## **Cross-Coupling Reactions of Organoboranes:**

An Easy Way for Carbon-Carbon Bonding

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## Conjugated Alkadienes

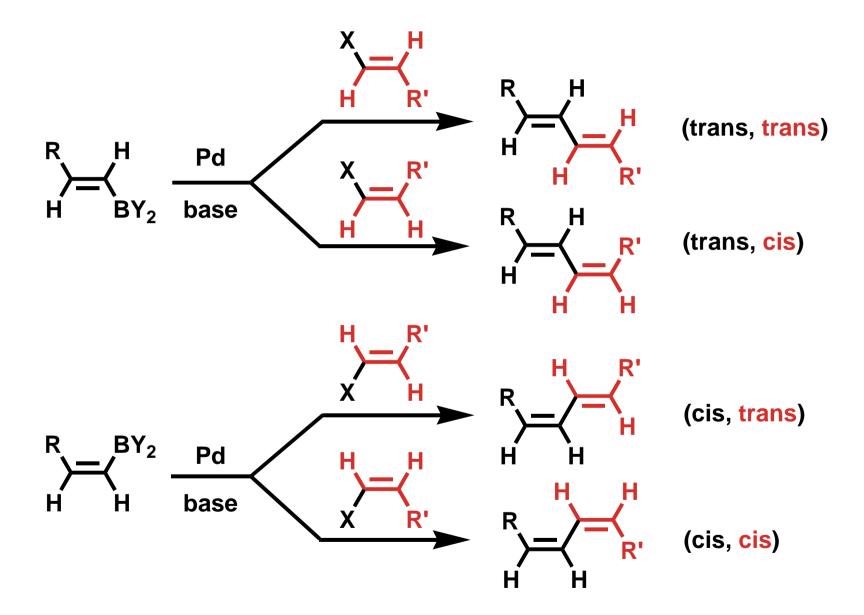
**M**: transition metal catalyst

#### Syntheses of (E)- and (Z)-1-Alkenylboranes

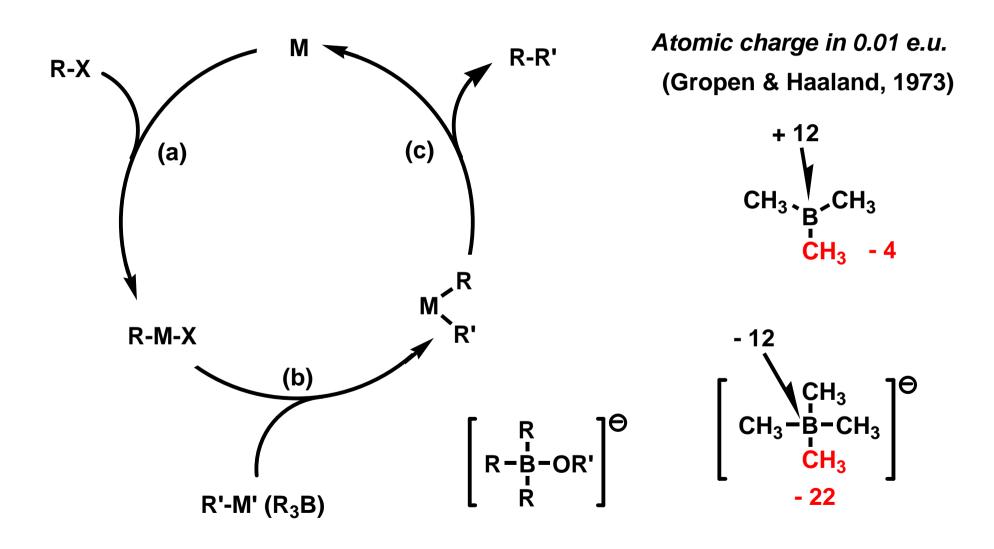
RC
$$\equiv$$
CH + HBY<sub>2</sub>  $\longrightarrow$   $\stackrel{R}{\longrightarrow}$   $\stackrel{H}{\longrightarrow}$  BY<sub>2</sub> trans > 99 % Y<sub>2</sub> = (Siamyl)<sub>2</sub>,

RC
$$\equiv$$
CX  $\xrightarrow{HBY_2}$   $\xrightarrow{R}$   $\xrightarrow{K}$   $\xrightarrow{H}$   $\xrightarrow{H}$   $\xrightarrow{H}$   $\xrightarrow{H}$   $\xrightarrow{R}$   $\xrightarrow{R}$   $\xrightarrow{BY_2}$   $\xrightarrow{R}$   $\xrightarrow{H}$   $\xrightarrow{R}$   $\xrightarrow{H}$   $\xrightarrow{R}$   $\xrightarrow{R}$   $\xrightarrow{H}$   $\xrightarrow{H}$   $\xrightarrow{R}$   $\xrightarrow{R}$ 

X = I or Br Y = Siamyl, Cyclohexyl



Common Catalytic Cycle Involving Sequential Oxidative Addition (a), Transmetalation (b), and Reductive Elimination (c)



$$Bu \nearrow_{BX_2} + Br \nearrow_{Ph} \longrightarrow Bu \nearrow_{Ph}$$

| <b>1</b> <sup>a)</sup> | Catalyst <sup>b)</sup><br>(mol %) | Base<br>(equiv / 2) | Solvent | React.<br>time (h) | Yield (%)<br>of 3 |
|------------------------|-----------------------------------|---------------------|---------|--------------------|-------------------|
| 1b                     | PdL <sub>4</sub> (3)              | None                | THF     | 6                  | 0                 |
| 1b                     | PdL <sub>4</sub> (3)              | None                | Benzene | 6                  | 0                 |
| 1a                     | PdL <sub>4</sub> (3)              | 2M NaOEt (2)-EtOH   | THF     | 2                  | 73                |
| 1b                     | PdL <sub>4</sub> (3)              | 2M NaOEt (2)-EtOH   | THF     | 4                  | 78                |
| 1b                     | PdL <sub>4</sub> (1)              | 2M NaOEt (2)-EtOH   | Benzene | 2                  | 86                |

a) 1a, 
$$X_2 = (Sia)_2$$
 1b,  $X_2 = \begin{pmatrix} 0 \\ 0 \end{pmatrix}$  b) L = PPh<sub>3</sub>

| 1-Alkenylborane  | 1-Alkenyl Bromide | Product | Yield (%)<br>[Purity (%)] |
|------------------|-------------------|---------|---------------------------|
| <b>Bu</b>        | Br Ph             | Bu      | 86 [98]                   |
| <b>Bu</b> _B< a) | Br Ph             | Bu      | <u>49</u> [99]            |
| <b>Bu</b> B< a)  | Br Ph             | BuPh    | <u>42</u> [89]            |
| <b>Bu</b>        | Br Hex            | Bu      | 88 [99]                   |
| <b>Bu</b> B< a)  | BrHex             | BuHex   | <u>49</u> [98]            |
| <b>Ph</b>        | Br Ph             | Ph Ph   | 89 [98]                   |

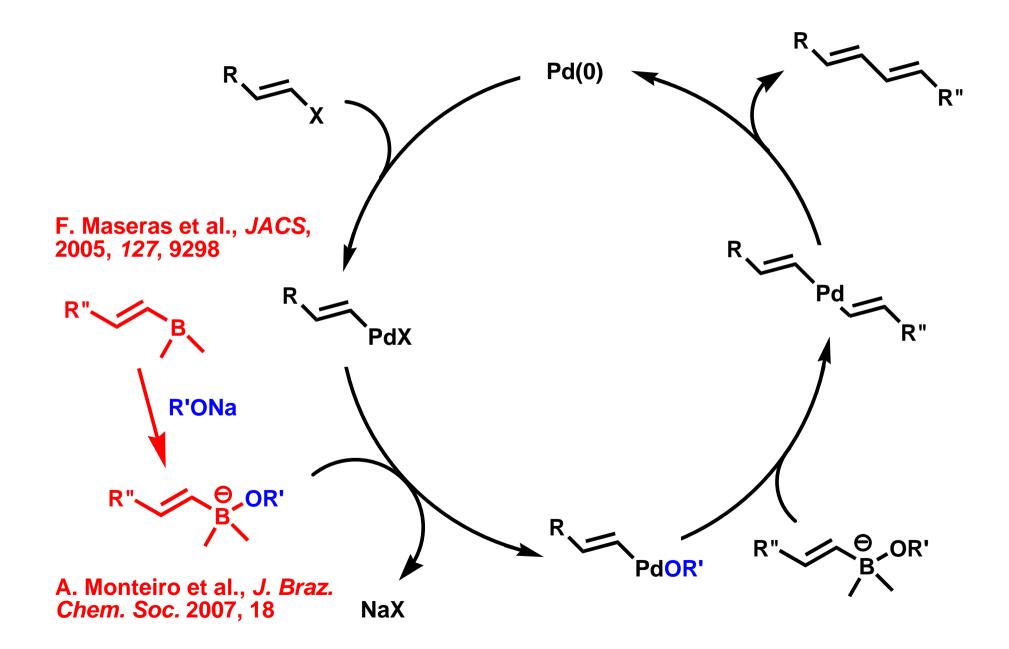
Reaction conditions: 1-3 mol % of Pd(PPh $_3$ ) $_4$  / NaOEt / Benzene / Reflux 2h a) Disiamyl b) 1,3,2-Benzodioxaboryl

| BY <sub>2</sub>  | RX     | Product | Yield (%) | Purity (%) |
|--|--------|---------|-----------|------------|
| B(Sia) <sub>2</sub><br>B(OPr <sup>i</sup> ) <sub>2</sub> | Br Hex | BuHex   | 49<br>87  | >98<br>>99 |
| B(Sia) <sub>2</sub>                                      |        |         | 58        | >94        |
| $B(\bigcirc)_2$  | PhI    | Bu Ph   | 49        | >83        |
| B(OPr <sup>i</sup> ) <sub>2</sub>                        |        |         | 98        | >97        |
| B(Sia) <sub>2</sub>                                      |        | Bu /    | 54        | >92        |
| B(OPr <sup>i</sup> ) <sub>2</sub>                        | · >=/  |         | 87        | >99        |

"Palytoxin" C<sub>129</sub>H<sub>223</sub>N<sub>3</sub>O<sub>54</sub> (MW. 2678.6)

Synthesis: Kishi et al., *J. Am. Chem. Soc*, 1989, 111, 7525, 7530

#### Reaction Mechanism:



### Reaction of B-Alkylboranes

$$R-B$$
 +  $R^4X$   $\longrightarrow$   $R-R^4$ 

R: Alkyl

#### Alkyl-Vinyl Coupling:

**Total Synthesis of Polycyclic Ether Natural Product** 

M. Sasaki, *Bull. Chem. Soc. Jpn.* 2007, *80*, 856

$$R^{1}O \xrightarrow{Q-BBN} R^{1}O \xrightarrow{Q-B$$

#### Polycyclic Ether Marine Natural Products:

### Aromatic-Aromatic Cross-Coupling Reactions

$$-B(OH)_2 + Br - Z$$

## Suzuki Coupling:

#### Valsartan (Novartis): Antihypertensive

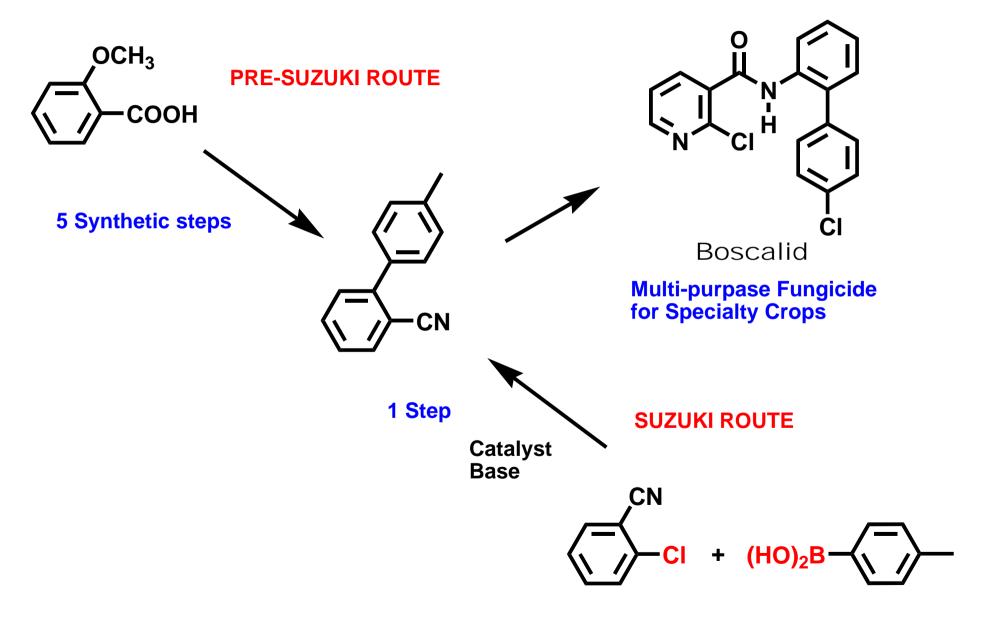
- 3.5 million users in Japan
- 22 million users in the whole world

Angiotensin II Receptor Antagonist (Losartan)

Merck , J. Org. Chem. 59, 6391 (1994)

Losartan (Antihypertensive)

# Suzuki coupling is a shortcut to biaryls (BASF's Boscalid Process)



## Boscalid; Agrochemicals (BASF, Germany)

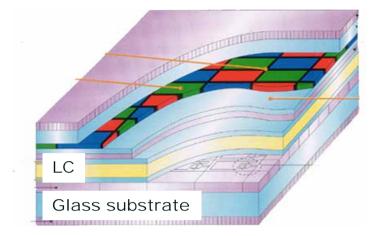


Boscalid

## Liquid crystal:

Chisso (Japan)

$$C_5H_{11}$$
  $\longrightarrow$   $B(OH)_2 + I$   $F$ 



$$C_5H_{11}$$
  $F$ 

Merck (Germany)

$$R - C$$
 $F$ 
 $F$ 
 $F$ 
 $OCF_3$ 

## EL Polymer materials

## Advantages of the Cross-Coupling Reaction between Organoboron Compounds and Organic Electrophiles:

- 1. Ready availability of reagents: hydroboration and transmetalation
- 2. Mild reaction conditions: base problem
- 3. Water stability
- 4. Easy use of the reaction both in aqueous and heterogeneous conditions
- 5. Toleration of a broad range of functional groups
- 6. High regio- and stereoselectivity of the reaction
- 7. Insignificant effect of the steric hindrance
- 8. Use of a small amount of catalysts
- 9. Application in one-pot synthesis
- 10. Nontoxic reaction
- 11. Easy separation of inorganic boron compounds