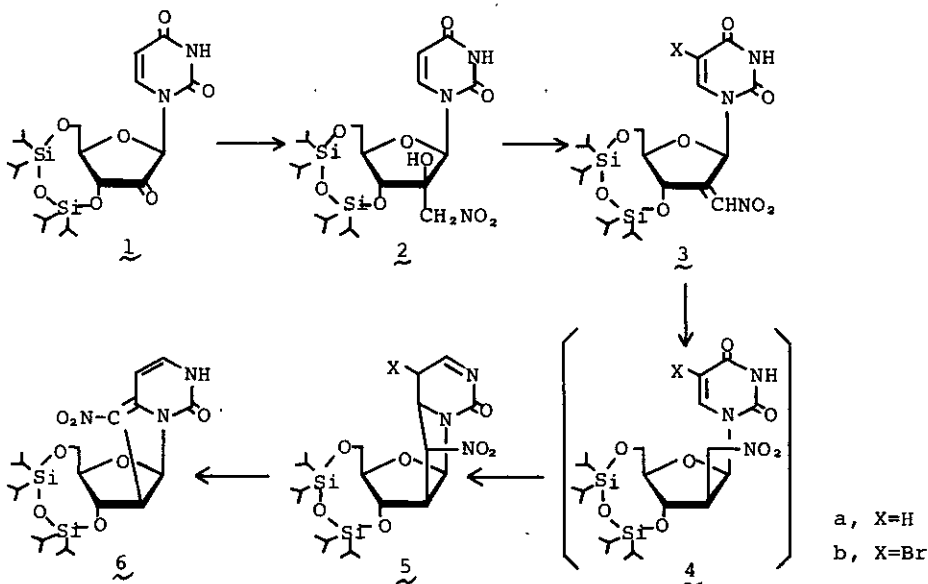


STRUCTURE OF A C-CYCLOPYRIMIDINE NUCLEOSIDE DERIVED FROM
A 2'-C-NITROMETHYLURIDINE

T. Ueda, S. Shuto and H. Inoue

Faculty of Pharmaceutical Sciences, Hokkaido University,
Sapporo 060, Japan

In the course of our studies on the transformation of nucleosides to carbon-branched sugar nucleosides, we found that 2'-carbon-branched uridine derivatives were easily prepared from protected 2'-ketouridine (1). For example, 6,2'-ethylene-cyclouridine was prepared from the Wittig product of 1 and ethoxycarbonylmethylene-triphenylphosphorane.¹ We envisioned that the synthesis of 6,2'-methylene-cyclouridines could be achieved by using a 2'-C-nitromethyluridine derivative (2) as the intermediate. For the preparation of 4, compound (3a) derived from 1 was allowed to react with NaBH₄, but the product obtained was 5,6-dihydro derivative (5a). To confirm its structure, 5-bromo derivative (3b) was treated with NaBH₄ in the same way. The product (5b) was then heated with DBU to afford a carbon-bridged 2-pyrimidinone cyclonucleoside (6).



1) T. Ueda, S. Shuto, T. Sano, H. Usui and H. Inoue, *Nucleic Acids Res. Symp. Series*, No. 11, 5 (1982).