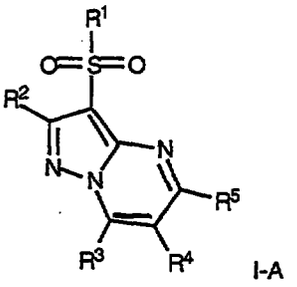
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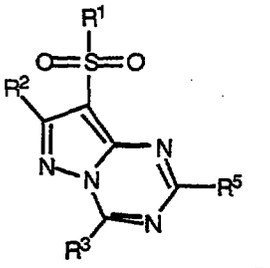
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The present invention is concerned with pyrazolopyrimidines and pyrazolotriazines of the general formulae



I-A

and



I-B

wherein

R^1	signifies phenyl, optionally substituted by one or more lower alkyl, halogen, lower alkoxy, tolyl, pyridyl, naphthyl or thiophenyl;
R^2	signifies hydrogen, lower alkyl, lower thioalkyl or hydroxy-lower-alkoxy;
R^3	signifies amino, lower alkylamino, di-lower-alkyl-amino, piperazinyl, optionally substituted by one or more lower alkyl, benzyl, phenyl or hydroxy-lower-alkyl, morpholinyl, imidazolyl, $(CH_3)_2N(CH_2)_nNH$-, $(CH_3)_2N(CH_2)_n)O$- or morpholinyl-$(CH_2)_nO$- in which n signifies 2 or 3;
R^4	signifies hydrogen, lower alkyl or hydroxy-lower-alkyl;
R^5	signifies hydrogen, halogen, lower alkyl, C_3-C_6-cycloalkyl, lower alkyl-lower-alkoxy, hydroxy-lower-alkyl-lower-alkoxy, $(CH_3)_2N(CH_2)_nNH$-, piperazinyl, optionally substituted by lower alkyl, methyl-piperazinyl, optionally substituted by lower alkyl, morpholinyl, methyl-morpholinyl, di-lower-alkylamino or di-lower-alkylamino-lower-alkyl, or
R^4 and R^5	together signify a group $-(CH_2)_m-$ or $-CH_2-S-CH_2-$ with m = 3 or 4,

as well as their pharmaceutically acceptable salts.

These compounds surprisingly have a selective affinity to 5HT-6 receptors. They are accordingly suitable for the treatment and prevention of central nervous disorders such as,

for example, psychoses, schizophrenia, manic depressions (Bryan L. Roth et al., J. Pharmacol. Exp. Ther., 268, pages 1403-1410 (1994)), depressions (David R. Sibley et al., Mol. Pharmacol., 43, pages 320-327 (1993)), neurological disorders (Anne Bourson et al., J. Pharmacol. Exp. Ther., 274, pages 173-180 (1995) ; R. P.Ward et al., Neuroscience, 64, pages 1105-1110 (1995)), memory disorders, Parkinson's disease, amyotrophic lateral sclerosis, Alzheimer's disease and Huntington's chorea (Andrew J. Sleight et al., Neurotransmissons, 11, pages 1-5 (1995)).

Objects of the present invention are the novel compounds of formulae I-A and I-B and their pharmaceutically useable salts per se, their use as therapeutically active substances, their manufacture, medicaments containing one or more of these compounds, optionally in the form of one of their pharmaceutically useable salts, as well as the production of such medicaments.

The following compounds are preferred for a use of the kind described above.

Especially preferred compounds of general formula I-A are those in which R^3 signifies amino, such as, for example

3-benzenesulphonyl-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ylamine,

3-(4-isopropyl-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]-pyrimidin-7-ylamine,

5-methyl-2-methylsulphanyl-3-(naphthalene-2-sulphonyl)-pyrazolo[1,5-a]-pyrimidin-7-ylamine,

3-(4-fluoro-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]-pyrimidin-7-ylamine,

3-benzenesulphonyl-5-cyclopropyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ylamine,

3-benzenesulphonyl-2-methylsulphanyl-6,7-dihydro-5H-cyclopenta[d]pyrazolo[1,5-a]pyrimidin-8-ylamine,

3-benzenesulphonyl-5-isopropyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ylamine,

5-methyl-2-methylsulphanyl-3-(toluene-2-sulphonyl)-pyrazolo[1,5-a]pyrimidin-7-ylamine,

5-methyl-2-methylsulphanyl-3-(toluene-3-sulphonyl)-pyrazolo[1,5-a]pyrimidin-7-ylamine,

3-benzenesulphonyl-5-methoxymethyl-2-methylsulphanyl-pyrazolo[1,5-a]-pyrimidin-6-ylamine,

3-benzenesulphonyl-N5,N5-dimethyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-5,7-diamine,

3-benzenesulphonyl-N5-(2-dimethylamino-ethyl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-5,7-diamine,

3-benzenesulphonyl-5-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ylamine and

3-benzenesulphonyl-5-dimethylamineomethyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ylamine.

Furthermore, compounds are preferred in which in formula I-A R^3 signifies piperazinyl, such as, for example,

3-benzenesulphonyl-5-methyl-2-methylsulphanyl-7-piperazin-1-yl-pyrazolo[1,5-a]pyrimidine,

3-(4-tert-butyl-benzenesulphonyl)-5-methyl-2-methylsulphanyl-7-piperazin-1-yl-pyrazolo[1,5-a]pyrimidine,

3-benzenesulphonyl-5,6-dimethyl-2-methylsulphanyl-7-piperazin-1-yl-pyrazolo-[1,5-a]pyrimidine,

3-benzenesulphonyl-2-methylsulphanyl-7-piperazin-1-yl-5-propyl-pyrazolo[1,5-a]pyrimidine,

3-benzenesulphonyl-5-cyclopropyl-2-methylsulphanyl-7-piperazin-1-yl-pyrazolo-[1,5-a]pyrimidine,

3-benzenesulphonyl-2-methylsulphanyl-8-piperazin-1-yl-6,7-dihydro-5H-cyclo-penta[d]pyrazolo[1,5-a]pyrimidine,

3-benzenesulphonyl-2-methylsulphanyl-8-piperazin-1-yl-5H,7H-pyrazolo[1,5-a]thieno[3,4-d]pyrimidine,

5-methyl-2-methylsulphanyl-7-piperazin-1-yl-3-(thiophen-2-sulphonyl)-pyrazolo-[1,5-a]pyrimidine,

3-benzenesulphonyl-2-ethyl-8-piperazin-1-yl-6,7-dihydro-5H-cyclopenta[d]-pyrazolo[1,5-a]pyrimidine and

5-methyl-2-methylsulphanyl-7-piperazin-1-yl-3-(toluol-2-sulphonyl)-pyrazolo[1,5-a]pyrimidine.

Moreover, of the already mentioned there are preferred compounds of formulae I-A and I-B in which R^3 signifies methylpiperazinyl, such as, for example,

3-benzenesulphonyl-5-cyclopropyl-2-methylsulphanyl-7-(4-methylpiperazin-1-yl)-pyrazolo[1,5-a]pyrimidine,

3-benzenesulphonyl-8-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-6,7-dihydro-5H-cyclopenta[d]pyrazolo[1,5-a]pyrimidine,

3-benzenesulphonyl-8-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-5H,7H-pyrazolo[1,5-a]thieno[3,4-d]pyrimidine,

3-benzenesulphonyl-5-isopropyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine and

2-[3-benzenesulphonyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-pyrazolo-[1,5-a]pyrimidine-5-yloxy]-ethanol.

8-Benzenesulphonyl-7-methylsulphanyl-pyrazolo[1,5-a][1,3,5]triazin-4-ylamine and 8-benzenesulphonyl-2-methyl-4-(4-methylpiperazin-1-yl)-7-methylsulphanyl-pyrazolo-[1,5-a][1,3,5]triazine are other preferred triazines of general formula I-B.

The term "lower alkyl" used in the present description denotes residues from 1 to 7, preferably from 1 to 4, carbon atoms, such as, for example, methyl, ethyl, n-propyl, i-propyl, n-butyl, i-butyl and t-butyl.

The term "lower alkoxy" denotes a lower alkyl residue in the sense of the foregoing definition bonded via an oxygen atom, such as, for example, methoxy, ethoxy, n-propoxy, i-propoxy, n-butoxy, i-butoxy and t-butoxy.

The term "lower alkylamino" denotes a lower alkyl residue in the sense of the foregoing definition bonded via a NH group, such as, for example, methylamino and ethylamino.

The term "di-lower-alkylamino" denotes to identical or different lower alkyl residues in the sense of the foregoing definition bonded via a nitrogen atom, such as, for example, dimethylamino, diethylamino or methyl-ethyl-amino.

The term "halogen" embraces fluorine, chlorine, bromine and iodine.

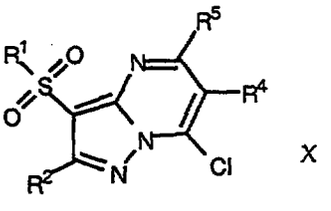
The term "lower thioalkyl" denotes a lower alkyl residue in the sense of the foregoing bonded via a sulphur atom.

The manufacture of the novel compounds can be effected in a manner known per se; that of formula I-A is described in Examples 1-123 and that of formula I-B is described in Examples 124-129. Moreover, Schemes 1-3 provide a general overview with respect to the possibilities available for the preparation of these novel compounds, with the starting compounds of formula III in Scheme 1 being known from the literature or being preparable in analogy to described methods.

The compounds of formulae I-A and I-B also embrace those compounds in which hydrogen can be replaced by tritium.

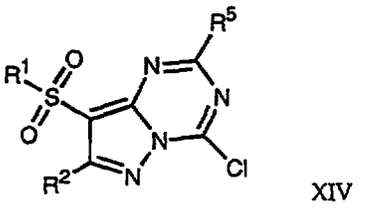
The compounds of formulae I-A and I-B can be manufactured by

reacting a compound of the formula



X

or a compound of the formula



XIV

with a compound of the formula

HR^3

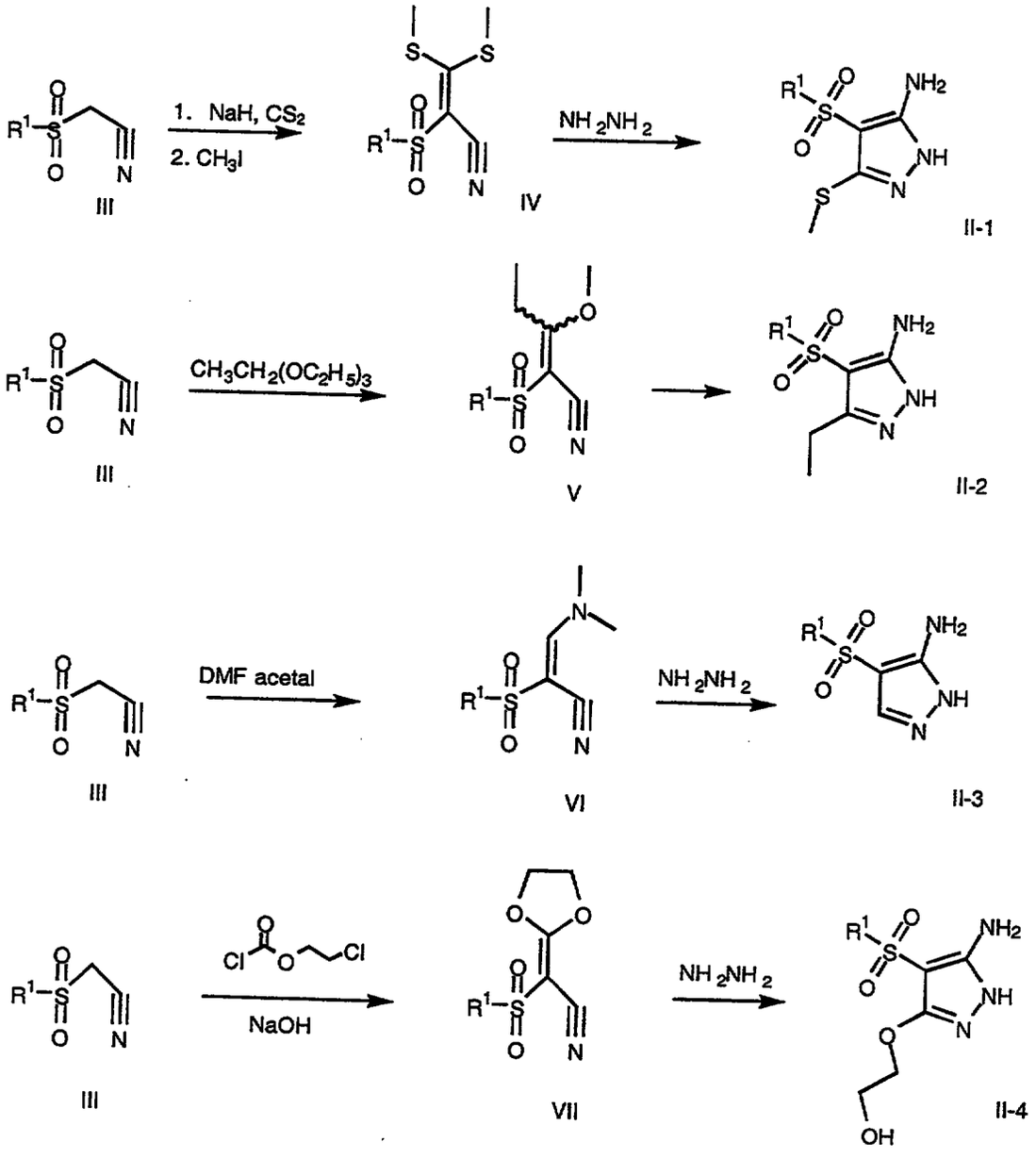
wherein R^1-R^5 have the significances described above, and optionally converting a compound of general formula I-A or I-B into a pharmaceutically acceptable salt.

The reaction of a compound of formulae X or XIV with a compound of the formula HR^3 is effected according to methods known per se. Conveniently, a compound

of formulae X or XIV is dissolved in DMF and depending on the reaction partner, which can be dissolved in DMF or an alcohol, reacted at room temperature or at the boiling temperature of the solvent used. Especially preferred reaction partners for the compounds of formulae X or XIV are piperazine, 1-methyl-piperazine, NH_3, methylamine, dimethylamine, morpholine, imidazole, N-benzylpiperidine, 1-(2-hydroxyethyl)-piperazine, 1-phenylpiperazine, 2-dimethylaminoethylamine or cis-2,6-dimethylpiperazine.

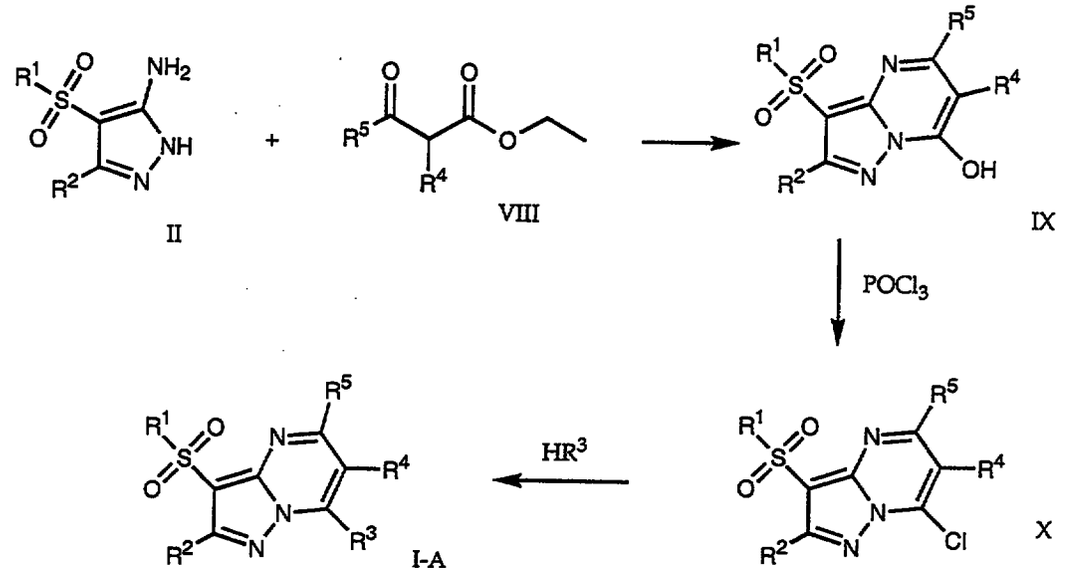
The compounds of formulae I-A and I-B can subsequently converted into pharmaceutically acceptable salts. Not only salts with inorganic acids, but also salts with organic acids come into consideration. Examples from the large number of such salts are hydrochlorides, hydrobromides, sulphates, nitrates, citrates, acetates, maleates, succinates, methanesulphonates, p-toluenesulphonates and the like. These salts can be manufactured according to methods which are known per se and which will be familiar to any person skilled in the art.

Scheme1



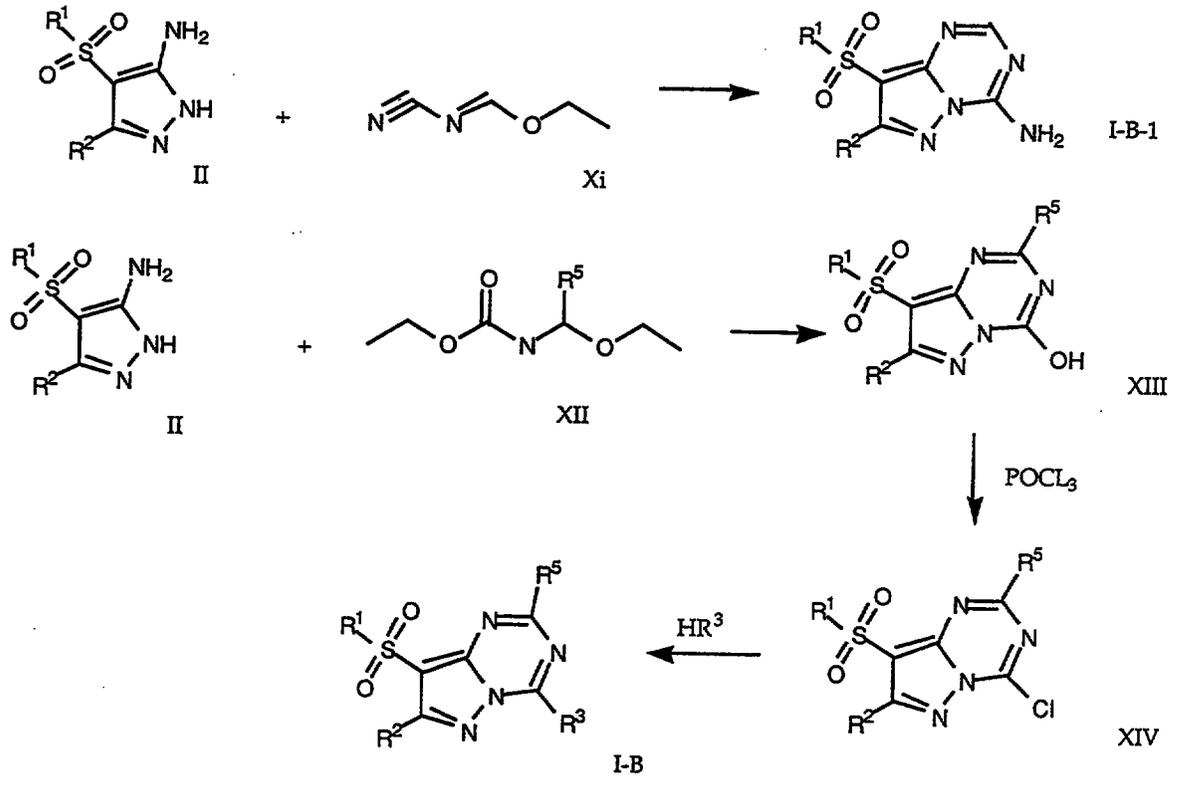
wherein R^1 has the significance given above.

Scheme 2



wherein R^1, R^2, R^3 and R^4 have the significance set forth above.

Scheme 3



wherein R^1, R^2, R^3 and R^5 have the significances set forth above.

The following compounds of formula I-A were manufactured according to Examples 1-123:

Example No.	R^1	R^2	R^3	R^4	R^5
1	Ph	SCH_3	Piperazinyl	H	CH_3
2	Ph	SCH_3	MePiperazinyl	H	CH_3
3	Ph	SCH_3	$NHCH_3$	H	CH_3
4	Ph	SCH_3	NH_2	H	CH_3
5	Ph	SCH_3	$N(CH_3)_2$	H	CH_3
6	Ph	SCH_3	Morpholinyl		CH_3
7	Ph	SCH_3	Imidazolyl	H	CH_3
8	Ph	SCH_3	Benzylpipe-razinyl	H	CH_3
9	Ph	SCH_3	Hydroxyethyl-piperazinyl	H	CH_3
10	Ph	SCH_3	Phenylpipe-razinyl	H	CH_3
11	Ph	SCH_3	$(CH_3)_2N$-CH_2CH_2NH	H	CH_3
12	Ph	SCH_3	2,6-Dimethyl-piperazinyl	H	CH_3
13	4-CH_3-Ph	SCH_3	Piperazinyl	H	CH_3
14	4-CH_3-Ph	SCH_3	MePiperazinyl	H	CH_3
15	4-CH_3-Ph	SCH_3	$NHCH_3$	H	CH_3
16	4-CH_3-Ph	SCH_3	NH_2	H	CH_3
17	4-CH_3-Ph	SCH_3	$N(CH_3)_2$	H	CH_3
18	4-$(CH_3)_2CHPh$	SCH_3	Piperazinyl	H	CH_3
19	4-$(CH_3)_2CHPh$	SCH_3	MePiperazinyl	H	CH_3
20	4-$(CH_3)_2CHPh$	SCH_3	NH_2	H	CH_3
21	4-$(CH_3)_2CHPh$	SCH_3	$N(CH_3)_2$	H	CH_3
22	4-t-Bu-Ph	SCH_3	Piperazinyl	H	CH_3
23	4-t-Bu-Ph	SCH_3	MePiperazinyl	H	CH_3
24	4-t-Bu-Ph	SCH_3	NH_2	H	CH_3

25	4-Cl-Ph	SCH_3	Piperazinyl	H	CH_3
26	4-Cl-Ph	SCH_3	MePiperazinyl	H	CH_3
27	4-Cl-Ph	SCH_3	NH_2	H	CH_3
28	4-Cl-Ph	SCH_3	$N(CH_3)_2$	H	CH_3
29	2,4-Di-Cl-Ph	SCH_3	Piperazinyl	H	CH_3
30	2,4-Di-Cl-Ph	SCH_3	MePiperazinyl	H	CH_3
31	2,4-Di-Cl-Ph	SCH_3	NH_2	H	CH_3
32	4-Br-Ph	SCH_3	Piperazinyl	H	CH_3
33	4-Br-Ph	SCH_3	MePiperazinyl	H	CH_3
34	4-Br-Ph	SCH_3	NH_2	H	CH_3
35	4-MeO-Ph	SCH_3	Piperazinyl	H	CH_3
36	4-MeO-Ph	SCH_3	MePiperazinyl	H	CH_3
37	4-MeO-Ph	SCH_3	NH_2	H	CH_3
38	2-Naphthyl	SCH_3	MePiperazinyl	H	CH_3
39	2-Naphthyl	SCH_3	$N(CH_3)_2$	H	CH_3
40	2-Naphthyl	SCH_3	NH_2	H	CH_3
41	4-F_3C-O-Ph	SCH_3	Piperazinyl	H	CH_3
42	4-F_3C-O-Ph	SCH_3	MePiperazinyl	H	CH_3
43	4-F-Ph	SCH_3	Piperazinyl	H	CH_3
44	4-F-Ph	SCH_3	MePiperazinyl	H	CH_3
45	4-F-Ph	SCH_3	NH_2	H	CH_3
46	4-I-Ph	SCH_3	NH_2	H	CH_3
47	Ph	SCH_3	Piperazinyl	CH_3	CH_3
48	Ph	SCH_3	MePiperazinyl	CH_3	CH_3
49	Ph	SCH_3	Piperazinyl	H	n-Propyl
50	Ph	SCH_3	MePiperazinvl	H	n-Propyl
51	Ph	SCH_3	Piperazinyl	H	Cyclo-propyl
52	Ph	SCH_3	MePiperazinvl	H	Cyclo-propyl
53	Ph	SCH_3	NH_2	H	Cyclo-propyl
54	Ph	SCH_3	Piperazinyl	$CH_2CH_2CH_2$	
55	Ph	SCH_3	MePiperazinyl	$CH_2CH_2CH_2$	

56	Ph	SCH_3	NH_2	$CH_2CH_2CH_2$	
57	Ph	SCH_3	Piperazinyl	$CH_2CH_2CH_2CH_2$	
58	Ph	SCH_3	MePiperazinyl	$CH_2CH_2CH_2CH_2$	
59	Ph	SCH_3	Piperazinyl	CH_2SCH_2	
60	Ph	SCH_3	MePiperazinyl	CH_2SCH_2	
61	Thiophenyl	SCH_3	Piperazinyl	H	CH_3
62	Thiophenyl	SCH_3	MePiperazinyl	H	CH_3
63	Thiophenyl	SCH_3	NH_2	H	CH_3
64	Ph	SCH_3	MePiperazinyl	H	i-Propyl
65	Ph	SCH_3	NH_2	H	i-Propyl
66	Ph	SCH_3	Piperazinyl	H	t-Butyl
67	Ph	SCH_3	Piperazinyl	H	CF_3
68	Ph	SCH_3	MePiperazinyl	H	CF_3
69	Ph	Ethyl	Piperazinyl	H	CH_3
70	Ph	Ethyl	MePiperazinyl	H	CH_3
71	Ph	Ethyl	$(CH_3)_2N$-CH_2CH_2NH	H	CH_3
72	4-Br-Ph	Ethyl	Piperazinyl	H	CH_3
73	4-Br-Ph	Ethyl	MePiperazinyl	H	CH_3
74	4-Br-Ph	Ethyl	$(CH_3)_2N$-CH_2CH_2NH	H	CH_3
75	4-MeO-Ph	Ethyl	Piperazinyl	H	CH_3
76	4-MeO-Ph	Ethyl	MePiperazinyl	H	CH_3
77	4-MeO-Ph	Ethyl	NH_2	H	CH_3
78	4-MeO-Ph	Ethyl	2,6-Dimethylpiperazinyl	H	CH_3
79	Ph	Ethyl	Piperazinyl	$CH_2CH_2CH_2$	
80	Ph	Ethyl	MePiperazinyl	$CH_2CH_2CH_2$	
81	Ph	Ethyl	NH_2	$CH_2CH_2CH_2$	
82	Ph	Ethyl	2,6-Dimethylpiperazinyl	$CH_2CH_2CH_2$	
83	Ph	H	Piperazinyl	H	CH_3
84	Ph	H	MePiperazinyl	H	CH_3

85	o-Tolyl	SCH_3	Piperazinyl	H	CH_3
86	o-Tolyl	SCH_3	MePiperazinyl	H	CH_3
87	o-Tolyl	SCH_3	NH_2	H	CH_3
88	m-Tolyl	SCH_3	Piperazinyl	H	CH_3
89	m-Tolyl	SCH_3	MePiperazinyl	H	CH_3
90	m-Tolyl	SCH_3	NH_2	H	CH_3
91	3-Pyridyl	SCH_3	Piperazinyl	H	CH_3
92	3-Pyridyl	SCH_3	MePiperazinyl	H	CH_3
93	3-Pyridyl	SCH_3	NH_2	H	CH_3
94	Ph	SCH_3	MePiperazinyl	CH_2CH_2OH	CH_3
95	Ph	OCH_2CH_2OH	MePiperazinyl	H	CH_3
96	Ph	SCH_3	Piperazinyl	H	CH_2CH_2O-CH_3
97	Ph	SCH_3	MePiperazinyl	H	CH_2CH_2O-CH_3
98	Ph	SCH_3	NH_2	H	CH_2CH_2O-CH_3
99	Ph	SCH_3	Piperazinyl	H	CH_2OCH_3
100	Ph	SCH_3	MePiperazinyl	H	CH_2OCH_3
101	Ph	SCH_3	NH_2	H	CH_2OCH_3
102	Ph	SCH_3	MePiperazinyl	H	Cl
103	Ph	SCH_3	MePiperazinyl	H	H
104	Ph	SCH_3	MePiperazinyl	H	OCH_2CH_2OH
105	Ph	SCH_3	NH_2	H	Cl
106 a	Ph	SCH_3	NH_2	H	$N(CH_3)_2$
106 b	Ph	SCH_3	NH_2	H	$NHCH_2CH_2$-$N(CH_3)_2$
107	Ph	SCH_3	NH_2	H	MePiperazinyl
108	Ph	Ethyl	MePiperazinyl	H	Cl
109	Ph	Ethyl	MePiperazinyl	H	H

110	Ph	Ethyl	MePiperazinyl	H	MePiperazinyl
111	Ph	Ethyl	MePiperazinyl	H	Morpholinyl
112	Ph	Ethyl	MePiperazinyl	H	OCH_2CH_2OH
113 a	Ph	Ethyl	MePiperazinyl	H	$N(CH_3)_2$
113 b	Ph	Ethyl	MePiperazinyl	H	$NHCH_2CH_2N(CH_3)_2$
114	Ph	Ethyl	NH_2	H	Cl
115	Ph	Ethyl	NH_2	H	MePiperazinyl
116 a	Ph	Ethyl	NH_2	H	$N(CH_3)_2$
116 b	Ph	Ethyl	NH_2	H	$NHCH_2CH_2N(CH_3)_2$
117	Ph	SCH_3	NH_2	H	$CH_2N(CH_3)_2$
118	Ph	SCH_3	NH_2	H	CH_2NHMePip
119	Ph	SCH_3	MePiperazinyl	H	CH_2NHMePip
120	Ph	SCH_3	NH_2	H	CH_2Morph
121	Ph	SCH_3	MePiperazinyl	H	CH_2Morph
122	Ph	SCH_3	$O(CH_2)N(CH_3)_2$	H	CH_3
123	Ph	SCH_3	$O(CH_2)_2$morpholinyl	H	CH_3

The following compounds of formula I-B were manufactured in accordance with Synthesis Examples 122-127:

Example No.	R^1	R^2	R^3	R^5
124	Ph	SCH_3	NH_2	H
125	Ph	SCH_3	$NHCH_3$	CH_3
126	Ph	SCH_3	$N(CH_3)_2$	CH_3
127	Ph	SCH_3	$NH(CH_2)_3N(CH_3)_2$	CH_3
128	Ph	SCH_3	MePiperazinyl	CH_3
129	Ph	SCH_3	Benzylpiperazinyl	CH_3

As mentioned earlier, the compounds of formulae I-A and I-B are novel. They have pharmacological properties and possess only a low toxicity. They have as a common feature they pronounced affinity to 5-HT_6 receptors and, on the basis of their action at this

receptor, are suitable for the treatment or prevention of central nervous disorders such as, for example, psychoses, schizophrenia, manic depressions, neurological disorders, memory disorders, Parkinson's disease, amyotrophic lateral sclerosis, Alzheimer's disease and Huntington's chorea.

The binding of the compounds of formulae I-A and I-B in accordance with the invention to 5-HT_6 receptors was determined as follows.

Membranes obtained from HEK 1293 cells which had been transfected with 5-HT_6 receptors from rats were used.

The cells were purified by two-fold centrifugation (10 minutes at 3000 g) in phosphate buffer-sodium chloride solution. The cell mass was suspended in an ice-cold solution consisting of 50 mm Tris-HCl buffer, 10 mm $MgCl_2$, 0.5 mm EDTA and 0.1 mm phenylmethylsulphonyl fluoride and homogenized (Polytron homogenizer, 15 seconds at maximum speed). The homogenizate was incubated at 37°C for 10 minutes and subsequently centrifuged (20 minutes at 20 000 g). The cell mass was again suspended in the aforementioned Tris buffer solution. The cell concentration obtained was 4×10^7 cells/ml. 1 ml aliquots of the homogenizate were frozen at -80°C.

Displacement experiments were carried out in order to determine the affinity of the test substance to the 5-HT_6 receptor. For the performance of the test, the homogenizate was thawed and suspended in a buffer solution (pH 7.4) consisting of 50 mm Tris-HCl buffer, 5 mm $MgCl_2$, 10^{-5} M pargyline and 0.1% ascorbic acid. 100 µl of membrane suspension, 50 µl of [^{3}H]-LSD (specific activity 85Ci/mMol, final concentration 1 NM) and 50 µl of test substance solution were incubated at 37°C for 1 hour. The respective test substance was investigated at 7 different concentrations of 10^{-10} M to 10^{-4} M. The binding reaction of the test substance was interrupted by rapid filtration through a Whatmann GF/B filter. The filters were washed with 2 x 2 ml of Tris-HCl buffer (50 mm pH 7.4) and the radioactivity of the filter was measured by scintillation spectroscopy in 2 ml of scintillation solution. All tests were carried out in triplicate and repeated three times. The pKi values ($pKi = -\log_{10}Ki$) of the test substances have been determined. The Ki value is defined by the following formula

$$Ki = \frac{IC_{50}}{1 + \frac{[L]}{K_D}}$$

in which the IC_{50} values are those concentrations of the test compounds in NM by which 50% of the ligands bonded to the receptor are displaced. [L] is the concentration of the ligand and the K_D value is the dissociation constant of the ligand.

The compounds in accordance with the invention have a selective affinity to 5-HT 6 receptors with a pKi value between 6.5 and 9.5.

The compounds of formulae I-A and I-B and the pharmaceutically acceptable salts of the compounds of formulae I-A and I-B can be used as medicaments, e.g. in the form of pharmaceutical preparations. The pharmaceutical preparations can be administered orally e.g. in the form of tablets, coated tablets, dragées, hard and soft gelatine capsules, solutions, emulsions or suspensions. However, the administration can also be effected rectally, e.g. in the form of suppositories, or parentally, e.g. in the form of injection solutions.

For the production of pharmaceutical preparations, the compounds of formulae I-A and I-B and the pharmaceutically acceptable salts of the compounds of formulae I-A and I-B can be processed with pharmaceutically inert, inorganic or organic carriers. Lactose, corn starch or derivatives thereof, talc, stearic acid or its salts and the like can be used as such carriers for tablets, coated tablets, dragées and hard gelatine capsules. Suitable carriers for soft gelatine capsules are, for example, vegetable oils, waxes, fats, semi-solid and liquid polyols and the like. Depending on the nature of the active ingredient no carriers are, however, usually required in the case of soft gelatine capsules. Suitable carriers for the production of solutions and syrups are, for example, water, polyols, glycerol, vegetable oils and the like. Suitable carriers for suppositories are, for example, natural or hardened oils, waxes, fats, semi-liquid or liquid polyols and the like.

Moreover, the pharmaceutical preparations can contain preservatives, solubilizers, stabilizers, wetting agents, emulsifiers, sweeteners, colorants, flavorants, salts for varying the osmotic pressure, buffers, masking agents or antioxidants. They can also contain still other pharmaceutically valuable substances.

Medicaments containing a compound of formula I-A or I-B or a pharmaceutically acceptable salt thereof and a therapeutically inert carrier are also an object of the present

invention, as is a process for their production, which comprises bringing one or more compounds of formula I and/or pharmaceutically acceptable acid addition salts thereof and, if desired, one or more other therapeutically valuable substances together with one or more therapeutically inert carriers into a galenical administration form in a manner known per se.

In accordance with the invention compounds of general formulae I-A and I-B as well as their pharmaceutically acceptable salts can be used in the treatment or prevention of central nervous disorders, such as depressions, psychoses, schizophrenia, neurological disorders, memory disorders, Parkinson's disease, amyotrophic lateral sclerosis, Alzheimer's disease, Huntington's chorea and for the production of corresponding medicaments. The dosage can vary within wide limits and will, of course, be fitted to the individual requirements in each particular case. In the case of oral administration the dosage lies in a range of about 0.1 mg per dose to about 1000 mg per day of a compound of general formula I-A or I-B or the corresponding amount of a pharmaceutically acceptable salt thereof, although the upper limit can also be exceeded when this is shown to be indicated.

The following Examples serve to illustrate the present invention in more detail. However, they are not intended to limit its scope in any manner.

Example 1

3-Benzenesulphonyl-5-methyl-2-methylsulphanyl-7-piperazin-1-yl-pyrazolo-[1,5-a]pyrimidine

a) A solution of 6.30 g (21 mmol) of 4-benzenesulphonyl-5-methylsulphanyl-2H-pyrazol-3-ylamine and 3.24 ml (25.8 mmol) of ethyl acetoacetate in 20 ml of acetic acid was heated at reflux for 1.5 hrs. The reaction solution was cooled to 0°C and stirred at this temperature for 30 min. The separated crystals were filtered off under suction and dried at 50°/10 Torr. There were thus obtained 6.10 g (87%) of 3-benzenesulphonyl-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ol as white crystals, m.p. > 220°.

b) A suspension of 3.0 g (8.94 mmol) of 3-benzenesulphonyl-5-methyl-2-methyl-sulphanyl-pyrazolo[1,5-a]pyrimidin-7-ol in 20 ml of $POCl_3$ was heated at reflux for 45 min. The reaction solution was cooled to RT and evaporated in a high vacuum. The residue was treated with 100 ml of ice-water and the pH value of the solution was adjusted

to 8 with sat. $NaHCO_3$ solution. The aqueous phase was extracted three times with 100 ml of CH_2Cl_2, and the organic phases were dried ($MgSO_4$), filtered and evaporated. Chromatography (SiO_2, CH_2Cl_2/AcOEt 19:1) of the residue yielded 3.0 g (94%) of 3-benzenesulphonyl-7-chloro-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine as pale yellow crystals, m.p. 163-165°C.

c) 0.3 g (3.4 mmol) of piperazine in 10 ml of DMF was added to a solution of 0.6 g (1.7 mmol) of 3-benzenesulphonyl-7-chloro-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine in 10 ml of DMF and the mixture was stirred at 60°C for 2 hrs. The reaction solution was cooled to RT and evaporated in a high vacuum. The residue was partitioned between 2N NaOH and CH_2Cl_2. The aqueous phase was extracted three times with CH_2Cl_2, and the combined organic phases were dried ($MgSO_4$), filtered and evaporated. Subsequent chromatography (SiO_2, CH_2Cl_2/MeOH 8:1) and crystallization from EtOH yielded 0.35 g (51%) of 3-benzenesulphonyl-5-methyl-2-methylsulphanyl-7-piperazin-1-yl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 201-202°.

Example 2

3-Benzenesulphonyl-5-methyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine

0.15 g (1.5 mmol) of 1-methyl-piperazine in 10 ml of DMF was added to a solution of 0.45 g (1.275 mmol) of 3-benzenesulphonyl-7-chloro-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine in 10 ml in DMF and stirred at 60° for 2 hrs. The reaction solution was cooled to RT and evaporated in a high vacuum. The residue was partitioned between 2N NaOH and CH_2Cl_2. The aqueous phase was extracted three times with CH_2Cl_2, and the combined organic phases were dried ($MgSO_4$), filtered and evaporated. Subsequent chromatography (SiO_2, CH_2Cl_2/MeOH 15:1) and crystallization from EtOH yielded 0.40 g (75%) of 3-benzenesulphonyl-5-methyl-7-(4-methyl-piperazine-1-yl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 209-210°.

Example 3

(3-Benzenesulphonyl-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-yl)-methyl-amine

10 ml of a 33% solution of methylamine in EtOH was added to a solution of 0.3 g (0.85 mmol) of 3-benzenesulphonyl-7-chloro-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine in 5 ml of DMF and stirred at RT for 2 hrs. The reaction solution was evaporated in a high vacuum and the residue was partitioned between 2N NaOH and CH_2Cl_2. The aqueous phase was extracted three times with CH_2Cl_2, and the combined organic phases were dried ($MgSO_4$), filtered and evaporated. Subsequent chromatography (SiO_2,CH_2Cl_2/AcOEt 15:1) and crystallization from EtOH yielded 0.18 g (60%) of (3-benzenesulphonyl-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-yl)-methyl-amine as colourless crystals, m.p. > 230°.

Example 4

(3-Benzenesulphonyl-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-yl)amine

10 ml of a 50% solution of NH_3 in MeOH were added to a solution of 0.40 g (1.13 mmol) of 3-benzenesulphonyl-7-chloro-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine in 10 ml of DMF and stirred at RT for 2 hrs. The reaction solution was evaporated in a high vacuum and the residue was partitioned between 2N NaOH and CH_2Cl_2. The aqueous phase was extracted three times with CH_2Cl_2, and the combined organic phases were dried ($MgSO_4$), filtered and evaporated. Subsequent chromatography (SiO_2, CH_2Cl_2/AcOEt 7:1) and crystallization from EtOH yielded 0.25 g (66%) of (3-benzenesulphonyl-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-yl)amine as colourless crystals, m.p. > 230°.

Example 5

(3-Benzenesulphonyl-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-yl)-dimethyl-amine added to a solution of 0.30 g (0.85 mmol) 3-benzenesulphonyl-7-chloro-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine in 10 ml of DMF and stirred at RT for 1 hr. The reaction solution was evaporated in a high vacuum and the residue was partitioned between 2N NaOH and CH_2Cl_2. The aqueous phase was extracted three times with CH_2Cl_2, and the combined organic phases were dried ($MgSO_4$), filtered and evaporated. Subsequent chromatography (SiO_2, CH_2Cl_2/AcOEt 15:1) and crystallization

from EtOH yielded 0.18 g (58%) of (3-benzene- sulphonyl-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-yl)-dimethyl-amine as colourless crystals, m.p. > 230°.

Example 6

3-Benzenesulphonyl-5-methyl-2-methylsulphanyl-7-morpholin-4-yl-pyrazolo-[1,5-a]pyrimidine

0.20 g (2.2 mmol) of morpholine was added to a solution of 0.25 g (0.85 mmol) of 3-benzenesulphonyl-7-chloro-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine in 10 l of DMF and stirred at 60° for 1 hr. The reaction solution was cooled to RT and evaporated in a high vacuum. The residue was partitioned between 2N NaOH and CH_2Cl_2. The aqueous phase was extracted three times with CH_2Cl_2, and the combined organic phases were dried ($MgSO_4$), filtered and evaporated. Subsequent chromatography (SiO_2, CH_2Cl_2/MeOH 25:1) and crystallization from EtOH yielded 0.20 g (70%) of 3-benzene- sulphonyl-5-methyl-2-methylsulphanyl-7-morpholin-4-yl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 206-208°.

Example 7

3-Benzenesulphonyl-7-imidazol-1-yl-5-methyl-2-methylsulphanyl-pyrazolo-[1,5-a]pyrimidine

0.08 g (1.5 mmol) of sodium methanolate was added to a suspension of 0.122 g (1.8 mmol) of imidazole in 30 ml of DMF and stirred at 60° for 15 min. Subsequently, a solution of 0.53 g (1.5 mmol) of 3-benzenesulphonyl-7-chloro-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]-pyrimidine in 10 ml of DMF was added to this suspension and stirred at 60° for 1 hr. The reaction solution was cooled to RT and evaporated in a high vacuum. The residue was partitioned between 2N NaOH and CH_2Cl_2. The aqueous phase was extracted three times with CH_2Cl_2, and the combined organic phases were dried ($MgSO_4$), filtered and evaporated. Subsequent chromatography (SiO_2, CH_2Cl_2/MeOH 15:1) and crystallization from EtOH yielded 0.30 g (51%) of 3-benzenesulphonyl-7-imidazol-1-yl-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine as colourless crystal, m.p. >230°.

Example 8

3-Benzenesulphonyl-7-(4-benzyl-piperazin-1-yl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine

0.35 g (2 mmol) of N-benzyl-piperidine was added to a solution of 0.35 g (1 mmol) of 3-benzenesulphonyl-7-chloro-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine in 5 ml of DMF and stirred at 60° for 2 hrs. The reaction solution was cooled to RT and evaporated in a high vacuum. The residue was partitioned between 2N NaOH and CH_2Cl_2. The aqueous phase was extracted three times with CH_2Cl_2, and the combined organic phases were dried ($MgSO_4$), filtered and evaporated. Subsequent chromatography (SiO_2, CH_2Cl_2/MeOH 19:1) and crystallization from EtOH yielded 0.36 g (73%) of 3-benzene- sulphonyl-7-(4-benzyl-piperazin-1-yl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine as colourless crystals m.p. 156-158°.

Example 9

2-[4-(3-Benzenesulphonyl-5-methyl-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-yl)-piperazin-1-yl]-ethanol

0.26 g (2 mmol) of 1(2-hydroxyethyl)-piperazine was added to a solution of 0.35 g (1 mmol) of 3-benzenesulphonyl-7-chloro-5-methyl-2-methylsulphanyl-pyrazolo[1,5 a]pyrimidine and stirred at 60° for 2 hrs. The reaction solution was cooled to RT and evaporated in a high vacuum. The residue was partitioned between 2N NaOH and CH_2Cl_2. The aqueous phase was extracted three times with CH_2Cl_2, and he combined organic phases were dried ($MgSO_4$), filtered and evaporated. Subsequent chromatography (SiO_2,CH_2Cl_2/MeOH 19:1) and crystallization from EtOH yielded 0.36 g (73%) of 2-[4-(3-benzenesulphonyl-5-methyl-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-yl)-piperazin-yl]-ethanol as colourless crystals, m.p. 189-190°.

Example 10

3-Benzenesulphonyl-5-methyl-2-methylsulphanyl-7-(4-phenyl-piperazine-1-yl)-pyrazolo-[1,5-a]pyrimidine

0.16 g (1 mmol) of N-phenyl-piperazine was added to a solution of 0.17 g (0.5 mmol) of 3-benzenesulphonyl-7-chloro-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine in 3 ml of DMF and stirred at 60° for 2 hrs. The reaction solution was cooled to RT and evaporated in a high vacuum. The residue was partitioned between 2N NaOH and CH_2Cl_2. The aqueous phase was extracted three times with CH_2Cl_2, and the combined organic phases were dried ($MgSO_4$), filtered and evaporated. Subsequent chromatography (SiO_2, CH_2Cl_2/MeOH 19:1) and crystallization from EtOH yielded 0.16 g (67%) of 3-benzenesulphonyl-5-methyl-2-methylsulphanyl-7-(4-phenyl-piperazin-1-yl)-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. > 230°.

Example 11

N-(3-Benzenesulphonyl-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-yl)-N',N'-dimethyl-ethane-1,2-diamine

0.088 g (1 mmol) of 2-dimethylaminoethylamine was added to a solution of 0.17 g (0.5 mmol) of 3-benzenesulphonyl-7-chloro-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine in 3 ml of DMF and stirred at 60° for 2 hrs. The reaction solution was cooled to RT and evaporated in a high vacuum. The residue was partitioned between 2N NaOH and CH_2Cl_2. The aqueous phase was extracted three times with CH_2Cl_2, and the combined organic phases were dried ($MgSO_4$), filtered and evaporated. Subsequent chromatography (SiO_2, CH_2Cl_2/MeOH 19:1) and crystallization from EtOH yielded 0.23 g (73%) of N-(3-benzenesulphonyl-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-yl)-N',N'-dimethyl-ethane-1,2-diamine as colourless crystals, m.p. 190-191°.

Example 12

(3R,5S)-3-Benzenesulphonyl-7-(3,5-dimethyl-piperazin-1-yl)-5-methyl-2-methyl-sulphanyl-pyrazolo[1,5-a]pyrimidine

0.28 g (2 mmol) of cis-2,6-dimethylpiperazine was added to a solution of 0.35 g (1 mmol) of 3-benzenesulphonyl-7-chloro-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine in 5 ml of DMF and stirred at 60° for 2 hrs. The reaction solution was cooled to RT and evaporated in a high vacuum. The residue was partitioned between 2N NaOH and CH_2Cl_2. The aqueous phase was extracted three times with CH_2Cl_2, and the combined organic phases were dried ($MgSO_4$), filtered and evaporated. Subsequent chromatography (SiO_2, CH_2Cl_2/MeOH 19:1) and crystallization from EtOH yielded 0.29 g (67%) of (3R,5S)-3-benzenesulphonyl-7-(3,5-dimethyl-piperazin-1-yl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 220-221°.

Example 13

3-Benzenesulphonyl-5-methyl-2-methylsulphanyl-7-piperazin-1-yl-pyrazolo[1,5-a]-pyrimidine

a) A solution of 7.08 g (25 mmol) of 5-methylsulphanyl-4-(toluene-4-sulphonyl)-2H-pyrazol-3-ylamine and 4.0 ml (31.25 mmol) of ethyl acetoacetate in 30 l of acetic acid was heated at reflux for 1.5 hrs. The reaction solution was cooled to 0° and stirred at this temperature for 30 min. The separated crystals were filtered off under suction and dried at 50°/10 Torr. There were thus obtained 7.20 g (82%) of 5-methyl-2-methylsulphanyl-3-(toluene-4-sulphonyl)-pyrazolo[1,5-a]pyrimidin-7-ol as white crystals, m.p. > 230°.

b) A suspension of 3.8 g (10.8 mmol) of 5-methyl-2-methylsulphanyl-3-(toluene-4-sulphonyl)-pyrazolo[1,5-a]pyrimidin-7-ol in 20 l of $POCl_3$ was heated at reflux for 1 hr. The reaction solution was cooled to RT and evaporated. The residue was treated with 100 ml of ice-water and the pH value of the solution was adjusted to 8 with sat. $NaHCO_3$ solution. The aqueous phase was extracted three times with CH_2Cl_2, and the organic phases were dried ($MgSO_4$), filtered and evaporated. Chromatography (SiO_2, CH_2Cl_2/AcOEt 25:1) of the residue yielded 3.8 g (95%) of 7-chloro-5-methyl-2-methylsulphanyl-3-(toluene-4-sulphonyl)-pyrazolo[1,5-a]pyrimidine as pale yellow crystals, m.p. 197-198°.

c) 0.3 g (3.4 mmol) of piperazine in 10 ml of DMF was added to a solution of 0.62 g (1.7 mmol) of 7-chloro-5-methyl-2-methyl-sulphanyl-3-(toluene-4-sulphonyl)-pyrazolo[1,5-a]pyrimidine in 10 ml of DMF and stirred at 60° for 2 hrs. The reaction solution was cooled to RT and evaporated in a high vacuum. The residue was partitioned between 2N NaOH and CH_2Cl_2. The aqueous phase was extracted three times with CH_2Cl_2, and the combined organic phases were dried ($MgSO_4$), filtered and evaporated. Subsequent chromatography (SiO_2, CH_2Cl_2/MeOH 6:1) and crystallization from EtOH yielded 0.30 g (42%) of 3-benzenesulphonyl-5-methyl-2-methylsulphanyl-7-piperazin-1-yl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 137-139°.

Example 14

5-Methyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-3-(toluene-4-sulphonyl)-pyrazolo[1,5-a]pyrimidine

In an analogous manner to that described in Example 2, from 7-chloro-5-methyl-2-methylsulphanyl-3-(toluene-4-sulphonyl)-pyrazolo[1,5-a]pyrimidine and 1-methyl-piperazine there was obtained 5-methyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-3-(toluene-4-sulphonyl)-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 221-223°.

Example 15

Methyl-[5-methyl-2-methylsulphanyl-3-(toluene-4-sulphonyl)-pyrazolo[1,5-a]pyrimidin-7-yl]-amine

In an analogous manner to that described in Example 3, from 7-chloro-5-methyl-2-methylsulphanyl-3-(toluene-4-sulphonyl)-pyrazolo[1,5-a]pyrimidine and methylamine in EtOH there was obtained methyl-[5-methyl-2-methylsulphanyl-3-(toluene-4-sulphonyl)-pyrazolo[1,5-a]pyrimidin-7-yl]-amine as colourless crystals, m.p. > 230°.

Example 16

5-Methyl-2-methylsulphanyl-3-(toluene-4-sulphonyl)-pyrazolo[1,5-a]pyrimidin-7-ylamine

In an analogous manner to that described in Example 4, from 7-chloro-5-methyl-2-methylsulphanyl-3-(toluene-4-sulphonyl)-pyrazolo[1,5-a]pyrimidine and NH_3 in MeOH

there was obtained 5-methyl-2-methylsulphanyl-3-(toluene-4-sulphonyl)-pyrazolo[1,5-a]-pyrimidin-7-ylamine as colourless crystals, m.p. > 230°.

Example 17

Dimethyl-[5-methyl-2-methylsulphanyl-3-(toluene-4-sulphonyl)-pyrazolo[1,5-a]-pyrimidin-7-yl]-amine

In an analogous manner to that described in Example 5, from 7-chloro-5-methyl-2-methylsulphanyl-3-(toluene-4-sulphonyl)-pyrazolo[1,5-a]pyrimidine and dimethylamine in EtOH there was obtained dimethyl-[5-methyl-2-methylsulphanyl-3-(toluene-4-sulphonyl)-pyrazolo[1,5-a]pyrimidin-7-yl]-amine as colourless crystals, m.p. > 230°.

Example 18

3-(4-Isopropyl-benzenesulphonyl)-5-methyl-2-methylsulphanyl-7-piperazin-1-yl-pyrazolo[1,5-a]pyrimidine

a) 1.8 g (44.6 mmol) of NaH (60% suspension in oil) were added portionwise to a solution of 5 g (22.3 mmol) of (4-isopropyl-benzenesulphonyl)-acetonitrile and 1.4 ml of CS_2 in 14 ml of DMSO and stirred at room temperature for 45 min. Subsequently, 2.9 ml (47 mmol) of methyl iodide were slowly added dropwise thereto and stirred at RT for 2 hrs. After the addition of 30 ml of H_2O the separated crystals were filtered off under suction and crystallized from $EtOH_2/CH_2Cl_2$. There were thus obtained 4.9 g (67%) of 2-(4-isopropyl-benzenesulphonyl)-3,3-bis-methylsulphanyl-acrylonitrile as pale yellow crystals, m.p. 87-.

b) 0.36 ml (7.3 ml) of NH_2NH_2 was added to a solution of 2.0 g (6.1 mmol) of 2-(4-isopropyl-benzenesulphonyl)-3,3-bis-methylsulphanyl-acrylonitrile in 11 ml of EtOH and heated at reflux for 30 min. The pale brown solution was evaporated and chromatographed (SiO_2, CH_2Cl_2/MeOH 9:1). There were thus obtained 1.82 g (69%) of 4-isopropyl-benzenesulphonyl)-5-methylsulphanyl-2H-pyrazol-3-ylamine as a beige foam.

c) A solution of 1.80 g (5.8 mmol) of 4-(4-isopropyl-benzenesulphonyl)-5-methyl-sulphanyl)-2H-pyrazol-3-ylamine and 1.13 ml (8.8 mmol) of ethyl acetoacetate in 10 ml of acetic acid was heated at reflux for 1.5 hrs. The reaction solution was cooled to 0° and

stirred at this temperature for 30 min. The separated crystals were filtered off and dried at 50°/10 Torr. There were thus obtained 1.82 g (83%) of 3-(4-isopropyl-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ol as white crystals, m.p. > 230°.

d) A suspension of 1.8 g (4.8 mmol) of of 3-(4-isopropyl-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ol in 30 ml of $POCl_3$ was heated at reflux for 45 min. The reaction solution was cooled to RT and evaporated. The residue was treated with 100 ml of ice-water and the pH value of the solution was adjusted to 8 with sat. $NaHCO_3$ solution. The aqueous phase was extracted three times with CH_2Cl_2, and the organic phases were dried ($MgSO_4$), filtered and evaporated. Chromatography (SiO_2, CH_2Cl_2/AcOEt 19:1) of the residue yielded 1.78 g (93%) of 3-(4-isopropyl-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ol as pale yellow crystals, m.p.183-184°.

e) In an analogous manner to that described in Example 1c), from 7-chloro-3-(4-isopropyl-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine and piperazine there was obtained 3-(4-isopropyl-benzenesulphonyl)-5-methyl-2-methyl-sulphanyl-7-piperazin-1-yl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. > 230°.

Example 19

3-(4-Isopropyl-benzenesulphonyl)-5-methyl-7-(4-methyl-piperazin-1-yl)-2-methyl-sulphanyl-pyrazolo[1,5-a]pyrimidine

In an analogous manner to that described in Example 2, from 7-chloro-3-(4-isopropyl-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine and 1-methyl-piperazine there was obtained 3-(4-isopropyl-benzenesulphonyl)-5-methyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 218-219°.

Example 20

3-(4-Isopropyl-benzenesulphonyl)-5-methyl-7-(4-methyl-piperazin-1-yl)-2-methyl-sulphanyl-pyrazolo[1,5-a]pyrimidine

In an analogous manner to that described in Example 4, from 7-chloro-3-(4-isopropyl-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine and NH_3 in MeOH there was obtained 3-(4-isopropyl-benzenesulphonyl)-5-methyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. >230°.

Example 21

[3-(4-Isopropyl-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]-pyrimidin-7-yl]-dimethyl-amine

In an analogous manner to that described in Example 5, from 7-chloro-3-(4-isopropyl-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine and dimethyl-amine in EtOH there was obtained [3-(4-isopropyl-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-yl]-dimethyl-amine as colourless crystals, m.p. 222-224°.

Example 22

3-(4-tert-Butyl-benzenesulphonyl)-5-methyl-2-methylsulphanyl-7-piperazin-1-yl-pyrazolo[1,5-a]pyrimidine

a) In an analogous manner to that described in Example 18 a) to d), starting from (4-tert-butyl-benzenesulphonyl)-acetonitrile there was obtained 3-(4-tert-butyl-benzene-sulphonyl)-7-chloro-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine as a beige foam.

b) In an analogous manner to that described in Example 1c), from 3-(4-tert-butyl-benzenesulphonyl)-7-chloro-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine and piperazine there was obtained 3-(4-tert-butyl-benzenesulphonyl)-5-methyl-2-methyl-sulphanyl-7-piperazin-1-yl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 178-180°.

Example 23

3-(4-tert-Butyl-benzenesulphonyl)-5-methyl-7-(4-methyl-piperazin-1-yl)-2-methyl-sulphanyl-pyrazolo[1,5-a]pyrimidine

In an analogous manner to that described in Example 2, from 3-(4-tert-butyl-benzene-sulphonyl)-7-chloro-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine and 1-methyl-piperazine there was obtained 3-(4-tert-butyl-benzenesulphonyl)-5-methyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 238-239°.

Example 24

3-(4-tert-Butyl-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]-pyrimidin-7-ylamine

In an analogous manner to that described in Example 4, from 3-(4-tert-butyl-benzene-sulphonyl)-7-chloro-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine and NH_3 in MeOH there was obtained 3-(4-tert-butyl-benzenesulphonyl)-5-methyl-2-methyl-sulphanyl-pyrazolo[1,5-a]pyrimidin-7-ylamine as colourless crystals, m.p. > 230°.

Example 25

3-(4-Chloro-benzenesulphonyl)-5-methyl-2-methylsulphanyl-7-piperazin-1-yl-pyrazolo[1,5-a]pyrimidine

(a) In an analogous manner to that described in Example 18 a) to d), starting from (4-chloro-benzenesulphonyl)-acetonitrile there was obtained 7-chloro-3-(4-chloro-benzene-sulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine as a colourless foam.

(b) In an analogous manner to that described in Example 1c), from 7-chloro-3-(4-chloro-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine and piperazine there was obtained 3-(4-chloro-benzenesulphonyl)-5-methyl-2-methyl-sulphanyl-7-piperazin-1-yl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 214-217°.

Example 26

3-(4-Chloro-benzenesulphonyl)-5-methyl-7-(4-methyl-piperazin-1-yl)-2-methyl-sulphanyl-pyrazolo[1,5-a]pyrimidine

In an analogous manner to that described in Example 2, from 7-chloro-3-(4-chloro-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine and 1-methyl-piperazine there was obtained 3-(4-chloro-benzenesulphonyl)-5-methyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 200-201°.

Example 27

3-(4-Chloro-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ylamine

In an analogous manner to that described in Example 4, from 7-chloro-3-(4-chloro-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine and NH_3 in MeOH there was obtained 3-(4-chloro-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ylamine as colourless crystals, m.p. >230°.

Example 28

[3-(4-Chloro-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-yl]-dimethyl-amine

In an analogous manner to that described in Example 5, from 7-chloro-3-(4-chloro-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine and dimethyl-amine in EtOH there was obtained [3-(4-chloro-benzenesulphonyl)-5-methyl-2-methyl-sulphanyl-pyrazolo[1,5-a]pyrimidin-7-yl]-dimethyl-amine as colourless crystals, m.p. 221-223°.

Example 29

3-(2,4-Dichloro-benzenesulphonyl)-5-methyl-2-methylsulphanyl-7-piperazin-1-yl-pyrazolo[1,5-a]pyrimidine

(a) In an analogous manner to that described in Example 18 a) to d), starting from (2,4-chloro-benzenesulphonyl)-acetonitrile there was obtained 7-chloro-3-(2,4-dichloro-

benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine as a colourless foam.

(b) In an analogous manner to that described in Example 1c), from 7-chloro-3-(2,4-dichloro-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine and piperazine there was obtained 3-(2,4-dichloro-benzenesulphonyl)-5-methyl-2-methyl-sulphanyl-7-piperazin-1-yl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. >230°.

Example 30

3-(2,4-Chloro-benzenesulphonyl)-5-methyl-7-(4-methyl-piperazin-1-yl)-2-methyl-sulphanyl-pyrazolo[1,5-a]pyrimidine

In an analogous manner to that described in Example 2, from 7-chloro-3-(2,4-dichloro-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine and 1-methyl-piperazine there was obtained 3-(2,4-chloro-benzenesulphonyl)-5-methyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. >230°.

Example 31

3-(2,4-Chloro-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ylamine

In an analogous manner to that described in Example 4, from 7-chloro-3-(2,4-chloro-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine and NH_3 in MeOH there was obtained 3-(2,4-chloro-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ylamine as colourless crystals, m.p. >230°.

Example 32

3-(4-Bromo-benzenesulphonyl)-5-methyl-2-methylsulphanyl-7-piperazin-1-yl-pyrazolo-[1,5-a]pyrimidine

(a) In an analogous manner to that described in Example 18 a) to d), starting from (4-bromo-benzenesulphonyl)-acetonitrile there was obtained 3-(4-bromo-benzene-sulphonyl)-7-chloro-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine as a colourless foam.

(b) In an analogous manner to that described in Example 1c), from 3-(4-bromo-benzenesulphonyl)-7-chloro-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine and

piperazine there was obtained 3-(4-bromo-benzenesulphonyl)-5-methyl-2-methyl-sulphanyl-7-piperazin-1-yl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 212-214°.

Example 33

3-(4-Bromo-benzenesulphonyl)-5-methyl-7-(4-methyl-piperazin-1-yl)-2-methyl-sulphanyl-pyrazolo[1,5-a]pyrimidine

In an analogous manner to that described in Example 2, from 3-(4-bromo-benzene-sulphonyl)-7-chloro-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine and 1-methyl-piperazine there was obtained 3-(4-bromo-benzenesulphonyl)-5-methyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 202-203°.

Example 34

3-(4-Bromo-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ylamine

In an analogous manner to that described in Example 4, from 3-(4-bromo-benzene-sulphonyl)-7-chloro-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine and NH_3 in MeOH there was obtained 3-(4-bromo-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ylamine as colourless crystals, m.p. >230°.

Example 35

3-(4-Methoxy-benzenesulphonyl)-5-methyl-2-methylsulphanyl-7-piperazin-1-yl-pyrazolo[1,5-a]pyrimidine

(a) In an analogous manner to that described in Example 18 a) to d), starting from (4-methoxy-benzenesulphonyl)-acetonitrile there was obtained 7-chloro-3-(4-methoxy-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine as a colourless foam.

(b) In an analogous manner to that described in Example 1c), from 7-chloro-3-(4-methoxy-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine and piperazine there was obtained 3-(4-methoxy-benzenesulphonyl)-5-methyl-2-methyl-sulphanyl-7-piperazin-1-yl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p.176-177°.

Example 36

3-(4-Methoxy-benzenesulphonyl)-5-methyl-7-(4-methyl-piperazin-1-yl)-2-methyl-sulphanyl-pyrazolo[1,5-a]pyrimidine

In an analogous manner to that described in Example 2, from 7-chloro-3-(4-methoxy-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine and 1-methyl-piperazine there was obtained 3-(4-methoxy-benzenesulphonyl)-5-methyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 212-213°.

Example 37

3-(4-Methoxy-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]-pyrimidin-7-ylamine

In an analogous manner to that described in Example 4, from 7-chloro-3-(4-methoxy-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine and NH_3 in MeOH there was obtained 3-(4-methoxy-benzenesulphonyl)-5-methyl-2-methyl-sulphanyl-pyrazolo[1,5-a]pyrimidin-7-ylamine as colourless crystals, m.p. >230°.

Example 38

5-Methyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-3-(naphthalene-2-sulphonyl)-pyrazolo[1,5-a]pyrimidine

(a) In an analogous manner to that described in Example 18 a) to d), starting from (naphthalene-2-sulphonyl)-acetonitrile there was obtained 7-chloro-5-methyl-2-methyl-sulphanyl-3-(naphthalene-2-sulphonyl)-pyrazolo[1,5-a]pyrimidine as a colourless foam.

(b) In an analogous manner to that described in Example 2), from 7-chloro-5-methyl-2-methylsulphanyl-3-(naphthalene-2-sulphonyl)-pyrazolo[1,5-a]pyrimidine and 1-methyl-piperazine there was obtained 5-methyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-3-(naphthalene-2-sulphonyl)-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p.> 230°

Example 39

Dimethyl-[5-methyl-2-methylsulphanyl-3-(naphthalene-2-sulphonyl)-pyrazolo[1,5-a]pyrimidin-7-yl]-amine

In an analogous manner to that described in Example 6, from 7-chloro-5-methyl-2-methylsulphanyl-3-(naphthalene-2-sulphonyl)-pyrazolo[1,5-a]pyrimidine and dimethylamine in EtOH there was obtained dimethyl-[5-methyl-2-methylsulphanyl-3-(naphthalene-2-sulphonyl)-pyrazolo[1,5-a]pyrimidin-7-yl]-amine as colourless crystals, m.p. >230°.

Example 40

5-Methyl-2-methylsulphanyl-3-(naphthalene-2-sulphonyl)-pyrazolo[1,5-a]pyrimidine-7-ylamine

In an analogous manner to that described in Example 4, from 7-chloro-5-methyl-2-methylsulphanyl-3-(naphthalene-2-sulphonyl)-pyrazolo[1,5-a]pyrimidine and NH_3 in MeOH there was obtained 5-methyl-2-methylsulphanyl-3-(naphthalene-2-sulphonyl)-pyrazolo[1,5-a]pyrimidine-7-ylamine as colourless crystals, m.p. >230°.

Example 41

5-Methyl-2-methylsulphanyl-7-piperazin-1-yl-3-(4-trifluoromethoxy-benzenesulphonyl)-pyrazolo[1,5-a]pyrimidine

(a) In an analogous manner to that described in Example 18 a) to d), starting from (4-trifluoromethoxy-benzenesulphonyl)-acetonitrile there was obtained 7-chloro-5-methyl-2-methylsulphanyl-3-(4-trifluoromethoxy-benzenesulphonyl)-pyrazolo[1,5-a]pyrimidine as a colourless foam.

(b) In an analogous manner to that described in Example 1c), from 7-chloro-5-methyl-2-methylsulphanyl-3-(4-trifluoromethoxybenzenesulphonyl)-pyrazolo[1,5a]pyrimidine and piperazine there was obtained 5-methyl-2-methylsulphanyl-7-piperazin-1-yl-3-(4-trifluoromethoxy-benzenesulphonyl)-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 203-204°.

Example 42

5-Methyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-3-(4-trifluoromethoxy-benzenesulphonyl)-pyrazolo[1,5-a]pyrimidine

In an analogous manner to that described in Example 2, from 7-chloro-5-methyl-2-methylsulphanyl-3-(4-trifluoromethoxybenzenesulphonyl)-pyrazolo[1,5a]pyrimidine and 1-methyl-piperazine there was obtained 5-methyl-7-(4-methyl-piperazin-1-yl)-2-methyl-sulphanyl-3-(4-trifluoromethoxy-benzenesulphonyl)-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 213-214°.

Example 43

3-(4-Fluoro-benzenesulphonyl)-5-methyl-2-methylsulphanyl-7-piperazin-1-yl-pyrazolo[1,5-a]pyrimidine

(a) In an analogous manner to that described in Example 18 a) to d), starting from (4-fluoro-benzenesulphonyl)-acetonitrile there was obtained 7-chloro-3-(4-fluoro-benzene-sulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine as a colourless foam.

(b) In an analogous manner to that described in Example 1c), from 7-chloro-3-(4-fluoro-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine and piperazine there was obtained 3-(4-fluoro-benzenesulphonyl)-5-methyl-2-methyl-sulphanyl-7-piperazin-1-yl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 180-181°.

Example 44

3-(4-Fluoro-benzenesulphonyl)-5-methyl-7-(4-methyl-piperazin-1-yl)-2-methyl-sulphanyl-pyrazolo[1,5-a]pyrimidine

In an analogous manner to that described in Example 2, from 7-chloro-3-(4-fluoro-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine and 1-methyl-piperazine there was obtained 3-(4-fluoro-benzenesulphonyl)-5-methyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 197-198°.

Example 45

3-(4-Fluoro-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ylamine

In an analogous manner to that described in Example 4, from 7-chloro-3-(4-fluoro-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine and NH_3 in MeOH there was obtained 3-(4-fluoro-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ylamine as colourless crystals, m.p. >230°.

Example 46

3-(4-Iodo-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine-7-ylamine

(a) In an analogous manner to that described in Example 18 a) to d), starting from (4-iodo-benzenesulphonyl)-acetonitrile there was obtained 7-chloro-3-(4-iodo-benzene-sulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine as a colourless foam.

(b) In an analogous manner to that described in Example 4), from 7-chloro-3-(4-iodo-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine and NH_3 in MeOH there was obtained 3-(4-iodo-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine-7-ylamine as colourless crystals, m.p. >230°.

Example 47

3-Benzenesulphonyl-5,6-dimethyl-2-methylsulphanyl-7-piperazin-1-yl-pyrazolo[1,5-a]pyrimidine

(a) In an analogous manner to that described in Example 1a, b), from 4-benzene-sulphonyl-5-methylsulphanyl-2H-pyrazol-3-ylamine and ethyl 2-methyl-acetoacetate there was obtained 3-benzenesulphonyl-7-chloro-5,6-dimethyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine as a colourless foam.

(b) In an analogous manner to that described in Example 1c), from 3-benzenesulphonyl-7-chloro-5,6-dimethyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine and piperazine there was obtained 3-benzenesulphonyl-5,6-dimethyl-2-methylsulphanyl-7-piperazin-1-yl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 159-160°.

Example 48

3-Benzenesulphonyl-5,6-dimethyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine

In an analogous manner to that described in Example 1c), from 3-benzenesulphonyl-7-chloro-5,6-dimethyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine and 1-methyl-piperazine there was obtained 3-benzenesulphonyl-5,6-dimethyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 183-185°.

Example 49

3-Benzenesulphonyl-2-methylsulphanyl-7-piperazin-1-yl-5-propyl-pyrazolo[1,5-a]-pyrimidine

(a) In an analogous manner to that described in Example 1a, b), from 4-benzene-sulphonyl-5-methylsulphanyl-2H-pyrazol-3-ylamine and ethyl butyrylacetate there was obtained 3-benzenesulphonyl-7-chloro-2-methylsulphanyl-5-propyl-pyrazolo[1,5-a]pyrimidine as a colourless foam.

(b) In an analogous manner to that described in Example 1c), from 3-benzene-sulphonyl-7-chloro-2-methylsulphanyl-5-propyl-pyrazolo[1,5-a]pyrimidine and piperazine there was obtained 3-benzenesulphonyl-2-methylsulphanyl-7-piperazin-1-yl-5-propyl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 197-199°.

Example 50

3-Benzenesulphonyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-5-propyl-pyrazolo[1,5-a]pyrimidine

In an analogous manner to that described in Example 2, from 3-benzenesulphonyl-7-chloro-2-methylsulphanyl-5-propyl-pyrazolo[1,5-a]pyrimidine and 1-methyl-piperazine there was obtained 3-benzenesulphonyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-5-propyl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 207-209°.

Example 51

3-Benzenesulphonyl-5-cyclopropyl-2-methylsulphanyl-7-piperazin-1-yl-pyrazolo[1,5-a]pyrimidine

a) In an analogous manner to that described in Example 1a, b), from 4-benzenesulphonyl-5-methylsulphanyl-2H-pyrazol-3-ylamine and ethyl 3-cyclopropyl-3-oxo-propionate there was obtained 3-benzenesulphonyl-7-chloro-5-cyclopropyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine as a colourless foam.

b) In an analogous manner to that described in Example 1c), from 3-benzenesulphonyl-7-chloro-5-cyclopropyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine and piperazine there was obtained 3-benzenesulphonyl-5-cyclopropyl-2-methylsulphanyl-7-piperazin-1-yl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 214-215°.

Example 52

3-Benzenesulphonyl-5-cyclopropyl-2-methylsulphanyl-7(4-methylpiperazin-1-yl)-pyrazolo[1,5-a]pyrimidine

In an analogous manner to that described in Example 2, from 3-benzenesulphonyl-7-chloro-5-cyclopropyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine and 1-methyl-piperazine there was obtained 3-benzenesulphonyl-5-cyclopropyl-2-methylsulphanyl-7(4-methylpiperazin-1-yl)-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 162-164°.

Example 53

3-Benzenesulphonyl-5-cyclopropyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine-7-ylamine

In an analogous manner to that described in Example 4, from 3-benzenesulphonyl-7-chloro-5-cyclopropyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine and NH_3 in MeOH there was obtained 3-benzenesulphonyl-5-cyclopropyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine-7-ylamine as colourless crystals, m.p. > 230°.

Example 54

3-Benzenesulphonyl-2-methylsulphanyl-8-piperazin-1-yl-6,7-dihydro-5H-cyclopenta-[d]pyrazolo[1,5-a]pyrimidine

a) In an analogous manner to that described in Example 1a, b), from 4-benzenesulphonyl-5-methylsulphanyl-2H-pyrazol-3-ylamine and ethyl cyclopentanone-2-carboxylate there was obtained 3-benzenesulphonyl-8-chloro-2-methylsulphanyl-6,7-dihydro-5H-cyclopenta[d]pyrazolo[1,5-a]pyrimidine as a colourless foam.

b) In an analogous manner to that described in Example 1c), from 3-benzenesulphonyl-8-chloro-2-methylsulphanyl-6,7-dihydro-5H-cyclopenta[d]pyrazolo[1,5-a]pyrimidine and piperazine there was obtained 3-benzenesulphonyl-2-methylsulphanyl-8-piperazin-1-yl-6,7-dihydro-5H-cyclopenta[d]pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 221-222.5°.

Example 55

3-Benzenesulphonyl-8-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-6,7-dihydro-5H-cyclopenta[d]pyrazolo[1,5-a]pyrimidine

In an analogous manner to that described in Example 2, from 3-benzenesulphonyl-8-chloro-2-methylsulphanyl-6,7-dihydro-5H-cyclopenta[d]pyrazolo[1,5-a]pyrimidine and 1-methyl-piperazine there was obtained 3-benzenesulphonyl-8-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-6,7-dihydro-5H-cyclopenta[d]pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 228-229.5°.

Example 56

3-Benzenesulphonyl-2-methylsulphanyl-6,7-dihydro-5H-cyclopenta[d]pyrazolo[1,5-a]pyrimidin-8-ylamine

In an analogous manner to that described in Example 4, from 3-benzenesulphonyl-8-chloro-2-methylsulphanyl-6,7-dihydro-5H-cyclopenta[d]pyrazolo[1,5-a]pyrimidine and NH_3 in MeOH there was obtained 3-benzenesulphonyl-2-methylsulphanyl-6,7-dihydro-5H-cyclopenta[d]pyrazolo[1,5-a]pyrimidin-8-ylamine as colourless crystals, m.p. >230°.

Example 57

3-Benzenesulphonyl-2-methylsulphanyl-9-piperazin-1-yl-5,6,7,8-tetrahydro-pyrazolo[5,1-b]quinazoline

a) In an analogous manner to that described in Example 1a, b), from 4-benzenesulphonyl-5-methylsulphanyl-2H-pyrazol-3-ylamine and ethyl cyclohexanone-2-carboxylate there was obtained 3-benzenesulphonyl-9-chloro-2-methylsulphanyl-5,6,7,8-tetrahydro-pyrazolo[5,1-b]quinazoline as a colourless foam.

b) In an analogous manner to that described in Example 1c), from 3-benzenesulphonyl-9-chloro-2-methylsulphanyl-5,6,7,8-tetrahydro-pyrazolo[5,1-b]quinazoline and piperazine there was obtained 3-benzenesulphonyl-2-methylsulphanyl-9-piperazin-1-yl-5,6,7,8-tetrahydro-pyrazolo[5,1-b]quinazoline as colourless crystals, m.p. 121-123°.

Example 58

3-Benzenesulphonyl-9-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-5,6,7,8-tetrahydro-pyrazolo[5,1-b]quinazoline

In an analogous manner to that described in Example 2, from 3-benzenesulphonyl-9-chloro-2-methylsulphanyl-5,6,7,8-tetrahydro-pyrazolo[5,1-b]quinazoline and 1-methyl-piperazine there was obtained 3-benzenesulphonyl-9-(4-methyl-piperazin-1-yl)-2-methyl-sulphanyl-5,6,7,8-tetrahydro-pyrazolo[5,1-b]quinazoline as colourless crystals, m.p. 198-200°.

Example 59

3-Benzenesulphonyl-2-methylsulphanyl-8-piperazin-1-yl-5H,7H-pyrazolo[1,5-a]-thieno[3,4-d]pyrimidine

a) In an analogous manner to that described in Example 1a, b), from 4-benzenesulphonyl-5-methylsulphanyl-2H-pyrazol-3-ylamine and methyl 4-oxo-tetrahydro-thiophene-3-carboxylate there was obtained 3-benzenesulphonyl-8-chloro-2-methylsulphanyl-5H,7H-pyrazolo[1,5-a]thieno[3,4-d]pyrimidine as a colourless foam.

b) In an analogous manner to that described in Example 1c), from 3-benzenesulphonyl-8-chloro-2-methylsulphanyl-5H,7H-pyrazolo[1,5-a]thieno[3,4-d]pyrimidine and piperazine there was obtained 3-benzenesulphonyl-2-methylsulphanyl-8-piperazin-1-yl-5H,7H-pyrazolo[1,5-a]thieno[3,4-d]pyrimidine as colourless crystals, m.p. >230°.

Example 60

3-Benzenesulphonyl-8-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-5H,7H-pyrazolo[1,5-a]thieno[3,4-d]pyrimidine

In an analogous manner to that described in Example 2, from 3-benzenesulphonyl-8-chloro-2-methylsulphanyl-5H,7H-pyrazolo[1,5-a]thieno[3,4-d]pyrimidine and 1-methyl-piperazine there was obtained 3-benzenesulphonyl-8-(4-methyl-piperazin-1-yl)-2-methyl-sulphanyl-5H,7H-pyrazolo[1,5-a]thieno[3,4-d]pyrimidine as colourless crystals, m.p. >230°.

Example 61

5-Methyl-2-methylsulphanyl-7-piperazin-1-yl-3-(thiophene-2-sulphonyl)-pyrazolo[1,5-a]pyrimidine

a) In an analogous manner to that described in Example 18 a) to d), starting from thien-2-ylsulphonylacetonitrile there was obtained 7-chloro-5-methyl-2-methylsulphanyl-3-(thiophene-2-sulphonyl)-pyrazolo[1,5-a]pyrimidine as a colourless solid.

b) In an analogous manner to that described in Example 1c), from 7-chloro-5-methyl-2-methylsulphanyl-3-(thiophene-2-sulphonyl)-pyrazolo[1,5-a]pyrimidine and piperazine there was obtained 5-methyl-2-methylsulphanyl-7-piperazin-1-yl-3-(thiophen-2-sulphonyl)-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 229-230°.

Example 62

5-Methyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-3-(thiophene-2-sulphonyl)-pyrazolo[1,5-a]pyrimidine

In an analogous manner to that described in Example 2, from 7-chloro-5-methyl-2-methylsulphanyl-3-(thiophene-2-sulphonyl)-pyrazolo[1,5-a]pyrimidine and 1-methyl-piperazine there was obtained 5-methyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-3-(thiophene-2-sulphonyl)-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. >230°.

Example 63

5-Methyl-2-methylsulphanyl-3-(thiophene-2-sulphonyl)-pyrazolo[1,5-a]pyrimidin-7-ylamine

In an analogous manner to that described in Example 4, from 7-chloro-5-methyl-2-methylsulphanyl-3-(thiophene-2-sulphonyl)-pyrazolo[1,5-a]pyrimidine and NH_3 in MeOH there was obtained 5-methyl-2-methylsulphanyl-3-(thiophene-2-sulphonyl)-pyrazolo[1,5-a]pyrimidin-7-ylamine as colourless crystals, m.p. >230°.

Example 64

3-Benzenesulphonyl-5-isopropyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine

a) In an analogous manner to that described in Example 1a, b), from 4-benzenesulphonyl-5-methylsulphanyl-2H-pyrazol-3-ylamine and ethyl isobutyrylacetate there was obtained 3-benzenesulphonyl-7-chloro-5-isopropyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine as a colourless foam.

b) In an analogous manner to that described in Example 2), from 3-benzenesulphonyl-7-chloro-5-isopropyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine and 1-methyl-piperazine there was obtained 3-benzenesulphonyl-5-isopropyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 212-214°.

Example 65

3-Benzenesulphonyl-5-isopropyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ylamine

In an analogous manner to that described in Example 4, from 3-benzenesulphonyl-7-chloro-5-isopropyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine and NH_3 in MeOH there was obtained 3-benzenesulphonyl-5-isopropyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ylamine as colourless crystals, m.p. 210-211°.

Example 66

3-Benzenesulphonyl-5-tert-butyl-2-methylsulphanyl-7-piperazin-1-yl-pyrazolo[1,5-a]pyrimidine

a) 2 g (7.4 mmol) of 4-benzenesulphonyl-5-methylsulphanyl-2H-pyrazol-3-ylamine and 2.91 ml (18.3 mmol) of ethyl pivaloylacetate were added to 27 g of polyphosphoric acid and heated to 120° for 5 hrs. After cooling 100 ml of water were slowly added thereto and the mixture was extracted three times with CH_2Cl_2. The organic phase was dried ($MgSO_4$), filtered and evaporated. Chromatography (SiO_2, CH_2Cl_2/EtOAc) yielded 1.34 g (48%) of 3-benzenesulphonyl-5-tert-butyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ol as a colourless foam.

b) A suspension of 1.34 g (3.5 mmol) of 3-benzenesulphonyl-5-tert-butyl-2-methyl-sulphanyl-pyrazolo[1,5-a]pyrimidin-7-ol in 20 ml of $POCl_3$ was heated at reflux for 30 min. The reaction solution was cooled to RT and evaporated. The residue was treated with 100 ml of ice-water and the pH value of the solution was adjusted to 8 with sat. $NaHCO_3$ solution. The aqueous phase was extracted three times with CH_2Cl_2, and the organic phases were dried ($MgSO_4$), filtered and evaporated. Chromatography (silica gel, CH_2Cl_2/AcOEt 19:1) of the residue yielded 1.23 g (78%) g of 3-benzenesulphonyl-5-tert-butyl-7-chloro-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine as a pale yellow foam.

c) 0.67 g (7.7 mmol) of piperazine in 10 ml of DMF was added to a solution of 1.23 g (3.1 mmol) of 3-benzenesulphonyl-5-tert-butyl-7-chloro-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine in 10 ml of DMF and stirred at RT for 2 hrs. The reaction solution was evaporated and the residue was partitioned between 2N NaOH and CH_2Cl_2. The aqueous phase was extracted three times with CH_2Cl_2, and the combined organic phases were dried ($MgSO_4$), filtered and evaporated. Subsequent chromatography (SiO_2, CH_2Cl_2/MeOH 19:1) and crystallization from EtOH yielded 0.25 g (18%) of 3-benzenesulphonyl-5-tert-butyl-2-methylsulphanyl-7-piperazin-1-yl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 236-237°.

Example 67

3-Benzenesulphonyl-2-methylsulphanyl-7-piperazin-1-yl-5-trifluoromethyl-pyrazolo[1,5-a]pyrimidine

a) In an analogous manner to that described in Example 66 a), b) from 4-benzene-sulphonyl-5-methylsulphanyl-2H-pyrazol-3-ylamine and ethyl 4,4,4-trifluoroacetate there was obtained 3-benzenesulphonyl-2-methylsulphanyl-5-trifluoromethyl-pyrazolo[1,5-a]-pyrimidin-7-ol as a colourless foam.

b) In an analogous manner to that described in Example 1c), from 3-benzenesulphonyl-2-methylsulphanyl-5-trifluoromethyl-pyrazolo[1,5-a]pyrimidin-7-ol and piperazine there was obtained 3-benzenesulphonyl-2-methylsulphanyl-7-piperazin-1-yl-5-trifluoromethyl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. >230°.

Example 68

3-Benzenesulphonyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-5-trifluoromethyl-pyrazolo[1,5-a]pyrimidine

In an analogous manner to that described in Example 2), from 3-benzenesulphonyl-2-methylsulphanyl-5-trifluoromethyl-pyrazolo[1,5-a]pyrimidin-7-ol and 1-methyl-piperazine there was obtained 3-benzenesulphonyl-7-(4-methyl-piperazin-1-yl)-2-methyl-sulphanyl-5-trifluoromethyl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. >230°.

Example 69

3-Benzenesulphonyl-2-ethyl-5-methyl-7-piperazin-1-yl-pyrazolo[1,5-a]pyrimidine

a) 0.2 ml of glacial acetic acid was added to a suspension of 5.0 g (27.6 mmol) of phenylsulphonylacetonitrile in 17.6 ml (88.6 mmol) of triethyl orthopropionate and subsequently heated to 140°. The EtOH formed was distilled off continuously. After 1.5 hr. the mixture was cooled to RT and evaporated to dryness in a high vacuum. There were thus obtained 7.3 g (100%) of a mixture of (E)- and (Z)-2-benzenesulphonyl-3-ethoxy-pent-2-enenitrile as a colourless oil.

b) A solution of 5.2 g (19.6 mmol) of (E)- and (Z)-2-benzenesulphonyl-3-ethoxy-pent-2-enenitrile and 1.24 ml (25.5 mol) of NH_2NH_2 in 50 ml of EtOH was heated at reflux for

1 hr. The brown reaction solution was cooled to RT, evaporated and chromatographed (SiO_2, CH_2Cl_2/MeOH 19:1). There were thus obtained 2.9 g (59%) of 4-benzene-sulphonyl-5-ethyl-2H-pyrazol-3-ylamine as a beige oil.

c) A solution of 2.9 g (11.5 mmol) of 4-benzenesulphonyl-5-ethyl-2H-pyrazol-3-ylamine and 1.8 ml (13.8 mmol) of ethyl acetoacetate in 10 ml of acetic acid was heated at reflux for 3 hrs. The reaction solution was cooled to RT and evaporated. The residue was partitioned between CH_2Cl_2 and H_2O and the aqueous phase was washed three times with 150 ml of CH_2Cl_2. The combined organic phases were dried ($MgSO_4$), filtered and evaporated. Crystallization from ethyl acetate yielded 2.6 g (71%) of 3-benzenesulphonyl-2-ethyl-5-methyl-pyrazolo[1,5-a]pyrimidin-7-ol as colourless crystals.

d) A suspension of 2.0 g (6.3 mmol) of 3-benzenesulphonyl-2-ethyl-5-methyl-pyrazolo-[1,5-a]pyrimidin-7-ol in 30 ml of $POCl_3$ was heated at reflux for 45 min. The reaction solution was cooled to RT and evaporated. The residue was treated with 100 ml of ice-water and the pH value of the solution was adjusted to 8 with sat. $NaHCO_3$ solution. The aqueous phase was extracted three times with 100 ml of CH_2Cl_2, and the organic phases were dried ($MgSO_4$), filtered and evaporated. Chromatography (SiO_2, CH_2Cl_2/MeOH 49:1) of the residue yielded 2.0 g (94%) of 3-benzenesulphonyl-7-chloro-2-ethyl-5-methyl-pyrazolo[1,5-a]pyrimidine as colourless crystals.

e) 0.64 g (7.4 mmol) of piperazine in 10 ml of DMF was added to a solution of 1.0 g (3 mmol) of 3-benzenesulphonyl-7-chloro-2-ethyl-5-methyl-pyrazolo[1,5-a]pyrimidine in 10 ml of DMF and stirred at 60° for 2 hrs. The DMF was evaporated in a high vacuum, the residue was partitioned between 2N NaOH and CH_2Cl_2. The aqueous phase was extracted three times with 50 ml of CH_2Cl_2 and the combined organic were dried ($MgSO_4$), filtered and evaporated. Subsequent chromatography (SiO_2, CH_2Cl_2/MeOH 9:1) and crystallization from EtOH yielded 0.32 g (27%) of 3-benzenesulphonyl-2-ethyl-5-methyl-7-piperazin-1-yl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 150-150.8°.

Example 70

3-Benzenesulphonyl-2-ethyl-5-methyl-7-(4-methyl-piperazin-1-yl)-pyrazolo[1,5-a]-pyrimidine

In an analogous manner to that described in Example 2, from 3-benzenesulphonyl-7-chloro-2-ethyl-5-methyl-pyrazolo[1,5-a]pyrimidine and 1-methyl-piperazine there was

obtained 3-benzenesulphonyl-2-ethyl-5-methyl-7-(4-methyl-piperazin-1-yl)-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 171-172°.

Example 71

N-(3-Benzenesulphonyl-2-ethyl-5-methyl-pyrazolo[1,5-a]pyrimidin-7-yl)-N',N'-dimethyl-ethane-1,2-diamine

In an analogous manner to that described in Example 11) from 3-benzenesulphonyl-7-chloro-2-ethyl-5-methyl-pyrazolo[1,5-a]pyrimidine and 2-dimethylaminoethylamine there was obtained N-(3-benzenesulphonyl-2-ethyl-5-methyl-pyrazolo[1,5-a]pyrimidin-7-yl)-N',N'-dimethyl-ethane-1,2-diamin

Example 72

3-(4-Bromo-benzenesulphonyl)-2-ethyl-5-methyl-7-piperazin-1-yl-pyrazolo[1,5-a]-pyrimidine

a) In an analogous manner to that described in Example 64 a) to d), from (4-bromo-benzenesulphonyl)-acetonitrile there was obtained 3-(4-bromo-benzenesulphonyl)-7-chloro-2-ethyl-5-methyl-pyrazolo[1,5-a]pyrimidine as colourless crystals.

b) In an analogous manner to that described in Example 1c), from 3-(4-bromo-benzenesulphonyl)-7-chloro-2-ethyl-5-methyl-pyrazolo[1,5-a]pyrimidine and piperazine there was obtained 3-(4-bromo-benzenesulphonyl)-2-ethyl-5-methyl-7-piperazin-1-yl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 196-197°.

Example 73

3-(4-Bromo-benzenesulphonyl)-2-ethyl-5-methyl-7-piperazin-1-yl-pyrazolo-[1,5-a]pyrimidine

In an analogous manner to that described in Example 2, from 3-(4-bromo-benzene-sulphonyl)-7-chloro-2-ethyl-5-methyl-pyrazolo[1,5-a]pyrimidine and 1-methyl-piperazine there was obtained 3-(4-bromo-benzenesulphonyl)-2-ethyl-5-methyl-7-piperazin-1-yl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 215-216°.

Example 74

N-[3-(4-Bromo-benzenesulphonyl)-2-ethyl-5-methyl-pyrazolo[1,5-a]pyrimidin-7-yl]-N',N'-dimethyl-ethane-1,2-diamine

In an analogous manner to that described in Example 11, from 3-(4-bromo-benzenesulphonyl)-7-chloro-2-ethyl-5-methyl-pyrazolo[1,5-a]pyrimidine and 2-dimethylamino-ethylamine there was obtained N-[3-(4-bromo-benzenesulphonyl)-2-ethyl-5-methyl-pyrazolo[1,5-a]pyrimidin-7-yl]-N',N'-dimethyl-ethane-1,2-diamine as colourless crystals, m.p. 210-211°.

Example 75

2-Ethyl-3-(4-methoxy-benzenesulphonyl)-5-methyl-7-piperazin-1-yl-pyrazolo[1,5-a]-pyrimidine

a) In an analogous manner to that described in Example 69 a) to d), from (4-methoxy-benzenesulphonyl)-acetonitrile there was obtained 7-chloro-2-ethyl-3-(4-methoxy-benzenesulphonyl)-5-methyl-pyrazolo[1,5-a]pyrimidine as colourless crystals.

b) In an analogous manner to that described in Example 1c), from 7-chloro-2-ethyl-3-(4-methoxy-benzenesulphonyl)-5-methyl-pyrazolo[1,5-a]pyrimidine and piperazine there was obtained 2-ethyl-3-(4-methoxy-benzenesulphonyl)-5-methyl-7-piperazin-1-yl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 196-197°.

Example 76

2-Ethyl-3-(4-methoxy-benzenesulphonyl)-5-methyl-7-(4-methyl-piperazin-1-yl)-pyrazolo[1,5-a]pyrimidine

In an analogous manner to that described in Example 2, from 7-chloro-2-ethyl-3-(4-methoxy-benzenesulphonyl)-5-methyl-pyrazolo[1,5-a]pyrimidine and 1-methyl-piperazine there was obtained 2-ethyl-3-(4-methoxy-benzenesulphonyl)-5-methyl-7-(4-methyl-piperazin-1-yl)-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 178-188°.

Example 77

2-Ethyl-3-(4-methoxy-benzenesulphonyl)-5-methyl-pyrazolo[1,5-a]pyrimidin-7-ylamine

In an analogous manner to that described in Example 4, from 7-chloro-2-ethyl-3-(4-methoxy-benzenesulphonyl)-5-methyl-pyrazolo[1,5-a]pyrimidine and NH_3 in MeOH there was obtained 2-ethyl-3-(4-methoxy-benzenesulphonyl)-5-methyl-pyrazolo[1,5-a]pyrimidin-7-ylamine as colourless crystals, m.p. 186-188°.

Example 78

(3R,5S)-7-(3,5-Dimethyl-piperazin-1-yl)-2-ethyl-3-(4-methoxy-benzenesulphonyl)-5-methyl-pyrazolo[1,5-a]pyrimidine

In an analogous manner to that described in Example12, from 7-chloro-2-ethyl-3-(4-methoxy-benzenesulphonyl)-5-methyl-pyrazolo[1,5-a]pyrimidine and cis-2,6-dimethyl-piperazine there was obtained (3R,5S)-7-(3,5-dimethyl-piperazin-1-yl)-2-ethyl-3-(4-methoxy-benzenesulphonyl)-5-methyl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 151-152°.

Example 79

3-Benzenesulphonyl-2-ethyl-8-piperazin-1-yl-6,7-dihydro-5H-cyclopenta[d]pyrazolo[1,5-a]pyrimidine

a) In an analogous manner to that described in Example 69 c) d), from 4-benzene-sulphonyl-5-ethyl-2H-pyrazol-3-ylamine and ethyl cyclopentanone-2-carboxylate there was obtained 3-benzenesulphonyl-8-chloro-2-ethyl-6,7-dihydro-5H-cyclopenta[d]-pyrazolo[1,5-a]pyrimidine as a colourless foam.

b) In an analogous manner to that described in Example 1c), from 3-benzenesulphonyl-8-chloro-2-ethyl-6,7-dihydro-5H-cyclopenta[d]pyrazolo[1,5-a]pyrimidine and piperazine there was obtained 3-benzenesulphonyl-2-ethyl-8-piperazin-1-yl-6,7-dihydro-5H-cyclopenta[d]pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 200-201°.

Example 80

3-Benzenesulphonyl-2-ethyl-8-(4-methyl-piperazin-1-yl)-6,7-dihydro-5H-cyclopenta[d]-pyrazolo[1,5-a]pyrimidine

In an analogous manner to that described in Example 2, from 3-benzenesulphonyl-8-chloro-2-ethyl-6,7-dihydro-5H-cyclopenta[d]pyrazolo[1,5-a]pyrimidine and 1-methyl-piperazine there was obtained 3-benzenesulphonyl-2-ethyl-8-(4-methyl-piperazin-1-yl)-6,7-dihydro-5H-cyclopenta[d]pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 235-236°.

Example 81

3-Benzenesulphonyl-2-ethyl-6,7-dihydro-5H-cyclopenta[d]pyrazolo[1,5-a]pyrimidin-8-ylamine

In an analogous manner to that described in Example 4, from 3-benzenesulphonyl-8-chloro-2-ethyl-6,7-dihydro-5H-cyclopenta[d]pyrazolo[1,5-a]pyrimidine and NH3 in MeOH there was obtained 3-benzenesulphonyl-2-ethyl-6,7-dihydro-5H-cyclopenta[d]-pyrazolo[1,5-a]pyrimidin-8-ylamine as colourless crystals, m.p. >230°.

Example 82

(3R,5S)-3-Benzenesulphonyl-8-(3,5-dimethyl-piperazin-1-yl)-2-ethyl-6,7-dihydro-5H-cyclopenta[d]pyrazolo[1,5-a]pyrimidine

In an analogous manner to that described in Example 12, from 3-benzenesulphonyl-8-chloro-2-ethyl-6,7-dihydro-5H-cyclopenta[d]pyrazolo[1,5-a]pyrimidine and cis-2,6-dimethylpiperazine there was obtained (3R,5S)-3-benzenesulphonyl-8-(3,5-dimethyl-piperazin-1-yl)-2-ethyl-6,7-dihydro-5H-cyclopenta[d]pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 220-221°.

Example 83

3-Benzenesulphonyl-5-methyl-7-piperazin-1-yl-pyrazolo[1,5-a]pyrimidine

a) 6.88 ml of N,N-dimethylformamide dimethyl acetal were added to a suspension of 7.0 g (38.6 mmol) of phenylsulphonylacetonitrile in 30 ml of hexane while cooling with ice and

subsequently stirred at RT for 12 hrs. The separated crystals were filtered off and therewere thus obtained 9.08 g (99%) of 2-benzenesulphonyl-3-dimethylamino-acrylonitrile as beige crystals, m.p. 108-110°.

b) 2.05 ml (40.9 mmol) of NH_2NH_2 were added to a solution of 9.08 g (38.3 mmol) of 2-benzenesulphonyl-3-dimethylamino-acrylonitrile in 60 ml of EtOH and stirred at 40° for 5 hrs. The reaction solution was evaporated and the residue was partitioned between H_2O and CH_2Cl_2. The aqueous phase was washed three times with CH_2Cl_2, and the combined organic phases were dried ($MgSO_4$), filtered and evaporated. Chromatography (SiO_2, CH_2Cl_2/MeOH 10:1) yielded 2.5 g (30%) of 4-benzenesulphonyl-1H-pyrazol-3-ylamine as a beige powder, m.p. 159-161°.

c) A solution of 1.0 g (4.47 mmol) of 4-benzenesulphonyl-1H-pyrazol-3-ylamine and 0.6 ml (5.37 mmol) of ethyl acetoacetate in 8 ml of acetic acid was heated at reflux for 1.5 hrs. The reaction solution was cooled to RT and evaporated. The residue was partitioned between CH_2Cl_2 and H_2O and the aqueous phase was washed three times with 150 ml of CH_2Cl_2. The combined organic phases were dried ($MgSO_4$), filtered and evaporated. Chromatography (SiO_2, CH_2Cl_2/MeOH 20:1) yielded 0.91 g (70%) of 3-benzenesulphonyl-5-methyl-pyrazolo[1,5-a]pyrimidine as beige crystals.

d) A suspension of 0.91 g (3.14 mmol) of 3-benzenesulphonyl-5-methyl-pyrazolo[1,5-a]pyrimidine in 15 ml of $POCl_3$ was heated at reflux for 1 hr. The reaction solution was cooled to RT and evaporated. The residue was treated with 30 ml of ice-water and the pH value of the solution was adjusted to 8 with sat. $NaHCO_3$ solution. The aqueous phase was extracted three times with 20 ml of CH_2Cl_2, and the organic phases were dried ($MgSO_4$), filtered and evaporated. Chromatography (SiO_2, CH_2Cl_2/AcOEt 19:1) of the residue yielded 0.84 g (87%) of 3-benzenesulphonyl-7-chloro-5-methyl-pyrazolo[1,5-a]pyrimidine as a beige solid.

e) In an analogous manner to that described in Example 1c), from 3-benzenesulphonyl-7-chloro-5-methyl-pyrazolo[1,5-a]pyrimidine and piperazine there was obtained 3-benzenesulphonyl-5-methyl-7-piperazin-1-yl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 180-181°.

Example 84

3-Benzenesulphonyl-5-methyl-7-(4-methyl-piperazin-1-yl)-pyrazolo[1,5-a]pyrimidine

In an analogous manner to that described in Example 2, from 3-benzenesulphonyl-7-chloro-5-methyl-pyrazolo[1,5-a]pyrimidine and 1-methyl-piperazine there was obtained 3-benzenesulphonyl-5-methyl-7-(4-methyl-piperazin-1-yl)-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. >230.

Example 85

5-Methyl-2-methylsulphanyl-7-piperazin-1-yl-3-(toluene-2-sulphonyl)-pyrazolo-[1,5-a]pyrimidine

a) In an analogous manner to that described in Example 18 a) to d), starting from (toluene-2-sulphonyl)-acetonitrile there was obtained 7-chloro-5-methyl-2-methylsulphanyl-3-(toluene-2-sulphonyl)-pyrazolo[1,5-a]pyrimidine as a colourless foam.

b) In an analogous manner to that described in Example 1c), from 7-chloro-5-methyl-2-methylsulphanyl-3-(toluene-2-sulphonyl)-pyrazolo[1,5-a]pyrimidine and piperazine there was obtained 5-methyl-2-methylsulphanyl-7-piperazin-1-yl-3-(toluene-2-sulphonyl)-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 215-215.5°

The (toluene-2-sulphonyl)-acetonitrile used was prepared as follows:

2.2 ml (34.5 mmol) of chloroacetonitrile were added to a suspension of 4.5 g (28.8 mmol) of toluene-2-sulphinic acid sodium salt in 100 ml of DMF and stirred at 100° for 1 hr. The reaction solution was evaporated and the residue was partitioned between H_2O and CH_2Cl_2. The aqueous phase was washed three times with CH_2Cl_2. The combined organic phases were washed once with H_2O, dried ($MgSO_4$). filtered and evaporated. Chromatography (SiO_2, CH_2Cl_2) yielded 2.9 g (50%) of (toluene-2-sulphonyl)-acetonitrile as a colourless oil.

Example 86

5-Methyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-3-(toluene-2-sulphonyl)-pyrazolo[1,5-a]pyrimidine

In an analogous manner to that described in Example 2), from 7-chloro-5-methyl-2-methylsulphanyl-3-(toluene-2-sulphonyl)-pyrazolo[1,5-a]pyrimidine and 1-methyl-piperazine there was obtained 5-methyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-3-(toluene-2-sulphonyl)-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 199-200°.

Example 87

5-Methyl-2-methylsulphanyl-3-(toluene-2-sulphonyl)-pyrazolo[1,5-a]pyrimidin-7-ylamine

In an analogous manner to that described in Example 4), from 7-chloro-5-methyl-2-methylsulphanyl-3-(toluene-2-sulphonyl)-pyrazolo[1,5-a]pyrimidine and NH_3 in MeOH there was obtained 5-methyl-2-methylsulphanyl-3-(toluene-2-sulphonyl)-pyrazolo[1,5-a]-pyrimidin-7-ylamine as colourless crystals, m.p. > 250°.

Example 88

5-Methyl-2-methylsulphanyl-7-piperazin-1-yl-3-(toluene-2-sulphonyl)-pyrazolo[1,5-a]-pyrimidine

a) In an analogous manner to that described in Example 18 a) to d), starting from (toluene-3-sulphonyl)-acetonitrile there was obtained 7-chloro-5-methyl-2-methyl-sulphanyl-3-(toluene-3-sulphonyl)-pyrazolo[1,5-a]pyrimidine as a colourless foam.

b) In an analogous manner to that described in Example 1c), from 7-chloro-5-methyl-2-methylsulphanyl-3-(toluene-2-sulphonyl)-pyrazolo[1,5-a]pyrimidine and piperazine there was obtained 5-methyl-2-methylsulphanyl-7-piperazin-1-yl-3-(toluene-2-sulphonyl)-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 165-165.5°.

The (toluene-3-sulphonyl)-acetonitrile used was prepared as follows:

2.65 ml (42 mmol) of chloroacetonitrile were added to a suspension of 7.5 g (42 mmol) of toluene-3-sulphinic acid sodium salt in 80 ml of DMF and stirred at 100° for 1 hr. The reaction solution was evaporated, the residue was partitioned between H_2O and CH_2Cl_2.

The aqueous phase was washed three times with CH_2Cl_2. The combined organic phases were washed once with H_2O, dried ($MgSO_4$). filtered and evaporated. Chromatography (SiO_2, CH_2Cl_2) yielded 2.06 g (25%) of (toluene-3-sulphonyl)-acetonitrile as a colourless oil.

5

Example 89

5-Methyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-3-(toluene-3-sulphonyl)-pyrazolo[1,5-a]pyrimidine

In an analogous manner to that described in Example 2), from 7-chloro-5-methyl-2-
10 methylsulphanyl-3-(toluene-3-sulphonyl)-pyrazolo[1,5-a]pyrimidine and 1-methyl-piperazine there was obtained 5-methyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-3-(toluene-3-sulphonyl)-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 183-184°.

Example 90

15 5-Methyl-2-methylsulphanyl-3-(toluene-3-sulphonyl)-pyrazolo[1,5-a]pyrimidin-7-ylamine

In an analogous manner to that described in Example 4), from 7-chloro-5-methyl-2-methylsulphanyl-3-(toluene-3-sulphonyl)-pyrazolo[1,5-a]pyrimidine and NH_3 in MeOH there was obtained 5-methyl-2-methylsulphanyl-3-(toluene-3-sulphonyl)-pyrazolo[1,5-a]-
20 pyrimidine-7-ylamine as colourless crystals, m.p. >250°.

Example 91

5-Methyl-2-methylsulphanyl-7-piperazin-1-yl-3-(pyridine-3-sulphonyl)-pyrazolo[1,5-a]pyrimidine

25 a) In an analogous manner to that described in Example 18 a) to d), starting from (pyridine-3-sulphonyl)-acetonitrile there was obtained 7-chloro-5-methyl-2-methyl-sulphanyl-3-(pyridine-3-sulphonyl)-pyrazolo[1,5-a]pyrimidine as a colourless foam.

b) In an analogous manner to that described in Example 1c), from 7-chloro-5-methyl-2-methylsulphanyl-3-(pyridine-3-sulphonyl)-pyrazolo[1,5-a]pyrimidine and piperazine
30 there was obtained 5-methyl-2-methylsulphanyl-7-piperazin-1-yl-3-(pyridine-3-sulphonyl)-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 222-223°.

The (pyridine-3-sulphonyl)-acetonitrile used was prepared as follows:

2.1 ml (33.4 mmol) of chloroacetonitrile were added to a solution of 4.6 g (42 mmol) of pyridine-3-sulphinic acid sodium salt in 50 ml of DMF and stirred at 90° for 1 hr. The reaction solution was evaporated, the residue was partitioned between H_2O and CH_2Cl_2. The aqueous phase was washed three times with CH_2Cl_2. The combined organic phases were washed once with H_2O, dried ($MgSO_4$) filtered and evaporated. Chromatography (SiO_2, AcOEt/hexane 2:1) yielded 4.1 g (80%) of (pyridine-3-sulphonyl)-acetonitrile as a beige solid.

Example 92

5-Methyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-3-(pyridine-3-sulphonyl)-pyrazolo[1,5-a]pyrimidine

In an analogous manner to that described in Example 2), from 7-chloro-5-methyl-2-methylsulphanyl-3-(pyridine-3-sulphonyl)-pyrazolo[1,5-a]pyrimidine and 1-methyl-piperazine there was obtained 5-methyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-3-(pyridine-3-sulphonyl)-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 188.4-189°.

Example 93

5-Methyl-2-methylsulphanyl-3-(pyridine-3-sulphonyl)-pyrazolo[1,5-a]pyrimidin-7-ylamine

In an analogous manner to that described in Example 4), from 7-chloro-5-methyl-2-methylsulphanyl-3-(pyridine-3-sulphonyl)-pyrazolo[1,5-a]pyrimidine and NH_3 in MeOH there was obtained 5-methyl-2-methylsulphanyl-3-(pyridine-3-sulphonyl)-pyrazolo[1,5-a]pyrimidin-7-ylamine as colourless crystals, m.p. 226.8 -227.5°.

Example 94

2-[3-Benzenesulphonyl-5-methyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-6-yl]-ethanol

a) A solution of 2.69 g (10 mmol) of 4-benzenesulphonyl-5-methylsulphanyl-2H-pyrazol-3-ylamine and 1.28 g (10 mmol) of 2-acetyl-butyrolactone in 10 ml of acetic acid was heated at reflux for 1.5 hrs. After cooling to RT the mixture was treated with 50 ml of H_2O

and extracted three times with CH_2Cl_2. The organic phases were dried ($MgSO_4$), filtered and evaporated. Chromatography (SiO_2, CH_2Cl_2/MeOH 20:1) of the residue yielded 1.5 g (36%) of ethyl 2-(3-benzenesulphonyl-7-hydroxy-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-6-yl)-acetate as a colourless foam.

b) A suspension of 1.5 g (3.56 mmol) of ethyl 2-(3-benzenesulphonyl-7-hydroxy-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-6-yl)-acetate in 30 ml of $POCl_3$ was heated at reflux for 4 hrs. The reaction solution was cooled to RT and evaporated. The residue was treated with 100 ml of ice-water and the pH value of the solution was adjusted to 8 with sat. $NaHCO_3$ solution. The aqueous phase was extracted three times with 70 ml of CH_2Cl_2, and the organic phases were dried ($MgSO_4$), filtered and evaporated.Chromatography (SiO_2, CH_2Cl_2 20:1) of the residue yielded 1.0 g (94%) of ethyl 2-(3-benzenesulphonyl-7-chloro-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-6-yl)-aceate as a pale yellow solid.

c) 0.1 g (1 mmol) of 1-methyl-piperazine in 5 ml of DMF was added to a solution of 0.35 g (0.8 mmol) of ethyl 2-(3-benzenesulphonyl-7-chloro-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-6-yl)-acetate in 15 ml of DMF and stirred at 60° for 2 hrs. The DMF was evaporated in a high vacuum, the residue was partitioned between 2N NaOH and CH_2Cl_2, the aqueous phase was extracted three times with CH_2Cl_2. The combined organic phases were dried ($MgSO_4$), filtered and evaporated. Subsequent chromatography (SiO_2, CH_2Cl_2/MeOH 10:1) yielded 0.34 g of a yellow foam, which was dissolved in a mixture of 50 ml of tetrahydrofuran/dioxan/H_2O 1:1:1. After the addition of 4 ml of 2N NaOH the mixture was stirred at 45° for 12 hrs., treated with 100 ml of H_2O and extracted three times with 60 ml of CH_2Cl_2. The combined organic phases were dried ($MgSO_4$), filtered and evaporated. Chromatography (SiO_2, CH_2Cl_2/MeOH 8:1) yielded 0.18 mg (48%) of 2-[3-benzenesulphonyl-5-methyl-7-(4-methyl-piperazin-1-yl)-2-methyl-sulphanyl-pyrazolo[1,5-a]pyrimidin-6-yl]-ethanol as a colourless foam.

Example 95

2-[3-Benzenesulphonyl-5-methyl-7-(4-methyl-piperazin-1-yl)-pyrazolo[1,5-a]pyrimidin-2-yloxy]-ethanol

a) 0.88 g (22 mmol) of powdered NaOH was added to a solution of 2 g (11 mmol) of phenysulphonylacetonitrile in 20 ml of acetonitrile and stirred at RT for 2 hrs.

Subsequently, a solution of 1.14 ml (11 mmol) of 2-chloroethyl chloroformate in 4 ml of acetonitrile was added dropwise thereto at 5° and the mixture was heated at reflux for 1 hr. After cooling to RT the precipitate was filtered off and the filtrate was evaporated. The thus-obtained brown oil was taken up in 50 ml of EtOH and treated with 0.54 ml (11 mmol) of NH_2NH_2 and heated at reflux for 1 hr. After evaporation of the reaction solution and subsequent chromatography (SiO_2, CH_2Cl_2/MeOH/NH_4OH 110:10:1) there were obtained 1.4 g (45%) of 2-(5-amino-4-benzenesulphonyl-1H-pyrazol-3-yloxy)-ethanol as a colourless solid.

b) 0.75 ml of ethyl acetoacetate was added to a solution of 1.14 g (4.9 mol) of 2-(5-amino-4-benzenesulphonyl-1H-pyrazol-3-yloxy)-ethanol in 10 ml of acetic acid and heated at reflux for 3 hrs. After cooling to RT the mixture was treated with 50 ml of H_2O and extracted three times with CH_2Cl_2. The organic phases were dried ($MgSO_4$), filtered and evaporated. Chromatography (SiO_2, CH_2Cl_2/MeOH 19:1) of the residue yielded 0.8 g (42%) of ethyl 2-(3-benzenesulphonyl-7-hydroxy-5-methyl-pyrazolo[1,5-a]pyrimidin-2-yloxy)-acetate as a colourless oil.

c) A suspension of 0.8 g (2 mmol) of ethyl 2-(3-benzenesulphonyl-7-hydroxy-5-methyl-pyrazolo[1,5-a]pyrimidin-2-yloxy)-acetate in 20 ml of $POCl_3$ was heated at reflux for 4 hrs. The reaction solution was cooled to RT and evaporated. The residue was treated with 80 ml of ice-water and the pH value of the solution was adjusted to 8 with sat. $NaHCO_3$ solution. The aqueous phase was extracted three times with 70 ml of CH_2Cl_2, and the organic phases were dried ($MgSO_4$), filtered and evaporated. Chromatography (SiO_2, CH_2Cl_2/MeOH 40:1) of the residue yielded 0.56 g (68%) of ethyl 2-[3-benzenesulphonyl-5-methyl-7-(4-methyl-piperazin-1-yl)-pyrazolo[1,5-a]pyrimidin-2-yloxy]-acetate as a colourless solid.

d) 0.38 ml (3.4 mmol) of 1-methyl-piperazine in 5ml of DMF was added to a solution of 0.56 g (1.4 mmol) of ethyl 2-[3-benzenesulphonyl-5-methyl-7-(4-methyl-piperazin-1-yl)-pyrazolo[1,5-a]pyrimidin-2-yloxy]-acetate in 10 ml of DMF and stirred at RT for 1.5 hrs. The DMF was evaporated in a high vacuum,and the residue was partitioned between 2N NaOH and CH_2Cl_2. The aqueous phase was extracted three times with CH_2Cl_2, and the combined organic phases were dried ($MgSO_4$), filtered and evaporated. Subsequent chromatography (silica gel, CH_2Cl_2/MeOH 19:1) yielded 0.61 g (92%) of ethyl 2-[3-benzenesulphonyl-5-methyl-7-(4-methyl-piperazin-1-yl)-pyrazolo[1,5-a]pyrimidin-2-yloxy]-acetate as a colourless foam.

e) A solution of 0.126 g of KOH in 5 ml of H_2O was added to a solution of 0.61 mg of ethyl 2-[3-benzenesulphonyl-5-methyl-7-(4-methyl-piperazin-1-yl)-pyrazolo[1,5-a]pyrimidin-2-yloxy]-acetate in 20 ml of dioxan/THF 1:1 and stirred at RT for 2.5 hrs. The reaction solution was evaporated and the residue was partitioned between H_2O and CH_2Cl_2. The aqueous phase was washed three times with 30 ml of CH_2Cl_2,and the combined organic phases were dried ($MgSO_4$), filtered and evaporated. Chromatography (SiO_2,CH_2Cl_2/MeOH 9:1) of the residue and crystallization from EtOH yielded 0.18 g (18%) of 2-[3-benzenesulphonyl-5-methyl-7-(4-methyl-piperazin-1-yl)-pyrazolo[1,5-a]pyrimidin-2-yloxy]-ethanol as colourless crystals. M.p. 177.5-178°.

Example 96

3-Benzenesulphonyl-5-(2-methoxy-ethyl)-2-methylsulphanyl-7-piperazin-1-yl-pyrazolo[1,5-a]pyrimidine

a) A solution of 2.69 g (10 mmol) of) 4-benzenesulphonyl-5-methylsulphanyl-2H-pyrazol-3-ylamine and 1.6 g (10 mmol) of methyl 5-methoxy-3-oxo-valerate in 10 ml of acetic acid was heated at reflux for 4 hrs. The reaction solution was evaporated and the residue was partitioned between H_2O and CH_2Cl_2. The aqueous phase was extracted three times with 80 ml of CH_2Cl_2. The combined organic phases were dried ($MgSO_4$), filtered and evaporated. Chromatography (SiO_2, CH_2Cl_2/MeOH 20:1) yielded 2.40 g (61%) of 3-benzenesulphonyl-5-(2-methoxy-ethyl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ol as a beige powder.

b) A suspension of 2.3 g (6.0 mmol) of 3-benzenesulphonyl-5-(2-methoxy-ethyl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ol in 40 ml of $POCl_3$ and 20 ml of diethyl-aniline was heated at reflux for 1 hr. The reaction solution was cooled to RT and evaporated. The residue was treated with 100 ml of ice-water and the pH value of the solution was adjusted to 8 with sat. $NaHCO_3$ solution. The aqueous phase was extracted three times with 100 ml of CH_2Cl_2, and the organic phases were dried ($MgSO_4$), filtered and evaporated. Chromatography (SiO_2, CH_2Cl_2/MeOH 200:1) of the residue yielded 1.8 g (75%) of 3-benzenesulphonyl-7-chloro-5-(2-methoxy-ethyl)-2-methylsuphanyl-pyrazolo-[1,5-a]pyrimidine as a pale yellow powder.

c) 0.4 g (4.8 mmol) of piperazine in 3 ml of DMF was added to a solution of 0.35 g (0.8 mmol) of 3-benzenesulphonyl-7-chloro-5-(2-methoxy-ethyl)-2-methylsuphanyl-pyrazolo-[1,5-a]pyrimidine in 10 ml of DMF and stirred at RT for 2 hrs. The reaction solution was evaporated, the residue was partitioned between 2N NaOH and CH_2Cl_2. The aqueous phase was extracted three times with CH_2Cl_2, the combined organic phases were dried ($MgSO_4$), filtered and evaporated. Subsequent chromatography (SiO_2, CH_2Cl_2/MeOH 8:1) and crystalliazation from EtOH yielded 0.25 g (62%) of 3-benzenesuphonyl-5-(2-methoxy-ethyl)-2-methylsulphanyl-7-piperazin-1-yl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 155-156°.

Example 97

3-Benzenesulphonyl-5-(2-methoxy-ethyl)-7-(4-methyl-piperazin-1-yl)-2-methyl-sulphanyl-pyrazolo[1,5-a]pyrimidine

0.5 g (5 mmol) of 1-methyl-piperazine in 3 ml of DMF was added to a solution of 0.45 g (1.13 mmol) of 3-benzenesulphonyl-7-chloro-5-(2-methoxy-ethyl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine in 20 ml of DMF and stirred at RT for 3 hrs. The reaction solution was evaporated, the residue was partitioned between 2N NaOH and CH_2Cl_2. The aqueous phase was extracted three times with CH_2Cl_2, the combined organic phases were dried ($MgSO_4$), filtered and evaporated. Subsequent chromatography (silica gel, CH_2Cl_2/MeOH 12:1) and crystallization from EtOH yielded 0.255 g (67%) of 3-benzenesulphonyl-5-(2-methoxy-ethyl)-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-pyrazolo[1,5-a]-pyrimidine as colourless crystals, m.p. 160-161°.

Example 98

3-Benzenesulphonyl-5-(2-methoxy-ethyl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ylamine

10 ml of a 50% solution of NH_3 in MeOH were added to a solution of 0.50 g (1.25 mmol) of 3-benzenesulphonyl-7-chloro-5-(2-methoxy-ethyl)-2-methylsulphanyl-pyrazolo[1,5-a]-pyrimidine in 10 ml of DMF and stirred at RT for 2 hrs. The reaction solution was evaporated in a high vacuum and the residue was partitioned between 2N NaOH and CH_2Cl_2. The aqueous phase was extracted three times with CH_2Cl_2 and the combined

organic phases were dried ($MgSO_4$), filtered and evaporated. Subsequent chromatography (SiO_2, CH_2Cl_2/MeOH 10:1) and crystallization from EtOH yielded 0.30 g (63%) of 3-benzene- sulphonyl-5-(2-methoxy-ethyl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ylamine as colourless crystals, m.p. 186-187°.

Example 99

3-Benzenesulphonyl-5-(2-methoxy-ethyl)-2-methylsulphanyl-7-piperazin-1-yl-pyrazolo[1,5-a]pyrimidine

a) A solution of 2.69 g (10 mmol) 4-benzenesulphonyl-5-methylsulphanyl-2H-pyrazol-3-ylamine and 1.46 g (10 mmol) of ethyl 4-methoxy-acetoacetate in 10 ml of acetic acid was heated at reflux for 3 hrs. The reaction solution was evaporated and the residue was partitioned between H_2O and CH_2Cl_2. The aqueous phase was extracted three times with 80 ml of CH_2Cl_2. The combined organic phases were dried ($MgSO_4$), filtered and evaporated. Chromatography (SiO_2, CH_2Cl_2/MeOH 25:1) yielded 3.1 g (85%) of 3-benzenesulphonyl-5-methoxymethyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ol as a beige powder, m.p. 175-155°.

b) A suspension of 2.5 g (6.8 mmol) of 3-benzenesulphonyl-5-methoxymethyl-2-methyl-sulphanyl-pyrazolo[1,5-a]pyrimidin-7-ol in 40 ml of $POCl_3$ and 20 ml of diethylaniline was heated at reflux for 1 hr. The reaction solution was cooled to RT and evaporated. The residue was treated with 100 ml of ice-water and the pH value of the solution was adjusted to 8 with sat. $NaHCO_3$ solution. The aqueous phase was extracted three times with 100 ml of CH_2Cl_2,and the organic phases were dried ($MgSO_4$), filtered and evaporated. Chromatography (SiO_2, CH_2Cl_2/MeOH 20:1) yielded 2.1 g (79%) of 3-benzenesulphonyl-7-chloro-5-methoxymethyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine as a pale yellow powder, m.p. 194-197°.

c) 0.4 g (4.8 mmol) of piperazine in 3 ml of DMF was added to a solution of 0.50 g (1.13 mmol) of 3-benzenesulphonyl-7-chloro-5-methoxymethyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine in 15 ml of DMF and stirred at RT for 2 hrs. The reaction solution was evaporated and the residue was partitioned between 2N NaOH and CH_2Cl_2. The aqueous phase was extracted three times with CH_2Cl_2, and the combined organic phases were dried ($MgSO_4$), filtered and evaporated. Subsequent chromatography (SiO_2, CH_2Cl_2/ MeOH 8:1) and crystallization from EtOH yielded 0.42 g (74%) of 3-

benzenesulphonyl-5-(2-methoxy-ethyl)-2-methylsulphanyl-7-piperazin-1-yl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 170-171°.

Example 100

3-Benzenesulphonyl-5-methoxymethyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine

0.50 g (5 mmol) of 1-methyl-piperazine in 3 ml of DMF was added to a solution of 0.50 g (1.3 mmol) of 3-benzenesulphonyl-7-chloro-5-methoxymethyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine in 15 ml of DMF and stirred at RT for 2 hrs. The reaction solution was evaporated and the residue was partitioned between 2N NaOH and CH_2Cl_2. The aqueous phase was extracted three times withCH_2Cl_2, and the combined organic phases were dried ($MgSO_4$), filtered and evaporated. Subsequent chromatography (SiO_2, CH_2Cl_2/ MeOH 10:1) and crystallization from EtOH yielded 0.42 g (72%) of 3-benzene-sulphonyl-5-methoxy- methyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-pyrazolo-[1,5-a]pyrimidine as colorless crystals, m.p. 207-208°.

Example 101

3-Benzenesulphonyl-5-methoxymethyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ylamine

10 ml of a 50% solution of NH_3 in MeOH was added to a solution of 0.50 g (1.3 mmol) of 3-benzenesulphonyl-7-chloro-5-methoxymethyl-2-methylsulphanyl-pyrazolo[1,5-a]-pyrimidine in 10 ml of DMF and stirred at RT for 2 hrs. The reaction solution was evaporated and the residue was partitioned between 2N NaOH and CH_2Cl_2. The aqueous phase was extracted three times with CH_2Cl_2, and the combined organic phases were dried ($MgSO_4$), filtered and evaporated. Subsequent chromatography (SiO_2, CH_2Cl_2/MeOH 10:1) and crystallization from EtOH yielded 0.38 g (80%) of 3-benzenesulphonyl-5-methoxymethyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ylamine as colourless crystals, m.p. >230°.

Example 102

3-Benzenesulphonyl-5-chloro-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-pyrazolo [1,5-a]pyrimidine

a) 5.38 g (20 mmol) of 4-benzensulphonyl-5-methylsulphanyl-2H-pyrazol-3-ylamine followed by 9 ml (60 mmol) of diethyl malonate were added to a freshly prepared solution of sodium ethanolate in EtOH (prepared from 0.89 g (77 mmol) of sodium in 100 ml of EtOH) and the mixture was heated at reflux for 48 hrs. After cooling to RT the mixture was subsequently poured on to 140 ml of ice-water. The resulting precipitate was filtered off and dried at 50° in a high vacuum. There were thus obtained 6.5 g (96%) of 3-benzene-sulphonyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine-5,7-diol as a beige powder, m.p. >230°.

b) A suspension of 3.0 g (8.89 mmol) of 3-benzenesulphonyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine-5,7-diol in 40 ml of $POCl_3$ was heated at reflux for 1 hr. The reaction solution was cooled to RT and evaporated. The residue was treated with 100 ml of ice-water and the pH value of the solution was adjusted to 8 with sat. $NaHCO_3$ solution. The aqueous phase was extracted three times with 90 ml of CH_2Cl_2, and the organic phases were dried ($MgSO_4$), filtered and evaporated. Chromatography (SiO_2, CH_2Cl_2/AcOEt 1.5:1) of the residue yielded 1.8 g (54%) of 3-benzenesulphonyl-5,7-dichloro-2-methyl-sulphanyl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 193-197°.

c) 0.1 g (1 mmol) of 1-methyl-piperazine in 3 ml of CH_2Cl_2 was added to a solution of 0.37 g (1 mmol) of 3-benzenesulphonyl-5,7-dichloro-2-methylsulphanyl-pyrazolo[1,5-a]-pyrimidine in 20 ml of CH_2Cl_2 and stirred at RT for 1 hr. The mixture was poured on to ice-water, adjusted to pH8 with $NaHCO_3$ solution and extracted three times with 30 ml of CH_2Cl_2. The combined organic phases were dried ($MgSO_4$), filtered and evaporated. Subsequent chromatography (silica gel, CH_2Cl_2/MeOH 4:1) yielded 0.38 g (86%) of 3-benzenesulphonyl-5-chloro-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. >230°.

Example 103

3-Benzenesulphonyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-pyrazolo[1,5-a]-pyrimidine

0.15 g of Pd/C (10%) and 0.3ml of NEt_3 were added to a solution of 0.189 g (0.4 mmol) of 3-benzenesulphonyl-5-chloro-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine in 30 ml of EtOH and hydrogenated at RT for 12 hrs. The reaction mixture was filtered over Dicalite and the filtrate was evaporated. Chromatography of the residue (SiO_2, CH_2Cl_2/MeOH 20:1) yielded 0.08 g (49%) of 3-benzenesulphonyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 107-109°.

Example 104

2-[3-Benzenesulphonyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-5-yloxy]-ethanol

0.115 g (5 mmol) of sodium was added to 20 ml of ethylene glycol and this solution was treated with 0.22 g (0.5 mmol) of 3-benzenesulphonyl-5-chloro-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine and subsequently stirred at 80° for 1 hr. After cooling to RT the reaction solution was poured on to 70 ml of ice-water and extracted three times with 50 ml of AcOEt. The combined organic phases were dried ($MgSO_4$), filtered and evaporated. Chromatography (SiO_2, CH_2Cl_2/MeOH/NH_4OH 110:10:1) of the residue and crystallization from EtOH yielded 0.16 g (69%) of 2-[3-benzenesulphonyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-5-yloxy]-ethanol as colourless crystals, m.p. 187-189°.

Example 105

3-Benzenesulphonyl-5-chloro-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ylamine

10 ml of a 50% solution of NH_3 in MeOH were added to a solution of 0.35 g (0.935 mmol) of 3-benzenesulphonyl-5,7-dichloro-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine in 10 ml of DMF and stirred at RT for 12 hrs. The DMF was evaporated in a high vacuum and the residue was partitioned between 2N NaOH and CH_2Cl_2. The aqueous phase was

extracted three times with CH_2Cl_2, and the combined organic phases were dried ($MgSO_4$), filtered and evaporated. Subsequent chromatography (silica gel, CH_2Cl_2/AcOEt 15:1) and crystallization from EtOH yielded 0.28 g (84%) of 3-benzenesulphonyl-5-chloro-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ylamine as colourless crystals, m.p. >230°.

Example 106

3-Benzenesulphonyl-N5,N5-dimethyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine-5,7-diamine and

3-benzenesulphonyl-N5-(2-dimethylamino-ethyl)-2-methylsulphanyl-pyrazolo[1,5-a] pyrimidine-5,7-diamine

0.26 g (3 mmol) of 2-dimethylaminoethylamine in 5 ml of DMF was added to a solution of 0.4 g (1 mmol) of 3-benzenesulphonyl-5-chloro-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ylamine in 10 ml of DMF and stirred at 90° for 1 hr. The reaction solution was evaporated and the residue was partitioned between H_2O and CH_2Cl_2. The aqueous phase was extracted three times with 50 ml of CH_2Cl_2. The combined organic phases were dried ($MgSO_4$), filtered and evaporated. Subsequent chromatography (silica gel, CH_2Cl_2/MeOH 8:1) yielded 0.20 g (48%) of 3-benzenesulphonyl-N5,N5-dimethyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine-5,7-diamine as colourless crystals, m.p. >230°, and 0.08 g (17%) of 3-benzenesulphonyl-N5-(2-dimethylamino-ethyl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine-5,7-diamine as colourless crystals, m.p. 210-212°.

Example 107

3-Benzenesulphonyl-5-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-pyrazolo[1,5-a] pyrimidin-7-ylamine

0.1 g (1 mmol) of 1-methyl-piperazine was added to a solution of 0.14 g (0.4 mmol) of 3-benzenesulphonyl-5-chloro-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ylamine in 5 ml of DMF and stirred at 90° for 1 hr. The reaction solution was evaporated and the residue was partitioned between H_2O and CH_2Cl_2. The aqueous phase was extracted three times with 50 ml of CH_2Cl_2. The combined organic phases were dried ($MgSO_4$), filtered and evaporated. Subsequent chromatography (silica gel CH_2Cl_2/MeOH 9:1) and

crystallization from EtOH yielded 0.1 g (59%) of 3-benzenesulphonyl-5-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ylamine as colourless crystals, m.p. >230°.

Example 108

3-Benzenesulphonyl-5-chloro-2-ethyl-7-(4-methyl-piperazin-1-yl)-pyrazolo[1,5-a]-pyrimidine

a) 7.90 g (31.4 mmol) of 4-benzenesulphonyl-5-ethyl-2H-pyrazol-3-ylamine followed by 14.3 ml (94.3 mmol) of diethyl malonate were added to a freshly prepared solution of sodium ethanolate in EtOH (prepared from 2.7 g (119.5 mmol) of sodium in 320 ml of EtOH) and heated at reflux for 48 hrs. After cooling to RT the mixture was subsequently poured into 140 ml of ice-water. The resulting precipitate was filtered off and dried at 50° in a high vacuum. There were thus obtained 4.8 g (48%) of 3-benzenesulphonyl-2-ethyl-pyrazolo- [1,5-a]pyrimidine-5,7-diol, m.p. >230°.

b) A suspension of 2.8 g (8.8 mmol) of 3-benzenesulphonyl-2-ethyl-pyrazolo1,5-a]pyrimidine-5,7-diol in 30 ml of $POCl_3$ was heated at reflux for 1 hr. The reaction solution was cooled to RT and evaporated. The residue was treated with 100 ml of ice-water and the pH value of the solution was adjusted to 8 with sat. $NaHCO_3$ solution. The aqueous phase was extracted three times with 100 ml of CH_2Cl_2, and the organic phases were dried ($MgSO_4$), filtered and evaporated. Chromatography (SiO_2, CH_2Cl_2) of the residue yielded 1.1 g (35%) of 3-benzenesulphonyl-5,7-dichloro-2-ethyl-pyrazolo[1,5-a]pyrimidine as a colourless solid.

c) 0.86 ml (7.7 mmol) of 1-methyl-piperazine in 3 ml of CH_2Cl_2 was added to a solution of 2.5 g (7 mmol) of 3-benzenesulphonyl-5,7-dichloro-2-ethyl-pyrazolo[1,5-a]pyrimidine in 20 ml of CH_2Cl_2 and stirred at RT for 2 hrs. The reaction mixture was poured on to ice-water, adjusted to pH 8 with $NaHCO_3$ solution and extracted three times with CH_2Cl_2. The combined organic phases were dried ($MgSO_4$), filtered and evaporated. Subsequent chromatography (silica gel, CH_2Cl_2/MeOH 4:1) and crystallization from EtOH yielded 2.5 g (85%) of 3-benzenesulphonyl-5,7-dichloro-2-ethyl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 166-167°.

Example 109

3-Benzenesulphonyl-2-ethyl-7-(4-methyl-piperazin-1-yl)-pyrazolo[1,5-a]pyrimidine

0.1 g of Pd/C (10%) was added to a solution of 0.267 g (0.63 mmol) of 3-benzene-sulphonyl-5-chloro-2-ethyl-7-(4-methyl-piperazin-1-yl)-pyrazolo[1,5-a]pyrimidine in 40 ml of EtOH and the mixture was hydrogenated at RT for 4 hrs. The reaction mixture was filtered over Dicalite and the filtrate was evaporated. The residue was partitioned between CH_2Cl_2 and sat. $NaHCO_3$ solution. The aqueous phase was extracted three times with CH_2Cl_2. The combined organic phases were dried ($MgSO_4$), filtered and evaporated. Chromatography of the residue ($SiO_2CH_2Cl_2$/MeOH 19:1) and crystallization from EtOH yielded 0.2 g (53%) of as pale beige crystals, m.p. 206-207°.

Example 110

3-Benzenesulphonyl-2-ethyl-5,7-bis-(4-methyl-piperazin-1-yl)-pyrazolo[1,5-a]pyrimidine

0.33 ml (3 mmol) of 1-methyl-piperazine in 5 ml of DMF was added to a solution of 0.50 g (1.2 mmol) of 3-benzenesulphonyl-5-chloro-2-ethyl-7-(4-methyl-piperazin-1-yl)-pyrazolo[1,5-a]pyrimidine in 15 ml of DMF and stirred at 100° for 1 hr. After cooling to RT the reaction solution was evaporated and the residue was partitioned between 2N NaOH and CH_2Cl_2. The aqueous phase was extracted three times with CH_2Cl_2, and the combined organic phases were dried ($MgSO_4$), filtered and evaporated. Subsequent chromatography (SiO_2, CH_2Cl_2/MeOH/NH_4OH 110:10:1) and crystallization from EtOH yielded 0.04 g (69%) of 3-benzenesulphonyl-2-ethyl-5,7-bis-(4-methyl-piperazin-1-yl)-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 232-235°.

Example 111

3-Benzenesulphonyl-2-ethyl-7-(4-methyl-piperazin-1-yl)-5-morpholin-4-yl-pyrazolo[1,5-a]-pyrimidine

0.26 ml (3 mmol) of morpholine in 5 ml of DMF was added to a solution of 0.50 g (1.2 mmol) of 3-benzenesulphonyl-5-chloro-2-ethyl-7-(4-methyl-piperazin-1-yl)-pyrazolo[1,5-a]pyrimidine in 15 ml of DMF and stirred at 100° for hr. After cooling to RT the reaction solution was evaporated and the residue was partitioned between 2N NaOH and CH_2Cl_2. The aqueous phase was extracted three times with CH_2Cl_2, and the combined organic phases were dried ($MgSO_4$), filtered and evaporated. Subsequent chromatography (SiO_2, CH_2Cl_2/MeOH 15:1) and crystallization from EtOH yielded 0.45 g (80%) of 3-

benzenesulphonyl-2-ethyl-7-(4-methyl-piperazin-1-yl)-5-morpholin-4-yl-pyrazolo[1,5-a]-pyrimidine as colourless crystals, m.p. >250°.

Example 112

2-[3-Benzenesulphonyl-2-ethyl-7-(4-methyl-piperazin-1-yl)-pyrazolo[1,5-a]pyrimidin-5-yl-oxy]-ethanol

0.274 g (12 mmol) of sodium was added to 40 ml of ethylene glycol and this solution was treated with 0.50 g (1.2 mmol) of 3-benzenesulphonyl-5-chloro-2-ethyl-7-(4-methyl-piperazin-1-yl)-pyrazolo[1,5-a]pyrimidine and subsequently stirred at 80° for 1 hr. After cooling to RT the reaction solution was poured on to 70 ml of ice-water and extracted three times with 50 ml of AcOEt. The combined organic phases were dried ($MgSO_4$), filtered and evaporated. Chromatography (SiO_2, CH_2Cl_2/MeOH 9:1) of the residue and crystallization from EtOH yielded 0.24 g (44%) of 2-[3-benzenesulphonyl-2-ethyl-7-(4-methyl-piperazin-1-yl)-pyrazolo[1,5-a]pyrimidin-5-yl-oxy]-ethanol as colourless crystals, m.p. 153-154°.

Example 113

[3-Benzenesulphonyl-2-ethyl-7-(4-methyl-piperazin-1-yl)-pyrazolo[1,5-a]pyrimidin-5-yl]-dimethyl-amine and

N-[3-benzenesulphonyl-2-ethyl-7-(4-methyl-piperazin-1-yl)-pyrazolo[1,5-a]pyrimidin-5-yl]-N',N'-dimethyl-ethane-1,2-diamine

0.72 g (6.5 mmol) of 2-dimethylaminoethylamine in 5 ml of DMF was added to a solution of 1.1 g (2.6 mmol) of 3-benzenesulphonyl-5-chloro-2-ethyl-7-(4-methyl-piperazin-1-yl)-pyrazolo[1,5-a]pyrimidine in 20 ml of DMF and stirred at 90° for 1 hr. After cooling to RT the reaction solution was evaporated and the residue was partitioned between 2N NaOH and CH_2Cl_2. The aqueous phase was extracted three times with 80 ml of CH_2Cl_2. The combined organic phases were dried ($MgSO_4$), filtered and evaporated. Subsequent chromatography (silica gel, CH_2Cl_2/MeOH/NH_4OH 65:10:1) yielded 0.20 g (13%) of [3-benzenesulphonyl-2-ethyl-7-(4-methyl-piperazin-1-yl)-pyrazolo[1,5-a]pyrimidin-5-yl]-dimethyl-amine as colourless crystals, m.p. 211-212°, and 0.30 g (24%) of N-[3-benzenesulphonyl-2-ethyl-7-(4-methyl-piperazin-1-yl)-pyrazolo[1,5-a]pyrimidin-5-yl]-N',N'-dimethyl-ethane-1,2-diamine as colourless crystals, m.p. 163-164°.

Example 114

3-Benzenesulphonyl-5-chloro-2-ethyl-pyrazolo[1,5-a]pyrimidin-7-ylamine

20 ml of a 50% solution of NH_3 in MeOH were added to a solution of 1.1 g (3.1 mmol) of 3-benzenesulphonyl-2-ethyl-pyrazolo[1,5-a]pyrimidine-5,7-diol in 10 ml of DMF and stirred at RT for 12 hrs. The reaction solution was evaporated and the residue was partitioned between 2N NaOH and CH_2Cl_2. The aqueous phase was extracted three times with CH_2Cl_2 and the combined organic phases were dried ($MgSO_4$), filtered and evaporated. The residue was treated with 5 ml of EtOH and the crystals obtained were filtered off. There was thus obtained 0.80 g (77%) of 3-benzenesulphonyl-5-chloro-2-ethyl-pyrazolo[1,5-a]- pyrimidin-7-ylamine as colourless crystals, m.p. >220°.

Example 115

3-Benzenesulphonyl-2-ethyl-5-(4-methyl-piperazin-1-yl)-pyrazolo[1,5-a]pyrimidin-7-yl-amine

0.33 ml (3 mmol) of 1-methyl-piperazine was added to a solution of 0.4 g (1.2 mmol) of 3-benzenesulphonyl-5-chloro-2-ethyl-pyrazolo[1,5-a]pyrimidin-7-ylamine in 5 ml of DMF and stirred at 90° for 1 hr. The reaction solution was evaporated and the residue was partitioned between H_2O and CH_2Cl_2. The aqueous phase was extracted three times with 50 ml of CH_2Cl_2. The combined organic phases were dried ($MgSO_4$), filtered and evaporated. Subsequent chromatography (silica gel, CH_2Cl_2/MeOH 9:1) and crystallization from EtOH yielded 0.24 g (50%) of 3-benzenesulphonyl-2-ethyl-5-(4-methyl-piperazin-1-yl)-pyrazolo[1,5-a]pyrimidin-7-yl-amine as colourless crystals, m.p. >250°.

Example 116

3-Benzenesulphonyl-2-ethyl-N5,N5-dimethyl-pyrazolo[1,5-a]pyrimidine-5,7-diamine and 3-benzenesulphonyl-N5-(2-dimethylamino-ethyl)-2-ethyl-pyrazolo[1,5-a]pyrimidine-5,7-diamine

0.73 g (6.68 mmol) of 2-dimethylaminoethylamine in 5 ml of DMF was added to a solution of 0.45 g (1.3 mmol) of 3-benzenesulphonyl-5-chloro-2-ethyl-pyrazolo[1,5-a]pyrimidin-7-ylamine in 10 ml of DMF and stirred at 90° for 1 hr. The reaction solution was evaporated and the residue was partitioned between H_2O and CH_2Cl_2. The aqueous phase was extracted three times with 80 ml of CH_2Cl_2. The combined organic phases were dried ($MgSO_4$), filtered and evaporated. Subsequent chromatography (silica gel, CH_2Cl_2/ MeOH/NH_4OH 65:10:1) yielded 0.12 g (26%) 3-benzenesulphonyl-2-ethyl-N5,N5-dimethyl-pyrazolo[1,5-a]pyrimidine-5,7-diamine as colourless crystals, m.p. 228-230°, and 0.09 g (17 %) 3-benzenesulphonyl-N5-(2-dimethylamino-ethyl)-2-ethyl-pyrazolo[1,5-a]pyrimidine-5,7-diamine as colourless crystals, m.p. 149.5-150.5°.

Example 117

3-Benzenesulphonyl-5-dimethylaminomethyl-2-methylsulphanyl-pyrazolo[1,5-a]-pyrimidin-7-ylamine

a) 6.8 ml (50 mmol) of ethyl 4-chloro-acetoacoacetate were added to a solution of 13.5 mmol (50 mmol) of 4-benzenesulphonyl-5-methylsulphanyl-2H-pyrazol-3-ylamine in 80 ml of acetic acid and heated at reflux for 1.5 hrs. After cooling to RT the crystals obtained were filtered off, washed with EtOH and dried in a high vacuum at 50°. There were thus obtained 10.5 g (56%) of 3-benzenesulphonyl-5-chloromethyl-2-methyl-sulphanyl-pyrazolo[1,5-a]pyrimidin-7-ol as a colourless powder, m.p. 215-217°.

b) 5 ml of a 33% solution of dimethylamine in EtOH were added to a solution of 1.4 g (3.7 mmol) of 3-benzenesulphonyl-5-chloromethyl-2-methyl- sulphanyl-pyrazolo[1,5-a]pyrimidin-7-ol in 20 ml of DMF and stirred at RT for 4 hrs. The reaction solution was evaporated, the residue was partitioned between 0.5N NaOH and CH_2Cl_2. The aqueous phase was extracted three times with CH_2Cl_2. The combined organic phases were dried ($MgSO_4$), filtered and evaporated. Chromatography (SiO_2, CH_2Cl_2/MeOH 4:1) yielded

1.3 g (92%) of 3-benzenesulphonyl-5-dimethylaminomethyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ol as a beige powder, m.p. >220°.

c) A suspension of 1.3 g (3.43 mmol) of 3-benzenesulphonyl-5-dimethylaminomethyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ol in 50 ml of $POCl_3$ was heated at reflux for 3 hrs. The reaction solution was cooled to RT and evaporated. The residue was treated with 100 ml of ice-water and the pH value of the solution was adjusted to 8 with sat. $NaHCO_3$ solution. The aqueous phase was extracted three times with 100 ml of CH_2Cl_2 and the organic phases were dried ($MgSO_4$), filtered and evaporated. Chromatography (SiO_2, CH_2Cl_2/AcOEt 1:1) of the residue yielded 1.2 g (88%) of (3-benzenesulphonyl-7-chloro-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-5-ylmethyl)-dimethyl-amine as a colourless foam.

d) 20 ml of a 50% solution of NH_3 in MeOH were added to a solution of 1.20 g (3 mmol) of (3-benzenesulphonyl-7-chloro-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-5-ylmethyl)-dimethyl-amine in 30 ml of DMF and stirred at RT for 4 hrs. The reaction solution was evaporated and the residue was partitioned between 2N NaOH and CH_2Cl_2. The aqueous phase was extracted three times with CH_2Cl_2, and the combined organic phases were dried ($MgSO_4$), filtered and evaporated. Subsequent chromatography (silica gel, CH_2Cl_2/MeOH 6:1) yielded 0.90 g (78%) of 3-benzenesulphonyl-5-dimethylaminomethyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ylamine as colourless crystals, m.p. 216-218°.

Example 118

3-Benzenesulphonyl-5-(4-methyl-piperazin-1-ylmethyl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ylamine

a) 1.57 ml (15.70 mml) of 1-methyl-piperazine were added to a solution of 2.9 g (7.84 mmol) of 3-benzenesulphonyl-5-chloromethyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ol in 20 ml of DMF and stirred at RT for 4 hrs. The reaction solution was evaporated and the residue was partitioned between 0.5N NaOH and CH_2Cl_2. The aqueous phase was extracted three times with CH_2Cl_2. The combined organic phases were dried ($MgSO_4$), filtered and evaporated. Chromatography (SiO_2, CH_2Cl_2/MeOH 4:1) yielded 1.85 g (53%) of 3-benzenesulphonyl-5-(4-methyl-piperazin-1-ylmethyl)-2-methyl-sulphanyl-pyrazolo[1,5-a]pyrimidin-7-ol as a beige powder, m.p. >220°.

b) A suspension of 4.0 g (9 mmol) of 3-benzenesulphonyl-5-(4-methyl-piperazin-1-ylmethyl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ol in 100 ml of $POCl_3$ was heated at reflux for 3 hrs. The reaction solution was cooled to RT and evaporated. The residue was treated with 100 ml of ice-water and the pH value of the solution was adjusted to 8 with sat. $NaHCO_3$ solution. The aqueous phase was extracted three times with 150 ml of CH_2Cl_2 and the organic phases were dried ($MgSO_4$), filtered and evaporated. Chromatography ($SiO_2CH_2Cl_2$/MeOH 10:1) of the residue yielded 4.0 g (98%) of 3-benzenesulphonyl-7-chloro-5-(4-methyl-piperazin-1-ylmethyl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine as a pale brown powder, m.p. 113-116°.

c) 10 ml of a 50% solution of NH_3 in MeOH were added to a solution of 0.8 g (1.77 mmol) of 3-benzenesulphonyl-7-chloro-5-(4-methyl-piperazin-1-ylmethyl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine in 20 ml of DMF and stirred at RT for 4 hrs. The reaction solution was evaporated and the residue was partitioned between 2N NaOH and CH_2Cl_2. The aqueous phase was extracted three times with CH_2Cl_2, and the combined organic phases were dried ($MgSO_4$), filtered and evaporated. Subsequent chromatography (SiO_2, CH_2Cl_2/ MeOH/NH4OH 90:10:1) yielded 0.59 g (77%) of 3-benzenesulphonyl-5-(4-methyl-piperazin-1-ylmethyl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ylamine as colourless crystals, m.p. 203-205°.

Example 119

3-Benzenesulphonyl-7-(4-methyl-piperazin-1-yl)-5-(4-methyl-piperazin-1-ylmethyl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine

0.4 g (4 mmol) of 1-methyl-piperazine was added to a solution of 0.8 g (1.77 mmol) of 3-benzenesulphonyl-7-chloro-5-(4-methyl-piperazin-1-ylmethyl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine in 20 ml of DMF and stirred at RT for 6 hrs. The reaction solution was evaporated and the residue was partitioned between 2N NaOH and CH_2Cl_2. The aqueous phase was extracted three times with CH_2Cl_2, and the combined organic phases were dried ($MgSO_4$), filtered and evaporated. Subsequent chromatography (SiO_2, CH_2Cl_2/MeOH/NH_4OH 80:10:1) and crystallization from EtOH yielded 0.75 g (82%) of 3-benzenesulphonyl-7-(4-methyl-piperazin-1-yl)-5-(4-methyl-piperazin-1-ylmethyl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p. 193-195°.

Example 120

3-Benzenesulphonyl-5-(4-methyl-piperazin-1-ylmethyl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ylamine

a) 1 ml (6 mmol) of morpholine was added to a solution of 2.0 g (5.40 mmol) of 3-benzenesulphonyl-5-chloromethyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ol in 20 ml of DMF and stirred at RT for 4 hrs. The reaction solution was evaporated and the residue was partitioned between 0.5N NaOH and CH_2Cl_2. The aqueous phase was extracted three times with CH_2Cl_2. The combined organic phases were dried ($MgSO_4$), filtered and evaporated. Chromatography (SiO_2, CH_2Cl_2/MeOH 10:1) yielded 2.0 g (88%) of 3-benzenesulphonyl-2-methylsulphanyl-5-morpholin-4-ylmethyl-pyrazolo[1,5-a]-pyrimidin-7-ol as a beige foam.

b) A suspension of 2.0 g (4.75 mmol) of 3-benzenesulphonyl-2-methylsulphanyl-5-morpholin-4-ylmethyl-pyrazolo[1,5-a]pyrimidin-7-ol in 30 ml of $POCl_3$ was heated at reflux for 3 hrs. The reaction solution was cooled to RT and evaporated. The residue was treated with 100 ml of ice-water and the pH value of the solution was adjusted to 8 with sat. $NaHCO_3$ solution. The aqueous phase was extracted three times with 70 ml of CH_2Cl_2 and the organic phases were dried ($MgSO_4$), filtered and evaporated. Chromatography (SiO_2, CH_2Cl_2/MeOH 10:1) of the residue yielded 1.8 g (86%) of 3-benzenesulphonyl-7-chloro-2-methylsulphanyl-5-morpholin-4-ylmethyl-pyrazolo[1,5-a]pyrimidine as a pale brown powder, m.p. 174-176°.

c) 10 ml of a 50% solution of NH_3 in MeOH were added to a solution of 0.9 g (2 mmol) of 3-benzenesulphonyl-7-chloro-2-methylsulphanyl-5-morpholin-4-ylmethyl-pyrazolo[1,5-a]pyrimidine in 20 ml of DMF and stirred at RT for 4 hrs. The DMF was evaporated in a high vacuum and the residue was partitioned between 2N NaOH and CH_2Cl_2. The aqueous phase was extracted three times with CH_2Cl_2 and the combined organic phases were dried ($MgSO_4$), filtered and evaporated. Subsequent chromatography (silica gel, CH_2Cl_2/MeOH/NH_4OH 80:10:1) and crystallization from EtOH yielded 0.70 g (83%) of 3-benzenesulphonyl-5-(4-methyl-piperazin-1-ylmethyl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ylamine as colourless crystals, m.p. 224-226°.

Example 121

3-Benzenesulphonyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-5-morpholin-4-ylmethyl-pyrazolo[1,5-a]pyrimidine

0.4 g (4 mmol) of 1-methyl-piperazine was added to a solution of 0.9 g (2 mmol) of 3-benzenesulphonyl-7-chloro-2-methylsulphanyl-5-morpholin-4-ylmethyl-pyrazolo[1,5-a]pyrimidine in 20 ml of DMF and stirred at RT for 4 hrs. The reaction solution was evaporated and the residue was partitioned between 2N NaOH and CH_2Cl_2. The aqueous phase was extracted three times with CH_2Cl_2 and the combined organic phases were dried ($MgSO_4$), filtered and evaporated. Subsequent chromatography (SiO_2, CH_2Cl_2/MeOH/NH_4OH 80:1:1) and crystallization from EtOH yielded 0.80 g (77%) of 3-benzene-sulphonyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-5-morpholin-4-ylmethyl-pyrazolo[1,5-a]pyrimidine as colourless crystals, m.p.199-201°.

Example 122

[2-(3-Benzenesulphonyl-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-yloxy)-ethyl]-dimethyl-amine

0.28 ml (2.38 mmol) of 2-dimethylaminoethanol and 3.68 g of Cs_2CO_3 were added to a solution of 0.8 g (2.26 mmol) of 3-benzenesulphonyl-7-chloro-5-methyl-2-methyl-sulphanyl-pyrazolo[1,5-a]pyrimidine in 40 ml of acetonitrile and the suspension was stirred at RT for 12 hrs. The reaction mixture was poured into semi-concentrated aqueous sodium chloride solution and extracted three times with ethyl acetate. The combined organic phases were dried ($MgSO_4$), filtered and evaporated. Subsequent chromatography yielded 0.65 g (70%) of 2-(3-benzenesulphonyl-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-yloxy)-ethyl]-dimethyl-amine as a pale yellow solid, m.p. 176-178°.

Example 123

3-Benzenesulphonyl-5-methyl-2-methylsulphanyl-7-(2-morpholin-4-yl-ethoxy)-pyrazolo[1,5-a]pyrimidine

In an analogous manner to that described in Example 122, from 3-benzenesulphonyl-7-chloro-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine and N-(2-hydroxy-ethyl)-

morpholine there was obtained 3-benzenesulphonyl-5-methyl-2-methylsulphanyl-7-(2-morpholin-4-yl-ethoxy)-pyrazolo[1,5-a]pyrimidine as a pale yellow solid.

Example 124

8-Benzenesulphonyl-7-methylsulphanyl-pyrazolo[1,5-a][1,3,5]triazin-4-ylamine

0.10 g (0.37 mmol) of 5-amino-3-methylthio-4-phenylsulphonyl-pyrazole was mixed with 0.25 g (2.55 mmol) of ethyl N-cyano-methanimate and stirred at 100°C for 16 hrs. The resulting pale beige paste was taken up in AcOEt/MeOH and treated in an ultrasound bath. The thus-obtained suspension was filtered. Chromatography (SiO_2, CH_2Cl_2/MeOH 95:5) yielded 0.052 g (44%) of 8-benzenesulphonyl-7-methylsulphanyl-pyrazolo[1,5-a][1,3,5]-triazin-4-ylamine as pale beige crystals, m.p. >280°C.

Example 125

(8-Benzenesulphonyl-2-methyl-7-methylsulphanyl-pyrazolo[1,5-a][1,3,5]triazin-4-yl)-methylamine

a) A solution of 0.54 g (2 mol) of 5-amino-3-methylthio-4-phenylsulphonyl-pyrazole and 0.54 g (3.4 mmol) of ethyl 1-ethoxy-ethylidene-carbamate in acetic acid was stirred at 100°C for 3 hrs. After cooling to RT the precipitate was filtered off, washed off with a copious amount of Et_2O and dried in a high vacuum at 45°C. There was obtained 0.41 g (61%) of 8-benzenesulphonyl-2-methyl-7-methylsulphanyl-3H-pyrazolo[1,5-a][1,3,5]-triazin-4-one as white crystals, m.p. >300°C.

b) A suspension of 0.36 g (1 mmol) of 8-benzenesulphonyl-2-methyl-7-methylsulphanyl-3H-pyrazolo[1,5-a][1,3,5]triazin-4-one in 20 ml of $POCl_3$ was treated with 0.12 ml (1.5 mmol) of pyridine and heated 110°C for 3 hrs. The reaction solution was cooled to RT and evaporated. The residue was dried azeotropically twice with 50 ml of toluene each time. The thus-obtained residue was taken in 10 ml of 2N methylamine in tetrahydrofuran and stirred at room temperature for 4 hrs. The reaction solution was evaporated and partitioned between H_2O and AcOEt. The organic phase was washed with H_2O and sat. NaCl solution, dried ($MgSO_4$), filtered and evaporated. Chromatography (SiO_2, AcOEt/hexane 1:1) yielded 0.28 g (80%) of (8-benzenesulphonyl-2-methyl-7-methyl-

sulphanyl-pyrazolo[1,5-a][1,3,5]triazin-4-yl)-methylamine as white crystals, m.p. 285° (dec).

Example 126

(8-Benzenesulphonyl-2-methyl-7-methylsulphanyl-pyrazolo[1,5-a][1,3,5]triazin-4-yl)-dimethylamine

In an analogous manner to that described in Example 123 b), from 8-benzenesulphonyl-2-methyl-7-methylsulphanyl-3H-pyrazolo[1,5-a][1,3,5]triazin-4-one and dimethylamine there was obtained (8-benzenesulphonyl-2-methyl-7-methylsulphanyl-pyrazolo[1,5-a]-[1,3,5]triazin-4-yl)-dimethylamine as pale pink coloured crystals, m.p. 228-230°.

Example 127

(8-Benzenesulphonyl-2-methyl-7-methylsulphanyl-pyrazolo[1,5-a][1,3,5]triazin-4-yl)-N',N'-dimethyl-propan-1,3-diamine

In an analogous manner to that described in Example 123 b), from 8-benzenesulphonyl-2-methyl-7-methylsulphanyl-3H-pyrazolo[1,5-a][1,3,5]triazin-4-one and 3-dimethylamino-1-propylamine there was obtained (8-benzenesulphonyl-2-methyl-7-methylsulphanyl-pyrazolo[1,5-a][1,3,5]triazin-4-yl)-N',N'-dimethyl-propan-1,3-diamine which was converted with HCl/diethyl ether into the corresponding hydrochloride (1:1.75), m.p. 249-250°.

Example 128

8-Benzenesulphonyl-2-methyl-4-(4-methylpiperazin-1-yl)-7-methylsulphanyl-pyrazolo-[1,5-a][1,3,5]triazine

In an analogous manner to that described in Example 123 b), from 8-benzenesulphonyl-2-methyl-7-methylsulphanyl-3H-pyrazolo[1,5-a][1,3,5]triazin-4-one and 1-methyl-piperazine there was obtained 8-benzenesulphonyl-2-methyl-4-(4-methyl–piperazin-1-yl)-7-methylsulphanyl-pyrazolo[1,5-a][1,3,5]triazine as yellow crystals, m.p. 169-171°.

Example 129

8-Benzenesulphonyl-2-methyl-4-(4-benzylpiperazin-1-yl)-7-methylsulphanyl-pyrazolo-[1,5-a][1,3,5]triazine

In an analogous manner to that described in Example 123 b), from 8-benzenesulphonyl-2-methyl-7-methylsulphanyl-3H-pyrazolo[1,5-a][1,3,5]triazin-4-one and 1-benzylpiperazine there was obtained 8-benzenesulphonyl-2-methyl-4-(4-benzylpiperazin-1-yl)-7-methyl-sulphanyl-pyrazolo[1,5-a][1,3,5]triazine as beige crystals, m.p. 190-192°.

Example A

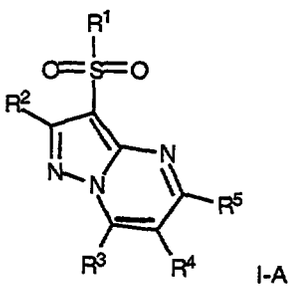
Tablets of the following composition are produced in the usual manner:

	mg/tablet
Active ingredient	100
Powd. lactose	95
White corn starch	35
Polyvinylpyrrolidone	8
Na carboxymethylstarch	10
Magnesium stearate	2
Tablet weight	250

* C L M S *

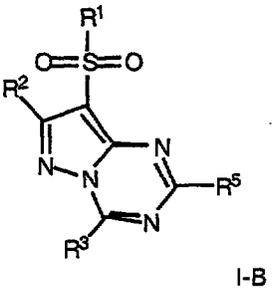
Claims

1. Compounds of the general formulae



I-A

and



I-B

wherein

R^1	signifies phenyl, optionally substituted by one or more lower alkyl, halogen, lower alkoxy, tolyl, pyridyl, naphthyl or thiophenyl;
R^2	signifies hydrogen, lower alkyl, lower thioalkyl or hydroxy-lower-alkoxy;
R^3	signifies amino, lower alkylamino, di-lower-alkyl-amino, piperazinyl, optionally substituted by one or more lower alkyl, benzyl, phenyl or hydroxy-lower-alkyl, morpholinyl, imidazolyl, $(CH_3)_2N(CH_2)_nNH$-, $(CH_3)_2N(CH_2)_nO$- or morpholinyl-$(CH_2)_n)O$- in which n signifies 2 or 3;
R^4	signifies hydrogen, lower alkyl or hydroxy-lower-alkyl;
R^5	signifies hydrogen, halogen, lower alkyl, C_3-C_6-cycloalkyl, lower alkyl-lower-alkoxy, hydroxy-lower-alkyl-lower-alkoxy, $(CH_3)_2N(CH_2)_nNH$-, piperazinyl, optionally substituted by lower alkyl, methyl-piperazinyl, optionally substituted by lower alkyl, morpholinyl, methyl-morpholinyl, di-lower-alkylamino or di-lower-alkylamino-lower-alkyl, or
R^4 and R^5	together signify a group $-(CH_2)_m-$ or $-CH_2-S-CH_2-$ with m = 3 or 4,

as well as their pharmaceutically acceptable salts.

2. Compounds of general formula I-A according to claim 1, wherein R^3 signifies amino.

* C L M S *

3. The following compounds in accordance with claim 2

3-benzenesulphonyl-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ylamine,

3-(4-isopropyl-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]-pyrimidin-7-ylamine,

5-methyl-2-methylsulphanyl-3-(naphthalene-2-sulphonyl)-pyrazolo[1,5-a]-pyrimidin-7-ylamine,

3-(4-fluoro-benzenesulphonyl)-5-methyl-2-methylsulphanyl-pyrazolo[1,5-a]-pyrimidin-7-ylamine,

3-benzenesulphonyl-5-cyclopropyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ylamine,

3-benzenesulphonyl-2-methylsulphanyl-6,7-dihydro-5H-cyclopenta[d]pyrazolo-[1,5-a]pyrimidin-8-ylamine,

3-benzenesulphonyl-5-isopropyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ylamine,

5-methyl-2-methylsulphanyl-3-(toluene-2-sulphonyl)-pyrazolo[1,5-a]pyrimidin-7-ylamine,

5-methyl-2-methylsulphanyl-3-(toluene-3-sulphonyl)-pyrazolo[1,5-a]pyrimidin-7-ylamine,

3-benzenesulphonyl-5-methoxymethyl-2-methylsulphanyl-pyrazolo[1,5-a]-pyrimidin-6-ylamine,

3-benzenesulphonyl-N5,N5-dimethyl-2-methylsulphanyl-pyrazolo[1,5-a]-pyrimidine-5,7-diamine,

3-benzenesulphonyl-N5-(2-dimethylamino-ethyl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine-5,7-diamine,

3-benzenesulphonyl-5-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ylamine and

3-benzenesulphonyl-5-dimethylaminomethyl-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidin-7-ylamine.

4. Compounds of general formula I-A in accordance with claim 1, wherein R^3 signifies piperazinyl.

5. The following compounds in accordance with claim 4

3-benzenesulphonyl-5-methyl-2-methylsulphanyl-7-piperazin-1-yl-pyrazolo[1,5-a]pyrimidine,

3-(4-tert-butyl-benzenesulphonyl)-5-methyl-2-methylsulphanyl-7-piperazin-1-yl-pyrazolo[1,5-a]pyrimidine,

3-benzenesulphonyl-5,6-dimethyl-2-methylsulphanyl-7-piperazin-1-yl-pyrazolo-[1,5-a]pyrimidine,

3-benzenesulphonyl-2-methylsulphanyl-7-piperazin-1-yl-5-propyl-pyrazolo[1,5-a]pyrimidine,

3-benzenesulphonyl-5-cyclopropyl-2-methylsulphanyl-7-piperazin-1-yl-pyrazolo-[1,5-a]pyrimidine,

3-benzenesulphonyl-2-methylsulphanyl-8-piperazin-1-yl-6,7-dihydro-5H-cyclo-penta[d]pyrazolo[1,5-a]pyrimidine,

3-benzenesulphonyl-2-methylsulphanyl-8-piperazin-1-yl-5H,7H-pyrazolo[1,5-a]-thieno[3,4-d]pyrimidine,

5-methyl-2-methylsulphanyl-7-piperazin-1-yl-3-(thiophene-2-sulphonyl)-pyrazolo[1,5-a]pyrimidine,

3-benzenesulphonyl-2-ethyl-8-piperazin-1-yl-6,7-dihydro-5H-cyclopenta[d]-pyrazolo[1,5-a]pyrimidine and

5-methyl-2-methylsulphanyl-7-piperazin-1-yl-3-(toluene-2-sulphonyl)-pyrazolo-[1,5-a]pyrimidine.

6. Compounds of general formula I-A in accordance with claim 1, wherein R^3 signifies methylpiperazinyl.

7. The following compounds in accordance with claim 6

3-benzenesulphonyl-5-cyclopropyl-2-methylsulphanyl-7-(4-methylpiperazin-1-yl)-pyrazolo[1,5-a]pyrimidine,

3-benzenesulphonyl-8-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-6,7-dihydro-5H-cyclopenta[d]pyrazolo[1,5-a]pyrimidine,

3-benzenesulphonyl-8-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-5H,7H-pyrazolo[1,5-a]thieno[3,4-d]pyrimidine,

3-benzenesulphonyl-5-isopropyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-pyrazolo[1,5-a]pyrimidine and

2-[3-benzenesulphonyl-7-(4-methyl-piperazin-1-yl)-2-methylsulphanyl-pyrazolo-[1,5-a]pyrimidine-5-yloxy]-ethanol.

8. Compounds of general formula I-B in accordance with claim 1, wherein R^3 signifies amino or methylpiperazinyl.

9. The following compounds in accordance with claim 8

8-benzenesulphonyl-7-methylsulphanyl-pyrazolo[1,5-a][1,3,5]triazin-4-ylamine and

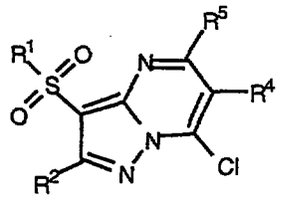
8-benzenesulphonyl-2-methyl-4-(4-methylpiperazin-1-yl)-7-methylsulphanyl-pyrazolo[1,5-a][1,3,5]triazine.

10. A medicament containing a compound according to any one of claims 1-9 and a therapeutically inert carrier material.

11. A medicament based on a compound of formula I-A or I-B according to any one of claims 1-9 and its pharmaceutically acceptable acid addition salts for the treatment or prevention of central nervous disorders such as psychoses, schizophrenia, manic depressions, depressions, neurological disorders, memory disorders, Parkinson's disease, amyotrophic lateral sclerosis, Alzheimer's disease and Huntington's chorea.

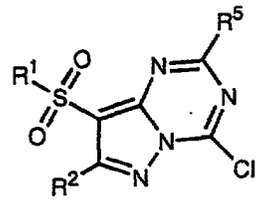
12. A process for the manufacture of compounds of formulae I-A and I-B according to claim 1, which process comprises

a) reacting a compound of the formula



X

or a compound of the formula



XIV

with a compound of the formula

HR^3

wherein R^1-R^5 have the significances given in claim,
and, if desired, converting a compound of general formula I-A or I-B into a pharmaceutically acceptable salt.

13. Compounds according to any one claims 1-9 insofar as they are manufactured according to the process defined in claim 12 or a process equivalent thereto.

14. Compounds according to any one of claims 1-9 for use as therapeutically active substances.

15. The use of compounds according to any one of claims 1-9 and of pharmaceutically useable salts thereof, especially for the treatment or prevention of central nervous disorders such as psychoses, schizophrenia, manic depressions, depressions, neurological disorders, memory disorders, Parkinson's disease, amyotrophic lateral sclerosis, Alzheimer's disease and Huntington's chorea or for the production of corresponding medicaments.

16. The invention as hereinbefore described.

This Sheet Has Been Replaced