A ONE-POT AND SELECTIVE PREPARATION OF 2-(N-ALKYL-4-CHLOROBUTYL-AMINO)-4-CHLOROQUINAZOLINES

Hideki Miki

Research Laboratories, Agricultural Chemicals Division, Takeda Chemical Industries, LTD., Juso-honmachi, Yodogawa-ku, Osaka, 532, Japan

<u>Abstract</u> — A new and facile synthesis of 2-(N-alkyl-4-chlorobutylamino)-4-chloroquinazolines by the reaction of 2,4(1H,3H)quinazolinedione with N-alkylpyrrolidines in phosphoryl chloride is described.

The chlorination of 2,4(1H,3H)-quinazolinedione (1) with phosphoryl chloride in the presence of triethylamine gave 4-chloro-2-diethylaminoquinazoline (3) instead of 2,4-dichloroquinazoline (2). On the other hand, when tripropylamine was used as a base in place of triethylamine, compound 1 was smoothly converted to $2^{(1)}$

To expand the studies of these reactions, N-alkylpyrrolidines (4) were used in place of triethyl- or tripropylamine as a base, and it was designed to convert the hydroxy group at 2-position in compound 1 to N-alkyl-4-chlorobutylamino group and to prepare 2-(N-alkyl-4-chlorobutylamino)-4-chloroquinazolines (5) in a one-pot selective preparation.

The reaction of 1 with phosphoryl chloride in the presence of N-methylpyrrolidine (4a) was carried out at 80-85°C for 20 min to give 4-chloro-2-(N-methyl-4chlorobutylamino)quinazoline (5a) as a yellow oil and 2 as colorless needles (mp 116-118°C). (Literature²); mp 118°C).

To make further investigation into this reaction, the study was expanded to the use of other N-alkylpyrrolidines such as N-ethylpyrrolidine (4b), N-propylpyrrolidine (4c), N-butylpyrrolidine (4d), N-sec-butylpyrrolidine (4e) and N-tertbutylpyrrolidine (4f). The results were summarized in Table I.

As shown in Table I, it has become apparent that the reaction of 1 with phosphoryl chloride in the presence of N-alkylpyrrolidines proceeds by a von Braun type reaction³ through the formation of a quaternary ammonium salt, which decomposes to give the new tertiary amines, 2-(N-alkyl-4-chlorobutylamino)-4-chloroquinazolines (5).

There are two possible reaction pathways to decompose the quaternary ammonium

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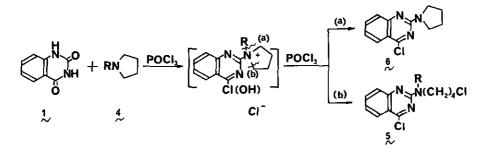
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R	4	2	5	
Me	4a.	3.3	(5a) 77.3	
Et	4b	17,9	(5b) 58 . 7	
Pr	4c	19.5	(5c) 59 . 4	
Bu	4d	48.5	(5d) 34.7	
sec-Bu	4e	84.6	(5e) trace	
tert-Bu	4f	88.5	(5f) trace	

Table I Yields (%) of Isolated Products from the Reaction of 1 with POCl3 in the Presence of 4.

salt, (a): the cleavage of N-alkyl group, and (b): the cleavage of pyrrolidine ring. On the results of these studies, the isolated product was mainly 5, 4-chloro-2-pyrrolidinoquinazoline (6) was not isolated.

When the bulky N-alkylpyrrolidine such as 4e or 4f was used as a base in the phosphoryl chloride chlorination of $\frac{1}{2}$, compound $\frac{2}{2}$ was mainly obtained. However, when the amine such as 4a, 4b, 4c or 4d was used, both of $\frac{2}{2}$ and $\frac{5}{2}$ were obtained.

It was found that the products ratio depended on the bulkiness of N-alkylpyrrolidines used. The introduction of N-alkyl-4-chlorobutylamino group to quinazoline ring has not been reported in the literature.



General procedure ---- N-Alkylpyrrolidine (15 ml) was added to a mixture of 1 (5.0 g, 0.031 mol) and phosphoryl chloride (70 ml) and the mixture was stirred at 80-85°C for 20 min. After the excess of phosphoryl chloride and N-alkylpyrrolidine were evaporated off <u>in vacuo</u>, the residue was disolved in 100 ml of chloroform. The chloroform layer was washed with water, satd. NaHCO3 aq., and then satd. NaCl aq. successively. After dring over MgSO4, the chloroform layer was concentrated to

give a yellow oil. The residue was subjected to the silica gel column chromatography. The elution with tetrachloromethane gave 2-(N-alkyl-4-chlorobutylamino)-4-chloroquinazoline 5 as a yellow oil and 2 as colorless needles. The results were summarized in Table II.

Compound No.	NZO	UV ^{EtOH} nm(E)	MS m/e(M ⁺)	PMR(8: ppm in CDCl ₃)
5a	1.6147	247(28700) 385(3300)	283,285,287.	1.57-1.92(4H,m,CH ₂),3.16(3H, s,CH ₃),3.37-3.19(4H,m,CH ₂), 6.85-7.95(4H,m,Ar-H).
5b	1.5987	247(31900) 384(3200)	297,299,301.	1.19(3H,t,CH ₃),1.59-1.98(4H, m,CH ₂),3.37-3.88(6H,m,CH ₂), 6.80-7.90(4H,m,Ar-H).
5c	1.5960	247(30600) 384(3100)	311,313,315.	0.91(3H,t,CH ₃),1.30-2.13(6H, m,CH ₂),3.34-3.72(6H,m,CH ₂), 6.80-7.90(4H,m,Ar-H).
5d	1.5862	247(29400) 384(3100)	325,327,329.	0.93(3H,t,CH3),1.40-2.15(8H, m,CH ₂),3.32-3.85(6H,m,CH ₂), 6.82-7.91(4H,m,Ar-H).

Table II 2-(N-Alkyl-4-chlorobutylamino)-4-chloroquinazoline

REFERENCES AND NOTES

1) H.C.Scarborough, B.C.Lawes, J.L.Minielli, and J.L.Compton, <u>J.Org.Chem</u>., 27, 957 (1962).

2) D.Libermann, and A.Rouaix, <u>Bull.Soc.Chim.France</u>, 1793(1959).

- 3) H.A.Hageman, Org.React., 7, 198(1953).
- 4) The structure of all new compounds was established from their analytical data of PMR, UV and MS spectra in addition to the elementalanalyses for C, H and N.

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