## SYNTHESIS OF DIARYLS FROM PHENYLBORIC ACID AND

ARYL IODIDES IN AN AQUEOUS MEDIUM

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The cross coupling of aryl halides is a convenient method for the synthesis of diaryls with virtually any functional groups. The reaction is usually carried out in a system containing benzene and aqueous alkali in the presence of 3 mole  $Pd(PPh_3)_4$  at reflux.

We have shown that the reaction of phenylboric acid with aryl iodides such as m-, p-, and  $o-IC_6H_4CO_2H$  and  $p-IC_6H_4OH$  dissolved in water in the presence of base proceeds in the absence of an organic solvent.  $Pd(OAc)_2$  may be used as the catalyst and sodium carbonate may be used as the base. The yields of asymmetrical diaryls vary from 70 to 95%

 $\operatorname{ArI} + \operatorname{PhB}(\operatorname{OH})_2 \frac{\operatorname{Pd}(\operatorname{OAc})_2, \operatorname{1mole} \%}{\operatorname{Na_2CC_3/H_2O, 20^\circ}} \operatorname{ArPh} 70-95\%$ 

 $ArI = m - IC_6H_4COOH; p - IC_6H_4COOH; o - IC_6H_4COOH; p - IC_6H_4OH.$ 

A mixture of 0.318 g (3 mmoles)  $Na_2CO_3$ , 0.248 g (1 mmole) m-IC<sub>6</sub>H<sub>4</sub>CO<sub>2</sub>H, 0.134 g (1.1 mmoles) PhB(OH)<sub>2</sub>, and 0.0022 g (0.01 mmole) Pd(OAc)<sub>2</sub> was dissolved in 4 ml water in an argon atmosphere and stirred for 90 min at ~20°C. Palladium black was filtered off. The filtrate was acidified by the addition of hydrochloric acid. The precipitate formed was filtered off, washed with water, and dried to give 0.185 g (93%) m-phenylbenzoic acid, mp 165-166°C [3].

## LITERATURE CITED

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