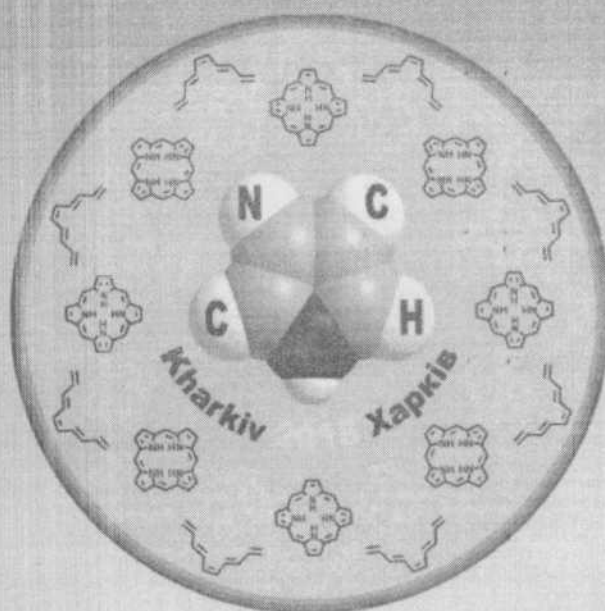


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Book of Abstracts

DECARBAMOYLATION OF N-ALKOXY-N-1-(4-DIMETHYLAMINOPYRIDINIUM)UREA'S CHLORIDES AS A ROUTE TO 1-N-ALKOXYAMINO-4-DIMETHYLAMINOPYRIDINIUM'S SALTS

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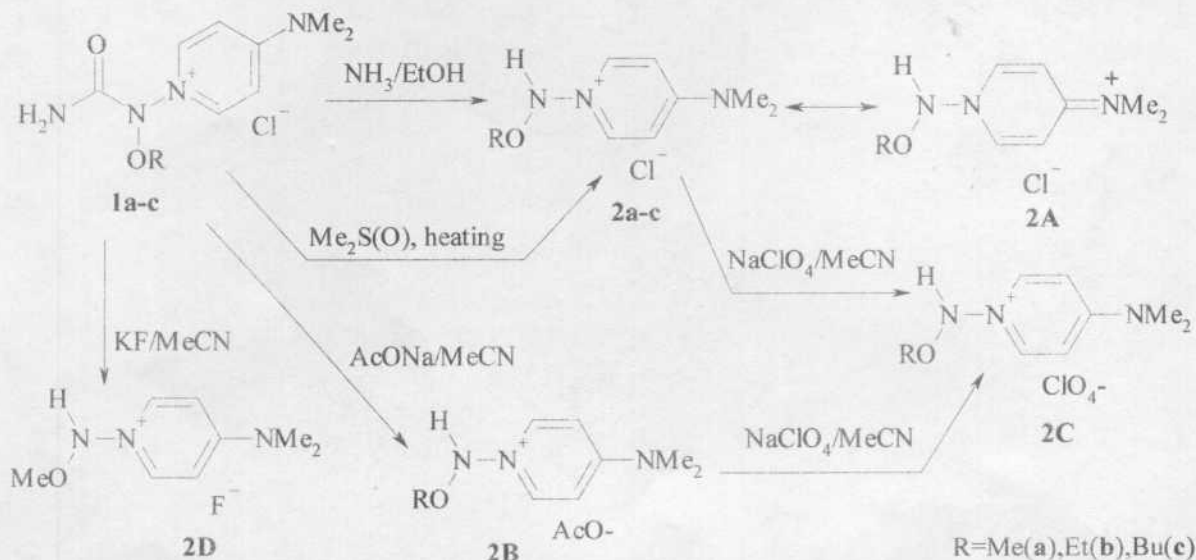
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1-N-Alkoxyamino-4-dimethylaminopyridium's salts **2a-c** can be easily synthesized from N-alkoxy-N-1-(4-dimethylaminopyridinium)urea's chlorides **1a-c** as by action of basic reagents (such as NH₃/EtOH, KF/MeCN or AcONa/MeCN) (route I), as by decarbamylation at heating in dimethylsulphoxide (route II). In the last case nature of N-alkoxy moiety takes great influence on the decarbamylation ease.



XRD study of compounds **2a** and **2b** revealed high degree of pyramidalicity of central nitrogen atom in O-N-N⁺ geminal system (the sum of bond angles is 312°). Lone pair of this nitrogen atom lies in the plane of pyridine moiety (the proper torsion angle is 4° (**2a**), 17° (**2b**)). It was found that quinonoid deformation of pyridine's ring (structure **2A**) had place. The positive charge is localized mainly on nitrogen atom of Me₂N-group.