

PAPER

View Article Online
View Journal | View IssueCite this: *RSC Adv.*, 2018, 8, 2267Received 6th December 2017
Accepted 29th December 2017

DOI: 10.1039/c7ra13080e

rsc.li/rsc-advances

Synthesis of benzoxazoles *via* an iron-catalyzed domino C–N/C–O cross-coupling reaction†

Bo Yang, Weiye Hu and Songlin Zhang *

An eco-friendly and efficient method has been developed for the synthesis of 2-arylbenzoxazoles *via* a domino iron-catalyzed C–N/C–O cross-coupling reaction. Some of the issues typically encountered during the synthesis of 2-arylbenzoxazoles in the presence of palladium and copper catalysts, including poor substrate scope and long reaction times have been addressed using this newly developed iron-catalyzed method.

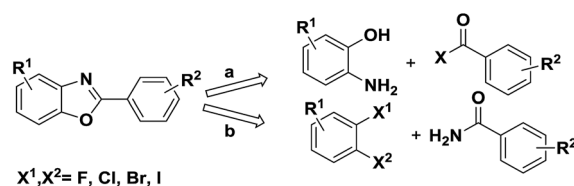
2-Arylbenzoxazoles are an important class of structures in natural products, and pharmaceuticals and has shown a wide range of biological activities, such as antitumor, antiviral, and antimicrobial activities.¹ In particular, they show a marvellous efficacy in the treatment of duchenne muscular dystrophy (DMD) which is one of the most common of the muscular dystrophies that is caused by a mutation in the gene DMD, located in humans on the X chromosome (Xp21).² So the synthesis of 2-arylbenzoxazoles has been intensively studied for use in organic and medicinal chemistry over the past few years.

Numerous methods have been reported to synthesise this motif, one of the common methods is transition-metal-catalyzed (like Pd,³ Ni,⁴ Cu,⁵ Mn⁶ *etc.*) cross-coupling from pre-existing benzoxazoles with aryl halide or arylboronic acid. And another method is the classic one employing a cyclocondensation approach between an aminophenol and either a carboxylic acid⁷ or benzaldehyde⁸ (Scheme 1, path a). In 2004, Frank Glorius' group reported a domino copper-catalyzed C–N and C–O cross-coupling for the conversion of primary amides into benzoxazoles⁹ (Scheme 1, path b) which is a new reaction type for the synthesis of benzoxazoles. Bunch *et al.* apply this domino reaction in the synthesis of planar heterocycles in 2014.¹⁰ In addition the cyclization of *o*-halobenzenamides to benzoxazoles has been reported several times.^{11,12} Nevertheless, some limitations in the reported methods need to be overcome, such as the use of palladium complexes and narrow substrate range.

In the last few years, there has been a significant increase in the number of reports pertaining to the development of iron-catalyzed reactions in organic synthesis, where iron has shown several significant advantages over other metals, such as

being more abundant, commercially inexpensive, environmentally friendly and drug safety.¹³ Compared with palladium and copper, the use of iron is particularly suitable for reactions involving the preparation of therapeutic agents for human consumption. With this in mind, it was envisaged that a new method should be developed for the synthesis of benzoxazoles *via* an iron-catalyzed domino C–N/C–O cross-coupling reaction.

The reaction of benzamide (**1a**) with 1-bromo-2-iodobenzene was used as model transformation to identify the optimum reaction conditions by screening a variety of different iron salts, bases, ligands and solvents (Table 1). Several iron salts were screened in this reaction, including FeCl₃, FeCl₂·4H₂O, FeSO₄·7H₂O, Fe(acac)₃, Fe₂O₃, Fe₃O₄, Fe₃O₄(nano), Fe₂O₃(nano), Fe₂(SO₄)₃ and Fe(NO₃)₃·9H₂O, Fe₂O₃ was found to give the best results with the desired product **3a** being formed in a yield of 15% while most of the iron salt have no effect on the reaction (Table 1, entries 1–10). Then, several other bases, including LiO^tBu, Na₂CO₃, NaOAc, KOH and K₂CO₃ were also evaluated under the same conditions using Fe₂O₃, but all of them failed to provided the desired product **3a** except K₂CO₃ with a yield of 37% (Table 1, entries 11–15). When the reaction was stirred for 24 h at 110 °C in the presence of 20% mol of Fe₂O₃, 20% mol *N,N'*-dimethylethanedi-amine (DMEDA) and 1 equiv. of K₂CO₃ in PhMe under nitrogen, (*N*-(2-bromophenyl)benzamide) was obtained as an intermediate which could be converted to the final product with a yield of 87% if extend the reaction time from 24 h to 48 h (Table 1, entries 15, 16). Several ligands were also screened in the model reaction, and the results revealed that the



Scheme 1 Classic method of benzoxazole formation.

Key Laboratory of Organic Synthesis of Jiangsu Province, College of Chemistry, Chemical Engineering and Materials Science, Soochow University, Suzhou 215123, P. R. China. E-mail: zhangsl@suda.edu.cn

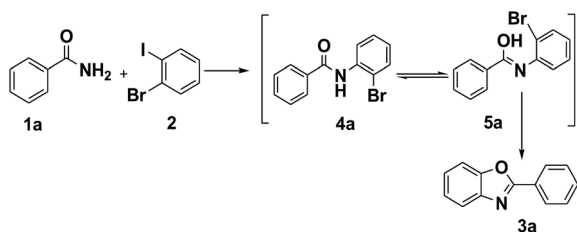
† Electronic supplementary information (ESI) available. See DOI: 10.1039/c7ra13080e

Table 1 Optimization of reaction conditions^a

Entry	Iron salt	Ligand	Base	Solvent	Y ^b (%)
1	FeCl ₃	DMEDA	KO ^t Bu	PhMe	Trace
2	FeCl ₂ ·4H ₂ O	DMEDA	KO ^t Bu	PhMe	Trace
3	FeSO ₄ ·7H ₂ O	DMEDA	KO ^t Bu	PhMe	0
4	Fe(acac) ₃	DMEDA	KO ^t Bu	PhMe	0
5	Fe ₂ O ₃	DMEDA	KO ^t Bu	PhMe	15
6	Fe ₃ O ₄	DMEDA	KO ^t Bu	PhMe	0
7	Fe ₃ O ₄ (nano)	DMEDA	KO ^t Bu	PhMe	10
8	Fe ₂ O ₃ (nano)	DMEDA	KO ^t Bu	PhMe	0
9	Fe ₂ (SO ₄) ₃	DMEDA	KO ^t Bu	PhMe	0
10	Fe(NO ₃) ₃ ·9H ₂ O	DMEDA	KO ^t Bu	PhMe	0
11	Fe ₂ O ₃	DMEDA	LiO ^t Bu	PhMe	0
12	Fe ₂ O ₃	DMEDA	Na ₂ CO ₃	PhMe	0
13	Fe ₂ O ₃	DMEDA	NaOAc	PhMe	0
14	Fe ₂ O ₃	DMEDA	KOH	PhMe	0
15	Fe ₂ O ₃	DMEDA	K ₂ CO ₃ (24 h)	PhMe	37
16	Fe ₂ O ₃	DMEDA	K ₂ CO ₃ (48 h)	PhMe	87
17	Fe ₂ O ₃	Phen	K ₂ CO ₃	PhMe	Trace
18	Fe ₂ O ₃	L-Proline	K ₂ CO ₃	PhMe	0
19	Fe ₂ O ₃	Dpy	K ₂ CO ₃	PhMe	0
20	Fe ₂ O ₃	DMEDA	K ₂ CO ₃	DMSO	0
21	Fe ₂ O ₃	DMEDA	K ₂ CO ₃	DMF	0
22	Fe ₂ O ₃	DMEDA	K ₂ CO ₃	PhMe ₂	0
23	—	DMEDA	K ₂ CO ₃	PhMe	0
24	Fe ₂ O ₃	DMEDA	K ₂ CO ₃	PhMe	86 ^c
25	Fe ₂ O ₃	DMEDA	K ₂ CO ₃	PhMe	58 ^d

^a Reaction conditions: benzamides (0.5 mmol), 1-bromo-2-iodobenzene (1.5 eq.), iron salt (20% mol), base (1 eq.), ligand (20%) were added to a solvent (2 mL) and react at 110 °C for 48 h under N₂. ^b Isolated yield based on **1a** after silica gel chromatography. ^c Fe₂O₃ and K₂CO₃ were applied in purity of 99.999% from Alfa. ^d with Fe₂O₃ in a dosage of 10 mmol%.

nature of the ligand has a dramatic impact on the yield of the reaction. For example, the use of DMEDA gave 2-phenylbenzo[d]oxazole in 85% yield, whereas 1,10-phenanthroline, dipyriddy and L-proline provided no product (Table 1, entries 16–19). The reaction was conducted in DMSO, DMF and PhMe₂ respectively and none of them provided a much higher of the desired product than toluene (Table 1, entries 20–22). Control experiments was taken in the absence of Fe₂O₃, no product was obtained (Table 1, entry 23). In view of the fact that the trace



Scheme 2 The pathway of the reaction.

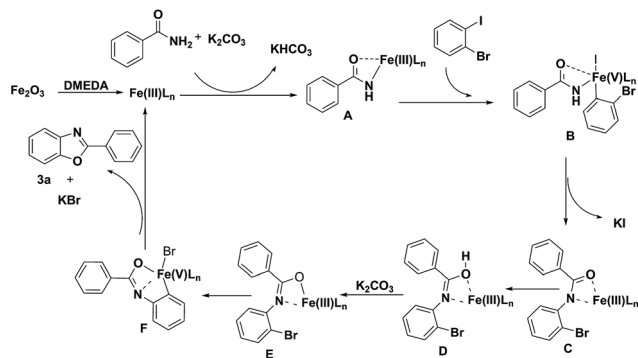
metals in catalytic, as is well-known, sometime could play an important role in the reaction,¹⁴ high-purity Fe₂O₃ (99.999%) and K₂CO₃ (99.999%) were applied in the reaction (Table 1, entry 24). The product was formed in a yield of 86% which was similar with the one of the entry 16.

Table 2 Reagent scope of reaction^a

R ¹		R ²		Y (%)
X=Br	R ¹ =H	R ² =H	Y=87%	3a
X=Br	R ¹ =H	R ² =Cl	Y=75%	3b
X=I	R ¹ =H	R ² =Cl	Y=62%	3b
X=Br	R ¹ =H	R ² =Me	Y=80%	3c
X=Br	R ¹ =F	R ² =H	Y=58%	3d
X=Br	R ¹ =F	R ² =Me	Y=92%	3e
X=Br	R ¹ =F	R ² =Cl	Y=85%	3f
X=Br	R ¹ =Me	R ² =H	Y=70%	3g
X=Br	R ¹ =Me	R ² =Cl	Y=65%	3h
X=Br	R ¹ =Me	R ² =Me	Y=45%	3i
X=Br	R ¹ =OMe	R ² =H	Y=50%	3j
X=Br	R ¹ =OMe	R ² =F	Y=54%	3n
X=Br	R ¹ =OMe	R ² =Cl	Y=35%	3o
X=Br	R ¹ =OMe	R ² =H	Y=0%	3p
X=Br	R ¹ =NO ₂	R ² =H	Y=0%	3q

^a Reaction conditions: **1a** (0.5 mmol), *o*-dihalo substrate (1.5 eq.), Fe₂O₃ (20% mol), K₂CO₃ (1 eq.), DMEDA (20%) were added to PhMe (2 mL) and react at 110 °C for 48 h under N₂.





Scheme 3 Possible catalytic cycle.

At last, the dosage of Fe_2O_3 was reduced to 10 mmol%, but only 58% yield was obtained (Table 1, entry 26). Taken together, the results of these screening experiments revealed that the optimal conditions for the reaction were Fe_2O_3 (20 mol%), DMEDA (20 mol%) and K_2CO_3 (1 eq.) in toluene at 110 °C for 48 h.

It is noteworthy that the intermediate product **4a** was formed under the optimized conditions *via* the C–N cross coupling reaction of benzamide (**1a**) with 1-bromo-2-iodobenzene (**2**). So a possible pathway of the reaction was proposed as shown in Scheme 2.

With the optimized reaction conditions in hand, we proceeded to investigate the substrate scope of the reaction using a variety of different 1,2-dihalobenzene substrates and aryl formamide (Table 2). Benzamide containing electron poor (**3d–f**, **3l**), electron-neutral (**3a–c**, **3k**), and electron-rich (**3g–j**, **3m–n**) substituents were all obtained in moderate to excellent yields. But some functional groups are intolerated in the reaction, like amino (**3p**) and nitro (**3q**).

Based on the results observed in the current study and Goldberg reaction,¹⁵ we have proposed a reaction mechanism for this transformation, which is shown in Scheme 3. The initial transmetalation of benzamide with $\text{Fe}_2\text{O}_3\text{L}_n$ in the presence of K_2CO_3 would give rise to the iron(III) species **A**. Complex **A** would then undergo an oxidative addition reaction with 1-bromo-2-iodobenzene to give the iron(V) species **B**, which would undergo a reductive elimination reaction to give iron(III) species **C** with the concomitant formation of a C–N bond. Followed the tautomerism of intermediate **C** to **D**, the intermediate iron(III) species **E** was formed in the presence of K_2CO_3 , which would undergo another oxidative addition reaction to afford iron(V) species **F**. Compound **3a** would then be obtained *via* a reductive elimination reaction from iron(V) species **F**.

In summary, we have demonstrated that the cheap and environmental friendly catalyst system composed of Fe_2O_3 and ligand DMEDA is highly effective for the synthesis of 2-aryl-benzoxazoles. The new catalyzed system can be effective for both C–N coupling and C–O coupling.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

We gratefully acknowledge A Project Funded by the Priority Academic Program Development of Jiangsu Higher Education Institutions, The Project of Scientific and Technologic Infrastructure of Suzhou (No. SZS201207) and the National Natural Science Foundation of China (No. 21072143) for financial support.

Notes and references

- (a) S. Tzanopoulou, M. Sagnou, M. Paravatou-Petsotas, E. Gourni, G. Loudos, S. Xanthopoulos, D. Lafkas, H. Kiaris, A. Varvarigou, I. C. Pirmettis, M. Papadopoulos and M. Pelecanou, *J. Med. Chem.*, 2010, **53**, 4633–4641; (b) M. C. V. Zandt, M. L. Jones, D. E. Gunn, L. S. Geraci, J. H. Jones, D. R. Sawicki, J. Sredy, J. L. Jacot, A. T. DiCioccio, T. Petrove, A. Mitschler and A. D. Podjarny, *J. Med. Chem.*, 2005, **48**, 3141–3152; (c) L. Katz, *J. Am. Chem. Soc.*, 1953, **75**, 712–714; (d) D. K. Dalvie, A. S. Kalgutkar, S. C. Khojasteh-Bakht, R. S. Obach and J. P. O'Donnell, *Chem. Res. Toxicol.*, 2002, **15**, 269; (e) H. Z. Boeini and K. H. Najafabadi, *Eur. J. Org. Chem.*, 2009, **2009**, 4926–4929.
- D. R. Chancellor, K. E. Davies, O. D. Moor, C. R. Dorgan, P. D. Johnson, A. G. Lambert, D. Lawrence, C. Lecci, C. Maillol, P. J. Middleton, G. Nugent, S. D. Poignant, A. C. Potter, P. D. Price, R. J. Pye, R. Storer, J. M. Tinsley, R. V. Well, R. Vickers, J. Vile, F. J. Wilkes, F. X. Wilson, S. P. Wren and G. M. Wynne, *J. Med. Chem.*, 2011, **54**, 3241–3250.
- (a) M. R. Kumar, K. Park and S. Lee, *Adv. Synth. Catal.*, 2010, **352**, 3255–3266; (b) A. M. Hamdy, N. Eleya, H. H. Mohammed, T. Patonay, A. Spannenberg and P. Langer, *Tetrahedron*, 2013, **69**, 2081–2086; (c) F. Shibahara, E. Yamaguchi and T. Murai, *Chem. Commun.*, 2010, **46**, 2471–2473; (d) J. Huang, J. Chan, Y. Chen, C. J. Borths, K. D. Baucom, R. D. Larsen and M. M. Faul, *J. Am. Chem. Soc.*, 2010, **132**, 3674–3675; (e) R. S. Sánchez and F. A. Zhuravlev, *J. Am. Chem. Soc.*, 2007, **129**, 5824–5825; (f) S. Ranjit and X.-G. Liu, *Chem.–Eur. J.*, 2011, **17**, 1105–1108.
- (a) T. Yamamoto, K. Muto, M. Komiyama, J. Canivet, J. Yamaguchi and K. Itami, *Chem.–Eur. J.*, 2011, **17**, 10113–10122; (b) J. Canivet, J. Yamaguchi, I. Ban and K. Itami, *Org. Lett.*, 2009, **11**, 1733–1736; (c) H. Hachiya, K. Hirano, T. Satoh and M. Miura, *ChemCatChem*, 2010, **2**, 140–1406.
- (a) H.-Q. Do and O. Daugulis, *J. Am. Chem. Soc.*, 2007, **129**, 12404–12405; (b) W. Zhang, Q.-L. Zeng, X.-M. Zhang, Y.-J. Tian, Y. Yue, Y.-J. Guo and Z.-H. Wang, *J. Org. Chem.*, 2011, **76**, 4741–4745; (c) W.-Y. Hu, P.-P. Wang and S.-L. Zhang, *Synthesis*, 2015, **47**, 42–48; (d) F.-Z. Yang, Z.-Q. Xu, Z. Wang, Z.-K. Yu and R. Wang, *Chem.–Eur. J.*, 2011, **17**, 6321–6325.
- S. K. Guchhait, M. Kashyap and S. Saraf, *Synthesis*, 2010, **2010**, 1166–1170.
- (a) Y. Wang, K. Sarris, D. R. Sauer and S. W. Djuric, *Tetrahedron Lett.*, 2006, **47**, 4823–4826; (b) C. O. Kangani,



- D. E. Kelley and B. W. Day, *Tetrahedron Lett.*, 2006, **47**, 6497–6499.
- 8 (a) Y. Riadi, R. Mamouni, R. Azzalou, M. E. Haddad, S. Routier and G. Guillaumet, *Tetrahedron Lett.*, 2011, **52**, 3492–3495; (b) S. Banerjee, S. Payra, A. Saha and G. Sereda, *Tetrahedron Lett.*, 2014, **55**, 5515–5520; (c) Y.-X. Chen, L.-F. Qian, W. Zhang and B. Han, *Angew. Chem., Int. Ed.*, 2008, **47**, 9330–9333; (d) L. Tang, X.-F. Guo, Y. Yang, Z.-G. Zha and Z.-Y. Wang, *Chem. Commun.*, 2014, **50**, 6145–6148; (e) A. K. Nezhad and F. Panahi, *ACS Catal.*, 2014, **4**, 1686–1692; (f) K. Osowska and O. S. Miljanic, *J. Am. Chem. Soc.*, 2011, **133**, 724–727.
- 9 G. Altenhoff and F. Glorius, *Adv. Synth. Catal.*, 2004, **346**, 1661–1664.
- 10 C. S. Demmer, J. C. Hansen, J. Kehler and L. Bunch, *Adv. Synth. Catal.*, 2014, **356**, 1047–1055.
- 11 (a) D. Xue and Y.-Q. Long, *J. Org. Chem.*, 2014, **79**, 4727–4734; (b) Q.-C. Xu, Z.-N. Li and H.-Y. Chen, *Chin. J. Chem.*, 2011, **29**, 925–932.
- 12 (a) J.-Y. Lu, X.-Y. Gong, H.-J. Yang and H. Fu, *Chem. Commun.*, 2010, **46**, 4172–4174; (b) P. Saha, M. Ashif Ali, P. Ghosh and T. Punniyamurthy, *Org. Biomol. Chem.*, 2010, **8**, 5692–5699; (c) N. Khatun, S. Guin, S. K. Rout and B. K. Patel, *RSC Adv.*, 2014, **4**, 10770–10778; (d) R. J. Perry and B. D. Wilson, *J. Org. Chem.*, 1992, **57**, 6351–6354; (e) G. Evindar and R. A. Batey, *J. Org. Chem.*, 2006, **71**, 1802–1808; (f) P. Saha, T. Ramana, N. Purkait, M. A. Ali, R. Paul and T. Punniyamurthy, *J. Org. Chem.*, 2009, **74**, 8719–8725; (g) Y.-T. Park, C. H. Jung and K.-W. Kim, *J. Org. Chem.*, 1999, **64**, 8546–8556; (h) J. Bonnamour and C. Bolm, *Org. Lett.*, 2008, **10**, 2665–2667.
- 13 (a) C. Bolm, J. Legros, J. L. Paih and L. Zani, *Chem. Rev.*, 2004, **104**, 6217–6254; (b) D. Damodara, R. Arundhathi and P. R. Likhar, *Catal. Sci. Technol.*, 2013, **3**, 797–802; (c) O. Bistri, A. Correa and C. Bolm, *Angew. Chem.*, 2008, **120**, 596–598; (d) C. L. Sun, B. J. Li and Z. J. Shi, *Chem. Rev.*, 2011, **111**, 1293–1314; (e) H. B. Wang, L. Wang, J. S. Shang, X. Li, H. Y. Wang, J. Guiand and A. W. Lei, *Chem. Commun.*, 2012, **48**, 76–78; (f) K. Gopalaiah, *Chem. Rev.*, 2013, **113**, 3248–3296; (g) I. Bauer and H.-J. Knolker, *Chem. Rev.*, 2015, **115**, 3170–3387.
- 14 (a) S. L. Buchwald and C. Bolm, *Angew. Chem., Int. Ed.*, 2009, **48**, 5586; (b) Z. Gonda, G. L. Tolnai and Z. Novak, *Chem.–Eur. J.*, 2010, **16**, 11822.
- 15 (a) A. Klapars, X.-H. Huang and S. L. Buchwald, *J. Am. Chem. Soc.*, 2002, **124**, 7421–7428; (b) E. R. Strieter, D. G. Blackmond and S. L. Buchwald, *J. Am. Chem. Soc.*, 2005, **127**, 4120–4121.

