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Cu(II)-catalyzed C–N coupling of 2-aminobenzothiazoles with boronic acids at room temperature†

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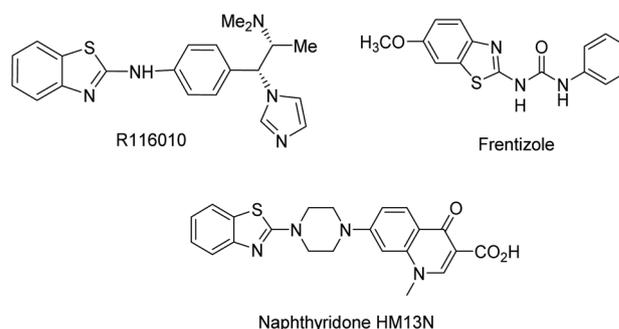
A Cu(II)-catalyzed, effective C–N coupling of 2-aminobenzothiazoles with boronic acids in acetonitrile under open vessel chemistry was achieved. This protocol demonstrates the *N*-arylation of 2-aminobenzothiazoles with a broad range of differently substituted phenylboronic acids at room temperature and accomplishes moderate to excellent yields of the desired products. Under the optimized condition, phenylboronic acids bearing halogen at the *para* and *meta* positions were found to be more fruitful.

1. Introduction

N-Arylated heterocyclic compounds are significant organic structural frameworks in various natural products¹ and drugs² and they also serve as precursors in aromatic heterocyclic synthesis.³ 2-Aminobenzothiazole derivatives have extensive applications in pharmaceutical chemistry due to their beneficial medicinal properties like anticancer,⁴ antidiabetic,⁵ anti-tubercular⁶ *etc.* and hence have generated inquisitiveness among researchers towards their synthesis. *N*-Substituted 2-aminobenzothiazoles containing molecules are employed as drugs (Scheme 1), for instance R116010,⁷ Frentizole⁸ and Naphthyridone HM13N,⁹ have been realized as a good inhibitor of *trans*-retinoic acid metabolism, as an immunosuppressive and as an efficient anti-HIV agent, respectively.

Copper catalyzed coupling strategies for carbon–heteroatom bond formation have emerged as an alternative to palladium chemistry, since they are less toxic, cheap, abundant and able to make milder reaction conditions. The comprehensive elaboration of copper based C–N coupling has grown from Ullmann's¹⁰ and Goldberg's¹¹ chemistry, but which needed drastic reaction conditions–stoichiometric amount of copper and temperature more than 200 °C. Later, Buchwald *et al.* reported the copper-catalyzed C–N coupling of amides with aryl halides using diamine ligands¹² whereas Chan and Lam disclosed copper mediated coupling of aryl boronic acid with nitrogen nucleophiles at room temperature (RT) under air with promising

functional group tolerance.¹³ The classical Chan–Lam coupling has faced some drawbacks such as requirement of stoichiometric quantity of copper catalyst, long reaction time, use of chlorinated solvent, narrow substrate scope, requirement of substrates with prefunctionalization and higher possibility for the presence of side products which make the purification of targeted products difficult. These downsides may provide some challenges for large-scale productions in industry. Therefore, a large group of researchers have tried to solve these problems and achieved substantial improvement in expanding copper catalyzed Chan–Lam coupling¹⁴ to deliver several methodologies for the formation of C–N bond, under relatively mild reaction conditions and with extended substrate scope.¹⁵ These methods used different types of copper-based catalytic systems like Cu(OAc)₂/amine bases or inorganic bases, Cu(OAc)₂/ligand/base, Cu₂O, [Cu(OH)·TMEDA]₂Cl₂, CuCl, CuCl/base, CuCl₂·2H₂O/ligand/base, CuOTf/ligand *etc.* to afford *N*-arylated compounds. Within these, C–N bond fabrication of heterocycles *via* Chan–Lam coupling was found effective on utilizing Cu(OAc)₂ in DCE for C–N coupling of 2-amino nitrogen containing heterocycles,¹⁴ Cu(OAc)₂/DMAP/KI in DME for the C–N



Scheme 1 Structure of drugs with 2-aminobenzothiazole scaffold.

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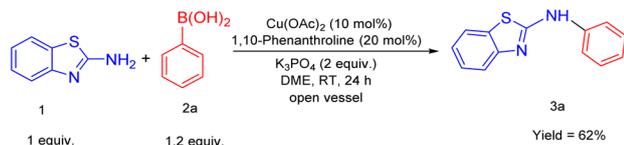
† Electronic supplementary information (ESI) available: A comprehensive report on the experimental approaches, procedures and materials employed *etc.* were included in the ESI. The characterization data with spectral details (¹H, ¹³C NMR) of the prepared compounds are also comprised in this section. See DOI: <https://doi.org/10.1039/d3ra02979d>



coupling of bidentate amides,¹⁶ Cu(OAc)₂ in methanol and Cu(OAc)₂/CsOPiv in DMF systems for the *N*-arylation of aminoazoles,¹⁷ CuCl₂·2H₂O/2,2'-bipyridine/K₂CO₃ in toluene or DMF system for the three-component amination involving boronic acids, cyanamides and amines as substrates,¹⁸ Cu₂O in DMSO for the amination of tetrazoles,¹⁹ CuOTf/1,10-phenanthroline in DMSO for the amination of nitrogen having heterocycles with tautomerism,²⁰ Cu(OAc)₂/CsCO₃ in DMF for the amination of 2-aminoheteroarenes,²¹ Cu(OAc)₂·H₂O in DCM for the amination of aminoindazoles,²² Cu(OAc)₂/decanoic acid/2,6-lutidine in toluene²³ for the amination of 2-

alkynylanilines, [Cu(OH)TMEDA]₂Cl₂/K₂CO₃ in DCM for the amination of tetrazoles²⁴ etc.

The detailed optimization studies and substrate scope exploration of Chan–Lam coupling of 2-aminobenzothiazoles under copper catalytic system was not reported. Motivated from the above mentioned approaches, we were enthusiastic to study the catalytic system for the C–N bond formation of 2-aminobenzothiazoles with aryl boronic acids. As an extension to our curiosity in transition metal catalysis,²⁵ herein we considered the Chan–Lam type coupling of 2-aminobenzothiazoles with arylboronic acid substrates under copper catalysis at room temperature.



Scheme 2 Preliminary reaction on Cu(II)-catalyzed C–N coupling of 2-aminobenzothiazole with phenylboronic acid.

2. Results and discussion

We have opted 2-aminobenzothiazole (**1**) and phenylboronic acid (**2a**) as our model substrates to optimize the reaction condition. Initially, the reaction of **1** and **2a** was executed using 10 mol% of Cu(OAc)₂ as catalyst, 20 mol% of 1,10-

Table 1 Optimizations of Cu catalysts, ligands, bases, solvents, time and temperature

S. no.	Cu salts	Ligands	Bases	Solvents	Temp (°C)	Time (h)	Yield (%) of 3a ^{a,b}
1	CuBr	L ₁	K ₃ PO ₄	DME	RT	24	28
2	Cu(OAc) ₂	L ₁	K ₃ PO ₄	DME	RT	24	62
3	CuBr ₂	L ₁	K ₃ PO ₄	DME	RT	24	51
4	—	L ₁	K ₃ PO ₄	DME	RT	24	nd
5	Cu(OAc) ₂	L ₂	K ₃ PO ₄	DME	RT	24	60
6	Cu(OAc) ₂	L ₃	K ₃ PO ₄	DME	RT	24	10
7	Cu(OAc) ₂	—	K ₃ PO ₄	DME	RT	24	Traces ^c
8	Cu(OAc) ₂	L ₁	Cs ₂ CO ₃	DME	RT	24	56
9	Cu(OAc) ₂	L ₁	(CH ₃) ₃ N	DME	RT	24	20
10	Cu(OAc) ₂	L ₁	C ₆ H ₁₃ N	DME	RT	24	36
11	Cu(OAc) ₂	L ₁	—	DME	RT	24	18 ^d
12	Cu(OAc)₂	L₁	K₃PO₄	CH₃CN	RT	24	76
13	Cu(OAc) ₂	L ₁	K ₃ PO ₄	DCE	RT	24	70
14	Cu(OAc) ₂	L ₁	K ₃ PO ₄	C ₆ H ₄ Cl ₂	RT	24	70
15	Cu(OAc) ₂	L ₁	K ₃ PO ₄	Water	RT	24	20 ^e
16	Cu(OAc) ₂	L ₁	K ₃ PO ₄	<i>t</i> -BuOH	RT	24	16
17	Cu(OAc) ₂	L ₁	K ₃ PO ₄	MeOH	RT	24	20
18	Cu(OAc) ₂	L ₁	K ₃ PO ₄	CH ₃ CN	45	24	35
19	Cu(OAc) ₂	L ₁	K ₃ PO ₄	CH ₃ CN	65	24	10
20	Cu(OAc) ₂	L ₁	K ₃ PO ₄	CH ₃ CN	RT	18	61
21	Cu(OAc) ₂	L ₁	K ₃ PO ₄	CH ₃ CN	RT	30	73

^a Reaction conditions: 2-aminobenzothiazole (1 equiv.), phenylboronic acid (1.2 equiv.), Cu salts (10 mol%), ligands (20 mol%), bases (2 equiv.), solvents (2 mL), reaction time: 18–30 h, temperature: RT–65 °C, air. Open vessel. ^b Isolated yield. nd = not detected. ^c Absence of ligands.

^d Absence of bases. ^e 1 equiv. of TBAB used.



phenanthroline as ligand and 2 equiv. of K_3PO_4 as base in DME at RT for 24 hours under open flask method. To our delight, the anticipated product *N*-phenylbenzo[d]thiazol-2-amine (**3a**) was noticed in 62% yield (Scheme 2, Table 1, entry 2). Encouraged from this result, we accomplished different optimizations for the C–N coupling of 2-aminobenzothiazoles with various boronic acids and are discussed in Table 1.

2.1 The choice of Cu-formed catalytic system

A blank test revealed that copper as a catalyst was indispensable to attain a reasonable result (Table 1, entry 4). So, we have studied the effect of different copper salts in the reaction as a catalyst. Cu(II) salts provided higher yields compared to Cu(I) salts, $Cu(OAc)_2$ showed the best catalytic activity with 62% yield of the expected product (Table 1, entry 2).

Subsequently, the influence of an array of different ligands, including *N,N*-type, *N,O*-type, *O,O*-type and phosphine ligands, were evaluated. *N,N*-planar ligands were found to be appropriate for the reaction (Table 1, entries 2 & 5) whereas *trans*-ligands disfavoured (Table 1, entry 6). 1,10-Phenanthroline was the ligand which offered the maximum outcome, and thus selected as the optimized ligand (Table 1, entry 2). Only traces of the required product **3a** was found for the experiment in the absence of ligand (Table 1, entry 7).

Later, the optimization of bases indicated that inorganic bases like potassium phosphate (Table 1, entry 2) displayed good results compared to the organic bases like triethyl amine and cyclohexylamine (Table 1, entries 9 & 10 respectively). A base-free condition was also carried out, but the yield was poor (Table 1, entry 11).

2.2 Catalytic properties of $Cu(OAc)_2/1,10$ -phenanthroline/ K_3PO_4 system

The impact of various solvents in the reaction was examined then. The replacement of DME with acetonitrile as solvent, improved the yield to 76% (Table 1, entry 12). Good yields were obtained with chlorinated solvents like DCE and dichlorobenzene also (Table 1, entries 13 & 14 respectively). But polar greener solvents like water, higher and lower alcohols were not applicable for the reaction because of their deprived yields (Table 1, entries 15–17). Since acetonitrile is well known for making coordination bonds with copper complexes, the increased rate of the reaction with the same may be due to its capability for coordination with $Cu(OAc)_2/1,10$ -phenanthroline/ K_3PO_4 catalytic system.

An appreciable depreciation in the yields of **3a** to 35% and 10% were noted, when the temperatures were elevated from RT to 45 °C and 65 °C respectively (Table 1, entries 18 & 19). The evaporation of solvents at these temperatures perhaps led to the non-availability of oxygen in the reaction. The duration of the reaction was then shortened to 18 hours and prolonged to 30 hours, which acquired the targeted products in 61% and 73% respectively (Table 1, entries 20 & 21). The significant decrease of yield in the previous situation could be ascribed to the reduced reaction rate for the lowered duration supported by the appearance of unreacted substrates in the reaction mixture. The

Table 2 Optimization of catalyst and ligand loading

S. no.	$Cu(OAc)_2$ (mol%)	1,10-Phenanthroline (mol%)	Yield (%) of 3a ^{a,b}
1	5	10	68
2	10	20	76
3	15	30	60
4	10	10	63
5	10	20	75 ^c
6	10	20	36 ^d
7	10	20	77 ^e
8	10	20	19 ^f

^a Reaction conditions: 2-aminobenzothiazole (1 equiv.), phenylboronic acid (1.2 equiv.), $Cu(OAc)_2$ (5–15 mol%), 1,10-phenanthroline (10–30 mol%), K_3PO_4 (2 equiv.), acetonitrile (2 mL), reaction time: 24 h, temperature: RT, air, Open vessel. ^b Isolated yield. ^c 1.5 equiv. of K_3PO_4 . ^d Excess of O_2 . ^e 1.6 equiv. of phenylboronic acid. ^f In N_2 atmosphere.

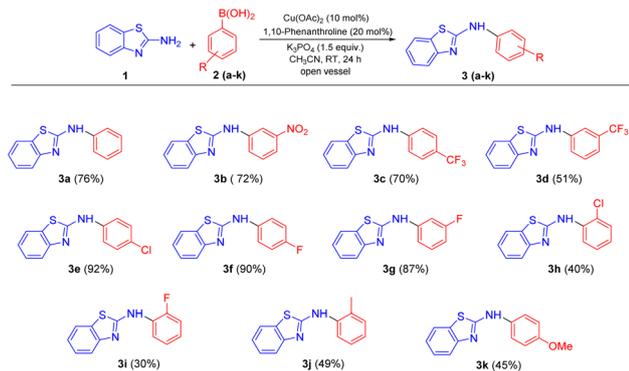
unaltered yield for the extended duration proved that 24 hours of the reaction is sufficient to form **3a** with maximum optimized yield.

The amount of copper catalyst was then reduced to 5 mol%, which offered 68% of **3a** (Table 2, entry 1). No enhancement in the yield of **3a** was observed when the amount of copper was raised to 15 mol% (Table 2, entry 3). Decreasing the quantity of 1,10-phenanthroline to 10 mol% (1:1 ratio of catalyst and ligand) did alter the yield appreciably to 63% (Table 2, entry 4). An additional trial using 1.5 equiv. of base gave similar yield of the reaction in comparison to that using 2 equiv. of base (Table 2, entry 5). Considerable increase in the yield was not detected, when the quantity of boronic acid was increased to 1.6 equiv. (Table 2, entry 7). Reactions carried out under excess of oxygen and in the presence of inert atmosphere (nitrogen) did not show drastic difference in the yield of the product (Table 2, entries 6 & 8). Thus, we have optimized the reaction condition, which utilized 10 mol% of $Cu(OAc)_2$, 20 mol% of 1,10-phenanthroline and 1.5 equiv. of K_3PO_4 in acetonitrile at RT for 24 hours for the coupling between 2-aminobenzothiazole and phenylboronic acid.

2.3 Efficiency of $Cu(OAc)_2/1,10$ -phenanthroline/ K_3PO_4 system

With the inspiring results in the optimized condition, we moved to the substrate scope studies by changing different phenylboronic acid substituents with 2-aminobenzothiazole derivatives and the observations are assembled in Scheme 3 and 4. A broad range of substituted phenylboronic acids underwent C–N coupling with 2-aminobenzothiazole (**1**) and afforded the corresponding *N*-arylated products in moderate to excellent yields (Scheme 3). *Para* and *meta* substituted halogen derivatives of phenylboronic acids provided the desired products in excellent yields with **1** (Scheme 3, **3e–g**). Electron-withdrawing group (EWG) containing substrates of phenylboronic acids having 3-nitro and 4- CF_3 substitution (Scheme 3, **3b** and **3c**) also reacted successfully and delivered products in good yields. The coupling of 3- CF_3 substituted phenylboronic acid with **1**

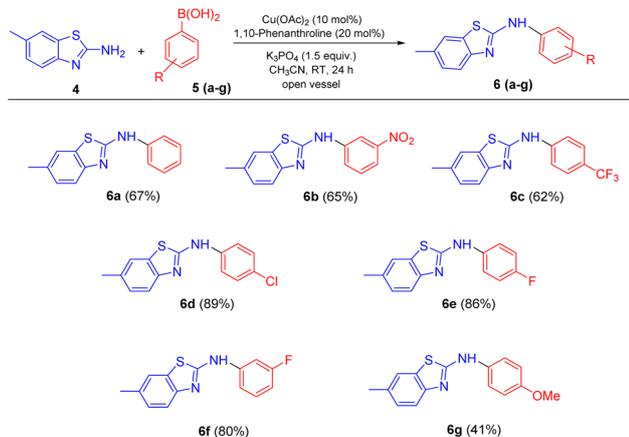




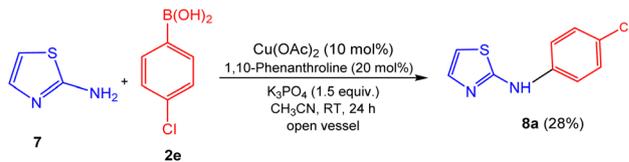
Scheme 3 Substrate scope of the reaction between substituted phenylboronic acids and 2-aminobenzothiazole.

furnished the targeted product in moderate yield (Scheme 3, **3d**). This may be explained on the basis of the reasonable nucleophilicity of the aromatic ring with EWG to follow transmetallation and reductive elimination in the mechanism. The EWG enhancing the yield of the reaction can be attributed mainly to the optimal formation of the complex C *via* transmetallation as well as the fast transformation of complex D to the targeted product. Fascinatingly, substrates containing electron-releasing substituents such as 2-methyl and 4-methoxy phenylboronic acids also coupled well affording 49% and 45% yield of the product respectively (Scheme 3, **3j** and **3k**). This methodology was also extended to *ortho* substituted halogen derivatives of phenylboronic acids, which afforded moderate yields, presumably due of steric reasons (Scheme 3, **3h** and **3i**).

To investigate the generality of the present coupling strategy, 6-methyl 2-aminobenzothiazole (**4**) was reacted with several phenylboronic acids (Scheme 4). 67% yield was obtained when phenylboronic acid was used as the coupling partner with **4** (Scheme 4, **6a**). 4-Cl, 4-F and 3-F substituted boronic acids tolerated well and yielded the corresponding products with 89%, 86% and 80% respectively (Scheme 4, **6d**, **6e** and **6f**). Electron deficient phenylboronic acids also afforded the coupling products in good yields (Scheme 4, **6b** and **6c**). The



Scheme 4 Scope of *N*-arylation with 6-methyl 2-aminobenzothiazole.

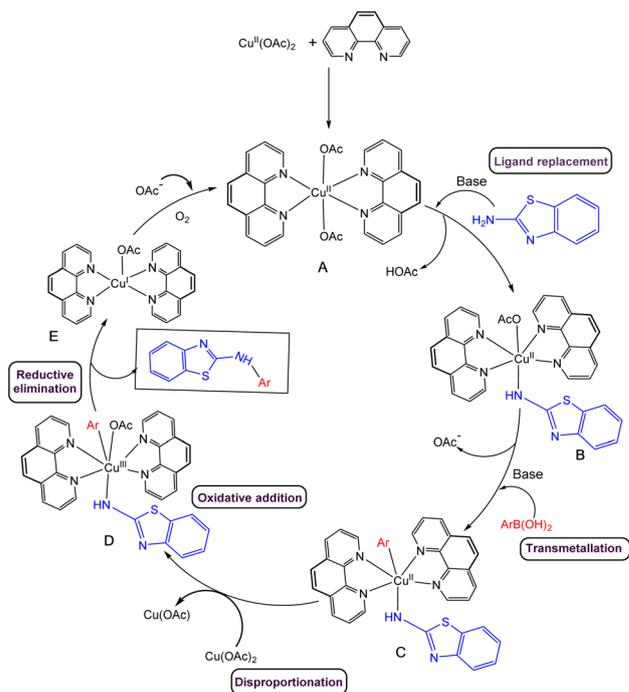


Scheme 5 C–N coupling of 2-aminobenzothiazole with 4-chlorophenylboronic acid.

anticipated *N*-arylation of *para*-methoxy phenylboronic acid was detected in 41% of the yield (Scheme 4, **6g**).

The reactivity of 2-aminobenzothiazole with *para*-chlorophenylboronic acid was also tried, which offered only 28% of the respective coupled product (Scheme 5, **8a**).

Based on the experimental studies and reported literature works,²⁶ we have suggested a possible mechanism for the C–N coupling approach (Scheme 6). The formation of an octahedral copper complex **A** from copper acetate and 1,10-phenanthroline could be considered as the first step. Acetate replacement of complex **A** with 2-aminobenzothiazole generates complex **B**. Transmetallation of boronic acid with complex **B** produces complex **C**, which undergoes oxidative addition to form a copper(III) complex **D**. **D** on reductive elimination furnishes the *N*-arylated product. Copper(I) complex **E** formed during this step experiences oxidation by O₂ and thus regenerates complex **A**. There is also a possibility for the solvent (acetonitrile) to coordinate with the copper²⁷ complexes mostly by the replacement of acetate ions, but this will be in equilibrium. Supposedly, K₃PO₄ base is found to be strong thermodynamically in aprotic solvents;²⁸ hence the rate of deprotonation from NH₂ group in **1** (or **4**) may be feasible by K₃PO₄ in acetonitrile.



Scheme 6 The possible mechanistic route.



3. Conclusions

In summary, Cu(II)/1,10-phenanthroline based catalytic system was established for the *N*-arylation of 2-aminobenzothiazoles with substituted phenylboronic acids at room temperature using acetonitrile as the solvent. This protocol is useful to synthesize a variety of *N*-arylated 2-aminobenzothiazole derivatives in moderate to excellent yields under simple and mild reaction conditions. Also, this method exhibited good functional group tolerance and particularly, proved to be highly efficient for the coupling with *meta* and *para* substituted halogen derivatives of phenylboronic acids. One of the chief benefits of our strategy include the use of less harmful acetonitrile as the reaction solvent at room temperature, though several Chan-Lam type C–N coupling reactions were reported in noxious organic solvents like DMF, THF *etc.* and chlorinated solvents like DCE at higher temperatures. Usually, addition of oxidants are required for Chan-Lam conditions at higher temperatures. Open flask chemistry made easier to monitor the development of the reaction, arrange the experiments, control the procedures and facilitates the substrates to react without adding oxidants. The optimized condition at room temperature with low copper loading, absence of additives like oxidant, favourable functional group compatibility are the other key advantages of our protocol. However, there are reports available in which researchers have performed their reactions under ligand- and base-free conditions.¹⁴ Since *N*-arylated heterocycles are unavoidable structural scaffolds in several drugs and natural products, novel discoveries in their synthetic routes might rise confidence among researchers in the scientific community.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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References

- R. G. S. Berlinck and M. H. Kossuga, *Modern Alkaloids*, Wiley-VCH Verlag GmbH & Co. KGaA, 2007, 305.
- A. Bikker, M. Brooijmans, A. Wissner and T. S. Mansour, *J. Med. Chem.*, 2009, **52**, 1493.
- M. A. McGowan, C. Z. McAvoy and S. L. Buchwald, *Org. Lett.*, 2012, **14**, 3800.
- (a) D. A. Ibrahim, D. S. Lasheen and M. Y. Zaky, *Bioorg. Med. Chem.*, 2015, **23**, 4989; (b) T. L. Dadmal, K. Appalanaidu, R. M. Kumbhare, T. Mondal, M. J. Ramaiah and M. P. Bhadra, *New J. Chem.*, 2018, **42**, 15546; (c) F. Sultana, M. A. Saifi, R. Syed, G. S. Mani, S. P. Shaik, E. G. Osas, C. Godugu, S. Shahjahanb and A. Kamal, *New J. Chem.*, 2019, **43**, 7150.
- (a) G. Sadhasivam and K. Kulanthai, *J. Chem. Pharm. Res.*, 2015, **7**, 425; (b) K. Harrouche, J.-F. Renard, N. Bouider, P. Tullio, E. Goffin, P. Lebrun, G. Faury, B. Pirotte and S. Khelili, *Eur. J. Med. Chem.*, 2016, **115**, 352.
- (a) F. M. Shaikh, N. B. Patel, G. Sanna, B. Busonera, P. L. Colla and D. P. Rajani, *Med. Chem. Res.*, 2015, **24**, 3129; (b) S. Radhika, M. B. Aleena and G. Anilkumar, *J. Catal.*, 2022, **416**, 233.
- J. Van Heudsen, R. Van Ginckel, H. Bruwiere, P. Moelans, B. Janssen, W. Floren, B. J. Van der Leede, J. van Dun, G. Sanz, M. Venet, L. Dillen, C. Van Hove, G. Willemsens, M. Janicot and W. Wouters, *Br. J. Cancer*, 2002, **86**, 605.
- M. Scheetz, D. Carlson and M. Schinitsky, *Infect. Immun.*, 1977, **15**, 145.
- S. Massari, D. Daelemans, M. L. Barreca, A. Knezevich, S. Sabatini, V. Cecchetti, A. Marcello, C. Pannecouque and O. Tabarrini, *J. Med. Chem.*, 2010, **53**, 641.
- F. Ullmann, *Ber. Dtsch. Chem. Ges.*, 1903, **36**, 2382.
- I. Goldberg, *Ber. Dtsch. Chem. Ges.*, 1906, **39**, 1691.
- (a) A. Klapars, J. C. Antilla, X. H. Huang and S. L. Buchwald, *J. Am. Chem. Soc.*, 2001, **123**, 7727; (b) A. Klapars, X. H. Huang and S. L. Buchwald, *J. Am. Chem. Soc.*, 2002, **124**, 7421.
- (a) D. M. T. Chan, K. L. Monaco, R.-P. Wang and M. P. Winters, *Tetrahedron Lett.*, 1998, **39**, 2933; (b) P. Y. S. Lam, C. G. Clark, S. Saubern, J. Adams, M. P. Winters, D. M. T. Chan and A. Combs, *Tetrahedron Lett.*, 1998, **39**, 2941.
- D. N. Rao, Sk. Rasheed, S. Aravinda, R. A. Vishwakarma and P. Das, *RSC Adv.*, 2013, **3**, 11472–11475.
- (a) A. Vijayan, D. N. Rao, K. V. Radhakrishnan, P. Y. S. Lam and P. Das, *Synthesis*, 2021, **53**, 805; (b) J.-Q. Chen, J.-H. Li and Z.-B. Dong, *Adv. Synth. Catal.*, 2020, **362**, 3311; (c) M. J. West, J. W. B. Fyfe, J. C. Vantourout and A. J. B. Watson, *Chem. Rev.*, 2019, **119**, 12491.
- H. Sahoo, S. Mukherjee, G. S. Grandhi, J. Selvakumar and M. Baidya, *J. Org. Chem.*, 2017, **82**, 2764.
- D. N. Rao, S. Rasheed, R. A. Vishwakarma and P. Das, *Chem. Commun.*, 2014, **50**, 12911.
- J. Li and L. Neuville, *Org. Lett.*, 2013, **15**, 6124.
- C. Y. Liu, Y. Li, J. Y. Ding, D. W. Dong and F. S. Han, *Chem.–Eur. J.*, 2014, **20**, 2373.
- K. A. Kumar, P. Kannaboina, C. K. Jaladanki, P. V. Bharatam and P. Das, *ChemistrySelect*, 2016, **3**, 601.
- S. Rasheed, D. N. Rao and P. Das, *J. Org. Chem.*, 2015, **80**, 9321.
- K. A. Kumar, P. Kannaboina, D. K. Dhaked, R. A. Vishwakarma, P. V. Bharatam and P. Das, *Org. Biomol. Chem.*, 2015, **13**, 1481.
- J. Gao, Y. Shao, J. Zhu, J. Zhu, H. Mao, X. Wang and X. Lv, *J. Org. Chem.*, 2014, **79**, 9000.
- T. Onaka, H. Umamoto, Y. Miki, A. Nakamura and T. Maegawa, *J. Org. Chem.*, 2014, **79**, 6703.



- 25 (a) S. Saranya, S. Radhika and G. Anilkumar, *ChemistrySelect*, 2021, **6**, 6847; (b) S. Radhika, T. Aneja, R. M. Philip and G. Anilkumar, *Appl. Organomet. Chem.*, 2021, **35**, e6217; (c) A. M. Thomas, S. Asha, K. S. Sindhu and G. Anilkumar, *Tetrahedron Lett.*, 2015, **56**, 6560.
- 26 D. N. Rao, Sk. Rasheed, K. A. Kumar, A. S. Reddy and P. Das, *Adv. Synth. Catal.*, 2016, **358**, 2126.
- 27 J. C. Vantourout, H. N. Miras, A. Isidro-Llobet, S. Sproules and A. J. B. Watson, *J. Am. Chem. Soc.*, 2017, **139**, 4769.
- 28 A. Klapars, X. Huang and S. L. Buchwald, *J. Am. Chem. Soc.*, 2002, **124**, 7421.

