

NOVEL C-NUCLEOSIDE SYNTHESSES

Haruo Ogura, Hiroshi Takahashi, and Masakazu Sakaguchi

School of Pharmaceutical Sciences, Kitasato University,

Shirokane, Minato-ku, Tokyo 108

Condensation of 5,6-diamino-1,3-dimethyluracil (1) with di-Q-benzylidene aldehydo-sugars (2a-c) and aldehydo-sugars afforded the corresponding Schiff base (3a-c, 5) in an excellent yield, and which were acetylated with acetic anhydride and pyridine to yield acetates (6). In the presence of NBS, the Schiff base (3a-c) and 6 were oxidized to form theophylline C-nucleosides (4a-c). On the other hand, on treatment of 6 with DDQ pteridine C-nucleosides (7) and furanose-type compounds (8), and the latter compounds were treated with ethyl orthoformate to yield pteridine C-nucleosides (9).

