POSTER PRESENTATION

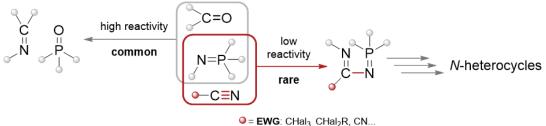
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α-EWG-CONTROLLED AZA-WITTIG REACTION WITH NITRILES

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The *aza*-Wittig reaction is a well-known rapid access to C=N bond formation. This process provides numerous opportunities for the synthesis of N-containing organic molecules. [1]. Among various *aza*-Wittig reactions, the interactions between *aza*-Wittig reagents, phosphazenes, and compounds containing polar double bonds (primarily in the carbonyl group) are the most studied. Oppositely, reactions involving the triple C=X bonds of acetylenes or nitriles are described scarcely. However, this challenging type of *aza*-Wittig reactions could allow for installing an additional nitrogen atom in the assembled molecules and, thus, synthesizing various heterocyclic systems with at least two nitrogen atoms.



 \mathbf{V} = **EWG**: CHal₃, CHal₂R, CN...

Here, we report an *aza*-Wittig reaction wherein the efficient control of the competition between the C=N and C=O groups provides for chemoselective assembly of N-heterocycles. α -EWGs were found to control chemoselectivity and, depending on their nature, act as CN group activators (*e.g.*, ester, amide or nitrile) or competitors (*e.g.*, ketone) in *aza*-Wittig reactions. To demonstrate the synthetic utility of the obtained iminophosphazenes **2** as *N*,*N*-binucleophiles, their transformations into pyrrole-fused systems **3**, pyrrolo[1,2-a]imidazoles and pyrrolo[1,2-a][1,3]diazepines, were carried out [2].



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