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Chloroformylation of Ketophosphonates

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Chloroformylation of Ketophosphonates

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Recently we have developed a reaction pathway for a novel type of phosphonyl chlorovinylaldehydes using general procedure of chloroformylation[1].

This reaction led stereoselectively to the products with (Z)-configuration along with a small amount of (E)-isomers. Z or E isomers was separated by column chromatography, and assigned by the ³J_{HP} coupling constant measurement in

1H NMR spectra[2]. Moreover, when the isolation pure Z or E isomers (R = H) are refluxed in ethyl acetate for about 10 h respectively, the equilibration of the geometrical isomers (Z = E) takes place, and the ratio of Z/E (93/7) is almost the same in each case.

The reactions of phosphonylchlorovinylaldehyde with phenylhydrazine or oxyammonia gave phosphonyl pyrazoles or phosphonyl isoxazole respectively in presence of sodium hydride. Hydrazone and oxime (R=H) could be separated in absence of sodium hydride.

$$(C_{2}H_{5}O)_{2}P$$

$$(C_{1}H_{5}O)_{2}P$$

$$(C_{2}H_{5}O)_{2}P$$

$$(C_{3}H_{5}O)_{2}P$$

$$(C_{3}H_{5}O)_{2}P$$

$$(C_{3}H_{5}O)_{2}P$$

$$(C_{3}H_{5}O)_{2}P$$

$$(C_{3}H_{5}O)_{2}P$$

$$(C_{3}H_{5}O)_{2}P$$

$$(C_{3}H_{5}O)_{2}P$$

$$(C_{3}H_{5}O)_{2}P$$

$$(C_{3}H_{5}O)_{3}P$$

$$(C_{3}H$$

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