

CYCLOADDITION REACTION OF A PYRIMIDINE-DIENOL

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During studies directed toward the development of new synthetic routes to fused pyrimidines employing 6-methyluracils, we have found that 5-acyl-1,3,6-trimethyluracil (I) causes base-catalyzed isomerization to the dienol(II) which is a reactive and versatile heterocyclic diene in the Diels-Alder reaction.

Thus, treatment of 5-formyl-1,3-dimethyluracil(I) ($R=H$) with lithium diisopropylamide(LDA) afforded a dimer (III) ($R=1,3,6$ -trimethyluracil-5-yl), which would be formed by the Diels-Alder reaction of the dienol(II) ($R=H$) and the aldehyde group of (I). The dimer was converted to a quinazoline (IV) ($R^1=H$, $R^2=CO_2Me$, $R^3=Me$) by refluxing in methanolic sodium hydroxide. 5-Acyl-1,3,6-trimethyluracil(I) ($R=H, Me, OEt$) reacted readily with dimethyl acetylenedicarboxylate in the presence of LDA yielding quinazolines (IV) ($R^1=H$, $R^2=R^3=CO_2Me$; $R^1=Me$, $R^2=R^3=CO_2Me$; $R^1=OH$, $R^2=R^3=CO_2Me$). Base-catalyzed cycloaddition of I ($R=H, Me$) with dimethyl maleate or dimethyl fumarate led to the formation of the 5,6,7,8-tetrahydroquinazoline (IV) ($R^1=H$, $R^2=R^3=CO_2Me$; $R^1=Me$, $R^2=R^3=CO_2Me$) with high stereoselectivity. In both the tetrahydroquinazolines, the relationship between the hydroxyl group at 5 position and the ester group at 6 position was determined to be *cis*. The adduct must be therefore formed by the Diels-Alder reaction of the (*E*)-dienol(II). Reaction of I ($R=H$) with other nucleophiles such as methyl vinyl ketone and *N*-phenylmaleimide afforded 7,8-dihydroquinazolines (IV) ($R^1=R^3=H$, $R^2=COMe$; $R^1=H$, $R^2=CONHPh$, $R^3=CO_2H$), respectively. Furthermore, the dienol (II) underwent cycloaddition reaction with aldehydes to yield pyrano[4,3-*d*]pyrimidines(III) ($R=H$, $Et, Ph, PhCH_2, 2$ -furfuryl), stereospecifically. Compound III ($R=Ph$) was epimerized to a 2:1 mixture with trifluoroacetic acid in DMSO and transformed to 1,3-dimethyl-6-(2-styryl)uracil(V) by refluxing in 10% HCl.

