A compound of formula I according to claim 1, substantially as hereinbefore described.

18. A composition for controlling and preventing pests according to claim 11, substantially as hereinbefore described with reference to the Examples.

19. Use according to claim 12, substantially as hereinbefore described with reference to the Examples.

20. Method of controlling or preventing infestation of cultivated plants by phytopathogenic microorganisms according to claim 13, substantially as hereinbefore described with reference to the Examples.

21. A method for the preparation of a compound of formula I according to claim 15, substantially as hereinbefore described with reference to the Examples.



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22. A compound of formula I prepared by the process of claims 15 or 21.

23. A compound of formula III according to claim 16, substantially as hereinbefore described with reference to the Examples.

5 DATED this 27th day of June, 2001
Novartis AG

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Patent Attorneys for the Applicants

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