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Rh(III)-catalyzed C–H olefination of *N*-pentafluoroaryl benzamides using air as the sole oxidant†

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Accepted 5th January 2015

DOI: 10.1039/c4sc03350g

www.rsc.org/chemicalscience

The oxidative olefination of a broad array of arenes and heteroarenes with a variety of activated and unactivated olefins has been achieved via a rhodium(III)-catalyzed C–H activation reaction. The use of an *N*-pentafluorophenyl benzamide directing group is crucial for achieving catalytic turnovers in the presence of air as the sole oxidant without using a co-oxidant.

Introduction

The direct coupling of unactivated aryl C–H bonds with olefins provides a step- and atom-economical method for the functionalization of arenes.¹ While a number of transition metals have been explored in this capacity, Rh(III) has emerged as an effective catalyst for the olefination of aryl C–H bonds under mild reaction conditions.^{2,3} Many research groups have sought to improve the utility of Rh(III)-catalyzed aryl C–H olefinations by increasing reactivity of the catalytic systems and improving site selectivity of the transformation. This has been achieved via the use of proximal directing groups to promote *ortho*-C–H bond cleavage. However, in the previously reported Rh(III)-catalyzed aryl C–H olefination reactions, stoichiometric oxidants such as peroxides, hypervalent iodonium salts, fluorinating agents, NXS (X = F, Cl, Br, or I) or inorganic salts are generally required to sustain the catalytic cycle.⁴ The use of any of these oxidants results in the undesired generation of stoichiometric reaction byproducts, reducing the utility and practicality of these transformations. An alternative approach has been the use of oxidizing directing groups CONHOR, which generally feature a cleavable N–O bond that can serve to reoxidize the Rh(I) catalyst.⁵ A greener and more environmentally benign approach is the use of molecular oxygen (O₂) or air as the terminal oxidant, which generates water as the sole reaction byproduct.⁶ The goal of using O₂ as the sole oxidant for C–H

olefination reactions has been achieved in several cases using palladium.⁷ More recently Huang and coworkers reported the Rh(III)-catalyzed C–H activation reactions using O₂ as the sole oxidant in alkynylation reactions for the synthesis of isoquinolinium salts⁸ and indoles⁹ and the direct C-2 olefination of electron-rich indoles.¹⁰ However, to date, we are unaware of any other instances of a Rh(III)-catalyzed C–H functionalization reaction of inert arenes using air as the sole oxidant without using a metal co-oxidant.^{3a}

While the use of molecular O₂ as oxidant is fundamentally important, the use of air as the sole oxidant for Rh(III)/Rh(I) catalysis is practically desirable in terms of operational safety. Herein we report a Rh(III)-catalyzed C–H activation/olefination reaction of aryl and heteroaryl benzamides under air at atmospheric pressure without the addition of an external oxidant. This reaction utilizes an *N*-pentafluorophenyl benzamide directing group, which has not previously been explored as a directing group for Rh(III)-catalyzed reactions. The transformation is compatible with a range of olefin coupling partners, including simple styrenes (Scheme 1).

Our group has recently described the effectiveness of *N*-polyfluoroaryl benzamide directing groups in Pd(II)-catalyzed C(sp³)-H activation/olefination reactions.¹¹ We hoped to extrapolate the use of this directing group to Rh(III)-catalyzed C–H olefination reactions. As shown in Table 1, treatment of pentafluorobenzamide **1a** and ethyl acrylate with 5 mol% [RhCp*Cl₂]₂ in the presence of AgOAc or Cu(OAc)₂ as a

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† Electronic supplementary information (ESI) available: Data for new compounds and experimental procedures. CCDC 1042327. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c4sc03350g



Scheme 1 Aerobic Rh(III)-catalyzed C–H olefination.



substituted substrates, which bear a handle for further chemical manipulation, are also tolerated in this transformation (Table 2, entries 2i–n). It is worth noting that the *meta*-substituted benzamides (Table 2, entries 2n and 2o) provide a regioisomeric mixture of olefination products. For *meta*-fluorinated substrate, the major product in most cases with Pd(OAc)₂-catalyzed C–H activation occurs *para* to F with one exceptional example.¹⁴ Interestingly, *meta*-fluorinated substrate affords the *ortho* position olefination product 2n, while *meta*-Me substrate affords the *para* position olefination product 2o, illustrating that the acidity of the C–H bond plays a role in this C–H olefination reaction. The reaction can be conducted on gram scale with no decrease in isolated yield (Table 2, entry 2l). The catalyst loading can also be lowered to 2 mol% [RhCp*Cl₂]₂ without significantly affecting the reactivity (Table 3).

In addition, we were pleased to find that heterocyclic amides, including thiophene, furan, and pyridine are competent substrates and can be olefinated in good yield (Table 3, entry 2p–2r) with just a slight increase in reaction temperature. Interestingly, although the *N*-aryl thiophene-2-carboxamide reacts to provide the cyclized product 2p in 81% yield (reaction time is increased to 48 hours), the *N*-aryl furan-2-carboxamide provides only the uncyclized product 2q (75% yield) even with higher reaction temperatures and an extended reaction time. While olefination of pyridine-4-carboxamide gave the desired product 2r in excellent yield, pyridine-3-carboxamide is less reactive, affording lower yield (Table 2, entry 2s, 48%). 4-(Pyridine-2-yl)benzamide was also subjected to the standard conditions and the olefination occurred *ortho* to the amide rather than the pyridyl, albeit affording low yield (28%, see ESI†). Not surprisingly, pyridine-2-carboxamide is not reactive under these

Table 4 Scope of olefin coupling partner^{a,b}

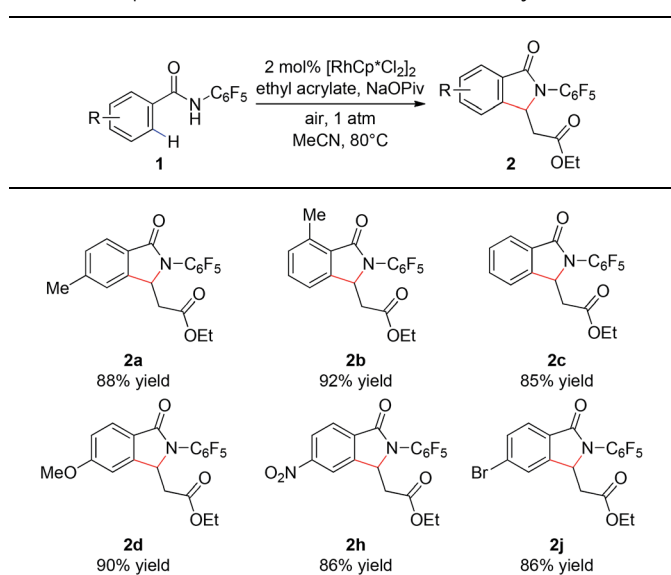
^a Reaction conditions: benzamide (0.2 mmol, 1.0 eq.), olefins (0.5 mmol, 2.5 eq.), [RhCp*Cl₂]₂ (0.01 mmol, 0.05 eq.), NaOPiv (0.2 mmol, 1.0 eq.), MeCN (2 mL), air, 1 atm, 80 °C, 24 h. ^b Isolated yield.

conditions due to the bis-dentate coordination of the substrate with Rh(III).

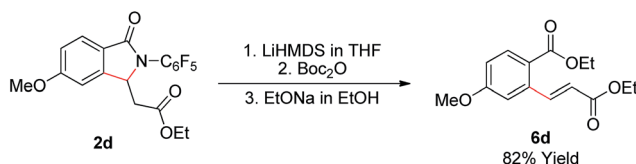
We subsequently analyzed the scope of the olefin coupling partner in this transformation. As shown in Table 4, electron-deficient olefins, including ethyl vinyl ketone and acrylonitrile, can be coupled in good yield to provide lactam products 4a–b. A variety of styrenes substituents are also coupled in good yields to provide the corresponding uncyclized products 5a–5c. In addition, the unactivated olefin pentene can also be coupled, albeit in modest yield, to provide amide 5d. However, di-substituted olefins are not reactive under these conditions.

As shown in Scheme 2, the γ -lactam products formed in this transformation are readily converted to the olefinated products. Treatment of lactam 2d with LiHMDS, Boc₂O, and EtONa step by step in one-pot provides enoate 6d in good yield.

Our proposed mechanism for this transformation is shown in Scheme 3. Thus coordination of the amide to the [Rh(III)] catalyst is followed by *ortho*-C–H bond activation to give a corresponding [Rh(III)–Ar] intermediate. Subsequent coordination to the olefin coupling partner and 1,2-migratory insertion provides a complex which can undergo β -hydride elimination to provide the uncyclized product 6. Reoxidation of the [Rh(I)] to [Rh(III)] by molecular oxygen would complete the catalytic cycle.

Table 3 Scope of benzamide substrate with 2% catalyst^{a,b}

^a Reaction conditions: benzamide (0.2 mmol, 1.0 eq.), ethyl acrylate (0.5 mmol, 2.5 eq.), [RhCp*Cl₂]₂ (0.004 mmol, 0.02 eq.), NaOPiv (0.2 mmol, 1.0 eq.), MeCN (2 mL), air, 1 atm, 80 °C, 48 h. ^b Isolated yield.



Scheme 2 Removal of the auxiliary.





Scheme 3 Proposed reaction mechanism.

This reoxidation step is likely facilitated by the acetate base, addition of which provides a substantial increase in reaction conversion. Subsequent base mediated 1,4-conjugate addition of the acidic *N*-pentafluorophenyl benzamide onto the pendant enoate provides the γ -lactam product **2**. The isolation of furan **2q** and alkenes **5a–5d**, which have undergone olefination but not cyclization, provide some evidence for the formation of **6** during the catalytic cycle.

In summary, we have developed a Rh(III)-catalyzed C–H olefination of aryl C–H bonds using the *N*-pentafluorophenyl amide auxiliary and air as the sole oxidant. The reaction conditions can be applied to a variety of both aryl and heteroaryl benzamides.

Acknowledgements

We gratefully acknowledge the National Natural Science Foundation of China (grant no. 21201100 and 21331002) and NSF under the CCI Center for Selective C–H Functionalization, CHE-1205646 for financial support. This work was also supported by a Project Funded by the Priority Academic Program Development of Jiangsu Higher Education Institutions.

Notes and references

‡ General procedure: to a 350 mL Schlenk-type sealed tube equipped with a magnetic stirring bar, were added the substrate (0.2 mmol, 1.0 eq.), [RhCp*Cl₂]₂ (0.01 mmol, 0.05 eq.), NaOPiv (0.2 mmol, 1.0 eq.), MeCN (2.0 mL) and olefin coupling partner (0.5 mmol, 2.5 eq.). The tube was capped and heated to 80 °C for 24 h. After cooling to room temperature, the reaction mixture was filtered through a pad of Celite. The filtrate was concentrated *in vacuo* to afford the crude product, which was purified by flash column chromatography (SiO₂) gel to provide the desired product.

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