



Synthesis of novel furo[3',2':5,6]pyrimido[2,1-c][1,2,4]triazines and furo[3,2-e][1,2,3,4]tetrazolo[1,5-a]pyrimidine

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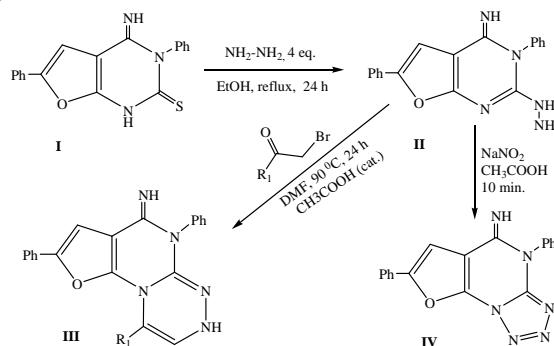
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Furo[2,3-*d*]pyrimidines have received much attention due to their biological activities. Antifungal, antibacterial, antiviral, antifolate, antitumor, and anti-HCMV (human cytomegalovirus) activities have been described for these compounds [1]. Recently, some furopyrimidines were shown to be potent LCK (lymphocyte-specific kinase) [2], PI3K (phosphoinositide 3-kinase) [3], VEGFR2 (vascular endothelial growth factor receptor2) and EGFR (epidermal growth factor receptor) inhibitors [4]. Furthermore, a wide range of biological activities has been attributed to fused triazoles, triazines and tetrazoles [5].

In connection with our interest in the synthesis of polyheterocyclic systems, we now report here the utility of 4-imino-3,6-diphenyl-3,4-dihydrofuro[2,3-*d*]pyrimidine-2(1*H*)-thione **I** for the synthesis of some novel tricyclic fused furo[3',2':5,6]pyrimido[2,1-*c*][1,2,4]triazines **III** and furo[3,2-*e*][1,2,3,4]tetrazolo[1,5-*a*]pyrimidine **IV** from α -haloketones and nitrous acid respectively in high yields.



References:

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