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Synthetic Applications of Phosphonium Diylides

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Abstract Phosphonium diylides 1 react with electrophiles leading to new monoylides which permit, by a Wittig reaction, the E stereoselective synthesis of various di- or tri-substituted α,β -unsaturated functionalized compounds.

Recently we have shown that phosphonium divlides 1, allowing the formation of various α,β -unsaturated functions, can be consider as very good tools in organic synthesis [1].

We report here a generalization of the method which permit, via the in situ formation of a new intermediate monoylide 2, the synthesis of α,β -unsaturated amides, amidines, esters, acids, phosphines, phosphonates and phosphine oxides.

In the case or R = H, alkyl, and when R^2R^3CO is an aldehyde, the products are isolated in good yields (60-90%) and excellent E stereoselectivity. When R = Ph, COPh or in presence of ketones, the yields are often lower or the reaction leads to the direct formation of the E alkene $R^2R^3C=CHR$.

REFERENCE

1. H. J. Cristau, Chem Rev 94, 1299 (1994) and references cited therein.