

Synthesis of Abnormal Acyclic Nucleosides in The Series of Triazoloazines

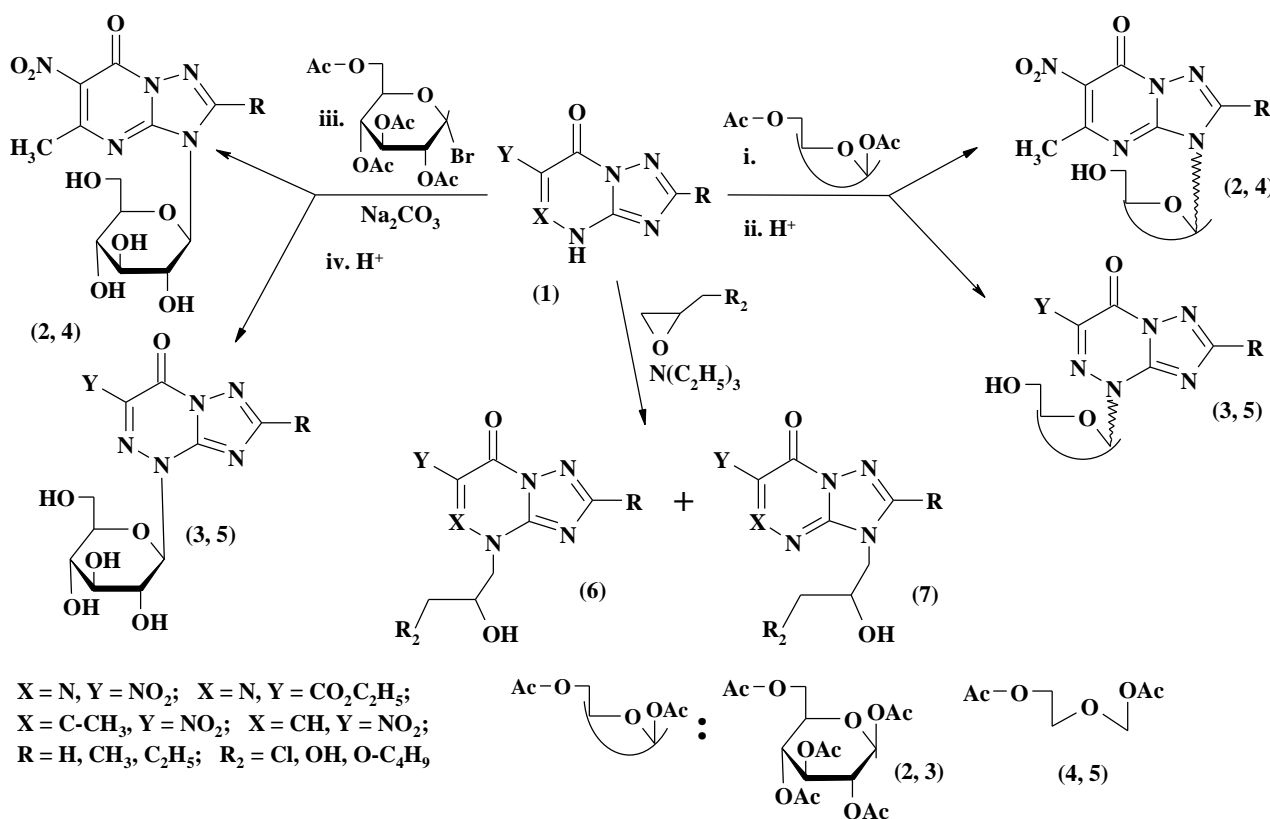
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Abnormal and acyclic nucleosides, derivatives of triazolo[1,5-a]pyrimidine and triazine-7-ones (1), which are structurally related to guanine have been obtained.

Acylated glucosides (2 and 3) have been obtained by two different ways. When heterocyclic bases were allowed to react with pentaacetyl glucose without of any catalyst the resulting nucleosides were obtained as mixtures of β - and α -anomers. The exclusive formation of β -glucosides can be reached by alkylation of triazoloazines with acetobromoglucose.

The synthesis of acyclovir analogues (4 and 5) has been performed by heating the diacetyl derivative of 2-oxabuta-1,4-diol with the corresponding NH-heterocycles. In order to remove the protecting acetyl groups all kind of acid catalysts proved to be the appropriate reagents.



Acyclic nucleosides (6,7) bearing the fragments of 3-chloropropan-2-ol and propane-1,2-diol were easily obtained by alkylation of NH-heterocycles with the corresponding in the presence of triethylamine. Sterical effects of substituents seem to be one of the main factors affecting the ratio of alkylation products (6,7).