Lipase-Catalyzed Alkoxycarbonylation of Alcohols

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Protection of functional groups are very important in multistep organic synthesis. The alkoxycarbonyl group has been used as a protecting group for alcohols because carbonates are generally more stable against a wide range of nucleophiles under basic conditions than esters.

Recently, enzymatic protecting techniques have increased importance in promoting green sustainable chemistry. We have previously developed the lipase-catalyzed *tert*-butoxycarbonylation of primary alcohols using Boc₂O.¹ In this research, we have investigated the scope of lipase-catalyzed alkoxycarbonylation of alcohols.

We explored the use of various reagents for alkoxycarbonylation of alcohols. For example, the reaction of benzyl alcohol with allyl phenyl carbonate was carried out in the presence of lipases such as *Aspergillus niger* lipase, *Pseudomonas fluorescens* lipase, *Candida rugosa* lipase, and *Burkholderia cepacia* lipase in hexane at 40 °C for 24 h (Scheme 1). From these results, we confirmed that *Pseudomonas fluorescens* lipase, *Candida rugosa* lipase, and *Burkholderia cepacia* lipase were suitable catalysts for the allyloxycarbonylation of benzyl alcohol (Table 1, entries 3–5).



Scheme 1.

Entry	Lipase	Yield (%) ^a	
1	No lipase	-	
2	Aspergillus niger lipase	6	
3	Pseudomonas fluorescens lipase	95	
4	Candida rugosa lipase	99	
5	Burkholderia cepacia lipase	99	

Table 1. Reaction of benzyl alcohol with allyl phenyl carbonate in the presence of lipase.

a) Determined by gas chromatography.

1) N. Kishi, H. Kojima, ChemistrySelect 2019, 4, 9570-9572.