PHOTOCHEMICAL ADDITION OF 6-CYANO-1,3-DIMETHYLURACIL TO SUBSTITUTED ACETYLENES AND OLEFINS

I<u>sao</u> S<u>aito</u>, K<u>oji</u> S<u>himozono</u> and T<u>eruo</u> M<u>atsuura</u> <u>Department of Synthetic Chemistry</u>, Faculty of <u>Engineering</u>, Kyoto <u>University</u> Sakyo-ku, Kyoto <u>606</u>, <u>Japan</u>

Photochemical reactions of 6-substituted 1,3-dimethyluracils in the presence of alkynes and alkenes have been investigated. Photolysis ($\lambda > 290$ nm) of 6-cyano-1,3-dimethyluracil (1) with diphenylacetylene or methylphenylacetylene in acetonitrile gave the corresponding cyclobutene-type adduct in a regiospecific fashion. A similar type of cycloaddition has been observed in the photoreaction of 6-carbomethoxy-1,3-dimethyluracil with methylphenylacetylene.

In contrast, photolysis of 1 in the presence of alkynes or alkenes gave rise to the formation of a novel 5-substituted uracil derivative. For example, irradiation of 1 with 1-hexyne in acetonitrile gave 5-(2-cyano-1-hexenyl)-1,3-dimethyluracil in high yield.

This novel photoreaction involves migration of cyano group and provides a convenient, useful method for forming carbon-carbon bonds at the C-5 position of uracil and uridine derivatives.