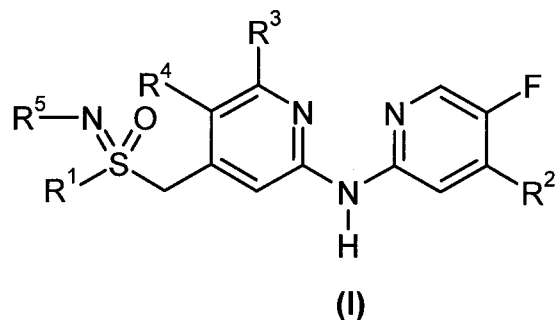


Patent claims

1. A compound of general formula (I)

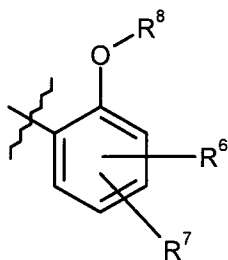


wherein

R^1 represents a C_1 - C_6 -alkyl group,

wherein said group is optionally substituted with one substituent, selected from the group of hydroxy, C_1 - C_3 -alkoxy, $-NH_2$, alkylamino-, dialkylamino-, or cyclic amines;

R^2 represents a group



R^3 represents a hydrogen atom, a fluoro atom or a chloro atom, or a methyl, methoxy, difluoromethyl or trifluoromethyl group;

R^4 represents a hydrogen atom or a bromo atom;

R^5 represents a group selected from a hydrogen atom, cyano, $-C(O)R^9$, $-C(O)OR^9$, $-C(O)NR^{10}R^{11}$;

R^6, R^7 represent, independently from each other, a group selected from a hydrogen atom, fluoro atom or chloro atom;

R^8 represents a C_1 - C_3 -alkyl group;

R^9 represents a C_1 - C_3 -alkyl group, a benzyl group, or trifluoromethyl;

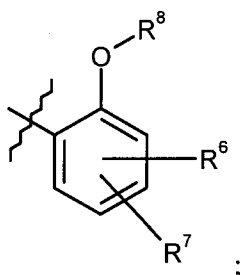
R^{10}, R^{11} represent, independently from each other, a group selected from hydrogen, C_1 - C_2 -alkyl-; and the enantiomers, diastereomers, salts, solvates or salts of solvates thereof.

2. The compound of general formula (I) according to claim 1, wherein

R^1 represents a C_1 - C_3 -alkyl group,

wherein said group is optionally substituted with one substituent, selected from the group of hydroxy, -NH_2 ;

R^2 represents a group



R^3 represents a hydrogen atom, a fluoro atom or a chloro atom, or a methyl, methoxy, difluoromethyl or trifluoromethyl group;

R^4 represents a hydrogen atom or a bromo atom;

R^5 represents a group selected from a hydrogen atom, cyano, -C(O)R^9 , -C(O)OR^9 , $\text{-C(O)NR}^{10}\text{R}^{11}$;

R^6 represents a fluoro atom;

R^7 represents hydrogen;

R^8 represents a methyl or ethyl group;

R^9 represents a methyl, ethyl or trifluoromethyl group;

R^{10} , R^{11} represent, independently from each other, a group selected from hydrogen, $\text{C}_1\text{-C}_2\text{-alkyl}$; and the enantiomers, diastereomers, salts, solvates or salts of solvates thereof.

3. The compound of general formula (I) according to claim 1, wherein

R^1 represents a methyl, ethyl, 2-hydroxyethyl or 2-aminoethyl group;

R^2 represents a 4-fluoro-2-methoxyphenyl or 4-fluoro-2-ethoxyphenyl group;

R^3 represents a hydrogen atom, a fluoro atom or a chloro atom, or a methyl, methoxy, difluoromethyl or trifluoromethyl group;

R^4 represents a hydrogen atom or a bromo atom;

R^5 represents a hydrogen atom, or a group selected from cyano, -C(O)CH_3 , -C(O)CF_3 , $\text{-C(O)OC}_2\text{H}_5$, $\text{-C(O)N(H)C}_2\text{H}_5$;

and the enantiomers, diastereomers, salts, solvates or salts of solvates thereof.

4. The compound according to claim 1, which is

- (*rac*)-5-Fluoro-4-(4-fluoro-2-methoxyphenyl)-*N*-{4-[(*S*-methylsulfonimidoyl)methyl]pyridin-2-yl}pyridin-2-amine;
- (+)-5-Fluoro-4-(4-fluoro-2-methoxyphenyl)-*N*-{4-[(*S*-methylsulfonimidoyl)methyl]pyridin-2-yl}pyridin-2-amine;
- (-)-5-Fluoro-4-(4-fluoro-2-methoxyphenyl)-*N*-{4-[(*S*-methylsulfonimidoyl)methyl]pyridin-2-yl}pyridin-2-amine;

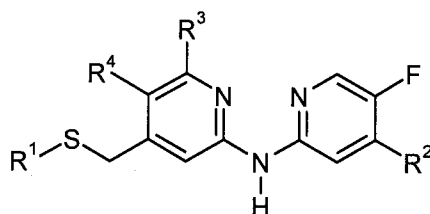
- (*rac*)-5-Fluoro-4-(4-fluoro-2-methoxyphenyl)-*N*-{6-methyl-4-[(*S*-methylsulfonimidoyl)methyl]pyridin-2-yl}pyridin-2-amine;
- (*rac*)-5-Bromo-*N*-[5-fluoro-4-(4-fluoro-2-methoxyphenyl)pyridin-2-yl]-6-methyl-4-[(*S*-methylsulfonimidoyl)methyl]pyridin-2-amine;
- (*rac*)-5-Fluoro-4-(4-fluoro-2-methoxyphenyl)-*N*-{6-methoxy-4-[(*S*-methylsulfonimidoyl)methyl]pyridin-2-yl}pyridin-2-amine;
- (*rac*)-*N*-{6-Chloro-4-[(*S*-methylsulfonimidoyl)methyl]pyridin-2-yl}-5-fluoro-4-(4-fluoro-2-methoxyphenyl)pyridin-2-amine;
- (*rac*)-2-{*S*-[(2-{[5-Fluoro-4-(4-fluoro-2-methoxyphenyl)pyridin-2-yl]amino}pyridin-4-yl)methyl]sulfonimidoyl}ethanol;
- (*rac*)-*N*-(4-{[*S*-(2-Aminoethyl)sulfonimidoyl]methyl}pyridin-2-yl)-5-fluoro-4-(4-fluoro-2-methoxyphenyl)pyridin-2-amine;
- {[2-{[5-Fluoro-4-(4-fluoro-2-methoxyphenyl)pyridin-2-yl]amino}pyridin-4-yl)methyl](methyl)oxido- λ^6 -sulfanylidene}cyanamide;
- (*rac*)-Ethyl {[2-{[5-fluoro-4-(4-fluoro-2-methoxyphenyl)pyridin-2-yl]amino}pyridin-4-yl)methyl](methyl)oxido- λ^6 -sulfanylidene}carbamate;
- (*rac*)-1-Ethyl-3-[[2-{[5-fluoro-4-(4-fluoro-2-methoxyphenyl)pyridin-2-yl]amino}pyridin-4-yl)methyl](methyl)oxido- λ^6 -sulfanylidene]urea;
- (*rac*)-*N*-[[2-{[5-Fluoro-4-(4-fluoro-2-methoxyphenyl)pyridin-2-yl]amino}pyridin-4-yl)methyl](methyl)oxido- λ^6 -sulfanylidene]acetamide;
- 5-Fluoro-4-(4-fluoro-2-methoxyphenyl)-*N*-{4-[(*S*-methylsulfonimidoyl)methyl]pyridin-2-yl}pyridin-2-amine hydrochloride; enantiomer 2;
- (*rac*)-Ethyl {[2-{[5-fluoro-4-(4-fluoro-2-methoxyphenyl)pyridin-2-yl]amino}-6-methoxypyridin-4-yl)methyl](methyl)oxido- λ^6 -sulfanylidene}carbamate;
- 5-Fluoro-4-(4-fluoro-2-methoxyphenyl)-*N*-{4-[(*S*-methylsulfonimidoyl)methyl]pyridin-2-yl}pyridin-2-amine hydrochloride; enantiomer 1;
- (*rac*)-5-Fluoro-4-(4-fluoro-2-methoxyphenyl)-*N*-{6-fluoro-4-[(*S*-methylsulfonimidoyl)methyl]pyridin-2-yl}pyridin-2-amine;
- (*rac*)-2,2,2-Trifluoro-*N*-{[(2-fluoro-6-{[5-fluoro-4-(4-fluoro-2-methoxyphenyl)pyridin-2-yl]amino}pyridin-4-yl)methyl](methyl)oxido- λ^6 -sulfanylidene}acetamide;
- (+)-5-Fluoro-4-(4-fluoro-2-methoxyphenyl)-*N*-{6-fluoro-4-[(*S*-methylsulfonimidoyl)methyl]pyridin-2-yl}pyridin-2-amine;
- (-)-5-Fluoro-4-(4-fluoro-2-methoxyphenyl)-*N*-{6-fluoro-4-[(*S*-methylsulfonimidoyl)methyl]pyridin-2-yl}pyridin-2-amine;
- (*rac*)-5-Fluoro-4-(4-fluoro-2-methoxyphenyl)-*N*-{4-[(*S*-methylsulfonimidoyl)methyl]-6-(trifluoromethyl)pyridin-2-yl}pyridin-2-amine;

- (+)-5-Fluoro-4-(4-fluoro-2-methoxyphenyl)-*N*-{4-[(*S*-methylsulfonimidoyl)methyl]-6-(trifluoromethyl)pyridin-2-yl}pyridin-2-amine;
- (-)-5-Fluoro-4-(4-fluoro-2-methoxyphenyl)-*N*-{4-[(*S*-methylsulfonimidoyl)methyl]-6-(trifluoromethyl)pyridin-2-yl}pyridin-2-amine;
- (+)-2,2,2-Trifluoro-*N*-[{2-[5-fluoro-4-(4-fluoro-2-methoxyphenyl)pyridin-2-yl]amino}-6-(trifluoromethyl)-pyridin-4-yl]methyl}(methyl)oxido- λ^6 -sulfanylidene]acetamide;
- (-)-2,2,2-Trifluoro-*N*-[{2-[5-fluoro-4-(4-fluoro-2-methoxyphenyl)pyridin-2-yl]amino}-6-(trifluoromethyl)-pyridin-4-yl]methyl}(methyl)oxido- λ^6 -sulfanylidene]acetamide;
- (*rac*)-4-(2-Ethoxy-4-fluorophenyl)-5-fluoro-*N*-{6-fluoro-4-[(*S*-methylsulfonimidoyl)methyl]pyridin-2-yl}pyridin-2-amine;
- (*rac*)-*N*-{[(2-[4-(2-Ethoxy-4-fluoro-phenyl)-5-fluoropyridin-2-yl]amino)-6-fluoropyridin-4-yl]methyl}-(methyl)oxido- λ^6 -sulfanylidene}-2,2,2-trifluoroacetamide;
- (+)-4-(2-Ethoxy-4-fluorophenyl)-5-fluoro-*N*-{6-fluoro-4-[(*S*-methylsulfonimidoyl)methyl]pyridin-2-yl}pyridin-2-amine;
- (-)-4-(2-Ethoxy-4-fluorophenyl)-5-fluoro-*N*-{6-fluoro-4-[(*S*-methylsulfonimidoyl)methyl]pyridin-2-yl}pyridin-2-amine;
- (*rac*)-*N*-{4-[(*S*-Ethylsulfonimidoyl)-methyl]pyridin-2-yl}-5-fluoro-4-(4-fluoro-2-methoxyphenyl)pyridin-2-amine;
- (*rac*)-*N*-{6-(Difluoromethyl)-4-[(*S*-methylsulfonimidoyl)methyl]pyridin-2-yl}-5-fluoro-4-(4-fluoro-2-methoxyphenyl)pyridin-2-amine;
- (+)-5-Fluoro-4-(4-fluoro-2-methoxyphenyl)-*N*-{6-methyl-4-[(*S*-methylsulfonimidoyl)methyl]pyridin-2-yl}pyridin-2-amine;
- (-)-5-Fluoro-4-(4-fluoro-2-methoxyphenyl)-*N*-{6-methyl-4-[(*S*-methylsulfonimidoyl)methyl]pyridin-2-yl}pyridin-2-amine,

and the enantiomers, diastereomers, salts, solvates or salts of solvates thereof.

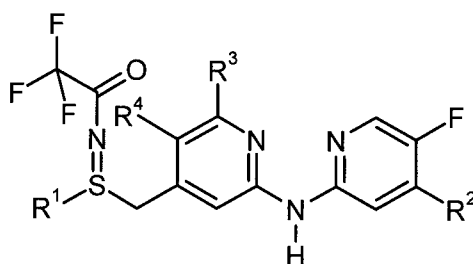
5. A compound of general formula (I) according to any one of claims 1 to 4 or an enantiomer, diastereomer, salt, solvate, or salt of the solvate thereof, for use in the treatment and/or prophylaxis of hyper-proliferative disorders, virally induced infectious diseases and/or of cardiovascular diseases.
6. A compound of general formula (I) according to any one of claims 1 to 4 or an enantiomer, diastereomer, salt, solvate, or salt of the solvate thereof, for use in the treatment and/or prophylaxis of lung carcinomas, prostate carcinomas, cervical carcinomas, colorectal carcinomas, melanomas, ovarian carcinomas or leukemias.
7. A pharmaceutical combination comprising a compound of general formula (I) according to any one of claims 1 to 4 or an enantiomer, diastereomer, salt, solvate, or salt of the solvate thereof, in combination with at least one or more further active ingredients.

8. A pharmaceutical composition comprising a compound of general formula (I) according to any one of claims 1 to 4 or an enantiomer, diastereomer, salt, solvate, or salt of the solvate thereof, in combination with an inert, nontoxic, pharmaceutically suitable adjuvant.
9. The pharmaceutical combination according to claim 7 for use in the treatment and/or prophylaxis of hyper-proliferative disorders, virally induced infectious diseases and/or of cardiovascular diseases.
10. The pharmaceutical composition according to claim 8 for use in the treatment and/or prophylaxis of hyper-proliferative disorders, virally induced infectious diseases and/or of cardiovascular diseases.
11. A compound of general formula (5)

**5**

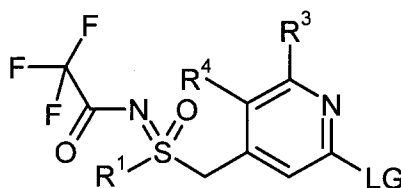
wherein R¹, R², R³ and R⁴ are as defined according to any one of claims 1 to 3 for the compounds of general formula (I),
or a salt, solvate, or salt of solvate thereof.

12. A compound of general formula (6)

**6**

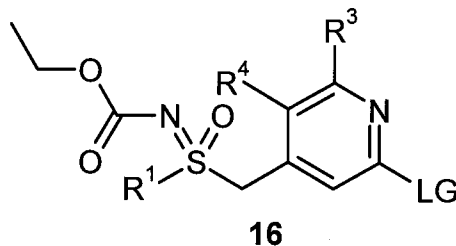
wherein R¹, R², R³ and R⁴ are as defined according to any one of claims 1 to 3 for the compounds of general formula (I),
or an enantiomer, diastereomer, salt, solvate, or salt of solvate thereof.

13. A compound of general formula (14)

**14**

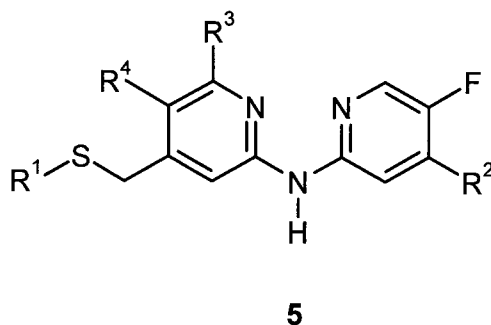
wherein R^1 , R^3 and R^4 are as defined according to any one of claims 1 to 3 for the compounds of general formula (I), and wherein LG stands for a leaving group,
or an enantiomer, diastereomer, salt, solvate, or salt of solvate thereof.

14. A compound of general formula (16)

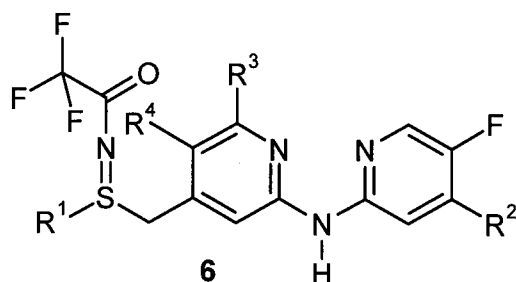


wherein R^1 , R^3 and R^4 are as defined according to any one of claims 1 to 3 for the compounds of general formula (I), and wherein LG stands for a leaving group,
or an enantiomer, diastereomer, salt, solvate, or salt of solvate thereof.

15. A method for the preparation of a compound of formula (6), in which R^1 , R^2 , R^3 , and R^4 are as defined for the compound of the formula (I) according to any one of claims 1 to 3, in which method a compound of formula (5), in which R^1 , R^2 , R^3 , and R^4 are as defined for the compound of the formula (I) according to any one of claims 1 to 3,

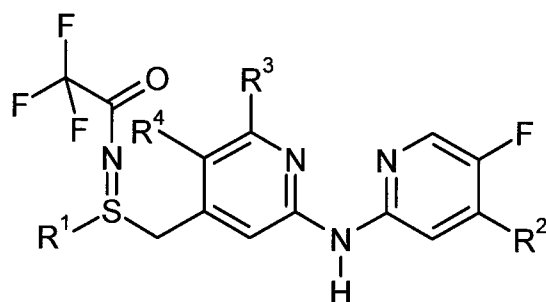


is reacted with trifluoroacetamide and 1,3-dibromo-5,5-dimethylhydantoin in the presence of an alkali salt of *tert*-butanol in a cyclic ether as a solvent, to give a compound of the formula (6),



and in which method the resulting compound is optionally, if appropriate, converted with the corresponding (i) solvents and/or (ii) bases or acids to the solvates, salts and/or solvates of the salts thereof.

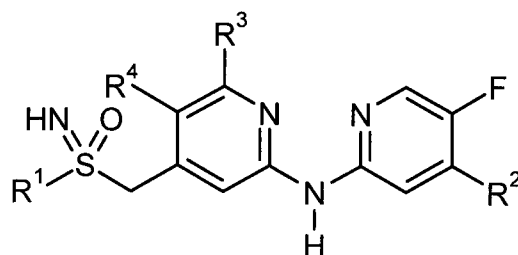
16. A method for the preparation of a compound of formula (I), according to any one of the claims 1 to 4, in which method a compound of formula (6), in which R^1 , R^2 , R^3 , and R^4 are as defined for the compound of the formula (I) according to any one of claims 1 to 3,



6

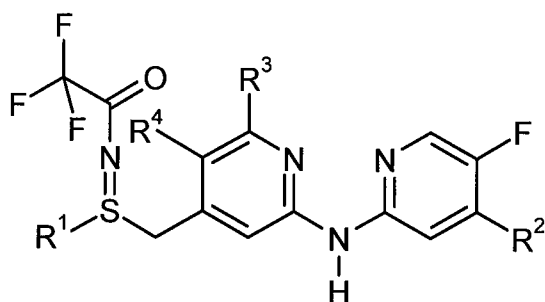
is oxidised with an alkali salt of permanganic acid in an aliphatic ketone of the formula $C_1-C_2-C(O)-C_1-C_2$ -alkyl as a solvent,

followed, if the trifluoroacetyl group present in the compound of formula (6) has not been cleaved off during abovementioned oxidation process, by the removal of said trifluoroacetyl group by treatment of the resulting intermediate with a suitable base in an alcoholic solvent, to give a compound of the formula (I), in which R^5 is hydrogen,

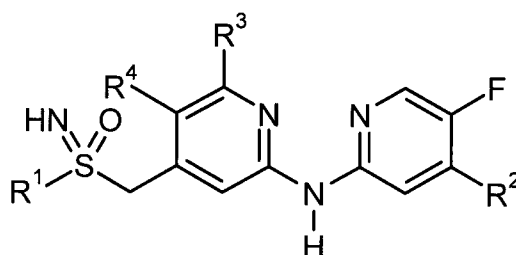
(I); $R^5 = H$

and in which method the resulting compound is optionally, if appropriate, converted with the corresponding (i) solvents and/or (ii) bases or acids to the solvates, salts and/or solvates of the salts thereof.

17. A method for the preparation of a compound of formula (I), according to any one of the claims 1 to 4, in which method a compound of formula (6), in which R^1 , R^2 , R^3 , and R^4 are as defined for the compound of the formula (I) according to any one of claims 1 to 3,

**6**

is oxidised with a peroxomonosulfate based oxidant in a solvent selected from an aliphatic alcohol of the formula C₁-C₃-alkyl-OH, water, and *N,N*-dimethylformamide, or a mixture thereof, to give a compound of the formula (I), in which R⁵ is hydrogen,

**(I); R⁵ = H**

and in which method the resulting compound is optionally, if appropriate, converted with the corresponding (i) solvents and/or (ii) bases or acids to the solvates, salts and/or solvates of the salts thereof.

18. Use of a compound of the formula (I) according to any one of claims 1 to 4, or an enantiomer, diastereomer, salt, solvate, or salt of the solvate thereof, in the manufacture of a medicament for the treatment and/or prophylaxis of a disorder relating to or mediated by CDK9 activity, wherein the disorder is a hyper-proliferative disorder, a virally induced infectious disease, or a cardiovascular disease.
19. Use according to claim 18, wherein the disorder is lung carcinoma, prostate carcinoma, cervical carcinoma, colorectal carcinoma, melanoma, ovarian carcinoma or leukemia.