## Category

Synthesis of Heterocycles

## Key words

oxazolines

oxazines

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palladium catalysis

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A General and Efficient Palladium-Catalyzed Carbonylative Synthesis of 2-Aryloxazolines and 2-Aryloxazines from Aryl Bromides

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## Palladium-Catalyzed Three-Component Synthesis of Oxazolines and Oxazines

$$Ar - Br + CO + CI \xrightarrow{NH_2 \cdot HCI} \frac{Pd(OAc)_2 \ (2 \text{ mol\%})}{PhMe, 110 \ ^\circ C, 16 \ h} Ar \xrightarrow{N} 27 \text{ examples } 50-89\% \text{ yield}$$
 
$$(10 \text{ bar}) \qquad (1 \text{ equiv}) \qquad (Ad = \text{adamantyI})$$
 
$$Ar = Ph, \text{ Tol}, 2 \cdot \dot{\ell} \text{ PrC}_6 H_4, 4 \cdot \text{PhC}_6 H_4, 4 \cdot \text{F}_3 \text{CC}_6 H_4, 4 \cdot \text{NCC}_6 H_4, 4 \cdot \text{OHCC}_6 H_4, 4 \cdot \text{CHC}_6 H_4, 4 \cdot \text{Me}(O) \text{CC}_6 H_4, 4 \cdot \dot{\ell} \text{BuO}_2 \text{CC}_6 H_4, 4 \cdot \dot{\ell} \text{Pr}_2 \text{N}(O) \text{CC}_6 H_4, 4 \cdot \text{CHC}_6 H_4, 4 \cdot \text{CHC}_6 H_4, 4 \cdot \text{Me}_2 \text{NC}_6 H_4, PMP, 3 \cdot \text{4-MeSC}_6 H_4, 1 \cdot \text{Naph}, 2 \cdot \text{Naph}, 6 \cdot \text{MeONaph-}2 \cdot \text{yl}, 2 \cdot \text{thiophenyI}, 3 \cdot \text{thiophenyI}, 3 \cdot \text{benzo}[b] \text{thiophenyI}, 3 \cdot \text{py}, 1 \cdot \text{methyI-}1 \cdot H \cdot \text{indole-}5 \cdot \text{yl}, 3 \cdot \text{quinolinyI}, 6 \cdot \text{quinolinyI}, 7 \cdot \text{quinoxalinyI}$$
 
$$Ar - Br + CO + CI \qquad NH_2 \cdot HCI \qquad Pd(OAc)_2 \ (2 \text{ mol\%}) \\ BuPAd_2 \ (6 \text{ mol\%}) \\ NEt_3 \ (3 \text{ equiv}) \qquad PhMe, 110 \ ^\circ \text{C}, 16 \ h} \qquad Ar \rightarrow N \qquad 11 \text{ examples} \\ 60 - 89\% \text{ yield}$$
 
$$Ar = Ph, 4 \cdot \text{NCC}_6 H_4, 4 \cdot \text{Me}(O) \text{CC}_6 H_4, 4 \cdot \text{Me}_2 \text{NC}_6 H_4, PMP, 4 \cdot \text{MeSC}_6 H_4, 1 \cdot \text{Naph}, 3 \cdot \text{thiophenyI}, 3 \cdot \text{benzo}[b] \text{thiophenyI}, 3 \cdot \text{Py}, 1 \cdot \text{methyI-}1 \cdot H \cdot \text{indole-}5 \cdot \text{yI}$$

Significance: Described is the synthesis of oxazolines via a three-component process, in which readily available aryl bromides, carbon monoxide and 2-chloroethylamine undergo a palladium-catalyzed carbonylation and a subsequent cyclization to afford 2-aryloxazolines in good yield. Replacing 2-chloroethylamine with 3-chloropropylamine also works well and in this case the corresponding 2-aryloxazine derivatives are formed. Notably, both electron-donating and electron-withdrawing groups are tolerated in this process. However, the scope of the three-component process was not well investigated, especially for *ortho-* and *meta*-substituted aryl bromides.

Comment: Oxazolines and oxazoles are important heterocycles for organic synthesis and materials chemistry (see Book below). A number of methodologies for the construction of the oxazoline ring from aryl aldehydes, nitriles, carboxylic acids and related derivatives have been developed (e.g., S. Takahashi, H. Togo *Synthesis* 2009, 2329). In comparison with those, the present carbonylation–cyclization strategy offers a straightforward way for the synthesis of oxazolines and their analogues from easily available starting materials. A drawback of this method is the high pressure required (10 bar).

**Book:** Oxazoles: synthesis, reactions, and spectroscopy, Part 2; D. C. Palmer, Ed.; Wiley: Hoboken, **2004**.

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