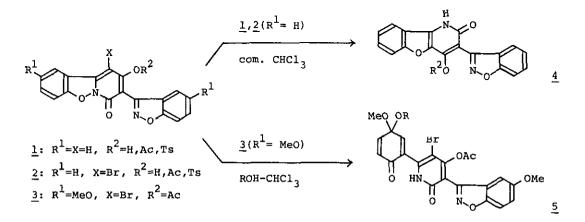
MECHANISMS OF PHOTO-INDUCED ISOMERIZATION OF 4-OXO-4H-BENZISOXAZOLO[2,3-a]PYRIDINE DERIVATIVES

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 $4-0xo-4\underline{H}$ -benzisoxazolo[2,3-<u>a</u>]pyridine derivatives <u>1</u>-<u>3</u> underwent novel photoinduced (a low pressure Hg-lamp or sun light) isomerizations to give benzofuro-[3,2-b]pyridines 4 or quinone ketal derivatives 5 in good to excellent yields.



A plausible mechanism for the ring transformation of $\underline{1}$ and $\underline{2}$ may involve the homolytic fission of the N₅-O₆ bond to give diradicals <u>6</u>, followed by isomerization and recyclization. In the case of the 9-methoxy derivative <u>3</u>, the first step may be the heterolytic cleavage of the N₅-O₆ bond to give a zwitter ion <u>7</u>, which is stabilized by both the methoxy and carbonyl groups; this stabilization effect is considered to enable the zwitter ion <u>7</u> to be formed, despite the fact that the electronegativity of oxygen is greater than that of nitrogen.

