



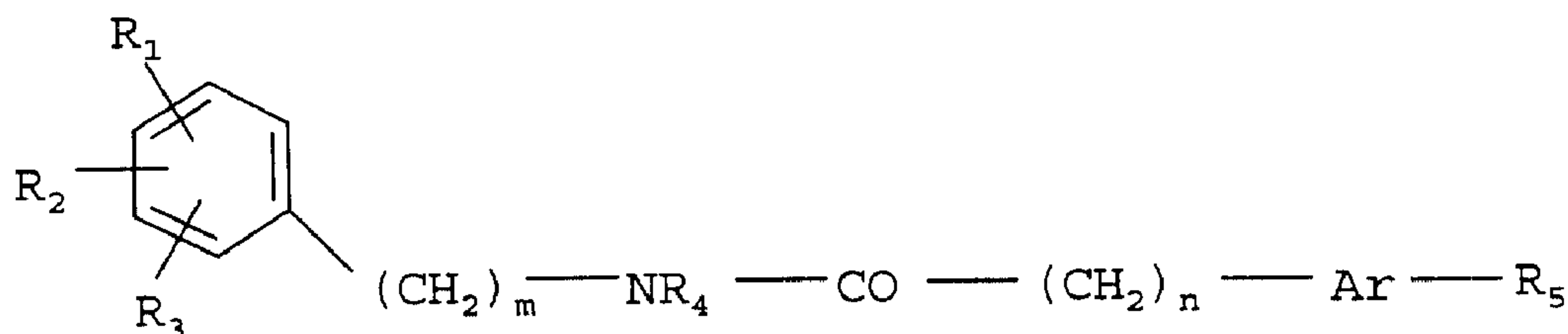
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(54) Titre : AMIDES D'ACIDE CARBOXYLIQUE, LEUR PRODUCTION ET LEUR UTILISATION COMME MEDICAMENTS
(54) Title: CARBOXYLIC ACID AMIDES, THE PREPARATION THEREOF AND THEIR USE AS PHARMACEUTICAL
COMPOSITIONS



(I)

(57) **Abrégé/Abstract:**

The present invention relates to carboxylic acid amides of general formula (see formula I), wherein R_1 to R_5 , Ar, m and n are defined as in claim 1, the tautomers, stereoisomers, mixtures thereof, the prodrugs and the salts thereof which have valuable properties. The compounds of the above general formula I wherein R_5 denotes a cyano group are valuable intermediate products for preparing the other compounds of general formula I, and the compounds of the above general formula I wherein R_5 denotes one of the amidino groups mentioned in claim 1 have valuable pharmacological properties, particularly an antithrombotic activity and a factor Xa-inhibiting activity.



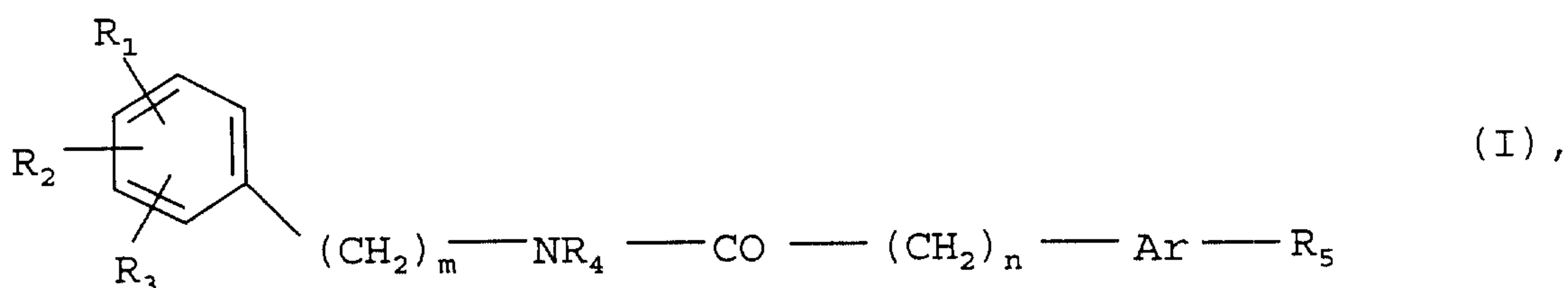
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Abstract

The present invention relates to carboxylic acid amides of general formula



wherein

R_1 to R_5 , Ar, m and n are defined as in claim 1, the tautomers, stereoisomers, mixtures thereof, the prodrugs and the salts thereof which have valuable properties.

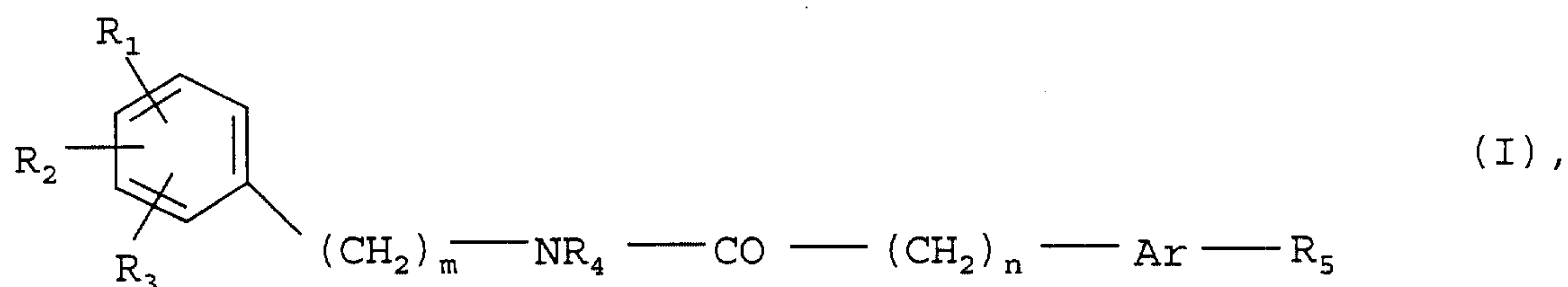
The compounds of the above general formula I wherein R_5 denotes a cyano group are valuable intermediate products for preparing the other compounds of general formula I, and the compounds of the above general formula I wherein R_5 denotes one of the amidino groups mentioned in claim 1 have valuable pharmacological properties, particularly an antithrombotic activity and a factor Xa-inhibiting activity.

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Carboxylic acid amides, the preparation thereof and their use
as pharmaceutical compositions

The present invention relates to carboxylic acid amides of
general formula



their tautomers, stereoisomers, mixtures thereof, the prodrugs thereof, the derivatives thereof which contain a group which is negatively charged under physiological conditions, and the salts thereof, particularly the physiologically acceptable salts thereof with inorganic or organic acids or bases which have valuable properties.

The compounds of the above general formula I wherein R_5 denotes a cyano group, are valuable intermediate products for preparing the above compounds of general formula I, and the compounds of the above general formula I wherein R_5 denotes one of the following amidino groups, as well as their tautomers, stereoisomers, mixtures thereof, the prodrugs thereof, the derivatives thereof which contain a group which is negatively charged under physiological conditions, and the salts thereof, particularly the physiologically acceptable salts thereof with inorganic or organic acids, and the stereoisomers thereof have valuable pharmacological properties, particularly an antithrombotic activity and a factor Xa-inhibiting activity.

The present application thus relates to the new compounds of the above general formula I and their preparation, pharmaceutical compositions containing the pharmacologically active compounds, the preparation and use thereof.

In the above general formula

one of the groups m or n denotes the number 0 and the other group m or n denotes the number 1,

Ar denotes a phenylene or naphthylene group optionally substituted by a fluorine, chlorine or bromine atom, by a trifluoromethyl, C₁₋₃-alkyl, hydroxy, C₁₋₃-alkoxy, phenyl-C₁₋₃-alkoxy, amino, C₁₋₃-alkylamino or di-(C₁₋₃-alkyl)-amino group, whilst the phenylene group may be substituted by another fluorine, chlorine or bromine atom or by another C₁₋₃-alkyl group,

a thienylene, thiazolylenes, pyridinylenes, pyrimidinylene, pyrazinylenes or pyridazinylene group optionally substituted in the carbon skeleton by a C₁₋₃-alkyl group,

R₁ denotes a C₁₋₃-alkyl group optionally substituted by an amino, C₁₋₃-alkylamino, di-(C₁₋₃-alkyl)-amino, phenyl, naphthyl, heteroaryl or 4- to 7-membered cycloalkyleneimino group,

a C₃₋₇-cycloalkyl group which is substituted in the 1 position by a 5- to 7-membered cycloalkyleneiminocarbonyl group,

an amino, C₁₋₅-alkylamino, C₅₋₇-cycloalkylamino or phenyl-C₁₋₃-alkylamino group which may in each case be substituted at the amino-nitrogen atom by a benzoyl or phenylsulphonyl group or by a C₁₋₃-alkyl or C₁₋₃-alkylcarbonyl group optionally substituted in the C₁₋₃-alkyl moiety by a carboxy group,

a 4- to 7-membered cycloalkyleneiminocarbonyl or cycloalkyleneiminosulphonyl group optionally substituted by a C₁₋₃-alkyl group,

an aminosulphonyl group optionally substituted by one or two C₁₋₃-alkyl groups,

a phenyl group optionally substituted by a fluorine, chlorine or bromine atom, by a trifluoromethyl, aminosulphonyl, C₁₋₃-alkyl or C₁₋₃-alkoxy group, which may additionally be substituted by a fluorine, chlorine or bromine atom or by a trifluoromethyl, C₁₋₃-alkyl or C₁₋₃-alkoxy group,

a C₁₋₃-alkoxy, phenyl-C₁₋₃-alkoxy, heteroaryloxy or heteroaryloxy-C₁₋₃-alkoxy group wherein the alkoxy moiety may be substituted in the 2 or 3 position in each case by an amino, C₁₋₃-alkylamino or di-(C₁₋₃-alkyl)-amino group,

a C₃₋₇-cycloalkoxy group, whilst the methylene group in the 3 or 4 position in a C₅₋₇-cycloalkoxy group may be replaced by an -NH group, whilst the -NH group may be substituted

by a C₁₋₃-alkyl group which may be substituted in the 2 or 3 position by an amino, C₁₋₃-alkylamino or di-(C₁₋₃-alkyl)-amino group, by a C₁₋₃-alkylcarbonyl, arylcarbonyl or arylsulphonyl group or

by an aminocarbonyl, C₁₋₃-alkylaminocarbonyl or di-(C₁₋₃-alkyl)-aminocarbonyl group, wherein in each case the oxygen atom of the carbonyl group is replaced by an imino group,

R₂ denotes a hydrogen, fluorine, chlorine or bromine atom, a C₁₋₃-alkyl, hydroxy or C₁₋₃-alkoxy group,

R₃ denotes a hydrogen atom or a C₁₋₃-alkyl group,

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R_4 denotes a hydrogen atom or a C_{1-3} -alkyl group optionally substituted by a carboxy group and

R_5 denotes a cyano group or an amidino group optionally substituted by one or two C_{1-3} -alkyl groups,

but particularly, if m , n , Ar and R_2 to R_5 are as hereinbefore defined,

R_1 denotes a C_{1-3} -alkyl group optionally substituted by an amino, C_{1-3} -alkylamino, di- $(C_{1-3}$ -alkyl)-amino, phenyl, naphthyl or heteroaryl group,

a C_{3-7} -cycloalkyl group which is substituted in the 1 position by a 5- to 7-membered cycloalkyleneiminocarbonyl group,

a 4- to 7-membered cycloalkyleneiminocarbonyl group,

a phenyl group optionally substituted by a fluorine, chlorine or bromine atom, by a trifluoromethyl, C_{1-3} -alkyl or C_{1-3} -alkoxy group,

a C_{1-3} -alkoxy, phenyl- C_{1-3} -alkoxy, heteroaryloxy or heteroaryloxy- C_{1-3} -alkoxy group wherein the alkoxy moiety in the 2 or 3 position may be substituted in each case by an amino, C_{1-3} -alkylamino or di- $(C_{1-3}$ -alkyl)-amino group,

a C_{3-7} -cycloalkoxy group, whilst the methylene group in the 3 or 4 position in a C_{5-7} -cycloalkoxy group may be replaced by an -NH group, whilst the -NH group may be substituted by an arylcarbonyl or arylsulphonyl group, by a C_{1-3} -alkylcarbonyl group wherein the oxygen atom of the carbonyl group may be replaced by an imino group and the alkanoyl moiety may be substituted by an amino, C_{1-3} -alkylamino or di- $(C_{1-3}$ -alkyl)-amino group, or by a C_{1-3} -alkyl group which may be substituted in the 2 or 3 position by an amino, C_{1-3} -alkylamino or di- $(C_{1-3}$ -alkyl)-amino group,

in particular

R_1 denotes a C_{1-3} -alkyl group substituted by a 4- to 7-membered cycloalkyleneimino group,

an amino, C_{1-5} -alkylamino, C_{5-7} -cycloalkylamino or phenyl- C_{1-3} -alkylamino group which may in each case be substituted at the amino-nitrogen atom by a benzoyl or phenylsulphonyl group or by a C_{1-3} -alkyl or C_{1-3} -alkylcarbonyl group optionally substituted in the C_{1-3} -alkyl moiety by a carboxy group,

a 4- to 7-membered cycloalkyleneiminocarbonyl group substituted by a C_{1-3} -alkyl group,

a 4- to 7-membered cycloalkyleneiminosulphonyl group optionally substituted by a C_{1-3} -alkyl group,

an aminosulphonyl group optionally substituted by one or two C_{1-3} -alkyl groups,

an aminosulphonylphenyl group,

a phenyl group substituted by a fluorine, chlorine or bromine atom, by a trifluoromethyl, aminosulphonyl, C_{1-3} -alkyl or C_{1-3} -alkoxy group, which may additionally be substituted by a fluorine, chlorine or bromine atom or by a trifluoromethyl, C_{1-3} -alkyl or C_{1-3} -alkoxy group,

a C_{3-7} -cycloalkoxy group, whilst the methylene group in the 3 or 4 position is replaced in a C_{5-7} -cycloalkoxy group by an -NH group, whilst the -NH group is substituted

by an aminocarbonyl, C_{1-3} -alkylaminocarbonyl or di-
(C_{1-3} -alkyl)-aminocarbonyl group, wherein in each case the oxygen atom of the carbonyl group is replaced by an imino group.

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By the abovementioned heteroaryl groups is meant a 5-membered heteroaryl group optionally substituted by a C₁₋₃-alkyl group which contains, in the heteroaromatic moiety,

an imino group optionally substituted by a C₁₋₃-alkyl group, or an oxygen or sulphur atom,

an imino group optionally substituted by a C₁₋₃-alkyl group and an oxygen, sulphur or nitrogen atom,

an imino group optionally substituted by a C₁₋₃-alkyl group and two nitrogen atoms or

an oxygen or sulphur atom and two nitrogen atoms,

or a 6-membered heteroarylene group optionally substituted by a C₁₋₃-alkyl group which contains one or two nitrogen atoms in the heteroaromatic moiety.

By the above mentioned heteroaryloxy groups is meant a heteroaryl group, as defined above, which is bound through an oxygen atom.

Moreover, the carboxy groups mentioned in the definition of the abovementioned groups may be replaced by a group which may be converted into a carboxy group *in vivo* or by a group which is negatively charged under physiological conditions or

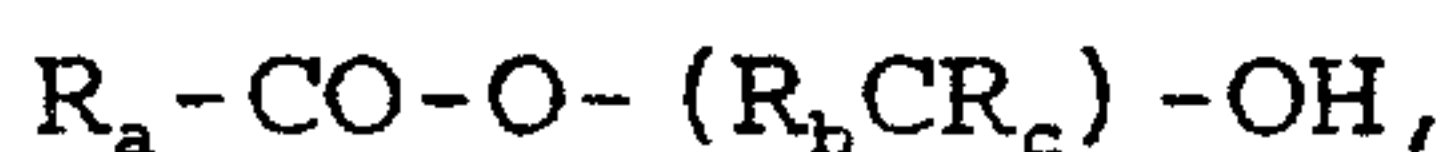
the amino and imino groups mentioned in the definition of the abovementioned groups may be replaced by a group which may be cleaved *in vivo*. Groups of this kind are described for example in WO 98/46576 and by N.M. Nielsen et al. in International Journal of Pharmaceutics 39, 75-85 (1987).

By a group which can be converted *in vivo* into a carboxy group is meant, for example, a hydroxymethyl group, a carboxy group esterified with an alcohol, wherein the alcoholic moiety preferably denotes a C₁₋₆-alkanol, a phenyl-C₁₋₃-alkanol, a C₃₋₉-cycloalkanol, whilst a C₅₋₈-cycloalkanol may additionally be substituted by one or two C₁₋₃-alkyl groups, a C₅₋₈-cycloalkanol

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wherein a methylene group in the 3 or 4 position is replaced by an oxygen atom or by an imino group optionally substituted by a C₁₋₃-alkyl, phenyl-C₁₋₃-alkyl, phenyl-C₁₋₃-alkoxycarbonyl or C₂₋₆-alkanoyl group and the cycloalkanol moiety may additionally be substituted by one or two C₁₋₃-alkyl groups, a C₄₋₇-cycloalkenol, a C₃₋₅-alkenol, a phenyl-C₃₋₅-alkenol, a C₃₋₅-alkynol or phenyl-C₃₋₅-alkynol, with the proviso that no bond to the oxygen atom starts from a carbon atom which carries a double or triple bond, a C₃₋₈-cycloalkyl-C₁₋₃-alkanol, a bicycloalkanol having a total of 8 to 10 carbon atoms which may additionally be substituted by one or two C₁₋₃-alkyl groups in the bicycloalkyl moiety, a 1,3-dihydro-3-oxo-1-isobenzfuranol or an alcohol of formula



wherein

R_a denotes a C₁₋₈-alkyl, C₅₋₇-cycloalkyl, phenyl or phenyl-C₁₋₃-alkyl group,

R_b denotes a hydrogen atom, a C₁₋₃-alkyl, C₅₋₇-cycloalkyl or phenyl group and

R_c denotes a hydrogen atom or a C₁₋₃-alkyl group,

by a group which is negatively charged under physiological conditions is meant, for example, a tetrazol-5-yl, phenylcarbonylaminocarbonyl, trifluoromethylcarbonylaminocarbonyl, C₁₋₆-alkylsulphonylamino, phenylsulphonylamino, benzylsulphonylamino, trifluoromethylsulphonylamino, C₁₋₆-alkylsulphonylaminocarbonyl, phenylsulphonylaminocarbonyl, benzylsulphonylaminocarbonyl or perfluoro-C₁₋₆-alkylsulphonylaminocarbonyl group

and by a group which can be cleaved in vivo from an imino or amino group is meant, for example, a hydroxy group, an acyl group such as a benzoyl group optionally mono- or

disubstituted by fluorine, chlorine, bromine or iodine atoms or by C₁₋₃-alkyl or C₁₋₃-alkoxy groups, whilst the substituents may be identical or different, a pyridinoyl group or a C₁₋₁₆-alkanoyl group such as the formyl, acetyl, propionyl, butanoyl, pentanoyl or hexanoyl group, a 3,3,3-trichloropropionyl or allyloxycarbonyl group, a C₁₋₁₆-alkoxycarbonyl or C₁₋₁₆-alkylcarbonyloxy group wherein hydrogen atoms may be wholly or partially replaced by fluorine or chlorine atoms, such as the methoxycarbonyl, ethoxycarbonyl, propoxycarbonyl, isopropoxycarbonyl, butoxycarbonyl, tert. butoxycarbonyl, pentoxycarbonyl, hexoxycarbonyl, octyloxycarbonyl, nonyloxycarbonyl, decyloxycarbonyl, undecyloxycarbonyl, dodecyloxycarbonyl, hexadecyloxycarbonyl, methylcarbonyloxy, ethylcarbonyloxy, 2,2,2-trichloroethylcarbonyloxy, propylcarbonyloxy, isopropylcarbonyloxy, butylcarbonyloxy, tert.butylcarbonyloxy, pentylcarbonyloxy, hexylcarbonyloxy, octylcarbonyloxy, nonylcarbonyloxy, decylcarbonyloxy, undecylcarbonyloxy, dodecylcarbonyloxy or hexadecylcarbonyloxy group, a phenyl-C₁₋₆-alkoxycarbonyl group such as the benzyloxycarbonyl, phenylethoxycarbonyl or phenylpropoxycarbonyl group, a 3-amino-propionyl group wherein the amino group may be mono or disubstituted by C₁₋₆-alkyl or C₃₋₇-cycloalkyl groups and the substituents may be identical or different, a C₁₋₃-alkylsulphonyl-C₂₋₄-alkoxycarbonyl, C₁₋₃-alkoxy-C₂₋₄-alkoxy-C₂₋₄-alkoxycarbonyl, R_a-CO-O-(R_bCR_c)-O-CO, C₁₋₆-alkyl-CO-NH-(R_dCR_e)-O-CO or C₁₋₆-alkyl-CO-O-(R_dCR_e)-(R_dCR_e)-O-CO group wherein R_a to R_c are as hereinbefore defined,

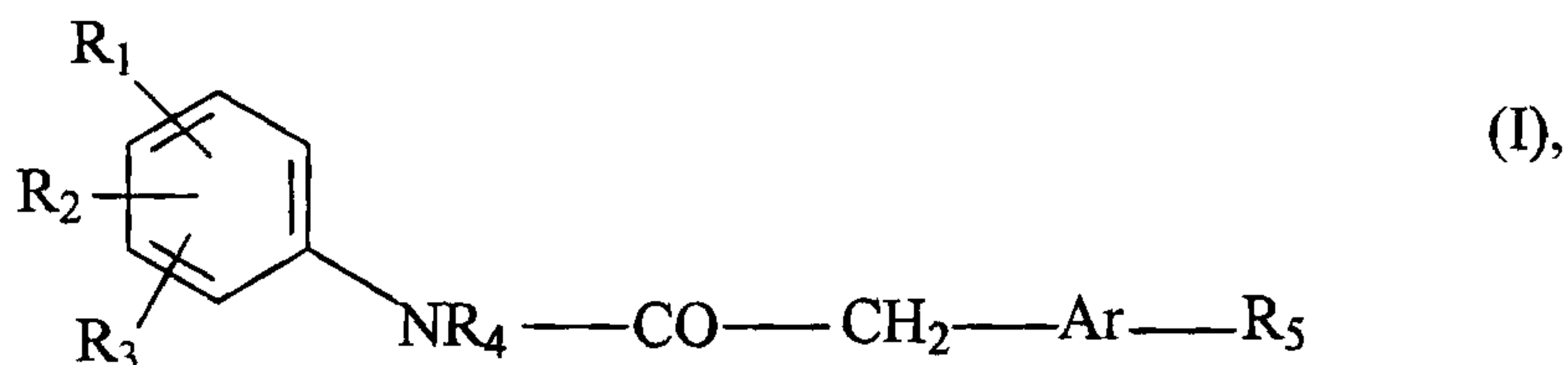
R_d and R_e, which may be identical or different, denote hydrogen atoms or C₁₋₃-alkyl groups.

Moreover, the saturated alkyl and alkoxy moieties containing more than 2 carbon atoms mentioned in the definitions given above also include the branched isomers thereof, such as the isopropyl, tert.butyl, isobutyl group, etc.

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According to one aspect of the present invention, there is provided carboxylic acid amides of general formula



wherein

5 Ar denotes a phenylene or naphthylene group optionally substituted by a fluorine, chlorine or bromine atom, by a trifluoromethyl, C₁₋₃-alkyl, hydroxy, C₁₋₃-alkoxy, phenyl-C₁₋₃-alkoxy, amino, C₁₋₃-alkylamino or di-(C₁₋₃-alkyl)-amino group, while the phenylene group is optionally further substituted
 10 by another fluorine, chlorine or bromine atom or by another C₁₋₃-alkyl group,

a thienylene, thiazolylen, pyridinylen, pyrimidinylene, pyrazinylen or pyridazinylene group optionally substituted in the carbon skeleton by a C₁₋₃-alkyl group,

15 R₁ denotes a C₁₋₃-alkyl group optionally substituted by an amino, C₁₋₃-alkylamino, di-(C₁₋₃-alkyl)-amino, phenyl, naphthyl, heteroaryl or 4- to 7-membered cycloalkyleneimino group,

a C₃₋₇-cycloalkyl group which is substituted in the
 20 1 position by a 5- to 7-membered cycloalkyleneiminocarbonyl group,

an amino, C₁₋₅-alkylamino, C₅₋₇-cycloalkylamino or phenyl-C₁₋₃-alkylamino group each of which is optionally substituted at the amino-nitrogen atom by a benzoyl or
 25 phenylsulphonyl group or by a C₁₋₃-alkyl or C₁₋₃-alkylcarbonyl

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group optionally substituted in the C₁₋₃-alkyl moiety by a carboxy group,

a 4- to 7-membered cycloalkyleneiminocarbonyl or cycloalkyleneiminosulphonyl group optionally substituted by
5 a C₁₋₃-alkyl group,

an aminosulphonyl group optionally substituted by one or two C₁₋₃-alkyl groups,

a phenyl group optionally substituted by a fluorine, chlorine or bromine atom, by a trifluoromethyl,
10 aminosulphonyl, C₁₋₃-alkyl or C₁₋₃-alkoxy group, wherein the substituent group is optionally substituted by a fluorine, chlorine or bromine atom or by a trifluoromethyl, C₁₋₃-alkyl or C₁₋₃-alkoxy group,

a C₁₋₃-alkoxy, phenyl-C₁₋₃-alkoxy, heteroaryloxy or
15 heteroaryloxy-C₁₋₃-alkoxy group wherein the alkoxy moiety is optionally substituted in the 2 or 3 position in each case by an amino, C₁₋₃-alkylamino or di-(C₁₋₃-alkyl)-amino group, or

a C₃₋₇-cycloalkoxy group, wherein the methylene group in the
20 3 or 4 position when the cycloalkoxy group is a C₅₋₇-cycloalkoxy group is optionally replaced by a -NH group, wherein the -NH group is optionally substituted

by a C₁₋₃-alkyl group which is optionally substituted in the 2 or 3 position by an amino,
25 C₁₋₃-alkylamino or di-(C₁₋₃-alkyl)-amino group, by a C₁₋₃-alkylcarbonyl, arylcarbonyl or arylsulphonyl group or

by an aminocarbonyl, C₁₋₃-alkylaminocarbonyl or di-(C₁₋₃-alkyl)-aminocarbonyl group, wherein in each case the oxygen atom of the carbonyl group is replaced by an imino
30 group,

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R₂ denotes a hydrogen, fluorine, chlorine or bromine atom, a C₁₋₃-alkyl, hydroxy or C₁₋₃-alkoxy group,

R₃ denotes a hydrogen atom or a C₁₋₃-alkyl group,

R₄ denotes a hydrogen atom or a methyl or ethyl group
5 optionally substituted by a carboxy or C₁₋₃-alkoxycarbonyl group and

R₅ denotes a cyano group or an amidino group optionally substituted by a C₁₋₆-alkoxycarbonyl or benzoyl group,

wherein

10 by the abovementioned heteroaryl groups is meant a 5-membered heteroaryl group optionally substituted by a C₁₋₃-alkyl group which comprises, in the heteroaromatic moiety,

an imino group optionally substituted by a C₁₋₃-alkyl group, or an oxygen or sulphur atom,

15 an imino group optionally substituted by a C₁₋₃-alkyl group and an oxygen, sulphur or nitrogen atom,

an imino group optionally substituted by a C₁₋₃-alkyl group and two nitrogen atoms or

an oxygen or sulphur atom and two nitrogen atoms,

20 or a 6-membered heteroarylene group optionally substituted by a C₁₋₃-alkyl group which comprises one or two nitrogen atoms in the heteroaromatic moiety,

by the abovementioned heteroaryloxy groups is meant a heteroaryl group, as defined above, which is bound through
25 an oxygen atom,

the carboxy groups mentioned in the definition of the abovementioned groups is optionally replaced by a group which

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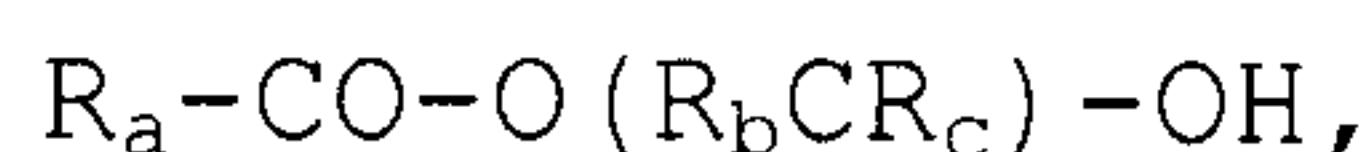
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is converted into a carboxy group *in vivo* or by a group which is negatively charged under physiological conditions or

the amino and imino groups mentioned in the definition of the abovementioned groups is optionally replaced by a group
5 which may be cleaved *in vivo*,

wherein said group which is converted into a carboxy group is selected from the group consisting of: a hydroxymethyl group, a carboxy group esterified with an alcohol wherein the alcoholic moiety denotes a C₁₋₆-alkanol, a
10 phenyl-C₁₋₃-alkanol, a C₃₋₉-cycloalkanol, wherein the cycloalkanol is optionally substituted by one or two C₁₋₃-alkyl groups when the cycloalkanol is a C₅₋₈-cycloalkanol, the C₅₋₈-cycloalkanol wherein a methylene group in the 3 or 4 position is replaced by an oxygen atom
15 or by an imino group optionally substituted by a C₁₋₃-alkyl, phenyl-C₁₋₃-alkyl, phenyl-C₁₋₃-alkoxycarbonyl or C₂₋₆-alkanoyl group and the cycloalkanol moiety is optionally further substituted by one or two C₁₋₃-alkyl groups, a C₄₋₇-cycloalkenol, a C₃₋₅-alkenol, a phenyl-C₃₋₅-alkenol, a
20 C₃₋₅-alkynol or phenyl;

C₃₋₅-alkynol, with the proviso that no bond to the oxygen atom starts from a carbon atom which carries a double or triple bond; a C₃₋₈-cycloalkyl-C₁₋₃-alkanol, a bicycloalkanol having a total of 8 to 10 carbon atoms optionally
25 substituted by one or two C₁₋₃-alkyl groups in the bicycloalkyl moiety, a 1,3-dihydro-3-oxo-1-isobenzofuranol and an alcohol of formula



wherein

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R_a denotes a C_{1-8} -alkyl, C_{5-7} -cycloalkyl, phenyl or phenyl- C_{1-3} -alkyl group,

R_b denotes a hydrogen atom, a C_{1-3} -alkyl, C_{5-7} -cycloalkyl or phenyl group and

5 R_c denotes a hydrogen atom or a C_{1-3} -alkyl group,

and wherein said group that is negatively charged under physiological conditions is selected from the group consisting of: a tetrazol-5-yl, phenylcarbonylaminocarbonyl, trifluoromethylcarbonylaminocarbonyl,

10 C_{1-6} -alkylsulphonylamino, phenylsulphonylamino, benzylsulphonylamino, trifluoromethylsulphonylamino, C_{1-6} -alkylsulphonylaminocarbonyl, phenylsulphonylaminocarbonyl, benzylsulphonylaminocarbonyl and perfluoro- C_{1-6} -alkylsulphonylaminocarbonyl group, and

15 and wherein said amino and imino groups are replaced by a group that is cleaved *in vivo* is selected from the group consisting of a hydroxy group, an acyl group, a pyridinoyl group, a C_{1-16} -alkanoyl group, a 3,3,3-trichloropropionyl, a allyloxycarbonyl group, a C_{1-16} -alkoxycarbonyl group, a

20 C_{1-16} -alkylcarbonyloxy group wherein hydrogen atoms are optionally wholly or partially replaced by fluorine or chlorine atoms, a phenyl- C_{1-6} -alkoxycarbonyl group; a 3-amino-propionyl group wherein the amino group is optionally mono or disubstituted by C_{1-6} -alkyl or

25 C_{3-7} -cycloalkyl groups wherein the substituents are identical or different; a C_{1-3} -alkylsulphonyl- C_{2-4} -alkoxycarbonyl, C_{1-3} -alkoxy- C_{2-4} -alkoxy- C_{2-4} -alkoxycarbonyl, R_a -CO-O- (R_bCR_c) -O-CO, C_{1-6} -alkyl-CO-NH- (R_dCR_c) -O-CO and C_{1-6} -alkyl-CO-O- (R_dCR_e) - (R_dCR_c) -O-CO group wherein R_a to R_c are as hereinbefore

30 defined,

and wherein

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R_d and R_c , which may be identical or different, denote hydrogen atoms or C_{1-3} -alkyl groups;

or an isomer or salt thereof.

Preferred compounds of the above general formula I are those wherein

one of the groups m or n denotes the number 0 and the other group m or n denotes the number 1,

Ar denotes a phenylene group optionally substituted by a fluorine, chlorine or bromine atom or by a methyl, hydroxy, methoxy or benzyloxy group, which may be substituted by another methyl group,

R₁ denotes a phenyl group optionally substituted by a fluorine, chlorine or bromine atom or by a trifluoromethyl, aminosulphonyl, C₁₋₃-alkyl or C₁₋₃-alkoxy group, which may additionally be substituted by a fluorine, chlorine or bromine atom or by a trifluoromethyl, C₁₋₃-alkyl or C₁₋₃-alkoxy group,

a methyl group substituted by a dimethylamino, pyrrolidino or imidazolyl group, wherein the imidazolyl moiety may be substituted by a methyl group,

an amino, C₁₋₅-alkylamino, cyclopentylamino or benzylamino group which may be substituted at the amino-nitrogen atom by a carboxy-C₁₋₂-alkyl, C₁₋₃-alkoxycarbonyl-C₁₋₂-alkyl, carboxy-C₁₋₂-alkylcarbonyl or C₁₋₃-alkoxycarbonyl-C₁₋₂-alkylcarbonyl group,

a benzoylamino or phenylsulphonylamino group,

a cyclopropyl group which is substituted in the 1 position by a 5- to 7-membered cycloalkyleneiminocarbonyl group,

an optionally methyl-substituted pyrrolidinocarbonyl, piperidinocarbonyl, pyrrolidinosulphonyl or piperidinosulphonyl group,

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a C₁₋₃-alkoxy group wherein the alkoxy moiety in the 2 or 3 position may be substituted in each case by an amino, C₁₋₃-alkylamino or di-(C₁₋₃-alkyl)-amino group,

a phenyl-C₁₋₃-alkoxy or pyridinyloxy group,

a C₅₋₇-cycloalkoxy group wherein the methylene group in the 3 or 4 position may be replaced by an -NH group, whilst the -NH group may be substituted

by a C₁₋₃-alkyl or C₂₋₃-alkanoyl group,

by a C₂₋₃-alkanoyl or aminocarbonyl group wherein in each case the oxygen atom of the carbonyl group is replaced by an imino group,

R₂ denotes a hydrogen, fluorine, chlorine or bromine atom, a methyl, hydroxy or methoxy group,

R₃ denotes a hydrogen atom or a methyl group,

R₄ denotes a hydrogen atom or a methyl or ethyl group optionally substituted by a carboxy or C₁₋₃-alkoxycarbonyl group and

R₅ denotes a cyano group or an amidino group optionally substituted by a C₁₋₆-alkoxycarbonyl or benzoyl group,

the isomers thereof and the salts thereof.

Particularly preferred compounds of general formula I are those wherein

one of the groups m or n denotes the number 0 and the other group m or n denotes the number 1,

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Ar denotes a phenylene group optionally substituted by a methyl, hydroxy, methoxy or benzyloxy group,

R₁ denotes a phenyl group optionally substituted by a fluorine, chlorine or bromine atom or by a trifluoromethyl, aminosulphonyl, C₁₋₃-alkyl or C₁₋₃-alkoxy group, which may additionally be substituted by a fluorine, chlorine or bromine atom or by a trifluoromethyl, C₁₋₃-alkyl or C₁₋₃-alkoxy group,

a cyclopropyl group which is substituted in the 1 position by a 5- to 7-membered cycloalkyleneiminocarbonyl group, or a 4- to 7-membered cycloalkyleneiminocarbonyl group,

an optionally methyl-substituted pyrrolidinocarbonyl, piperidinocarbonyl or pyrrolidinosulphonyl group,

R₂ denotes a hydrogen, fluorine, chlorine or bromine atom or a methyl group,

R₃ denotes a hydrogen atom or a methyl group,

R₄ denotes a hydrogen atom or a methyl or ethyl group substituted by a carboxy, methoxycarbonyl or ethoxycarbonyl group and

R₅ denotes an amidino group optionally substituted by a C₁₋₆-alkoxycarbonyl or benzoyl group,

the isomers thereof and the salts thereof.

The following compounds may be mentioned by way of example:

(a) 2-(5-carbamimidoyl-2-hydroxy-phenyl)-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide,

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(b) 2-(2-benzyloxy-5-carbamimidoyl-phenyl)-N-(2-ethoxycarbonyl-ethyl)-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide,

(c) 2-(2-hydroxy-5-carbamimidoyl-phenyl)-N-(2-ethoxycarbonylethyl)-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide,

(d) 2-(2-hydroxy-5-carbamimidoyl-phenyl)-N-(2-carboxy-ethyl)-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide,

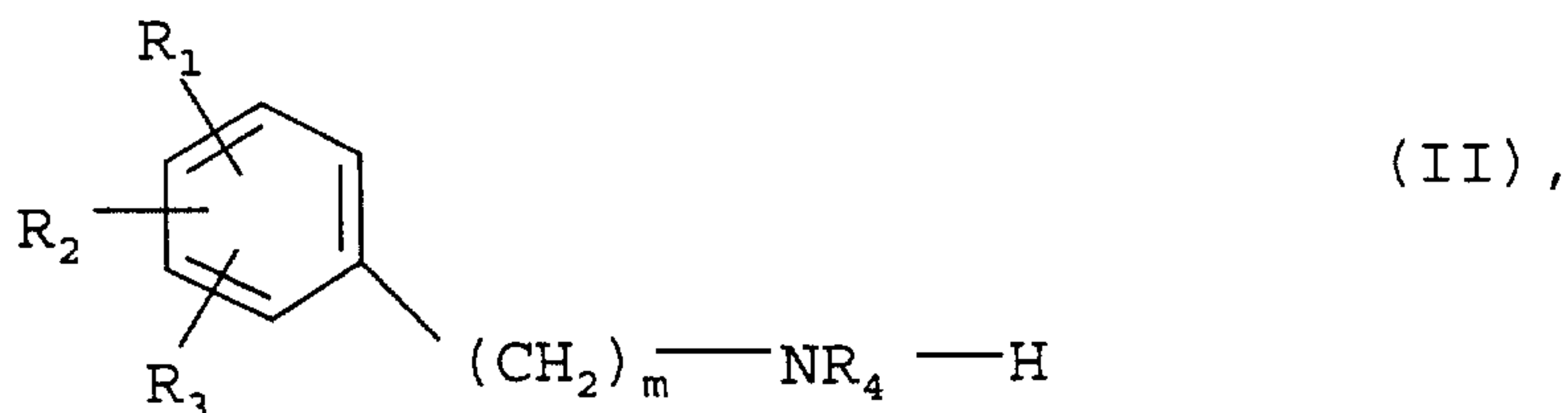
(e) 2-(5-carbamimidoyl-2-hydroxy-phenyl)-N-[3-methyl-4-(piperidin-1-yl-carbonyl)-phenyl]-acetamide and

(f) 2-(5-carbamimidoyl-2-hydroxy-phenyl)-N-[3-methyl-4-(2-aminosulphonyl-phenyl)-phenyl]-acetamide,

wherein the amidino group may additionally be substituted by a C_{1-6} -alkoxycarbonyl or benzoyl group, and the salts thereof.

According to the invention, the compounds of general formula I are obtained by methods known per se, e.g. by the following processes:

a) acylation of a compound of general formula



wherein

R_1 to R_4 and m are as hereinbefore defined, with a carboxylic acid of general formula



wherein

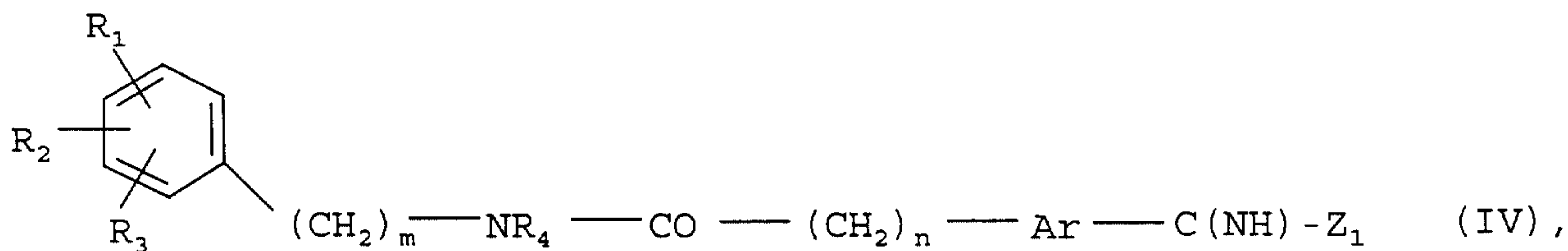
Ar, R₅ and n are as hereinbefore defined, or with the reactive derivatives thereof.

The acylation is conveniently carried out with a corresponding halide or anhydride in a solvent such as methylene chloride, chloroform, carbon tetrachloride, ether, tetrahydrofuran, dioxane, benzene, toluene, acetonitrile or sulpholane, optionally in the presence of an inorganic or organic base at temperatures between -20 and 200°C, but preferably at temperatures between -10 and 160°C.

The acylation may, however, also be carried out with the free acid, optionally in the presence of an acid-activating agent or a dehydrating agent, e.g. in the presence of isobutyl chloroformate, thionyl chloride, trimethylchlorosilane, hydrogen chloride, sulphuric acid, methanesulphonic acid, p-toluenesulphonic acid, phosphorus trichloride, phosphorus pentoxide, N,N'-dicyclohexylcarbodiimide, N,N'-dicyclohexylcarbodiimide/N-hydroxysuccinimide or 1-hydroxy-benzotriazole, N,N'-carbonyldiimidazole or N,N'-thionyl-diimidazole or triphenylphosphine/carbon tetrachloride, at temperatures between -20 and 200°C, but preferably at temperatures between -10 and 160°C.

b) In order to prepare a compound of general formula I wherein R₅ denotes an amidino group which may be substituted by one or two C₁₋₃-alkyl groups:

reaction of a compound of general formula



optionally formed in the reaction mixture wherein.

R_1 to R_4 , Ar and n are as hereinbefore defined and Z_1 denotes an alkoxy or aralkoxy group such as the methoxy, ethoxy, n-propoxy, isopropoxy or benzyloxy group or an alkylthio or aralkylthio group such as the methylthio, ethylthio, n-propylthio or benzylthio group, with an amine of general formula



wherein

R_6 and R_7 , which may be identical or different, each denote a hydrogen atom or a C_{1-3} -alkyl group, or with the salts thereof.

The reaction is conveniently carried out in a solvent such as methanol, ethanol, n-propanol, tetrahydrofuran or dioxane at temperatures between 0 and 150°C, preferably at temperatures between 0 and 80°C, with an amine of general formula V or with a corresponding acid addition salt such as, for example, ammonium carbonate or ammonium acetate.

A compound of general formula IV is obtained, for example, by reacting a corresponding cyano compound with a corresponding alcohol such as methanol, ethanol, n-propanol, isopropanol or benzylalcohol in the presence of an acid such as hydrochloric acid or by reacting a corresponding amide with a trialkyloxonium salt such as triethyloxonium tetrafluoroborate in a solvent such as methylene chloride, tetrahydrofuran or dioxane at temperatures between 0 and 50°C, but preferably at 20°C, or a corresponding nitrile with hydrogen sulphide, conveniently in a solvent such as pyridine or dimethylformamide and in the presence of a base such as triethylamine, and subsequently alkylating the thioamide formed with a corresponding alkyl or aralkyl halide.

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If according to the invention a compound of general formula I is obtained which contains an amino or imino group, this can subsequently be converted with a corresponding acyl derivative into a corresponding acyl compound of general formula I, and/or

if a compound of general formula I is obtained which contains an esterified carboxy group, this can be converted by hydrolysis into a corresponding carboxylic acid of general formula I, and/or

if a compound of general formula I is obtained which contains a carboxy group, this can be converted by esterification into a corresponding ester.

The subsequent acylation is conveniently carried out with a corresponding halide or anhydride in a solvent such as methylene chloride, chloroform, carbon tetrachloride, ether, tetrahydrofuran, dioxane, benzene, toluene, acetonitrile or sulpholane, optionally in the presence of an inorganic or organic base at temperatures between -20 and 200°C, but preferably at temperatures between -10 and 160°C. This may also, however, be carried out with the free acid, optionally in the presence of an acid-activating agent or a dehydrating agent, e.g. in the presence of isobutyl chloroformate, thionyl chloride, trimethylchlorosilane, hydrogen chloride, sulphuric acid, methanesulphonic acid, p-toluenesulphonic acid, phosphorus trichloride, phosphorus pentoxide, N,N'-dicyclohexylcarbodiimide, N,N'-dicyclohexylcarbodiimide/N-hydroxysuccinimide or 1-hydroxy-benzotriazole, N,N'-carbonyldiimidazole or N,N'-thionyl diimidazole or triphenylphosphine/carbon tetrachloride, at temperatures between -20 and 200°C, but preferably at temperatures between -10 and 160°C.

The subsequent hydrolysis is conveniently carried out either in the presence of an acid such as hydrochloric acid,

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sulphuric acid, phosphoric acid, acetic acid, trichloroacetic acid, trifluoroacetic acid or mixtures thereof or in the presence of a base such as lithium hydroxide, sodium hydroxide or potassium hydroxide in a suitable solvent such as water, water/methanol, water/ethanol, water/isopropanol, methanol, ethanol, water/tetrahydrofuran or water/dioxane and the subsequent decarboxylation is carried out in the presence of an acid as hereinbefore described at temperatures between -10 and 120°C, e.g. at temperatures between ambient temperature and the boiling temperature of the reaction mixture.

The subsequent esterification is carried out with a corresponding alcohol, conveniently in a solvent or mixture of solvents such as methylene chloride, benzene, toluene, chlorobenzene, tetrahydrofuran, benzene/tetrahydrofuran or dioxane, but preferably in an excess of the alcohol used, optionally in the presence of an acid such as hydrochloric acid or in the presence of a dehydrating agent, e.g. in the presence of isobutyl chloroformate, thionyl chloride, trimethylchlorosilane, hydrochloric acid, sulphuric acid, methanesulphonic acid, p-toluenesulphonic acid, phosphorus trichloride, phosphorus pentoxide, N,N'-dicyclohexylcarbodiimide, N,N'-dicyclohexylcarbodiimide/N-hydroxysuccinimide, N,N'-carbonyldiimidazole or N,N'-thionyl-diimidazole, triphenylphosphine/carbon tetrachloride or triphenylphosphine/diethyl azodicarboxylate optionally in the presence of a base such as potassium carbonate, N-ethyl-diisopropylamine or N,N-dimethylamino-pyridine conveniently at temperatures between 0 and 150°C, preferably at temperatures between 0 and 80°C, or with a corresponding halide in a solvent such as methylene chloride, tetrahydrofuran, dioxane, dimethylsulphoxide, dimethylformamide or acetone optionally in the presence of a reaction accelerator such as sodium or potassium iodide and preferably in the presence of a base such as sodium carbonate or potassium carbonate or in the presence of a tertiary organic base such as N-ethyl-diisopropylamine or

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N-methyl-morpholine, which may simultaneously serve as solvent, or optionally in the presence of silver carbonate or silver oxide at temperatures between -30 and 100°C, but preferably at temperatures between -10 and 80°C.

In the reactions described hereinbefore, any reactive groups present such as hydroxy, carboxy, amino, alkylamino or imino groups may be protected during the reaction by conventional protecting groups which are cleaved again after the reaction.

For example, a protecting group for a hydroxy group may be a methoxy, benzyloxy, trimethylsilyl, acetyl, benzoyl, tert.butyl, trityl, benzyl or tetrahydropyranyl group,

protecting groups for a carboxy group may be a trimethylsilyl, methyl, ethyl, tert.butyl, benzyl or tetrahydropyranyl group and

protecting groups for an amino, alkylamino or imino group may be an acetyl, trifluoroacetyl, benzoyl, ethoxycarbonyl, tert.butoxycarbonyl, benzyloxycarbonyl, benzyl, methoxybenzyl or 2,4-dimethoxybenzyl group and additionally, for the amino group, a phthalyl group.

Any protecting group used is optionally subsequently cleaved for example by hydrolysis in an aqueous solvent, e.g. in water, isopropanol/water, tetrahydrofuran/water or dioxan/water, in the presence of an acid such as trifluoroacetic acid, hydrochloric acid or sulphuric acid or in the presence of an alkali metal base such as lithium hydroxide, sodium hydroxide or potassium hydroxide or by ether splitting, e.g. in the presence of iodotrimethylsilane, at temperatures between 0 and 100°C, preferably at temperatures between 10 and 50°C.

However, a benzyl, methoxybenzyl or benzyloxycarbonyl group is cleaved, for example, hydrogenolytically, e.g. with hydrogen in the presence of a catalyst such as palladium/charcoal in a

solvent such as methanol, ethanol, ethyl acetate, dimethylformamide, dimethylformamide/acetone or glacial acetic acid, optionally with the addition of an acid such as hydrochloric acid at temperatures between 0 and 50°C, but preferably at ambient temperature, and at a hydrogen pressure of 1 to 7 bar, but preferably 3 to 5 bar.

A methoxybenzyl group may also be cleaved in the presence of an oxidising agent such as cerium(IV) ammonium nitrate in a solvent such as methylene chloride, acetonitrile or acetonitrile/water at temperatures between 0 and 50°C, but preferably at ambient temperature.

A methoxy may conveniently be cleaved in the presence of boron tribromide in a solvent such as methylene chloride at temperatures between -35 and -25°C.

A 2,4-dimethoxybenzyl group, however, is preferably cleaved in trifluoroacetic acid in the presence of anisole.

A tert.butyl or tert.butyloxycarbonyl group is preferably cleaved by treating with an acid such as trifluoroacetic acid or hydrochloric acid or by treating with iodotrimethylsilane, optionally using a solvent such as methylene chloride, dioxane, methanol or diethylether.

A phthalyl group is preferably cleaved in the presence of hydrazine or a primary amine such as methylamine, ethylamine or n-butylamine in a solvent such as methanol, ethanol, isopropanol, toluene/water or dioxane at temperatures between 20 and 50°C.

An allyloxycarbonyl group is cleaved by treating with a catalytic amount of tetrakis-(triphenylphosphine)-palladium(0), preferably in a solvent such as tetrahydrofuran and preferably in the presence of an excess of a base such as

morpholine or 1,3-dimedone at temperatures between 0 and 100°C, preferably at ambient temperature and under inert gas, or by treating with a catalytic amount of tris-(triphenylphosphine)-rhodium(I)chloride in a solvent such as aqueous ethanol and optionally in the presence of a base such as 1,4-diazabicyclo[2.2.2]octane at temperatures between 20 and 70°C.

The compounds of general formulae II to V used as starting materials, some of which are known from the literature, are obtained by methods known from the literature, and furthermore their preparation is described in the Examples.

The chemistry of the compounds of general formula II is described, for example, by Schröter in Stickstoffverbindungen [Nitrogen compounds] II, pages 341-730, Methoden der organischen Chemie (Houben-Weyl), 4th edition, published by Thieme, Stuttgart 1957, and those of general formula III are described by J.F. Hartwig in Angew. Chem. 110, 2154-2157 (1998).

Moreover, the compounds of general formula I obtained may be resolved into their enantiomers and/or diastereomers.

Thus, for example, the compounds of general formula I obtained which occur as racemates may be separated by methods known per se (cf. Allinger N. L. and Eliel E. L. in "Topics in Stereochemistry", Vol. 6, Wiley Interscience, 1971) into their optical enantiomers and compounds of general formula I with at least 2 asymmetric carbon atoms may be resolved into their diastereomers on the basis of their physical-chemical differences using methods known per se, e.g. by chromatography and/or fractional crystallisation, and, if these compounds are obtained in racemic form, they may subsequently be resolved into the enantiomers as mentioned above.

The enantiomers are preferably separated by column separation on chiral phases or by recrystallisation from an optically active solvent or by reacting with an optically active substance which forms salts or derivatives such as e.g. esters or amides with the racemic compound, particularly acids and the activated derivatives or alcohols thereof, and separating the diastereomeric mixture of salts or derivatives thus obtained, e.g. on the basis of their differences in solubility, whilst the free antipodes may be released from the pure diastereomeric salts or derivatives by the action of suitable agents. Optically active acids in common use are e.g. the D- and L-forms of tartaric acid or dibenzoyltartaric acid, di-o-tolyltartaric acid, malic acid, mandelic acid, camphorsulphonic acid, glutamic acid, aspartic acid or quinic acid. An optically active alcohol may be for example (+) or (-)-menthol and an optically active acyl group in amides, for example, may be a (+)- or (-)-menthyloxycarbonyl.

Furthermore, the compounds of formula I may be converted into the salts thereof, particularly for pharmaceutical use into the physiologically acceptable salts with inorganic or organic acids. Acids which may be used for this purpose include for example hydrochloric acid, hydrobromic acid, sulphuric acid, methanesulphonic acid, phosphoric acid, fumaric acid, succinic acid, lactic acid, citric acid, tartaric acid or maleic acid.

Moreover, if the new compounds of formula I contain a carboxy group, they may subsequently, if desired, be converted into the salts thereof with inorganic or organic bases, particularly for pharmaceutical use into the physiologically acceptable salts thereof. Suitable bases for this purpose include for example sodium hydroxide, potassium hydroxide, cyclohexylamine, ethanolamine, diethanolamine and triethanolamine.

As already mentioned, the new compounds of general formula I and the salts thereof have valuable properties. Thus, the compounds of general formula I wherein R₅ denotes a cyano group

are valuable intermediate products for preparing the other compounds of general formula I and the compounds of general formula I wherein R₅ denotes one of the abovementioned amidino groups and the tautomers, the stereoisomers and the physiologically acceptable salts thereof have valuable pharmacological properties, particularly an antithrombotic activity which is preferably based on an effect on thrombin or factor Xa, on a prolonging effect on aPTT time and on an inhibitory effect on related serine proteases such as e.g. trypsin, urokinase factor VIIa, factor IX, factor XI and factor XII.

For example, the compounds

A = 2-(5-carbamimidoyl-2-hydroxy-phenyl)-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide-hydrochloride,

B = 2-(2-benzyloxy-5-carbamimidoyl-phenyl)-N-(2-ethoxycarbonyl-ethyl)-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide-hydrochloride,

C = 2-(2-hydroxy-5-carbamimidoyl-phenyl)-N-(2-ethoxycarbonyl-ethyl)-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide-hydrochloride,

D = 2-(2-hydroxy-5-carbamimidoyl-phenyl)-N-(2-carboxy-ethyl)-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide-hydrochloride

E = 2-(5-carbamimidoyl-2-hydroxy-phenyl)-N-[3-methyl-4-(piperidin-1-yl-carbonyl)-phenyl]-acetamide-hydrochloride and

F = 2-(5-carbamimidoyl-2-hydroxy-phenyl)-N-[3-methyl-4-(2-aminosulphonyl-phenyl)-phenyl]-acetamide-hydrochloride,

were investigated for their effect on the inhibition of factor Xa as follows:

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Method: Enzyme-kinetic measurement with chromogenic substrate. The quantity of anp-nitroaniline (pNA) released from the colourless chromogenic substrate by human factor Xa is determined photometrically at 405 nm. It is proportional to the activity of the enzyme used. The inhibition of the enzyme activity by the test substance I (in relation to the solvent control) is determined at various concentrations of test substance and from this the IC_{50} is calculated, as the concentration which inhibits the factor Xa used by 50 %.

Material:

Tris(hydroxymethyl)-aminomethane buffer (100 mmol) of and sodium chloride (150 mMol), pH 8.0

Factor Xa (Roche), spec. activity: 10 U/0.5 ml, final concentration: 0.175 U/ml per reaction mixture

Substrate Chromozym X (Roche), final concentration: 200 μ Mol/l per reaction mixture

Test substance: final concentration 100, 30, 10, 3, 1, 0.3, 0.1, 0.03, 0.01, 0.003, 0.001 μ Mol/l

Procedure: 10 μ l of a 23.5-times concentrated starting solution of the test substance or solvent (control), 175 μ l of tris(hydroxymethyl)-aminomethane buffer and 25 μ l of Factor Xa working solution of 1.65 U/ml are incubated for 10 minutes at 37°C. After the addition of 25 μ l of Chromozym X working solution (1.88 μ Mol/l) the sample is measured in a photometer (SpectraMax 250) at 405 nm for 150 seconds at 37°C.

Evaluation:

1. Determining the maximum increase ($\Delta OD/\text{minutes}$) over 3 measuring points.

2. Determining the % inhibition based on the solvent control.
3. Plotting a dosage/activity curve (% inhibition vs substance concentration).
4. Determining the IC_{50} by interpolating the X value (substance concentration) of the dosage/activity curve at Y = 50 % inhibition.

The following Table shows the results obtained:

substance	inhibition of factor Xa (IC_{50} in μM)
A	0.030
B	0.680
C	0.120
D	0.850
E	0.085
F	0.260

The compounds prepared according to the invention are well tolerated, as no toxic side effects could be observed at therapeutic doses.

In view of their pharmacological properties the new compounds and the physiologically acceptable salts thereof are suitable for the prevention and treatment of venous and arterial thrombotic diseases, such as for example the treatment of deep leg vein thrombosis, for preventing reocclusions after bypass operations or angioplasty (PT(C)A), and occlusion in peripheral arterial diseases such as pulmonary embolism, disseminated intravascular coagulation, for preventing coronary thrombosis, stroke and the occlusion of shunts. In addition, the compounds according to the invention are suitable for antithrombotic support in thrombolytic treatment,

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such as for example with rt-PA or streptokinase, for preventing long-term restenosis after PT(C)A, for preventing metastasis and the growth of clot-dependent tumours and fibrin-dependent inflammatory processes, e.g. in the treatment of pulmonary fibrosis.

The dosage required to achieve such an effect is appropriately 0.1 to 30 mg/kg, preferably 0.3 to 10 mg/kg by intravenous route, and 0.1 to 50 mg/kg, preferably 0.3 to 30 mg/kg by oral route, in each case administered 1 to 4 times a day. For this purpose, the compounds of formula I prepared according to the invention may be formulated, optionally together with other active substances, with one or more inert conventional carriers and/or diluents, e.g. with corn starch, lactose, glucose, microcrystalline cellulose, magnesium stearate, polyvinylpyrrolidone, citric acid, tartaric acid, water, water/ethanol, water/glycerol, water/sorbitol, water/polyethyleneglycol, propyleneglycol, cetylstearyl alcohol, carboxymethylcellulose or fatty substances such as hard fat or suitable mixtures thereof, to produce conventional galenic preparations such as plain or coated tablets, capsules, powders, suspensions or suppositories.

The Examples which follow are intended to illustrate the invention:

Example 12-(3-carbamimidoyl-phenyl)-N-[2-chloro-5-(1-(pyrrolidin-1-yl-carbonyl)-cyclopropyl)-phenyl]-acetamide-hydrochloridea. 1-(4-chloro-3-nitro-phenyl)-cyclopropanecarboxylic acid

350 ml of fuming nitric acid are combined at -25 to -30°C with 50.0 g (0.21 mol) of 1-(4-chloro-phenyl)-cyclopropanecarboxylic acid in batches. After it has all been added the mixture is stirred for a further 15 minutes at -25°C and then poured onto ice. The substance precipitated is suction filtered, washed with water and dried.

Yield: 58.5 g (95 % of theory),

R_f value: 0.43 (silica gel; methylene chloride/methanol = 9.5:0.5)

b. 5-[1-(pyrrolidin-1-yl-carbonyl)-cyclopropyl]-2-chloro-nitrobenzene

2.4 g (0.01 mol) of 1-(4-chloro-3-nitro-phenyl)-cyclopropanecarboxylic acid are dissolved in 25 ml of tetrahydrofuran and after the addition of 3.2 g (0.01 mol) of O-(benzotriazol-1-yl)-N,N,N',N'-tetramethyluronium tetrafluoroborate, 1.1 ml of (0.01 mol) of N-methyl-morpholine and 1.0 ml of (0.012 mol) of pyrrolidine stirred for 16 hours at ambient temperature. The solvent is distilled off, the residue is poured onto ice water, made alkaline with ammonia and extracted with ethyl acetate. The organic phase is dried and evaporated down.

Yield: 2.5 g (85 % of theory),

R_f value: 0.18 (silica gel; cyclohexane/ethyl acetate = 1:1)

c. 5-[1-(pyrrolidin-1-yl-carbonyl)-cyclopropyl]-2-chloro-aniline

1.8 g (8.14 mmol) of 5-[1-(pyrrolidin-1-yl-carbonyl)-cyclopropyl]-2-chloro-nitrobenzene are dissolved in 30 ml of ethyl acetate and 30 ml of ethanol and after the addition of 0.8 g palladium on active charcoal (10%) hydrogenated for 3

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hours at ambient temperature with hydrogen. Then the catalyst is filtered off and the filtrate is evaporated down.

Yield: 2.0 g (92.8 % of theory),

R_f value: 0.24 (silica gel; cyclohexane/ethyl acetate/ammonia = 1:1:0.01)

C₁₄H₁₇ClN₂O (264.75)

mass spectrum: M⁺ = 264/6 (Cl)

d. 2-(3-cyano-phenyl)-N-[2-chloro-5-(1-(pyrrolidin-1-yl-carbonyl)-cyclopropyl)-phenyl]-acetamide

Prepared analogously to Example 1b from 5-[1-(pyrrolidin-1-yl-carbonyl)-cyclopropyl]-2-chloro-aniline, O-(benzotriazol-1-yl)-N,N,N',N'-tetramethyluronium tetrafluoroborate, N-methyl-morpholine and 3-cyanophenylacetic acid in dimethylformamide. Yield: 43 % of theory,

R_f value: 0.21 (silica gel; cyclohexane/ethyl acetate = 1:2)

e. 2-(3-carbamimidoyl-phenyl)-N-[2-chloro-5-(1-(pyrrolidin-1-yl-carbonyl)-cyclopropyl)-phenyl]-acetamide-hydrochloride

400 mg (0.1 mmol) of 2-(3-cyano-phenyl)-N-[2-chloro-5-(1-(pyrrolidin-1-yl-carbonyl)-cyclopropyl)-phenyl]-acetamide are dissolved in 60 ml of saturated ethanolic hydrochloric acid and stirred for 17 hours at ambient temperature. The solvent is distilled off, the residue is dissolved in 50 ml of absolute ethanol and mixed with 1.5 g (15.6 mmol) of ammonium carbonate. After 22 hours at ambient temperature the mixture is evaporated to dryness. The residue is chromatographed on silica gel, eluting with methylene chloride/methanol/glacial acetic acid (9:1:0.01).

Yield: 50 mg (11 % of theory),

R_f value: 0.59 (silica gel; methylene chloride/methanol/ammonia = 4:1:0.01)

C₂₃H₂₅ClN₄O₂ x HCl (424.94/461.4)

mass spectrum: (M+H)⁺ = 425/7 (Cl)

Example 23-carbamimidoyl-N-[3-(1-(pyrrolidin-1-yl-carbonyl)-cyclopropyl)-benzyl]-benzamide-hydrochloridea. 1-(3-bromo-phenyl)-1-cyclopropane-nitril

25 g (0.13 mol) of 3-bromo-benzylcyanide are taken up in 32 ml (0.38 mol) of 1-bromo-2-chloro-ethane and combined with 0.6 g (2.6 mmol) of benzyltriethylammonium chloride. Then a solution of 105.8 g (2.65 mol) of sodium hydroxide in 106 ml of water is added dropwise at 10 to 25°C. After 20 hours at 55°C the reaction solution is poured onto ice water and extracted with ethyl acetate. The organic extracts are dried and evaporated down. The residue is triturated with petroleum ether, suction filtered and dried.

Yield: 19.3 g (68 % of theory),

R_f value: 0.69 (petroleum ether/ethyl acetate = 4:1)

b. 1-(3-bromo-phenyl)-cyclopropanecarboxylic acid

7.6 g (0.135 mol) of potassium hydroxide are dissolved in 60 ml of ethyleneglycol, combined batchwise with 10.0 g (0.045 mol) of 1-(3-bromo-phenyl)-1-cyclopropane-nitrile and after the addition of 30 ml of water heated to 140°C for 4.5 hours. After cooling it is poured onto 600 ml of ice water and extracted with ether. The aqueous phase is poured onto ice/conc. hydrochloric acid, the product precipitated is suction filtered and dried.

Yield: 10.1 g (93 % of theory),

R_f value: 0.85 (silica gel; cyclohexane/ethyl acetate/glacial acetic acid = 1:1:0.01)

c. 3-[1-(pyrrolidin-1-yl-carbonyl)-cyclopropyl]-bromo-benzene

Prepared analogously to Example 1b from 1-(3-bromo-phenyl)-cyclopropanecarboxylic acid, pyrrolidine, O-(benzotriazol-1-yl)-N,N,N',N'-tetramethyluronium tetrafluoroborate and N-methyl-morpholine in tetrahydrofuran.

Yield: 98 % of theory,

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R_f value: 0.55 (silica gel; cyclohexane/ethyl acetate/glacial acetic acid = 1:1:0.01)

d. 3-[1-(pyrrolidin-1-yl-carbonyl)-cyclopropyl]-benzonitrile

6 g (20.4 mmol) of 3-[1-(pyrrolidin-1-yl-carbonyl)-cyclopropyl]-bromo-benzene are dissolved in 25 ml of dimethylformamide and after the addition of 2.7 g (30.6 mmol) of copper-I-cyanide, 0.3 g (0.216 mmol) of tetrakis-triphenylphosphine-palladium-(0) and 5 g of aluminium oxide stirred for 30 hours at 140°C. The insoluble matter is filtered off and the solution is evaporated down. The residue is chromatographed on silica gel, eluting with cyclohexane/ethyl acetate (1:2).

Yield: 1.8 g (36 % of theory),

R_f value: 0.32 (silica gel; cyclohexane/ethyl acetate/glacial acetic acid = 1:1:0.01)

e. 3-[1-(pyrrolidin-1-yl-carbonyl)-cyclopropyl]-benzylamine

1.8 g (7.5 mmol) of 3-[1-(pyrrolidin-1-yl-carbonyl)-cyclopropyl]-benzonitrile are hydrogenated with hydrogen in 50 ml of methanolic ammonia with the addition of 300 mg Raney nickel for 3 hours at 70°C. Then the catalyst is filtered off and the filtrate is evaporated down.

Yield: 1.8 g (98 % of theory),

R_f value: 0.94 (silica gel; methylene chloride /methanol/ammonia = 4:1:0.01)

f. 3-cyano-N-[3-(1-(pyrrolidin-1-yl-carbonyl)-cyclopropyl)-benzyl]-benzamide

Prepared analogously to Example 1b from 3-[1-(pyrrolidin-1-yl-carbonyl)-cyclopropyl]-benzylamine, 3-cyanobenzoic acid, O-(benzotriazol-1-yl)-N,N,N',N'-tetramethyluronium tetrafluoroborate and N-methyl-morpholine in tetrahydrofuran. Yield: 96 % of theory,

R_f value: 0.56 (silica gel; ethyl acetate/ethanol = 9:1)

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g. 3-carbamimidoyl-N-[3-(1-(pyrrolidin-1-yl-carbonyl)-cyclopropyl)-benzyl]-benzamide-hydrochloride

Prepared analogously to Example 1e from 3-cyano-N-[3-(1-(pyrrolidin-1-yl-carbonyl)-cyclopropyl)-benzyl]-benzamide and hydrochloric acid/ammonium carbonate in ethanol.

Yield: 58 % of theory,

R_f value: 0.19 (Reversed phase RP 8; 5% sodium chloride solution/methanol = 1:1)

C₂₃H₂₆N₄O₂ x HCl (390.48/426.95)

mass spectrum: (M+H)⁺ = 391
(M-H+HCl)⁻ = 425/7 (Cl)

The following compounds are prepared analogously to Example 2:

(1) 3-carbamimidoyl-N-[4-(1-(pyrrolidin-1-yl-carbonyl)-cyclopropyl)-benzyl]-benzamide-hydrochloride

Yield: 68 % of theory,

C₂₃H₂₆N₄O₂ x HCl (390.48/426.95)

mass spectrum: (M+H)⁺ = 391
(M+2H)⁺⁺ = 196

(2) 5-carbamimidoyl-2-hydroxy-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-benzyl]-benzamide-hydrochloride

Yield: 34% of theory,

R_f value: 0.1 (Reversed phase RP8; 5% saline solution/methanol = 1:1)

C₂₁H₂₄N₄O₃ x HCl (380.46/416.91)

mass spectrum: (M+H)⁺ = 381
(M-H)⁻ = 379

Example 3

2-(5-carbamimidoyl-2-hydroxy-phenyl)-N-[2-methyl-5-(1-(pyrrolidin-1-carbonyl)-cyclopropyl)-phenyl]-acetamide-hydrochloride

a. 5-cyano-2-methoxy-phenylacetic acid

Prepared analogously to Example 2d from 5-bromo-2-methoxy-phenylacetic acid, copper-I-cyanide, tetrakis-triphenylphosphine-palladium-(0) and aluminium oxide in dimethylformamide.

Yield: 37 % of theory,

R_f value: 0.26 (silica gel; cyclohexane/ethyl acetate/glacial acetic acid = 1:1:0.01)

b. 2-(5-cyano-2-methoxy-phenyl)-N-[2-methyl-5-(1-(pyrrolidin-1-carbonyl)-cyclopropyl)-phenyl]-acetamide

0.6 g (3.3 mmol) of 5-cyano-2-methoxy-phenylacetic acid are dissolved in 10 ml of dimethylformamide and after the addition of 0.5 g (3.3 mmol) of N,N-carbonyldiimidazole stirred for 10 minutes at ambient temperature. Then 0.8 g (3.3 mmol) of 5-(pyrrolidin-1-yl-carbonyl)-cyclopropyl-2-methyl-aniline are added. The reaction mixture is stirred for 4 hours at 80°C, cooled to ambient temperature, combined with ice water, made alkaline with ammonia and extracted several times with ethyl acetate. The combined organic extracts are dried and evaporated down. The residue is chromatographed on silica gel, eluting with cyclohexane/ethyl acetate (7:3).

Yield: 73 % of theory,

R_f value: 0.30 (silica gel; ethyl acetate)

c. 2-(5-cyano-2-hydroxy-phenyl)-N-[2-methyl-5-(1-(pyrrolidin-1-carbonyl)-cyclopropyl)-phenyl]-acetamide

0.7 g (1.67 mmol) of 2-(5-cyano-2-methoxy-phenyl)-N-[2-methyl-5-(1-(pyrrolidin-1-carbonyl)-cyclopropyl)-phenyl]-acetamide are dissolved in 35 ml of methylene chloride and at -35 to -25°C 10 ml of a 1-molar solution of boron tribromide in

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methylene chloride (10 mmol) are added dropwise. After 1 hour's stirring at 20°C, first ice is added, then 20 ml of 2N hydrochloric acid. The aqueous phase is extracted several times with methylene chloride, the combined organic extracts are dried and evaporated down. The residue is chromatographed on silica gel, eluting with methylene chloride/ethanol (100:1).

Yield: 81 % of theory,

R_f value: 0.14 (silica gel; methylene chloride/ethanol = 49:1)

d. 2-(5-carbamimidoyl-2-hydroxy-phenyl)-N-[2-methyl-5-(1-(pyrrolidin-1-carbonyl)-cyclopropyl)-phenyl]-acetamide-hydrochloride

Prepared analogously to Example 1e from 2-(5-cyano-2-hydroxy-phenyl)-N-[2-methyl-5-(1-(pyrrolidin-1-carbonyl)-cyclopropyl)-phenyl]-acetamide, and hydrochloric acid/ammonium carbonate in ethanol.

Yield: 80 % of theory,

R_f value: 0.39 (silica gel; methylene chloride/methanol/glacial acetic acid = 4:1:0.01)

C₂₄H₂₈N₄O₃ x HCl (420.51/456.98)

mass spectrum: (M+H)⁺ = 421

(M+Cl)⁻ = 455/7 (Cl)

The following compound is prepared analogously to Example 3:

(1) 2-(5-carbamimidoyl-2-methoxy-phenyl)-N-[2-methyl-5-(1-(pyrrolidin-1-carbonyl)-cyclopropyl)-phenyl]-acetamide-hydrochloride

Yield: 92 % of theory,

R_f value: 0.33 (silica gel; methylene chloride/methanol/glacial acetic acid = 4:1:0.01)

C₂₅H₃₀N₄O₃ x HCl (434.55/471.01)

mass spectrum: (M+H)⁺ = 435

Example 4

2-(5-carbamimidoyl-2-hydroxy-phenyl)-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide-hydrochloride

a. 2-(5-cyano-2-methoxy-phenyl)-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide

0.4 g (2 mmol) of 3-methyl-4-(pyrrolidin-1-yl-carbonyl)-aniline are dissolved in 15 ml of tetrahydrofuran and after the addition of 0.3 ml (2 mmol) of triethylamine and 0.4 g (2 mmol) of 5-cyano-3-methoxy-phenylacetic acid chloride stirred for 48 hours at ambient temperature. Then the mixture is combined with water, made alkaline with ammonia and extracted with ethyl acetate. The combined organic extracts are washed with 1N hydrochloric acid, dried and evaporated down.

Yield: 0.45 g (59 % of theory),

R_f value: 0.18 (silica gel; ethyl acetate)

b. 2-(5-carbamimidoyl-2-methoxy-phenyl)-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide-hydrochloride

Prepared analogously to Example 1e from 2-(5-cyano-2-methoxyphenyl)-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide and hydrochloric acid/ammonium carbonate in ethanol.

Yield: 36 % of theory,

R_f value: 0.33 (Reversed phase RP 8; 5% sodium chloride solution/methanol = 1:1)

C₂₂H₂₆N₄O₃ x HCl (394.48/430.94)

mass spectrum: (M+H)⁺ = 395

(M-H+HCl)⁻ = 429

c. 2-(5-carbamimidoyl-2-hydroxy-phenyl)-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide-hydrochloride

Prepared analogously to Example 3c from 2-(5-carbamimidoyl-2-methoxy-phenyl)-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide-hydrochloride and boron tribromide in dichloromethane.

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Yield: 19 % of theory,

R_f value: 0.38 (Reversed phase RP 8; 5% sodium chloride solution/methanol = 1:1)

C₂₁H₂₄N₄O₃ x HCl (380.45/416.91)

mass spectrum: (M+H)⁺ = 381

(M-H)⁻ = 379

The following compounds are prepared analogously to Example 4:

(1) 2-(3-carbamimidoyl-phenyl)-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide-hydrochloride

Yield: 12 % of theory,

C₂₁H₂₄N₄O₂ x HCl (364.45/400.92)

mass spectrum: (M+H)⁺ = 365

(2) 2-(5-carbamimidoyl-2-hydroxy-phenyl)-N-methyl-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide-hydrochloride

Yield: 99 % of theory,

C₂₂H₂₆N₄O₃ x HCl (394.48/430.94)

mass spectrum: (M+H)⁺ = 395

(M-H)⁻ = 393

(3) 2-(5-carbamimidoyl-2-benzyloxy-phenyl)-N-methyl-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide-hydrochloride

Yield: 90 % of theory,

C₂₉H₃₂N₄O₃ x HCl (484.60/521.06)

mass spectrum: (M+H)⁺ = 485

(M-H+HCl)⁻ = 519/21 (Cl)

(4) 2-(5-carbamimidoyl-2-hydroxy-phenyl)-N-[3-methyl-4-(N-(3-ethoxycarbonyl-propionyl)-N-cyclopentyl-amino)-phenyl]-acetamide-hydrochloride

Yield: 74% of theory,

C₂₇H₃₄N₄O₅ x HCl (494.61/531.06)

R_f value: 0.36 (Reversed phase RP8; 5% saline solution/methanol = 4:6)

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mass spectrum: $(M+H)^+ = 495$
 $(M+Cl)^- = 529/531$ (Cl)
 $(M-H)^- = 493$

(5) 2-(5-carbamimidoyl-2-benzyloxy-phenyl)-N-[3-methyl-4-(N-(3-ethoxycarbonyl-propionyl-N-(2-methyl-propyl)-amino)-phenyl]-acetamide-hydrochloride

Yield: 74% of theory,

R_f value: 0.21 (silica gel; methylene chloride/ethanol = 4:1)

$C_{33}H_{40}N_4O_5 \times HCl$ (572.71/609.18)

mass spectrum: $(M+H)^+ = 573$
 $(M-H)^- = 571$

(6) 2-(5-carbamimidoyl-2-hydroxy-phenyl)-N-[3-methyl-4-(N-(3-ethoxycarbonyl-propionyl)-N-(2-methyl-propyl)-amino)-phenyl]-acetamide-hydrochloride

Yield: 100% of theory,

R_f value: 0.33 (Reversed phase RP8; 5% saline solution/methanol = 4:6)

$C_{26}H_{34}N_4O_5 \times HCl$ (482.58/519.05)

mass spectrum: $(M+H)^+ = 483$
 $(M-H)^- = 481$
 $(M+Cl)^- = 517/519$ (Cl)

(7) 2-(5-carbamimidoyl-2-hydroxy-phenyl)-N-[3-methyl-4-(N-ethoxycarbonylacetyl-N-cyclopentyl-amino)-phenyl]-acetamide-hydrochloride

(8) 2-(5-carbamimidoyl-2-hydroxy-phenyl)-N-[3-methyl-4-(pyrrolidin-1-yl-sulphonyl)-phenyl]-acetamide-hydrochloride

Example 5

2-(5-carbamimidoyl-2-hydroxy-phenyl)-N-[2,5-dimethyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide-hydrochloride

a. 2,5-dimethyl-4-(pyrrolidin-1-yl-carbonyl)-bromobenzene

Prepared analogously to Example 1b from 4-bromo-3,5-dimethylbenzoic acid, pyrrolidine, O-(benzotriazol-1-yl)-N,N,N',N'-tetramethyluronium tetrafluoroborate and triethylamine in dimethylformamide.

Yield: 63 % of theory,

R_f value: 0.45 (silica gel; methylene chloride/ethanol = 19:1)

b. 2,5-dimethyl-4-(pyrrolidin-1-yl-carbonyl)-benzylaniline

2.3 g (0.01 mol) of 2,5-dimethyl-4-(pyrrolidin-1-yl-carbonyl)-bromobenzene and 1.3 g (0.012 mol) of benzylamine are dissolved in 25 ml of toluene and after the addition of 4.6 g of caesium carbonate, 100 mg palladium-II-acetate and 200 mg 2,2'-bis(-diphenylphosphino)-1,1'-binaphthyl stirred for 7 hours under an argon atmosphere at 100°C. After cooling the mixture is diluted with ice water and extracted with ethyl acetate. The combined organic extracts are dried and evaporated down. The residue is chromatographed on silica gel, eluting with methylene chloride/ethanol (50:1 and 25:1).

Yield: 60 % of theory,

R_f value: 0.30 (silica gel; methylene chloride/ethanol = 9:1)

c. 2,5-dimethyl-4-(pyrrolidin-1-yl-carbonyl)-aniline

Prepared analogously to Example 1c from 2,5-dimethyl-4-(pyrrolidin-1-yl-carbonyl)-benzylaniline and palladium on active charcoal in methanol.

Yield: 94 % of theory,

R_f value: 0.30 (silica gel; ethyl acetate/petroleum ether = 1:1)

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d. 2-benzyloxy-5-bromo-phenylacetic acid

A solution of 12.4 g (0.053 mol) of 2-hydroxy-5-bromo-phenylacetic acid in 125 ml of dimethylformamide is combined with 14 g (0.125 mol) of potassium tert.butoxide. After 15 minutes at ambient temperature 18.5 g (0.108 mol) of benzylbromide are added. The reaction solution is stirred for 3 hours at ambient temperature, poured onto ice water and extracted with ethyl acetate. The combined organic extracts are dried and evaporated down. The residue is dissolved in 100 ml of ethanol and after the addition of 50 ml of 2N sodium hydroxide solution stirred for 3 hours at ambient temperature. The solvent is distilled off, the residue is adjusted to pH 4 with 2N hydrochloric acid. After extraction with ethyl acetate the organic phases are dried and evaporated down. The residue is chromatographed on silica gel and eluted with petroleum ether/ethyl acetate (8:2).

Yield: 6.7 g (38 % of theory),

R_f value: 0.50 (silica gel; ethyl acetate/petroleum ether = 1:1)

e. 2-benzyloxy-5-cyano-phenylacetic acid

Prepared analogously to Example 2d from 2-benzyloxy-5-bromo-phenylacetic acid, copper-I-cyanide, tetrakis-triphenylphosphine-palladium-(0) and aluminium oxide in dimethylformamide.

Yield: 26 % of theory,

R_f value: 0.45 (silica gel; methylene chloride/ethanol = 19:1)

f. 2-(5-cyano-2-benzyloxy-phenyl)-N-[2,5-dimethyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide

Prepared analogously to Example 1b from 2-benzyloxy-5-cyano-phenylacetic acid, 2,5-dimethyl-4-(pyrrolidin-1-yl-carbonyl)-aniline, O-(benzotriazol-1-yl)-N,N,N',N'-tetramethyluronium-tetrafluoroborate and N-methylmorpholine in tetrahydrofuran.

Yield: 44 % of theory,

R_f value: 0.75 (silica gel; ethyl acetate/ethanol = 9:1)

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g. 2-(5-carbamimidoyl-2-benzyloxy-phenyl)-N-[2,5-dimethyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide

Prepared analogously to Example 1e from 2-(5-cyano-2-benzyloxy-phenyl)-N-[2,5-dimethyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide and hydrochloric acid/ammonium carbonate in ethanol.

Yield: 86 % of theory,

R_f value: 0.20 (silica gel; methylene chloride/ethanol/glacial acetic acid = 8:2:0.01)

h. 2-(5-carbamimidoyl-2-hydroxy-phenyl)-N-[2,5-dimethyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide-hydrochloride

355 mg (0.68 mmol) of 2-(5-carbamimidoyl-2-benzyloxy-phenyl)-N-[2,5-dimethyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide are dissolved in 40 ml of methanol and after the addition of 250 mg of palladium on active charcoal the mixture is hydrogenated with hydrogen for 15 minutes. Then the catalyst is filtered off and the filtrate is evaporated down.

Yield: 145 mg (49 % of theory),

R_f value: 0.10 (silica gel; methylene chloride/ethanol/glacial acetic acid = 8:2:0.01)

C₂₂H₂₆N₄O₃ x HCl (394.48/430.94)

mass spectrum: (M+H)⁺ = 395

(M-H)⁻ = 393

The following compounds are prepared analogously to Example 5:

(1) 2-(5-carbamimidoyl-2-hydroxy-phenyl)-N-[3-methyl-4-(piperidin-1-yl-carbonyl)-phenyl]-acetamide-hydrochloride

Yield: 98% of theory,

R_f value: 0.75 (Reversed phase RP8; 5% saline solution/methanol = 1:4)

C₂₂H₂₆N₄O₃ x HCl (394.49/430.94)

mass spectrum: M⁺ = 395

(M+Cl)⁻ = 429/431 (Cl)

(M-H)⁻ = 393

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(2) 2-(5-carbamimidoyl-2-hydroxy-phenyl)-N-[3-methyl-4-(2-methyl-pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide-hydrochloride

Yield: 100% of theory,

R_f value: 0.7 (Reversed phase RP8; 5% saline solution/methanol = 1:4)

C₂₂H₂₆N₄O₃ x HCl (394.49/430.94)

mass spectrum: M⁺ = 395

(M+Cl)⁺ = 429/431 (Cl)

(M-H)⁺ = 393

Example 6

2-(2-benzyloxy-5-carbamimidoyl-phenyl)-N-(2-ethoxycarbonyl-ethyl)-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide-hydrochloride

a. N-(2-methoxycarbonyl-ethyl)-3-methyl-4-(pyrrolidin-1-yl-carbonyl)-aniline

1.5 g (7.3 mmol) of 3-methyl-4-(pyrrolidin-1-yl-carbonyl)-aniline, 20 ml of (220 mmol) of methyl acrylate, 1 ml (2.2 mmol) of Triton B and 60 mg (0.27 mmol) of 2,5-di-tert.butyl-hydroquinone are stirred for 22 hours at 85°C. Then the reaction mixture is evaporated down, the residue is chromatographed on silica gel, eluting with methylene chloride + 0 to 5 % ethanol.

Yield: 1.6 g (76 % of theory),

R_f value: 0.70 (silica gel; methylene chloride/ethanol = 9:1)

b. 2-(2-benzyloxy-5-cyano-phenyl)-N-(2-ethoxycarbonyl-ethyl)-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide

0.8 g (2.88 mmol) of N-(2-methoxycarbonyl-ethyl)-3-methyl-4-(pyrrolidin-1-yl-carbonyl)-aniline are dissolved in 50 ml of tetrahydrofuran and after the addition of 1.1 ml (7.86 mmol) of triethylamine and 0.8 g (2.62 mmol) of 2-benzyloxy-5-cyano-phenylacetic acid chloride stirred for 8 hours at ambient temperature. Then the mixture is diluted with water and

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extracted with methylene chloride. The combined organic extracts are dried and evaporated down. The residue is chromatographed on silica gel, eluting with methylene chloride.

Yield: 1.0 g (71 % of theory),

R_f value: 0.72 (silica gel; methylene chloride/ethanol = 9:1)

c. 2-(2-benzyloxy-5-carbamimidoyl-phenyl)-N-(2-ethoxycarbonyl-ethyl)-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide-hydrochloride

Prepared analogously to Example 1e from 2-(2-benzyloxy-5-cyano-phenyl)-N-(2-ethoxycarbonyl-ethyl)-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide and hydrochloric acid/ammonium carbonate in ethanol.

Yield: 44 % of theory,

R_f value: 0.17 (silica gel; methylene chloride/ethanol = 4:1)

C₃₃H₃₈N₄O₅ x HCl (570.69/607.16)

mass spectrum: (M+H)⁺ = 571

Example 7

2-(2-hydroxy-5-carbamimidoyl-phenyl)-N-(2-ethoxycarbonyl-ethyl)-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide-hydrochloride

Prepared analogously to Example 5h from 2-(2-benzyloxy-5-carbamimidoyl-phenyl)-N-(2-ethoxycarbonyl-ethyl)-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide-hydrochloride and palladium on active charcoal in methanol.

Yield: 96 % of theory,

R_f value: 0.45 (Reversed phase RP 8; methanol/5% sodium chloride solution = 6:4)

C₂₆H₃₂N₄O₅ x HCl (480.57/517.04)

mass spectrum: (M+H)⁺ = 481

Example 8

2-(2-hydroxy-5-carbamimidoyl-phenyl)-N-(2-carboxy-ethyl)-
N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide-
hydrochloride

0.3 g (0.58 mmol) of 2-(2-hydroxy-5-carbamimidoyl-phenyl)-
N-(2-ethoxycarbonyl-ethyl)-N-[3-methyl-4-(pyrrolidin-1-yl-
carbonyl)-phenyl]-acetamide-hydrochloride are stirred into a
mixture of 3.2 ml (3.2 mmol) of 1-molar lithium hydroxide
solution, 6.2 ml of water and 7.6 ml of tetrahydrofuran for 2
hours at ambient temperature. After the addition of 74 mg
ammonium chloride the solution is evaporated down. The residue
is chromatographed on the reversed phase and eluted with
water.

Yield: 0.2 g (71 % of theory),

R_f value: 0.62 (Reversed phase RP 8; methanol/5% sodium
chloride solution = 6:4)

C₂₄H₂₈N₄O₅ x HCl (452.52/488.97)

mass spectrum: (M+H)⁺ = 453

(M-H)⁻ = 451

The following compounds are prepared analogously to Example 8:

(1) 2-(5-carbamimidoyl-2-hydroxy-phenyl)-N-[3-methyl-
4-(N-(3-carboxypropionyl)-N-cyclopentyl-amino)-phenyl]-
acetamide-hydrochloride

Yield: 83% of theory,

C₂₅H₃₀N₄O₅ x HCl (466.55/503.01)

R_f value: 0.84 (Reversed phase RP8; 5% saline solution/metha-
nol = 6:4)

(2) 2-(5-carbamimidoyl-2-hydroxy-phenyl)-N-[3-methyl-
4-(N-carboxyacetyl-N-cyclopentyl-amino)-phenyl]-acetamide-
hydrochloride

Example 93-carbamimidoyl-N-[4-(pyrrolidin-3-yl-oxy)-benzyl]-benzamide-dihydrochloridea. N-tert.butyloxycarbonyl-3-pyrrolidinol

5.8 g (66.5 mmol) of 3-pyrrolidinol and 6.7 g (67 mmol) of triethylamine are dissolved in 80 ml of methylene chloride and a solution of 15.3 g (70 mmol) of di-tert.butyl-dicarbonate in 40 ml of methylene chloride is added dropwise. After 16 hours at ambient temperature the mixture is stirred with water, the organic phase is dried and evaporated down.

Yield: 12.4 g (100 % of theory),

R_f value: 0.75 (silica gel; ethyl acetate/methanol = 9:1)

b. 4-[(N-tert.butyloxycarbonyl)-pyrrolidin-3-yl-oxy]-benzonitrile

3.8 g (20 mmol) of N-tert.butyloxycarbonyl-3-pyrrolidinol are dissolved in 100 ml of tetrahydrofuran and after the addition of 2.4 g (20 mmol) of 4-hydroxybenzonitrile, 5.7 g (22 mmol) of triphenylphosphine and 3.9 g (22 mmol) of diethyl diethylazodicarboxylate the mixture is stirred for 18 hours at ambient temperature. The solvent is distilled off and the residue is chromatographed on silica gel, eluting with cyclohexane/ethyl acetate (10:5).

Yield: 4.5 g (78 % of theory),

R_f value: 0.40 (silica gel; cyclohexane/ethyl acetate = 10:5)

c. 4-[(N-tert.butyloxycarbonyl)-pyrrolidin-3-yl-oxy]-benzylamine

4.5 g (15.6 mmol) of 4-[(N-tert.butyloxycarbonyl)-pyrrolidin-3-yl-oxy]-benzonitrile are dissolved in 100 ml of methanol and 50 ml of methanolic ammonia and after the addition of 1 g of Raney nickel hydrogenated for 2 hours at 50°C with hydrogen. Then the catalyst is filtered off and the filtrate is evaporated down.

Yield: 4.2 g (92 % of theory),

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R_f value: 0.08 (silica gel; ethyl acetate/methanol = 4:1)

d. 3-cyano-N-[4-(N'-tert.butyloxycarbonyl-pyrrolidin-3-yl-oxy)-benzyl]-benzamide

1.1 g (3.8 mmol) of 4-[(N-tert.butyloxycarbonyl)-pyrrolidin-3-yl-oxy]-benzylamine are dissolved in 30 ml of methylene chloride and after the addition of 0.9 g (9 mmol) of triethylamine, 1.6 g (3.8 mmol) of 3-cyanobenzoic acid chloride are added batchwise. After 4 hours at ambient temperature the solution is combined with water, the organic phase is dried and evaporated down.

Yield: 1.5 g (94 % of theory),

R_f value: 0.27 (silica gel; methylene chloride/ethyl acetate = 9:1)

e. 3-carbamimidoyl-N-[4-(pyrrolidin-3-yl-oxy)-benzyl]-benzamide-dihydrochloride

Prepared analogously to Example 1e from 3-cyano-N-[4-(N'-tert.butyloxycarbonyl-pyrrolidin-3-yl-oxy)-benzyl]-benzamide and hydrochloric acid/ammonium chloride in ethanol.

Yield: 100 % of theory,

Melting point: from 180°C (decomposition)

C₁₉H₂₂N₄O₂ x 2 HCl (338.41/411.41)

mass spectrum: (M+H)⁺ = 339

The following compounds are prepared analogously to Example 9:

(1) 3-carbamimidoyl-N-[4-(cyclopentyloxy)-benzyl]-benzamide-hydrochloride

Yield: 86 % of theory

R_f value: 0.42 (silica gel; methylene chloride/ethanol = 8:2)

C₂₀H₂₃N₃O₂ x HCl (337.43/373.89)

mass spectrum: (M+H)⁺ = 338

(2) 3-carbamimidoyl-N-[4-(benzyloxy)-benzyl]-benzamide-hydrochloride

Yield: 63 % of theory

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R_f value: 0.28 (silica gel: methylene chloride/ethanol = 17:1)

C₂₂H₂₁N₃O₂ x HCl (359.43/395.89)

mass spectrum: (M+H)⁺ = 360

(3) 3-carbamimidoyl-N-[4-(N'-acetyl-pyrrolidin-3-yl-oxy)-benzyl]-benzamide-hydrochloride

Yield: 100 % of theory,

R_f value: 0.08 (silica gel; methylene chloride/ethanol = 9:1)

C₂₁H₂₄N₄O₃ x HCl (380.45/416.91)

mass spectrum: (M+H)⁺ = 381

(4) 3-carbamimidoyl-N-[4-(N'-methyl-pyrrolidin-3-yl-oxy)-benzyl]-benzamide-hydrochloride

Yield: 29 % of theory,

R_f value: 0.07 (silica gel; methylene chloride/ethanol = 7:3)

C₂₀H₂₄N₄O₂ x HCl (352.44/388.91)

mass spectrum: (M+H)⁺ = 353

(5) 3-carbamimidoyl-N-[4-(N'-(aminomethylcarbonyl)-pyrrolidin-3-yl-oxy)-benzyl]-benzamide-dihydrochloride

Yield: 82 % of theory,

Melting point: from 160°C (decomposition)

C₂₁H₂₅N₅O₃ x 2 HCl (395.54/468.46)

mass spectrum: (M+H)⁺ = 396

(6) 3-carbamimidoyl-N-[4-(N'-(2-aminoethyl-carbonyl)-pyrrolidin-3-yl-oxy)-benzyl]-benzamide-dihydrochloride

Yield: 88 % of theory,

Melting point: from 165°C (decomposition)

C₂₂H₂₇N₅O₃ x 2 HCl (409.48/482.48)

mass spectrum: (M+H)⁺ = 410

(7) 3-carbamimidoyl-N-[4-(3-amino-propyloxy)-benzyl]-benzamide-dihydrochloride

Yield: 82 % of theory,

Melting point: from 122°C (decomposition)

C₁₈H₂₂N₄O₂ x 2 HCl (326.40/399.4)

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mass spectrum: $(M+H)^+ = 327$

(8) 3-carbamimidoyl-N-[4-(2-dimethylamino-ethyloxy)-benzyl]-benzamide-dihydrochloride

Yield: 85 % of theory,

Melting point: from 65°C (decomposition)

$C_{19}H_{24}N_4O_2 \times 2 HCl$ (340.43/413.43)

mass spectrum: $(M+H)^+ = 341$

(9) 3-carbamimidoyl-N-[4-(pyridin-4-yl-oxy)-benzyl]-benzamide-hydrochloride

Yield: 66 % of theory,

Melting point: 115°C (decomposition)

$C_{20}H_{18}N_4O_2 \times HCl$ (346.39/382.89)

mass spectrum: $(M+H)^+ = 347$

(10) 3-carbamimidoyl-N-[4-(piperidin-4-yl-oxy)-benzyl]-benzamide-hydrochloride

Yield: 62 % of theory

Melting point: from 170°C (decomposition)

$C_{20}H_{24}N_4O_2 \times HCl$ (352.44/388.89)

mass spectrum: $(M+H)^+ = 353$

Example 10

3-carbamimidoyl-N-[4-(1-(1-imino-ethyl)-pyrrolidin-3-yl-oxy)-benzyl]-benzamide-dihydrochloride

a. 3-cyano-N-[4-(pyrrolidin-3-yl-oxy)-benzyl]-benzamide

2.4 g (5.7 mmol) of 3-cyano-N-[4-(N-tert.butyloxycarbonyl-pyrrolidin-3-yl-oxy)-benzyl]-benzamide are dissolved in 30 ml of methylene chloride and at 0°C combined with 8 ml of trifluoroacetic acid. After 1 hour at ambient temperature the solvent is distilled off, the residue is taken up in methylene chloride, made alkaline with ammonia and water is added. The combined organic extracts are dried and evaporated down. Yield: 1.4 g (76 % of theory),

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R_f value: 0.29 (silica gel; methylene chloride/methanol/ammonia = 7:3:0.2)

b. 3-cyano-N-[4-(1-(1-imino-ethyl)-pyrrolidin-3-yl-oxy)-benzyl]-benzamide

0.7 g (2.17 mmol) of 3-cyano-N-[4-(pyrrolidin-3-yl-oxy)-benzyl]-benzamide, 0.4 g (3.2 mmol) of ethyl acetimidate hydrochloride and 1 g (10 mmol) of triethylamine are dissolved in 70 ml of ethanol and the mixture is stirred for 6 days at ambient temperature. The solvent is distilled off, the residue is taken up in water and made alkaline with sodium carbonate. Then it is extracted with methylene chloride, the combined organic extracts are dried and evaporated down. The residue is triturated with ether and suction filtered.

Yield: 0.6 g (76 % of theory),

R_f value: 0.37 (silica gel; methylene chloride/methanol/ammonia = 7:3:0.2)

Melting point: from 80°C (decomposition)

c. 3-carbamimidoyl-N-[4-(1-(1-imino-ethyl)-pyrrolidin-3-yl-oxy)-benzyl]-benzamide-dihydrochloride

Prepared analogously to Example 1e from 3-cyano-N-[4-(1-(1-imino-ethyl)-pyrrolidin-3-yl-oxy)-benzyl]-benzamide and hydrochloric acid/ammonium carbonate in ethanol.

Yield: 64 % of theory,

Melting point: from 100°C (decomposition)

C₂₁H₂₅N₅O₂ x 2 HCl (379.46/452.46)

mass spectrum: (M+H)⁺ = 380

The following compound is prepared analogously to Example 10:

(1) 3-carbamimidoyl-N-[4-(1-carbamimidoyl-pyrrolidin-3-yl-oxy)-benzyl]-benzamide-dihydrochloride

Yield: 88 % of theory,

Melting point: from 160°C (decomposition)

C₂₀H₂₄N₆O₂ x 2 HCl (380.45/453.38)

mass spectrum: (M+2H)⁺⁺ = 191

Example 113-carbamimidoyl-N-[4-(benzoylamino)-benzyl]-benzamide-hydrochloride

a. 3-cyano-N-(4-amino-benzyl)-benzamide

6 g (0.05 mol) of 4-aminobenzylamine and 10 g (0.1 mol) of triethylamine are dissolved in 150 ml of methylene chloride and at ambient temperature a solution of 8.3 g (0.05 mol) of 3-cyanobenzoylchloride in 20 ml of methylene chloride is added dropwise. After one hour 150 ml of water and 20 ml of methanol are added. After extraction the combined organic extracts are dried and evaporated down. The residue is chromatographed on silica gel and eluted with ethyl acetate. Yield: 4.4 g (35 % of theory),
R_f value: 0.69 (silica gel; ethyl acetate)

b. 3-cyano-N-[4-(benzoylamino)-benzyl]-benzamide

A solution of 0.6 g (4.2 mmol) of benzoylchloride in 10 ml of methylene chloride is added dropwise to a solution of 1 g (4 mmol) of 3-cyano-N-(4-amino-benzyl)-benzamide and 0.6 g (6 mmol) of triethylamine in 30 ml of methylene chloride at ambient temperature. After 8 hours at ambient temperature the product which has crystallised out is dissolved in methylene chloride and methanol. After extraction with water the combined organic extracts are dried and evaporated down. Yield: 1.2 g (84 % of theory),
Melting point: 210°C

c. 3-carbamimidoyl-N-[4-(benzoylamino)-benzyl]-benzamide-hydrochloride

Prepared analogously to Example 1e from 3-cyano-N-[4-(benzoylamino)-benzyl]-benzamide and hydrochloric acid/ammonium carbonate in ethanol.

Yield: 65 % of theory,
Melting point: 190-215°C

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$C_{22}H_{20}N_4O_2 \times HCl$ (372.43/408.93)

mass spectrum: $(M+H)^+ = 373$

The following compounds are prepared analogously to Example 11:

(1) 3-carbamimidoyl-N-[4-(phenylsulphonylamino)-benzyl]-benzamide-hydrochloride

Yield: 80 % of theory,

Melting point: 266°C

$C_{21}H_{20}N_4O_3S \times HCl$ (408.48/444.98)

mass spectrum: $(M+H)^+ = 409$

(2) 3-carbamimidoyl-N-[4-(benzylamino)-benzyl]-benzamide-hydrochloride

Yield: 69 % of theory,

$C_{22}H_{22}N_4O \times HCl$ (358.44/394.94)

mass spectrum: $(M+H)^+ = 359$

(3) 3-carbamimidoyl-N-[4-(N-benzyl-N-ethoxycarbonylmethyl-amino)-benzyl]-benzamide-hydrochloride

Yield: 79 % of theory,

Melting point: from 100°C

$C_{26}H_{28}N_4O_3 \times HCl$ (444.54/481.04)

mass spectrum: $(M+H)^+ = 445$

(4) 3-carbamimidoyl-N-[4-biphenyl-methyl]-benzamide

Yield: 79 % of theory,

Melting point: from 160°C (decomposition)

$C_{21}H_{19}N_3O$ (329.40)

mass spectrum: $(M+H)^+ = 330$

(5) 3-carbamimidoyl-N-[4-(cyclopentylamino)-benzyl]-benzamide-hydrochloride

Yield: 80 % of theory,

Melting point: from 135°C (decomposition)

$C_{20}H_{24}N_4O \times HCl$ (336.44/372.94)

mass spectrum: $M^+ = 336$

Example 12

3-carbamimidoyl-N-(4-dimethylaminomethyl-benzyl)-benzamide-dihydrochloride

a. 4-cyano-N,N-dimethyl-benzylamine

A solution of 7.3 g (0.16 mol) of dimethylamine in 100 ml of ether is added dropwise at -5°C to a solution of 10 g (0.05 mol) of 4-cyanobenzylbromide in 400 ml of ether. Then the reaction mixture is stirred for 2 hours at -5°C and for 20 hours at ambient temperature. After the addition of 200 ml of water and 200 ml of conc. hydrochloric acid the aqueous solution is separated off, made alkaline with sodium hydroxide solution and extracted with methylene chloride. The combined organic extracts are dried and evaporated down.

Yield: 8 g (100 % of theory),

R_f value: 0.58 (silica gel; methylene chloride/ethanol = 9:1)

b. 4-dimethylaminomethyl-benzylamine

Prepared analogously to Example 9c from 4-cyano-N,N-dimethyl-benzylamine, methanolic ammonia and Raney nickel/hydrogen.

Yield: 94 % of theory,

R_f value: 0.13 (silica gel; methylene chloride/ethanol = 9:1)

c. 3-cyano-N-(4-dimethylaminomethyl-benzyl)-benzamide

Prepared analogously to Example 9d from 4-dimethylaminomethyl-benzylamine, 3-cyanobenzoylchloride and triethylamine in methylene chloride.

Yield: 73 % of theory,

Melting point: 100°C

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d. 3-carbamimidoyl-N-(4-dimethylaminomethyl-benzyl)-benzamide-dihydrochloride

Prepared analogously to Example 1e from 3-cyano-N-(4-dimethylaminomethyl-benzyl)-benzamide and hydrochloric acid/ammonium carbonate in ethanol.

Yield: 100 % of theory,

Melting point: from 101°C (decomposition)

$C_{18}H_{22}N_4O \times 2 HCl$ (310.40/383.40)

mass spectrum: $(M+H)^+ = 311$

The following compounds are prepared analogously to Example 12:

(1) 3-carbamimidoyl-N-[4-(imidazol-1-yl)-methyl-benzyl]-benzamide-hydrochloride

Yield: 86 % of theory,

Melting point: from 152°C (decomposition)

$C_{19}H_{19}N_5O \times HCl$ (333.39/369.89)

mass spectrum: $(M+H)^+ = 334$

(2) 2-(5-carbamimidoyl-2-hydroxy-phenyl)-N-[3-methyl-4-(pyrrolidin-1-yl-methyl)-phenyl]-acetamide-dihydrochloride

Yield: 60 % of theory

R_f value (reversed phase RP8; 5% saline solution / methanol = 2:3): 0.7

$C_{21}H_{26}N_4O_2 \times 2 HCl$ (366.47/439.38)

mass spectrum: $(M+H)^+ = 367$

$(M-H)^- = 365$

(3) 2-(5-carbamimidoyl-2-hydroxy-phenyl)-N-[3-methyl-4-(imidazol-1-yl-methyl)-phenyl]-acetamide-dihydrochloride

Yield: 57 % of theory

R_f value (reversed phase RP8; 5% saline solution / methanol = 2:3): 0.7

$C_{20}H_{21}N_5O_2 \times 2 HCl$ (363.42/436.33)

mass spectrum: $(M+H)^+ = 364$

$(M-H)^- = 362$

(4) 2-(5-carbamimidoyl-2-hydroxy-phenyl)-N-[3-methyl-4-(2-methyl-imidazol-1-yl-methyl)-phenyl]-acetamide-dihydrochloride

Example 13

2-(5-carbamimidoyl-2-hydroxy-phenyl)-N-[3-methyl-4-(2-amino-sulphonyl-phenyl)-phenyl]-acetamide-hydrochloride

a. 3-Allyl-4-hydroxy-benzonitrile

82.3 g (0.52 mol) of 4-allyloxy-benzonitrile are heated to 200°C for 2 hours under a nitrogen atmosphere. After cooling, the crude product is purified on silica gel, eluting first with petroleum ether, later with petroleum ether/ethyl acetate (9:1, 8:2, 7:3 and 1:1). The uniform fractions are combined and evaporated down, the residue is washed with petroleum ether and dried.

Yield: 43 g (52% of theory),

R_f value: 0.45 (silica gel; petroleum ether/ethyl acetate = 1:1)

C₁₀H₉NO (159.19)

mass spectrum: (M-H)⁻ = 158

(2M+Na)⁺ = 341

b. 3-Allyl-4-benzyloxy-benzonitrile

Prepared analogously to Example 5d from 3-allyl-4-hydroxy-benzonitrile and benzylbromide/potassium carbonate in dimethylformamide.

Yield: 90% of theory,

Melting point: 59-60°C

R_f value: 0.55 (silica gel; petroleum ether/ethyl acetate = 4:1)

c. 2-benzyloxy-5-cyano-phenylacetic acid

30 g (0.12 mol) of 3-allyl-4-benzyloxy-benzonitrile are dissolved in 450 ml of acetonitrile and at 40°C 0.7 g of ruthenium trichloride hydrate and a solution of 179.7 g (0.84

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mol) of sodium periodate in 1 litre of water is added. After it has all been added, the reaction mixture is heated to 40°C for a further 30 minutes and then extracted 3 x with 1 litre of ethyl acetate. The organic phases are washed with saline solution and dried over sodium sulphate. The crude product is recrystallised from petroleum ether/ethyl acetate (7:3) with the addition of activated charcoal.

Yield: 18.4 g (58% of theory),

Melting point: 144-145°C

R_f value: 0.2 (silica gel; petroleum ether/ethyl acetate = 1:1)

C₁₆H₁₃NO₃ (267.29)

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mass spectrum: $(M-H)^- = 266$
 $(M+Na)^+ = 290$

d. 2-(5-cyano-2-benzyloxy-phenyl)-N-[3-methyl-4-(2-tert.butylaminosulphonyl-phenyl)-phenyl]-acetamide

Prepared analogously to Example 1b from 2-benzyloxy-5-cyano-phenylacetic acid and 4'-amino-2'-methyl-biphenyl-2-sulphonic acid-tert.-butylamide/O-(benzotriazol-1-yl)-N,N,N',N'-tetramethyluronium tetrafluoroborate/triethylamine in dimethylformamide.

Yield: 60% of theory,

R_f value: 0.5 (silica gel; methylene chloride/ethanol = 19:1)

e. 2-(5-carbamimidoyl-2-benzyloxy-phenyl)-N-[3-methyl-4-(2-aminosulphonyl-phenyl)-phenyl]-acetamide-hydrochloride

Prepared analogously to Example 1e from 2-(5-cyano-2-benzyloxy-phenyl)-N-[3-methyl-4-(2-tert.butylaminosulphonyl-phenyl)-phenyl]-acetamide and hydrochloric acid/ammonium carbonate in ethanol.

Yield: 70% of theory,

R_f value: 0.3 (silica gel; methylene chloride/ethanol = 9:1 + 1% glacial acetic acid)

C₂₉H₂₈N₄O₄S x HCl (528.63/565.08)

mass spectrum: $(M-H)^- = 527$
 $(M+H)^+ = 529$

f. 2-(5-carbamimidoyl-2-hydroxy-phenyl)-N-[3-methyl-4-(2-aminosulphonyl-phenyl)-phenyl]-acetamide-hydrochloride

Prepared analogously to Example 5h from 2-(5-carbamimidoyl-2-benzyloxy-phenyl)-N-[3-methyl-4-(2-aminosulphonyl-phenyl)-phenyl]-acetamide-hydrochloride and hydrogen/palladium on activated charcoal.

Yield: 62% of theory,

R_f value: 0.45 (Reversed phase RP8; 5% saline solution/methanol = 1:1)

C₂₂H₂₂N₄O₄S x HCl (438.52/474.97)

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mass spectrum: $(M+H)^+ = 439$
 $(M+Cl)^- = 473/5$ (Cl)

The following compounds are prepared analogously to Example 13:

(1) 2-(5-carbamimidoyl-2-hydroxy-phenyl)-N-(3-methyl-4-phenyl-phenyl]-acetamide-hydrochloride

Yield: 13% of theory,

R_f value: 0.15 (silica gel; methylene chloride/ethanol = 4:1)

C₂₂H₂₁N₃O₂ x HCl (359.45/395.9)

mass spectrum: $(M+H)^+ = 360$
 $(M-H)^- = 358$

(2) 2-(5-carbamimidoyl-2-hydroxy-phenyl)-N-[3-methyl-4-(2-aminosulphonyl-5-methyl-phenyl)-phenyl]-acetamide-hydrochloride

Yield: 23 % of theory

R_f value (silica; methylene chloride / ethanol = 7:3): 0.3

C₂₃H₂₄N₄O₄S x HCl (452.54/488.99)

mass spectrum: $(M+H)^+ = 453$
 $(M-H)^- = 451$

Example 14

2-[5-(N-benzoyl-carbamimidoyl)-2-hydroxy-phenyl]-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide

a. 2-[5-(N-benzoyl-carbamimidoyl)-2-benzyloxy-phenyl]-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide

350 mg (0.69 mmol) of 2-(5-carbamimidoyl-2-benzyloxy-phenyl)-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide-hydrochloride are suspended in 40 ml of methylene chloride and combined with 1.0 ml of triethylamine and 300 mg (1.3 mmol) of 4-nitrophenylbenzoate. The reaction mixture is refluxed for 4 hours. After the addition of 100 ml of saline solution the aqueous phase is extracted 3 x with methylene chloride. The

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combined organic phases are dried over sodium sulphate and concentrated by evaporation. The crude product is purified on silica gel, eluting first with methylene chloride, then with methylene chloride/ethanol (50:1 and 19:1). The uniform fractions are combined, concentrated by evaporation and stirred with petroleum ether/ether (1:1). The solid formed is suction filtered and dried.

Yield: 280 mg (71% of theory),

R_f value: 0.2 (silica gel; petroleum ether/ethyl acetate = 1:1)

C₃₅H₃₄N₄O₄ (574.69)

mass spectrum: (M-H)⁻ = 573

(M+H)⁺ = 575

b. 2-[5-(N-benzoyl-carbamimidoyl)-2-hydroxy-phenyl]-

N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide

Prepared analogously to Example 5h from 2-[5-(N-benzoyl-carbamimidoyl)-2-benzyloxy-phenyl]-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide and hydrogen/palladium on activated charcoal.

Yield: 31% of theory,

R_f value: 0.3 (silica gel; methylene chloride/ethanol = 19:1)

C₂₈H₂₈N₄O₄ (484.56)

mass spectrum: (M+H)⁺ = 485

(M+Na)⁺ = 507

The following compounds are prepared analogously to Example 14:

(1) 2-[5-(N-n-hexyloxycarbonyl-carbamimidoyl)-2-hydroxy-phenyl]-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide

Yield: 17% of theory,

R_f value: 0.3 (silica gel; methylene chloride/ethanol = 4:1)

C₂₈H₃₆N₄O₅ (508.62)

mass spectrum: (M+H)⁺ = 509

(M-H)⁻ = 507

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(2) 2-[5-(N-benzoyl-carbamimidoyl)-2-methoxy-phenyl]-
N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide

Yield: 40% of theory,

R_f value: 0.3 (silica gel; methylene chloride/ethanol = 19:1)

C₂₉H₃₀N₄O₄ (498.59)

mass spectrum: (M+H)⁺ = 499

(M-H)⁻ = 497

(3) 2-[5-(N-n-hexyloxycarbonyl-carbamimidoyl)-2-methoxy-
phenyl]-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-
acetamide

Yield: 35% of theory,

R_f value: 0.25 (silica gel; methylene chloride/ethanol = 19:1)

C₂₉H₂₈N₄O₅ (522.65)

mass spectrum: (M+H)⁺ = 523

(M-H)⁻ = 521

(M+Na)⁺ = 545

(4) 2-[5-(N-ethyloxycarbonyl-carbamimidoyl)-2-methoxy-phenyl]-
N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide

Yield: 32% of theory,

R_f value: 0.45 (silica gel; methylene chloride/ethanol = 9:1)

C₂₅H₃₀N₄O₅ (466.54)

mass spectrum: (M+H)⁺ = 467

(M-H)⁻ = 465

(M+Na)⁺ = 489

Example 15

2-[5-(N-hydroxy-carbamimidoyl)-2-hydroxy-phenyl]-N-[3-methyl-
4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide-acetate

a. 2-[5-(N-hydroxy-carbamimidoyl)-2-benzyloxy-phenyl]-
N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide-
acetate

1.1 g (2.5 mmol) of 2-(5-cyano-2-benzyloxy-phenyl)-N-[3-
methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide are

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dissolved in 100 ml of methanol and combined with a solution of 300 mg (5 mmol) of hydroxylamine hydrochloride in 2.0 ml of water. After the addition of 800 mg (2.5 mmol) of caesium carbonate and 300 mg (3.0 mmol) of sodium carbonate the reaction mixture is refluxed for 6 hours. After cooling and the addition of 0.5 l of ice water the crude product obtained is suction filtered and purified on silica gel, eluting first with methylene chloride and methylene chloride/ethanol (19:1), then with methylene chloride/ethanol (9:1 + 1% glacial acetic acid and 4:1 + 1% glacial acetic acid). The uniform fractions are combined and concentrated by evaporation.

Yield: 620 mg (51% of theory),

R_f value: 0.3 (silica gel; methylene chloride/ethanol = 9:1)

C₂₈H₃₀N₄O₄ (486.58)

mass spectrum: (M-H)⁻ = 485

(M+H)⁺ = 487

(M+Na)⁺ = 509

b. 2-[5-(N-hydroxy-carbamimidoyl)-2-hydroxy-phenyl]-
N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide-
acetate

Prepared analogously to Example 5h from 2-[5-(N-hydroxy-carbamimidoyl)-2-benzyloxy-phenyl]-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide-acetate and hydrogen/palladium on activated charcoal.

Yield: 50% of theory,

R_f value: 0.25 (silica gel; methylene chloride/ethanol = 4:1 + 1% glacial acetic acid)

C₂₁H₂₄N₄O₄ x CH₃COOH (396.45/456.5)

mass spectrum: (M+H)⁺ = 397

(M-H)⁻ = 395

The following compound is prepared analogously to Example 15:

(1) 2-[5-(N-hydroxy-carbamimidoyl)-2-methoxy-phenyl]-
N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide-
acetate

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Yield: 7% of theory,

R_f value: 0.28 (silica gel; methylene chloride/ethanol = 4:1 + 1% glacial acetic acid)

C₂₂H₂₆N₄O₄ x CH₃COOH (410.48/470.52)

mass spectrum: (M+H)⁺ = 411

(M-H)⁻ = 409

(M+Na)⁺ = 433

Example 16

Dry ampoule containing 75 mg of active substance per 10 ml

Composition:

Active substance	75.0 mg
Mannitol	50.0 mg
water for injections	ad 10.0 ml

Preparation:

Active substance and mannitol are dissolved in water. After packaging the solution is freeze-dried. To produce the solution ready for use, the product is dissolved in water for injections.

Example 17

Dry ampoule containing 35 mg of active substance per 2 ml

Composition:

Active substance	35.0 mg
Mannitol	100.0 mg
water for injections	ad 2.0 ml

Preparation:

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Active substance and mannitol are dissolved in water. After packaging, the solution is freeze-dried.

To produce the solution ready for use, the product is dissolved in water for injections.

Example 18

Tablet containing 50 mg of active substance

Composition:

(1) Active substance	50.0 mg
(2) Lactose	98.0 mg
(3) Maize starch	50.0 mg
(4) Polyvinylpyrrolidone	15.0 mg
(5) Magnesium stearate	<u>2.0 mg</u>
	215.0 mg

Preparation:

(1), (2) and (3) are mixed together and granulated with an aqueous solution of (4). (5) is added to the dried granulated material. From this mixture tablets are pressed, biplanar, faceted on both sides and with a dividing notch on one side. Diameter of the tablets: 9 mm.

Example 19

Tablet containing 350 mg of active substance

Composition:

(1) Active substance	350.0 mg
----------------------	----------

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(2) Lactose	136.0 mg
(3) Maize starch	80.0 mg
(4) Polyvinylpyrrolidone	30.0 mg
(5) Magnesium stearate	<u>4.0 mg</u>
	600.0 mg

Preparation:

(1), (2) and (3) are mixed together and granulated with an aqueous solution of (4). (5) is added to the dried granulated material. From this mixture tablets are pressed, biplanar, faceted on both sides and with a dividing notch on one side. Diameter of the tablets: 12 mm.

Example 20

Capsules containing 50 mg of active substance

Composition:

(1) Active substance	50.0 mg
(2) Dried maize starch	58.0 mg
(3) Powdered lactose	50.0 mg
(4) Magnesium stearate	<u>2.0 mg</u>
	160.0 mg

Preparation:

(1) is triturated with (3). This trituration is added to the mixture of (2) and (4) with vigorous mixing.

This powder mixture is packed into size 3 hard gelatine capsules in a capsule filling machine.

Example 21

Capsules containing 350 mg of active substance

Composition:

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(1) Active substance	350.0 mg
(2) Dried maize starch	46.0 mg
(3) Powdered lactose	30.0 mg
(4) Magnesium stearate	<u>4.0 mg</u>
	430.0 mg

Preparation:

(1) is triturated with (3). This trituration is added to the mixture of (2) and (4) with vigorous mixing.

This powder mixture is packed into size 0 hard gelatine capsules in a capsule filling machine.

Example 22

Suppositories containing 100 mg of active substance

1 suppository contains:

Active substance	100.0 mg
Polyethyleneglycol (M.W. 1500)	600.0 mg
Polyethyleneglycol (M.W. 6000)	460.0 mg
Polyethylenesorbitan monostearate	<u>840.0 mg</u>
	2,000.0 mg

Preparation:

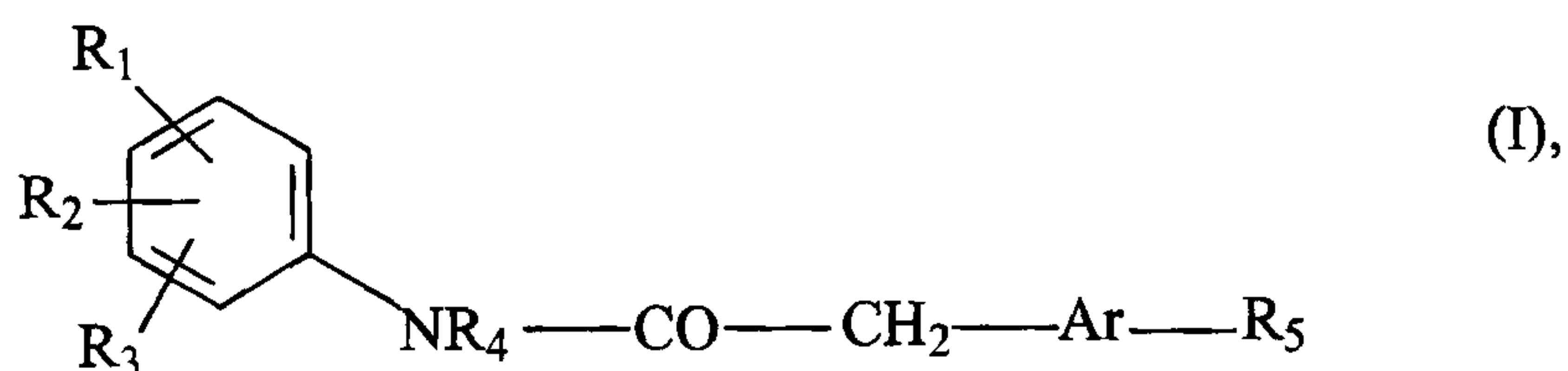
The polyethyleneglycol is melted together with polyethylenesorbitan monostearate. At 40°C the ground active substance is homogeneously dispersed in the melt. This is then cooled to 38°C and poured into slightly chilled suppository moulds.

27400-234

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CLAIMS:

1. A compound of general formula



wherein

5 Ar denotes a phenylene or naphthylene group optionally substituted by a fluorine, chlorine or bromine atom, by a trifluoromethyl, C₁₋₃-alkyl, hydroxy, C₁₋₃-alkoxy, phenyl-C₁₋₃-alkoxy, amino, C₁₋₃-alkylamino or di-(C₁₋₃-alkyl)-amino group, while the phenylene group is optionally further substituted
 10 by another fluorine, chlorine or bromine atom or by another C₁₋₃-alkyl group,

a thienylene, thiazolylene, pyridinylene, pyrimidinylene, pyrazinylene or pyridazinylene group optionally substituted in the carbon skeleton by a C₁₋₃-alkyl group,

15 R₁ denotes a C₁₋₃-alkyl group optionally substituted by an amino, C₁₋₃-alkylamino, di-(C₁₋₃-alkyl)-amino, phenyl, naphthyl, heteroaryl or 4- to 7-membered cycloalkyleneimino group,

a C₃₋₇-cycloalkyl group which is substituted in the
 20 1 position by a 5- to 7-membered cycloalkyleneiminocarbonyl group,

an amino, C₁₋₅-alkylamino, C₅₋₇-cycloalkylamino or phenyl-C₁₋₃-alkylamino group each of which is optionally substituted at the amino-nitrogen atom by a benzoyl or

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phenylsulphonyl group or by a C₁₋₃-alkyl or C₁₋₃-alkylcarbonyl group optionally substituted in the C₁₋₃-alkyl moiety by a carboxy group,

a 4- to 7-membered cycloalkyleneiminocarbonyl or
5 cycloalkyleneiminosulphonyl group optionally substituted by a C₁₋₃-alkyl group,

an aminosulphonyl group optionally substituted by one or two C₁₋₃-alkyl groups,

a phenyl group optionally substituted by a fluorine,
10 chlorine or bromine atom, by a trifluoromethyl, aminosulphonyl, C₁₋₃-alkyl or C₁₋₃-alkoxy group, wherein the substituent group is optionally substituted by a fluorine, chlorine or bromine atom or by a trifluoromethyl, C₁₋₃-alkyl or C₁₋₃-alkoxy group,

15 a C₁₋₃-alkoxy, phenyl-C₁₋₃-alkoxy, heteroaryloxy or heteroaryloxy-C₁₋₃-alkoxy group wherein the alkoxy moiety is optionally substituted in the 2 or 3 position in each case by an amino, C₁₋₃-alkylamino or di-(C₁₋₃-alkyl)-amino group, or

20 a C₃₋₇-cycloalkoxy group, wherein the methylene group in the 3 or 4 position when the cycloalkoxy group is a C₅₋₇-cycloalkoxy group is optionally replaced by a -NH group, wherein the -NH group is optionally substituted

by a C₁₋₃-alkyl group which is optionally
25 substituted in the 2 or 3 position by an amino, C₁₋₃-alkylamino or di-(C₁₋₃-alkyl)-amino group, by a C₁₋₃-alkylcarbonyl, arylcarbonyl or arylsulphonyl group or

by an aminocarbonyl, C₁₋₃-alkylaminocarbonyl or di-(C₁₋₃-alkyl)-aminocarbonyl group, wherein in each case the

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oxygen atom of the carbonyl group is replaced by an imino group,

R₂ denotes a hydrogen, fluorine, chlorine or bromine atom, a C₁₋₃-alkyl, hydroxy or C₁₋₃-alkoxy group,

5 R₃ denotes a hydrogen atom or a C₁₋₃-alkyl group,

R₄ denotes a hydrogen atom or a methyl or ethyl group optionally substituted by a carboxy or C₁₋₃-alkoxycarbonyl group and

R₅ denotes a cyano group or an amidino group optionally
10 substituted by a C₁₋₆-alkoxycarbonyl or benzoyl group,

wherein

by the abovementioned heteroaryl groups is meant a 5-membered heteroaryl group optionally substituted by a C₁₋₃-alkyl group which comprises, in the heteroaromatic moiety,

15 an imino group optionally substituted by a C₁₋₃-alkyl group, or an oxygen or sulphur atom,

an imino group optionally substituted by a C₁₋₃-alkyl group and an oxygen, sulphur or nitrogen atom,

an imino group optionally substituted by a
20 C₁₋₃-alkyl group and two nitrogen atoms or

an oxygen or sulphur atom and two nitrogen atoms,

or a 6-membered heteroarylene group optionally substituted by a C₁₋₃-alkyl group which comprises one or two nitrogen atoms in the heteroaromatic moiety,

25 by the abovementioned heteroaryloxy groups is meant a heteroaryl group, as defined above, which is bound through an oxygen atom,

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the carboxy groups mentioned in the definition of the abovementioned groups is optionally replaced by a group which is converted into a carboxy group *in vivo* or by a group which is negatively charged under physiological conditions or

5 the amino and imino groups mentioned in the definition of the abovementioned groups is optionally replaced by a group which may be cleaved *in vivo*,

wherein said group which is converted into a carboxy group is selected from the group consisting of: a hydroxymethyl
 10 group, a carboxy group esterified with an alcohol wherein the alcoholic moiety denotes a C₁₋₆-alkanol, a phenyl-C₁₋₃-alkanol, a C₃₋₉-cycloalkanol, wherein the cycloalkanol is optionally substituted by one or two C₁₋₃-alkyl groups when the cycloalkanol is a
 15 C₅₋₈-cycloalkanol, the C₅₋₈-cycloalkanol wherein a methylene group in the 3 or 4 position is replaced by an oxygen atom or by an imino group optionally substituted by a C₁₋₃-alkyl, phenyl-C₁₋₃-alkyl, phenyl-C₁₋₃-alkoxycarbonyl or C₂₋₆-alkanoyl group and the cycloalkanol moiety is optionally further
 20 substituted by one or two C₁₋₃-alkyl groups, a C₄₋₇-cycloalkenol, a C₃₋₅-alkenol, a phenyl-C₃₋₅-alkenol, a C₃₋₅-alkynol or phenyl;

C₃₋₅-alkynol, with the proviso that no bond to the oxygen atom starts from a carbon atom which carries a double or
 25 triple bond; a C₃₋₈-cycloalkyl-C₁₋₃-alkanol, a bicycloalkanol having a total of 8 to 10 carbon atoms optionally substituted by one or two C₁₋₃-alkyl groups in the bicycloalkyl moiety, a 1,3-dihydro-3-oxo-1-isobenzfuranol and an alcohol of formula

30
$$R_a-CO-O(R_bCR_c)-OH,$$

wherein

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R_a denotes a C_{1-8} -alkyl, C_{5-7} -cycloalkyl, phenyl or phenyl- C_{1-3} -alkyl group,

R_b denotes a hydrogen atom, a C_{1-3} -alkyl, C_{5-7} -cycloalkyl or phenyl group and

5 R_c denotes a hydrogen atom or a C_{1-3} -alkyl group,

and wherein said group that is negatively charged under physiological conditions is selected from the group consisting of: a tetrazol-5-yl, phenylcarbonylaminocarbonyl, trifluoromethylcarbonylaminocarbonyl,

10 C_{1-6} -alkylsulphonylamino, phenylsulphonylamino, benzylsulphonylamino, trifluoromethylsulphonylamino, C_{1-6} -alkylsulphonylaminocarbonyl, phenylsulphonylaminocarbonyl, benzylsulphonylaminocarbonyl and perfluoro- C_{1-6} -alkylsulphonylaminocarbonyl group, and

15 and wherein said amino and imino groups are replaced by a group that is cleaved *in vivo* is selected from the group consisting of a hydroxy group, an acyl group, a pyridinoyl group, a C_{1-16} -alkanoyl group, a 3,3,3-trichloropropionyl, a allyloxycarbonyl group, a C_{1-16} -alkoxycarbonyl group, a

20 C_{1-16} -alkylcarbonyloxy group wherein hydrogen atoms are optionally wholly or partially replaced by fluorine or chlorine atoms, a phenyl- C_{1-6} -alkoxycarbonyl group; a 3-amino-propionyl group wherein the amino group is optionally mono or disubstituted by C_{1-6} -alkyl or

25 C_{3-7} -cycloalkyl groups wherein the substituents are identical or different; a C_{1-3} -alkylsulphonyl- C_{2-4} -alkoxycarbonyl, C_{1-3} -alkoxy- C_{2-4} -alkoxy- C_{2-4} -alkoxycarbonyl, R_a -CO-O- (R_bCR_c) -O-CO, C_{1-6} -alkyl-CO-NH- (R_dCR_c) -O-CO and C_{1-6} -alkyl-CO-O- (R_dCR_e) - (R_dCR_c) -O-CO group wherein R_a to R_c are as hereinbefore

30 defined,

and wherein

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R_d and R_c, which may be identical or different, denote hydrogen atoms or C₁₋₃-alkyl groups;

or an isomer or salt thereof.

2. A compound, isomer or salt according to claim 1,
5 wherein

Ar denotes a phenylene group optionally substituted by a fluorine, chlorine or bromine atom or by a methyl, hydroxy, methoxy or benzyloxy group, which is optionally further substituted by a methyl group,

10 R₁ denotes a phenyl group optionally substituted by a fluorine, chlorine or bromine atom or by a trifluoromethyl, aminosulphonyl, C₁₋₃-alkyl or C₁₋₃-alkoxy group, which is optionally further substituted by a fluorine, chlorine or bromine atom or by a trifluoromethyl, C₁₋₃-alkyl or
15 C₁₋₃-alkoxy group,

a methyl group substituted by a dimethylamino, pyrrolidino or imidazolyl group, wherein the imidazolyl moiety is optionally substituted by a methyl group,

an amino, C₁₋₅-alkylamino, cyclopentylamino or benzylamino
20 group which is optionally substituted at the amino-nitrogen atom by a carboxy-C₁₋₂-alkyl, C₁₋₃-alkoxycarbonyl-C₁₋₂-alkyl, carboxy-C₁₋₂-alkylcarbonyl or C₁₋₃-alkoxycarbonyl-C₁₋₂-alkylcarbonyl group,

a benzoylamino or phenylsulphonylamino group,

25 a cyclopropyl group which is substituted in the 1 position by a 5- to 7-membered cycloalkyleneiminocarbonyl group,

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an optionally methyl-substituted pyrrolidinocarbonyl,
piperidinocarbonyl, pyrrolidinosulphonyl or
piperidinosulphonyl group,

5 a C₁₋₃-alkoxy group wherein the alkoxy moiety in the
2 or 3 position is optionally substituted in each case by an
amino, C₁₋₃-alkylamino or di-(C₁₋₃-alkyl)-amino group,

a phenyl-C₁₋₃-alkoxy or pyridinyloxy group, or

a C₅₋₇-cycloalkoxy group wherein the methylene group in the
3 or 4 position is optionally replaced by a -NH group,
10 wherein the -NH group is optionally substituted

by a C₁₋₃-alkyl or C₂₋₃-alkanoyl group,

by a C₂₋₃-alkanoyl or aminocarbonyl group wherein
in each case the oxygen atom of the carbonyl group is
replaced by an imino group,

15 R₂ denotes a hydrogen, fluorine, chlorine or bromine atom, a
methyl, hydroxy or methoxy group,

R₃ denotes a hydrogen atom or a methyl group,

R₄ denotes a hydrogen atom or a methyl or ethyl group
optionally substituted by a carboxy or C₁₋₃-alkoxycarbonyl
20 group and

R₅ denotes a cyano group or an amidino group optionally
substituted by a C₁₋₆-alkoxycarbonyl or benzoyl group.

3. A compound, isomer or salt according to claim 1,
wherein

25 Ar denotes a phenylene group optionally substituted by a
methyl, hydroxy, methoxy or benzyloxy group,

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R₁ denotes a phenyl group optionally substituted by a fluorine, chlorine or bromine atom or by a trifluoromethyl, aminosulphonyl, C₁₋₃-alkyl or C₁₋₃-alkoxy group, which is optionally further substituted by a fluorine, chlorine or bromine atom or by a trifluoromethyl, C₁₋₃-alkyl or C₁₋₃-alkoxy group,

a cyclopropyl group which is substituted in the 1 position by a 5- to 7-membered cycloalkyleneiminocarbonyl group, or a 4- to 7-membered cycloalkyleneiminocarbonyl group, or
 10 an optionally methyl-substituted pyrrolidinocarbonyl, piperidinocarbonyl or pyrrolidinosulphonyl group,

R₂ denotes a hydrogen, fluorine, chlorine or bromine atom or a methyl group,

R₃ denotes a hydrogen atom or a methyl group,

15 R₄ denotes a hydrogen atom or a methyl or ethyl group substituted by a carboxy, methoxycarbonyl or ethoxycarbonyl group and

R₅ denotes an amidino group optionally substituted by a C₁₋₆-alkoxycarbonyl or benzoyl group.

20 4. 2-(5-Carbamimidoyl-2-hydroxy-phenyl)-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide; wherein the amidino group is optionally substituted by a C₁₋₆-alkoxycarbonyl or benzoyl group, or a salt thereof.

5. 2-(2-Benzyloxy-5-carbamimidoyl-phenyl)-N-
 25 (2-ethoxycarbonyl-ethyl)-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide; wherein the amidino group is optionally substituted by a C₁₋₆-alkoxycarbonyl or benzoyl group, or a salt thereof.

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6. 2-(2-Hydroxy-5-carbamimidoyl-phenyl)-N-(2-ethoxycarbonylethyl)-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide; wherein the amidino group is optionally substituted by a C₁₋₆-alkoxycarbonyl or benzoyl group, or a salt thereof.

7. 2-(2-Hydroxy-5-carbamimidoyl-phenyl)-N-(2-carboxy-ethyl)-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide; wherein the amidino group is optionally substituted by a C₁₋₆-alkoxycarbonyl or benzoyl group, or a salt thereof.

8. 2-(5-Carbamimidoyl-2-hydroxy-phenyl)-N-[3-methyl-4-(piperidin-1-yl-carbonyl)-phenyl]-acetamide; wherein the amidino group is optionally substituted by a C₁₋₆-alkoxycarbonyl or benzoyl group, or a salt thereof.

9. 2-(5-Carbamimidoyl-2-hydroxy-phenyl)-N-[3-methyl-4-(2-aminosulphonyl-phenyl)-phenyl]-acetamide; wherein the amidino group is optionally substituted by a C₁₋₆-alkoxycarbonyl or benzoyl group, or a salt thereof.

10. 2-(5-Carbamimidoyl-2-hydroxy-phenyl)-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide or a salt thereof.

11. A physiologically acceptable salt of the compound defined in any one of claims 1 to 3, wherein R₅ denotes one of the amidino groups defined in any one of claims 1 to 3.

12. A pharmaceutical composition comprising the compound defined in any one of claims 1 to 3, wherein R₅ denotes one of the amidino groups defined in any one of claims 1 to 3, the salt according to claim 11 or the compound or salt as defined in any one of claims 4 to 10 and a pharmaceutically acceptable carrier or diluent.

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13. The pharmaceutical composition according to claim 12 for treating thrombosis.

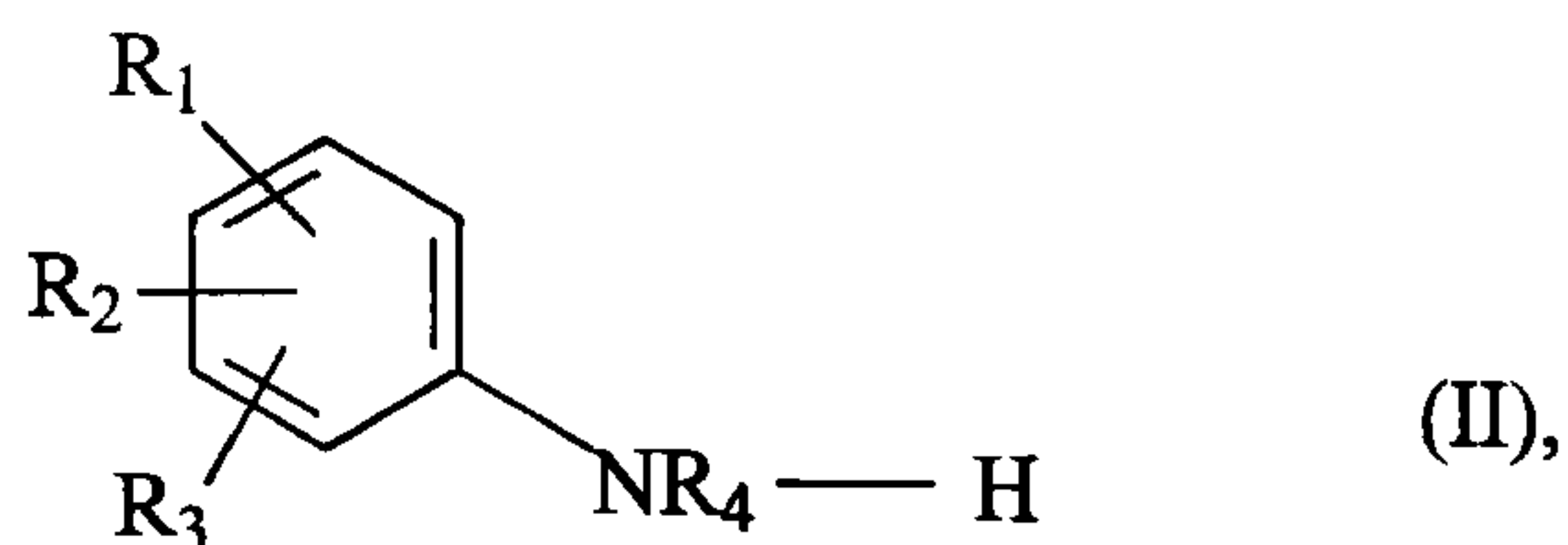
14. Use of the compound defined in any one of claims 1 to 3, wherein R_5 denotes one of the amidino groups mentioned in any one of claims 1 to 3, the salt according to claim 11, or the compound or salt defined in any one of claims 4 to 10, for preparing a pharmaceutical composition having an antithrombotic activity.

15. Use of the compound defined in any one of claims 1 to 3, wherein R_5 denotes one of the amidino groups mentioned in any one of claims 1 to 3, the salt according to claim 11, or the compound or salt defined in any one of claims 4 to 10, for treating thrombosis.

16. Process for preparing the pharmaceutical composition defined in claim 12 wherein the compound according to any one of claims 1 to 3, wherein R_5 denotes one of the amidino groups mentioned in any one of claims 1 to 3, the salt according to claim 11 or the compound or salt defined in any one of claims 4 to 10 is admixed with a pharmaceutically acceptable carrier or diluent.

17. A process for preparing the compound defined in any one of claims 1 to 3, wherein:

a) a compound of general formula

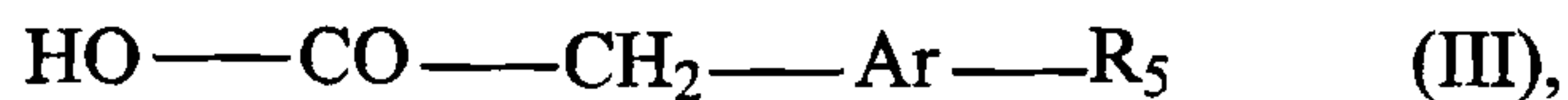


25 wherein

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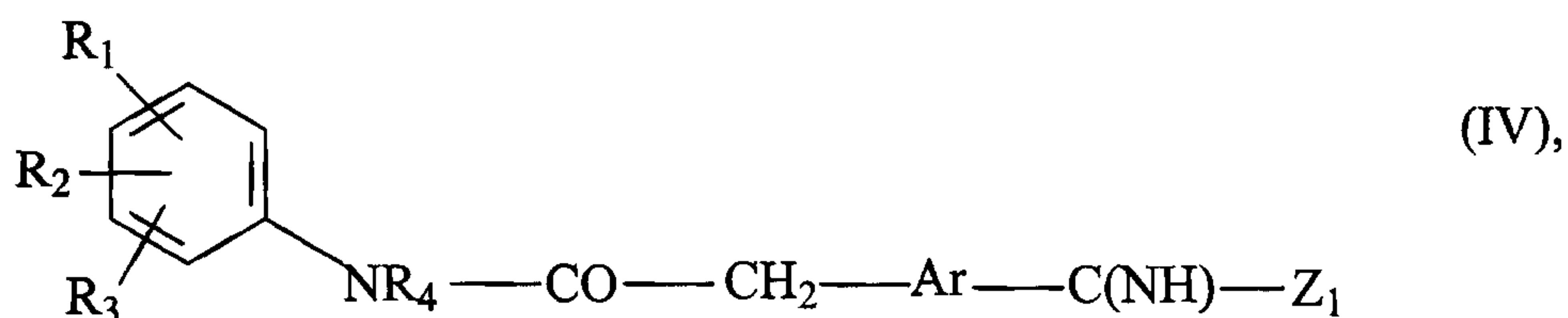
R₁ to R₄ are defined as in any one of claims 1 to 3, is acylated with a carboxylic acid of general formula



wherein

5 Ar and R₅ are defined as in any one of claims 1 to 3, or with a reactive derivatives thereof, or

b) in order to prepare a compound of general formula I wherein R₅ denotes an amidino group which is optionally substituted by one or two C₁₋₃-alkyl groups, a compound of
10 general formula



optionally formed in a reaction mixture,

wherein

R₁ to R₄ and Ar are defined as in any one of claims 1 to 3
15 and

Z₁ denotes an alkoxy, aralkoxy, alkylthio or aralkylthio group, is reacted with an amine of general formula



wherein

20 R₆ and R₇, which are identical or different, each denote a hydrogen atom or a C₁₋₃-alkyl group, or with the salts thereof, and

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subsequently, one or more steps is optionally performed, wherein the one or more steps are selected from the group consisting of:

the compound of general formula I thus obtained which
5 comprises an amino or imino group is converted by means of a corresponding acyl derivative into a corresponding acyl compound of general formula I;

the compound of general formula I thus obtained which
comprises an esterified carboxy group is converted by
10 hydrolysis into a corresponding carboxylic acid of general formula I;

the compound of general formula I thus obtained which
contains a carboxy group is converted by esterification into
a corresponding ester;

15 a protecting group used to protect a reactive group during the reactions is cleaved;

the compound of general formula I thus obtained is resolved into the stereoisomers thereof;

the compound of general formula I thus obtained which
20 comprises a cyano group is converted into a corresponding compound of general formula I which comprises an amidino group; and

the compound of general formula I thus obtained is converted into a salt thereof.

25 18. The process according to claim 17, wherein the compound of general formula I is converted into a physiologically acceptable salt thereof with an inorganic or organic acid or base.

