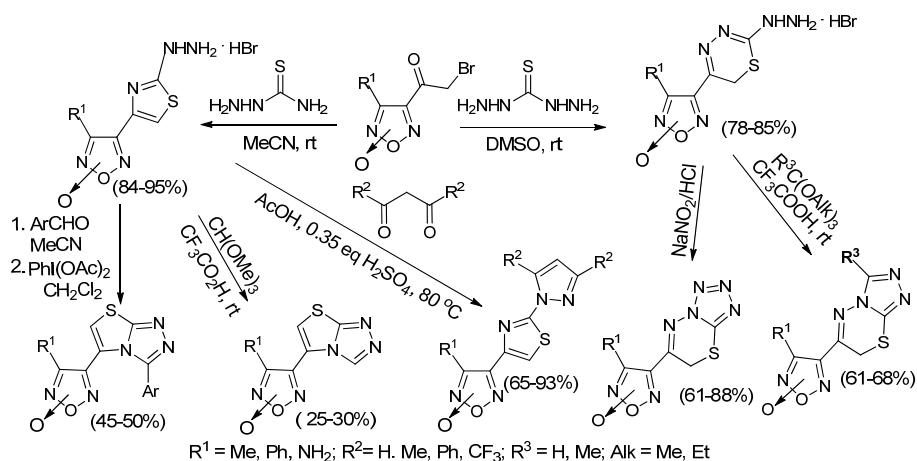


SYNTHESIS OF NEW PHARMACOLOGICALLY ORIENTED NO-DONOR FUROXAN-BASED HETEROCYCLIC ENSEMBLES

N. N. Makhova, L. L. Fershtat, M. A. Epishina, A. S. Kulikov, A. I. Churakov

N.D. Zelinsky Institute of Organic Chemistry, Russian Academy of Sciences
47 Leninsky prosp., 119991 Moscow, Russian Federation

The design of potential drugs with improved pharmacokinetic profile has been focused in recent years on the molecular hybridization of diverse compounds with known pharmacological activity [1]. Special efforts were directed to the synthesis of pharmacologically oriented structures comprising a framework capable of nitric oxide (NO) release, including 1,2,5-oxadiazole 2-oxides (furoxans) [2]. In this work simple, effective and regioselective methods for the synthesis of pharmacologically oriented polyheterocyclic ensembles containing furoxan motif as NO-donor fragment linked to various pharmacophoric nitrogen-containing heterocycles and their annulated derivatives (thiazoles [3], 1,3,4-thiadiazines [4], thiazolo[2,3-*c*][1,2,4]triazoles [3], 1,2,4-triazolo[3,4-*b*][1,3,4]thiadiazines [5], tetrazolo[5,1-*b*][1,3,4]thiadiazines [4], pyrazolylthiazoles [6]) have been presented.



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