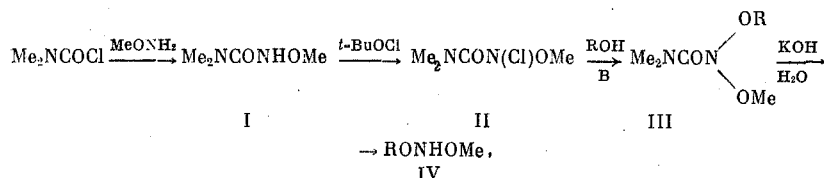


N,N-DIALKOXYUREAS AND NH-DIALKOXYAMINES

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For the first time we have obtained N,N-dialkoxyureas (III) and NH-dialkoxyamines (IV), particularly the simplest representative dimethoxyamine (IVa) (Table 1):



where B = RNa or 2,4,6-trimethylpyridine, and R = Me (a), Et (b), and i-Bu (c).

It was found that in II, as in N-chloro-N-alkoxy-N-tert-alkylamines [1, 2], nucleophilic substitution of the Cl on the N takes place smoothly. This is achieved by lowering the electron-acceptor effect of Me₂NCO with respect to N(Cl)OMe owing to the competitive, stronger amide conjugation. An increase in the electron-acceptor activity of the ROCO group causes the homolysis of the N-Cl bond to occur in N-chloro-N-alkoxyurethylanes when they are reacted with alcohols owing to the captodative stabilization of the aminyl radical [3]. N-Chloro-N-alkylamides, as we know, are chlorinating reagents owing to the effective stabilization of the N-anion by the electron-acceptor RCO group.

The structure of I-IV was confirmed by spectral data (see Table 1) and elemental analysis.

TABLE 1

Compound	Yield, %	Bp, °C (p, mm Hg)	PMR spectrum (60 MHz, in CCl ₄ relative to HMDS; δ, ppm, J, Hz)		
			Me ₂ N	MeO	other groups
(I)	81,3	91-92(1)	2,85	3,56	8,35 (NH)
(II)	~100	—	2,99	3,79	—
(IIIa)	84,5	61(1)	2,85	3,53	—
(IIIb)	59,7	75(2)	2,93	3,57	1,20, 3,86 (Et, J=6,8)
(IIIc)	50,7	97-98(2)	2,88	3,54	0,91 (Me ₂ CH, J=6,8), 1,8 (CH), 3,57 (CH ₂ CH, J=6)
(IVa)	61,5	83,5(760)	—	3,59	7,74 (NH)
(IVb)	41,7	95-97(760)	—	3,54	1,15, 3,79 (Et, J=6,8), 7,63 (NH)
(IVc)	44,4	95(160)	—	3,57	0,9 (Me ₂ CH, J=6,8), 1,86 (CH), 3,61 (CH ₂ CH, J=6), 7,67 (NH)

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