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Tertiary Carbamoylmethylphosphine Oxides and their Analogs

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Tertiary Carbamoylmethylphosphine Oxides and their Analogs

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Dialkyl(diaryl)carbamoylmethylphosphine oxides (I) were prepared by interaction of tervalent phosphorous acid esters with chloracetic amides:

$$\begin{array}{c}
\mathbb{R}^{\mathbf{I}} & \mathbb{P}(0) - \mathbb{C}\mathbb{H}_{\mathbf{Z}^{\mathbf{C}}}(0)\mathbb{N} \\
\mathbb{R}^{\mathbf{Z}}
\end{array} \tag{1}$$

 $R=R^{I}=n-Tol$, Ph, $c-C_{6}H_{II}$, Bu, Et, BuO; $R^{2}=R^{3}=Et$. $R=R^{I}=n-Tol$, Ph; $R^{2}=R^{3}=Bu$. R=Ph; $R^{I}=Bu$, BuO, $R^{2}=R^{3}=Et$, Bu. R=Ph, $R^{I}=Ph$, BuO; $R^{2}=Et$, $R^{3}=H$. R=Ph; $R^{I}=R^{2}=R^{3}=Et$. $R=R^{I}=R^{2}=R^{3}=Ph$.

The structure of (I) were established. Some of their chemical properties were investigated: Horner's reaction, formation of kalium salts, methylene group alkylation.

The structure of complexes (I) with perchloric and nitric acids were studied by potenciometric and thermometric titration methods and IR-spectra.

Complexation (I) and their analogs with Li^J were studied. The complexes of some compounds (I) with uranyl nitrate were synthesized. The structure of complexes (I) with PF₅ and TaF₅ were confirmed by means of the NMR ¹⁹F and ³¹P data.

Extraction and concentration by (I) of TPE were studied. (I), as it has been shown, are efficient extractants.

The work was carried out in collaboration with K.B.Yatsimirsky, E.I.Sinyavskaya (Kiev), Yu.A.Buslaev, E.G.Il'in (Moscow), B.F.Myasoedov, M.K.Chmutova (Moscow)