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C-5 Substituted Pyrimidine Nucleosides, Synthesis of 5-Alkylglycoside Deoxyuridine and Uridine Analogs VIA Organopalladium Intermediates

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C-5 SUBSTITUTED PYRIMIDINE NUCLEOSIDES. SYNTHESIS OF 5-ALKYLGLYCOSIDE DEOXYURIDINE AND URIDINE ANALOGS VIA ORGANOPALLADIUM INTERMEDIATES

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In this communication, we report a chemical synthesis of 5-alkyl glycoside (β -D-2-acetamido-2-deoxyglucopyranoside and α -D-manno pyranoside) uridine derivatives. These compounds could have importance in molecular genetics as affinity reagents, since it is expected that these modified nucleosides and nucleotides have strong non-bonding interactions with lectins and thus could be selectively retained on such a column.

In a preliminary study, the glycosides $\underline{4}$ and $\underline{5}$ were prepared by condensation of the activated aglycon $\underline{1}$ with the oxazoline $\underline{2}$ and perbenzoylated bromomannose $\underline{3}$, respectively. The resulting glycosides $\underline{4}$ and $\underline{5}$ on reaction with organomercurial pyrimidine in presence of palladium as catalyst gave the corresponding 5-alkylglycoside deoxyuridine derivatives $\underline{6}$ and $\underline{7}$.

This reaction was extended to the synthesis of monophosphate (dUMP and UMP) and triphosphate (dUPT and UTP) derivatives.