



 Cite this: *RSC Adv.*, 2021, **11**, 32106

# Convenient one-pot synthesis of 1,2,4-oxadiazoles and 2,4,6-triarylpyridines using graphene oxide (GO) as a metal-free catalyst: importance of dual catalytic activity†

 Puja Basak, Sourav Dey and Pranab Ghosh \*

A convenient and efficient process for the synthesis of 3,5-disubstituted 1,2,4-oxadiazoles and 2,4,6-triarylpyridines has been described using an inexpensive, environmentally benign, metal-free heterogeneous carbocatalyst, graphene oxide (GO). GO plays a dual role of an oxidizing agent and solid acid catalyst for synthesizing 1,2,4-oxadiazoles and triarylpyridines. This dual catalytic activity of GO is due to the presence of oxygenated functional groups which are distributed on the nanosheets of graphene oxide. A broad scope of substrate applicability and good sustainability is offered in this developed protocol. The results of a few control experiments reveal a plausible mechanism and the role of GO as a catalyst was confirmed by FTIR, XRD, SEM, and HR-TEM analysis.

 Received 21st August 2021  
 Accepted 16th September 2021

DOI: 10.1039/d1ra06331f

[rsc.li/rsc-advances](http://rsc.li/rsc-advances)

## Introduction

Nitrogen-containing heterocyclic compounds are valuable due to their potential application as a key intermediate in the synthesis of numerous drugs.<sup>1</sup> 3,5-Disubstituted 1,2,4-oxadiazoles are a remarkably important class of nitrogen-containing heterocyclic scaffold as they are widely used as pharmacophores, bioactive molecules, and functional materials.<sup>1,2</sup> Among the oxadiazole derivatives, the 1,2,4-oxadiazole motif has received interest due to its application as a stable bioisostere in place of an amide, ester, or urea functionality.<sup>3</sup> These compounds when selectively functionalized, have performed as various muscarinic agonists,<sup>4</sup> benzodiazepine receptor partial agonists,<sup>5</sup> serotonergic (5-HT<sub>3</sub>) antagonists,<sup>6</sup> dopamine transporters,<sup>7</sup> antischistosomal drugs,<sup>8</sup> G-quadruplex ligands for probing DNA superstructure in antitumor research.<sup>9,10</sup> Another nitrogen-containing heterocycle, pyridines are ubiquitous and have attracted much attention due to their unique biological, medicinal, and pharmaceutical properties.<sup>11–13</sup> 2,4,6-Triarylpyridines are frequently used as a synthon in supramolecular chemistry owing to their  $\pi$ -stacking ability.<sup>14</sup> In addition, pyridines have received a growing interest as monomeric building blocks in thin films and organometallic polymers.<sup>15</sup>

It is noteworthy that, in the last decade many efficient protocols have been developed to synthesize these significant heterocyclic moieties. Among the known synthetic strategies of

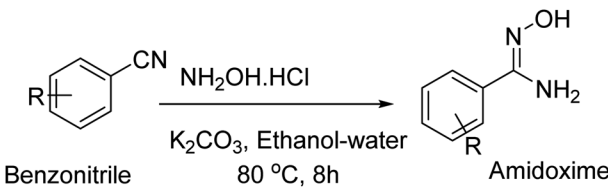
1,2,4-oxadiazoles, the most conventional approach involves the use of amidoximes as starting materials or intermediates. Other common approaches involve *O*-acylation of amidoximes by an activated carboxylic acid derivative, followed by cyclodehydration,<sup>16</sup> the 1,3-dipolar cycloaddition of nitrile oxide to nitriles, and intermolecular cyclodehydration reaction of amidoximes with aldehydes followed by oxidative dehydrogenation.<sup>17,18</sup> Besides this, base-mediated one-pot synthesis, MnO<sub>2</sub>/GO based synthesis, microwave-assisted efficient synthesis of oxadiazoles using PTSA and ZnCl<sub>2</sub> have also been reported.<sup>17,19–21</sup> On the other hand, efficient protocols for the synthesis of another important heterocycle 2,4,6-triarylpyridines involve condensation reaction between benzaldehydes, acetophenones, and ammonium acetate in presence of different acid catalysts<sup>22–24</sup> e.g. pentafluorophenylammonium triflate,<sup>23</sup> heteropolyacid,<sup>25</sup> HClO<sub>4</sub>-SiO<sub>2</sub>,<sup>26</sup> Brønsted-acidic ionic liquid,<sup>27</sup> and nano-metal catalyst.<sup>23,28,29</sup> Nevertheless, most of the traditional synthetic method requires harsh reaction condition, prolonged heating, and use of toxic transition metal catalyst. However, only a few protocols have shown greener context and high atom economy. Multicomponent reaction (MCR) is considered to be an effective and straightforward approach for the synthesis of heterocycles in an atom economical way. Considering the efficiency of MCRs and the aspects of green chemistry,<sup>30–33</sup> there is a need for new methods which involve metal-free, environmentally friendly catalytic protocol to synthesize 1,2,4-oxadiazoles and 2,4,6-triarylpyridines.

Recently, carbonaceous nanomaterials have gained considerable attention in green chemistry, especially in the development of metal-free sustainable heterogeneous catalysts.<sup>34–36</sup> Among the carbonaceous nanomaterials, graphene oxide (GO)

Department of Chemistry, University of North Bengal, Darjeeling, West Bengal, India.  
 E-mail: pizy12@yahoo.com; Fax: +91-353-2699001; Tel: +91-353-2776381

† Electronic supplementary information (ESI) available: The scanned copies of <sup>1</sup>H and <sup>13</sup>C NMR are included in this section. See DOI: 10.1039/d1ra06331f



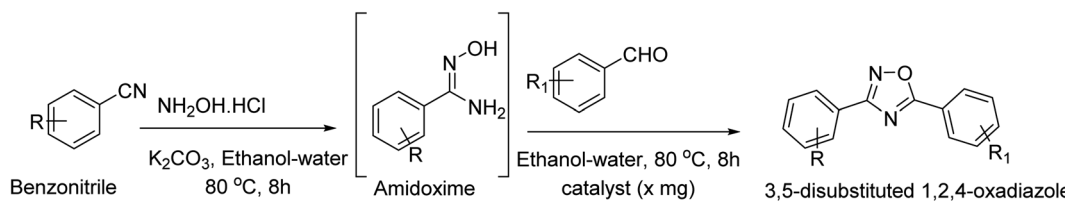
**Table 1** Optimization of reaction condition for the synthesis of amidoxime (intermediate)<sup>a</sup>


Entry	Solvent	Temp (°C)	Base	Yield <sup>b</sup> (%)
1	Water	100	K <sub>2</sub> CO <sub>3</sub>	68
2	Water	100	Cs <sub>2</sub> CO <sub>3</sub>	72
3	Ethanol	80	K <sub>2</sub> CO <sub>3</sub>	66
4	Ethanol	80	TEA	70
5	Ethanol-water	80	TEA	80
6	Ethanol-water	80	—	<50 <sup>c</sup>
7	Ethanol-water	80	K <sub>2</sub> CO <sub>3</sub>	91
8	Ethanol-water	80	K <sub>2</sub> CO <sub>3</sub>	94 <sup>d</sup>
9	Ethanol-water	80	Cs <sub>2</sub> CO <sub>3</sub>	93
10	THF	120	K <sub>2</sub> CO <sub>3</sub>	54
11	Toluene	110	K <sub>2</sub> CO <sub>3</sub>	<50
12	CH <sub>3</sub> CN	82	K <sub>2</sub> CO <sub>3</sub>	68
13	DMF	120	K <sub>2</sub> CO <sub>3</sub>	76

<sup>a</sup> Reaction condition: benzonitrile (1.5 mmol), hydroxylamine hydrochloride (1.5 mmol), base (1.5 mmol) and solvent (5 mL).

<sup>b</sup> Isolated yield. <sup>c</sup> No base was added. <sup>d</sup> The reaction was carried out for 24 h.

has been reported to accelerate several organic transformation reactions replacing different hazardous chemical reagents. GO, a thin two-dimensional unique nanomaterial contains different oxygen functionalities like carbonyl (C=O), carboxyl (COOH), epoxy (O), and hydroxyl (OH) on its edges and basal plane.<sup>37–39</sup> On account of the presence of large surface area, and diverse oxygen functionalities, GO has been identified as a heterogeneous solid acid catalyst (pH 4.5 at 0.1 mg mL<sup>-1</sup>) as well as a benign oxidizing agent.<sup>40–42</sup> Its abundance from low-cost natural carbon sources, low toxicity, reusability, and metal-free catalytic activity makes this heterogeneous carbon material (GO) as a promising carbocatalyst. Due to the inherent acidic and oxidation property of GO, it is explored as a catalyst in different organic transformations like C–H oxidations,<sup>43</sup> oxidative coupling of amines<sup>44</sup> to the imines, oxidation of thioanisole,<sup>45</sup> glutaraldehyde to glutaric acid,<sup>46</sup> 5-hydroxymethylfurfural,<sup>47</sup> benzylpyrazolyl coumarins,<sup>48</sup> Fisher esterification,<sup>49</sup> and transamidation.<sup>50–54</sup> The versatility and sustainability of GO as a catalyst leads us to employ GO as a metal-free catalyst for the synthesis of substituted 1,2,4-oxadiazoles and 2,4,6-triarylpyridines to overcome the drawbacks of the reported protocols and reduce environmental hazards. Our present study explores the role of GO as an acid catalyst as well as an oxidizing agent using the surface-bound oxygen-containing functional groups. To unleash the dual catalytic activity of GO, a plausible oxidative cyclization pathway to the

**Table 2** Optimization of reaction condition for the synthesis of 3,5-disubstituted 1,2,4-oxadiazole from amidoxime<sup>a</sup>


Entry	Catalyst (mg)	Solvent	Temperature	Time (h)	Yield%
1	—	Ethanol	80	12	Trace
2	15 (GO)	Ethanol	80	12	73
3	15 (GO)	Water	100	12	77
4	15 (GO)	DMF	100	12	60
5	15 (GO)	Ethanol-water	80	12	79
6	15 (GO)	Ethanol-water	80	24	83
7	25 (GO)	Ethanol-water	80	12	89
8	25 (GO)	Ethanol-water	80	8	88
9	25 (GO)	Ethanol-water	RT	12	52
10	25 (graphite)	Ethanol-water	80	8	40 <sup>b</sup>
11	25 (rGO)	Ethanol-water	80	8	45 <sup>c</sup>
12	25 (GO)/oxidant	Ethanol-water	80	8	67 <sup>d</sup>
13	Oxidant	Ethanol-water	80	8	<40 <sup>e</sup>
14	25 (GO)	Neat	80	8	69
15	25 (GO)	Ethanol-water	80	8	85 <sup>f</sup>
16	—	Ethanol-water	80	8	Nil <sup>f</sup>

<sup>a</sup> Reaction condition: benzaldehyde (1 mmol), amidoxime (1 mmol) and ethanol-water (5 mL), pristine GO (25 mg). <sup>b</sup> Graphite powder was used.

<sup>c</sup> Reduced graphene oxide (rGO). <sup>d</sup> GO and extra oxidant 30% H<sub>2</sub>O<sub>2</sub> (1 mmol) were used. <sup>e</sup> Only H<sub>2</sub>O<sub>2</sub> was used. <sup>f</sup> Under inert atmospheric condition.



Table 3 Synthesis of diversely functionalised 3,5-disubstituted 1,2,4-oxadiazole<sup>a</sup>

Entry	R	R <sub>1</sub>	Product	Yield <sup>b</sup> (%)
1	4-H	4-H		83
2	4-H	4-CH <sub>3</sub>		81
3	4-H	4-OCH <sub>3</sub>		80
4	4-H	4-F		78
5	4-H	3-NO <sub>2</sub>		75
6 <sup>c</sup>	4-H	4-N(CH <sub>3</sub> ) <sub>2</sub>	No 1,2,4-oxadiazole, only imine formation	—
7	4-H	1-Napthaldehyde		62
8	4-H	Furan-2-carbaldehyde		72
9	4-H	Thiophene-2-carbaldehyde		70
10	4-CH <sub>3</sub>	4-H		80



Table 3 (Contd.)

Entry	R	R <sub>1</sub>	Product	Yield <sup>b</sup> (%)
11	4-OCH <sub>3</sub>	4-H		78
12	4-OCH <sub>3</sub>	4-Cl		82
13	4-Pyridinecarbonitrile	4-H		68
14	4-H	CH <sub>3</sub> CHO		75
15 <sup>d</sup>	4-H	Heptaldehyde	NR	—
16 <sup>e</sup>	CH <sub>3</sub> CN	4-H	NR	—

<sup>a</sup> In the first step, benzonitrile (1 mmol), hydroxylamine hydrochloride (1.5 mmol), K<sub>2</sub>CO<sub>3</sub> (1.5 mmol), and ethanol–water (5 mL) were stirred for 8 h and in the 2<sup>nd</sup> step benzaldehyde (1 mmol) and GO (*x* mg) were added and stirred for another 8 h. <sup>b</sup> Isolated yield after purification through column chromatography. <sup>c</sup> 4-(Dimethylamino)benzaldehyde (1 mmol) was used. <sup>d</sup> Heptaldehyde was used. <sup>e</sup> Acetonitrile (1 mmol) was used.

synthesis of oxadiazoles and triarylpyridines under benign conditions has also been established.

## Results and discussion

For screening the reaction parameter benzonitrile (1.5 mmol), hydroxylamine hydrochloride (1.5 mmol), and base (1.5 mmol) were taken as model substrates to find out suitable conditions for the synthesis of amidoxime (intermediate). To satisfy our curiosity, the reaction was performed in different solvents *e.g.* polar protic, polar aprotic, and nonpolar. However, in absence of a base, a low yield was obtained (Table 1, entry 6). Gratifyingly, the reaction results showed (Table 1) the formation of amidoxime is highly favored in mixed solvent ethanol–water (1 : 3) using K<sub>2</sub>CO<sub>3</sub> as a base. To control the reaction conditions, after completion of the reaction, the solvent was removed by a rotary evaporator to separate the intermediate. While monitoring the TLC, only one spot was observed other than the reactant. After workup and purification by column

chromatography, 91% yield of the intermediate (amidoxime) was obtained (Table 1, entry 7). Although other bases were also employed (Table 1, entries 2, 5 and 9), K<sub>2</sub>CO<sub>3</sub> exerted the best result in an ethanol–water solvent. The synthesized amidoxime was characterized by NMR (300 MHz) and the spectral data was shown in ESI.†

In the second step of the reaction, benzaldehyde (1 mmol) and the catalyst were added to the reaction mixture to prioritize the synthesis of 3,5-disubstituted 1,2,4-oxadiazole. In presence of a small amount of GO, 73% yield of the product was obtained at 80 °C temperature (entry 2). Further increase in the amount of GO, proved to be favorable in the formation of 1,2,4-oxadiazole. No product was obtained when the reaction was carried out in absence of GO (Table 2, entry 1). High yield of the product was observed in aqueous ethanolic solution with a ratio ethanol–water (1 : 3). The outstanding catalytic activity of GO in ethanol–water (1 : 3) is revealed due to its better dispersibility. To establish the catalytic activity of GO, few controlled experiments were carried out using various catalysts. Other



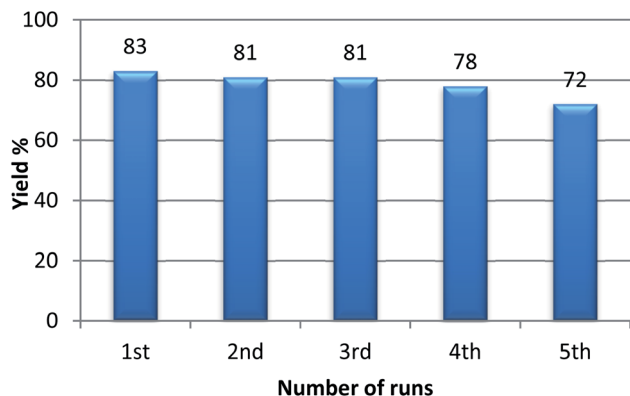


Fig. 1 Recyclability study of GO for the synthesis of 3,5-disubstituted 1,2,4-oxadiazole.

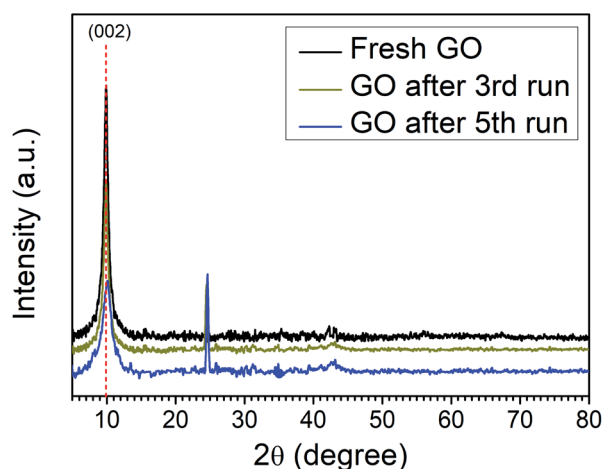


Fig. 2 XRD spectra of fresh GO, after 3<sup>rd</sup> run and 5<sup>th</sup> run.

carbonaceous nanomaterials *e.g.* powdered graphite, reduced graphene oxide (rGO) showed less catalytic activity than GO because they do not contain as many hydroxyl and carboxylic groups, indicating oxygen-containing functional groups in graphene oxide have a profound effect in catalyzing the synthesis of 3,5-disubstituted 1,2,4-oxadiazole. The reaction was also carried out in presence of GO and an oxidant H<sub>2</sub>O<sub>2</sub>, the reason for the low yield may be due to the oxidation of benzaldehyde to benzoic acid in presence of H<sub>2</sub>O<sub>2</sub> (Table 2, entry 12). The yield was not improved when only an H<sub>2</sub>O<sub>2</sub> oxidant was used (entry 13). These control experiments infer the significant catalytic role of GO in the reaction.

The scope and the substrate applicability of the reaction were also examined and results were summarized in Table 3.

With the optimized condition in hand, we have extended the substrate scope in organic transformations and a series of diversely substituted aldehydes and benzonitriles are subjected to the synthesis of 3,5-disubstituted 1,2,4-oxadiazole (Table 3). Both the electron-donating (Table 3, entries 2, 3, 10 and 11) and electron-withdrawing groups (entries 4 and 5) in the substituents afforded the corresponding product in good to excellent

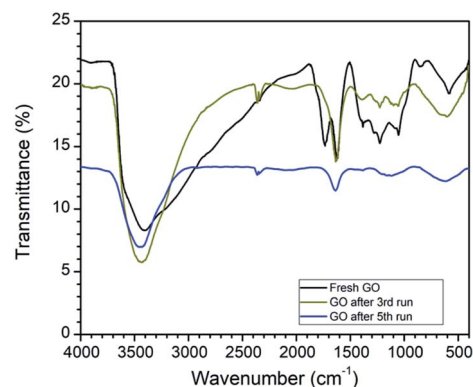


Fig. 3 Comparative FTIR of fresh GO, after 3<sup>rd</sup> run and 5<sup>th</sup> run.

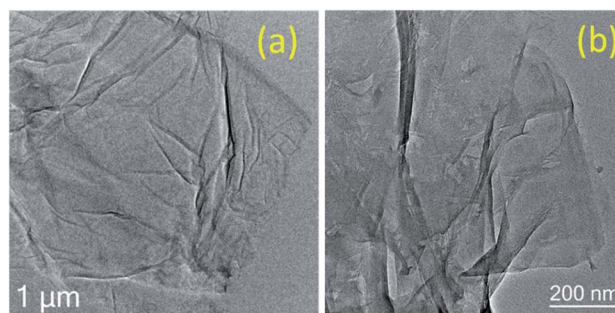


Fig. 4 HR-TEM images of (a) GO and (b) GO after the 5<sup>th</sup> run.

yield which indicates that the electronic nature of the substituents is not much influential to determine the yield of the reaction. 1-Naphthaldehyde offered the product with low yield and the reason may be due to steric hindrance (Table 3, entry 7). In the case of 4-*N,N*-(dimethylamino) benzaldehyde, the reaction was stopped at amidoxime, no desired oxadiazole is obtained (Table 3, entry 6). The present catalytic condition showed a wide tolerance to heterocyclic aldehydes (Table 3 entries 8, 9) and they were found to be highly effective to afford the corresponding product. The generality of the reaction was examined in the case of aliphatic aldehydes also. Interestingly, acetaldehyde was equally effective to yield the product with excellent quantity (entry 14). However, no product was found with increasing the side chain of aliphatic aldehydes (entry 15). It was disappointing that acetonitrile did not exert the corresponding product (entry 16). Due to the heterogeneous nature

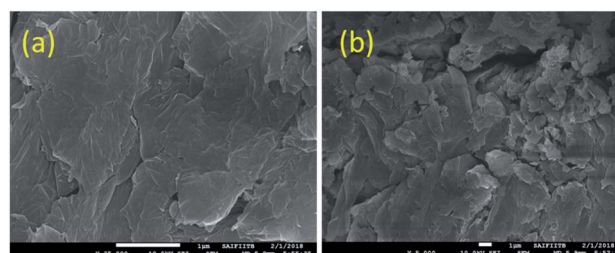


Fig. 5 SEM images of (a) GO and (b) GO after the 5<sup>th</sup> run.



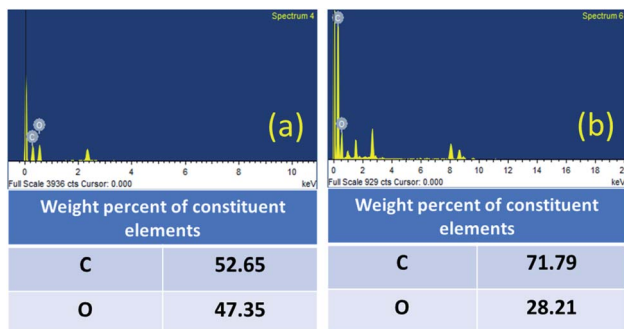
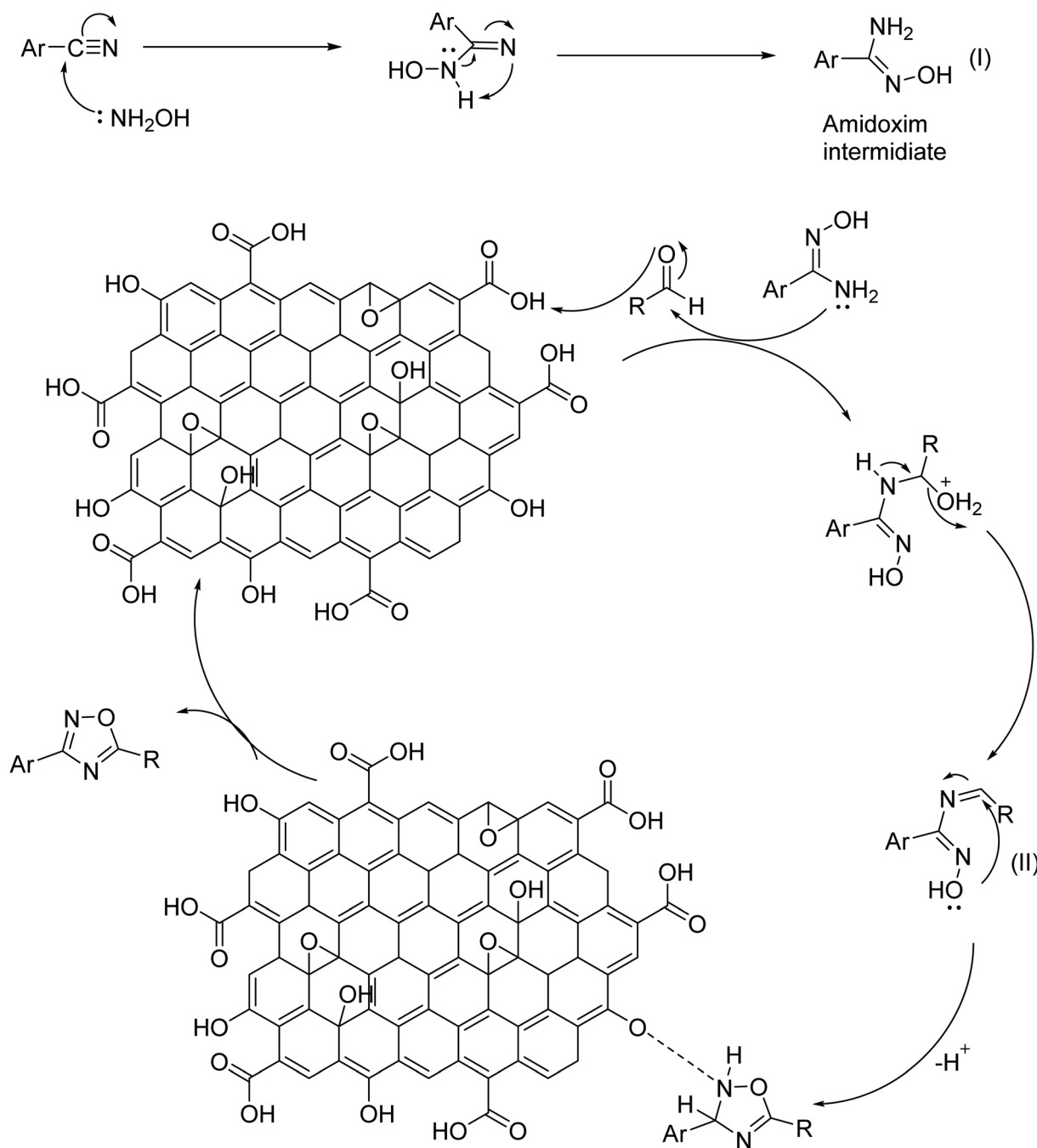


Fig. 6 EDX spectra of (a) GO and (b) GO after the 5<sup>th</sup> run.

of GO, it can be easily isolated from the reaction mixture and reused. The catalytic activity of GO was examined for five consecutive cycles for the synthesis of 3,5-disubstituted 1,2,4-oxadiazole from benzaldehyde and amidoxime under reflux conditions for 8 h to ascertain the recyclability potential of graphene oxide. The catalyst was separated after each recycle and washed thoroughly with ethanol and reused. A marginal decrease in the yield of oxadiazole is observed after each cycle which indicates a slight loss of catalytic activity of GO with recycling (Fig. 1).



Scheme 1 A plausible route to the synthesis of 3,5-disubstituted 1,2,4-oxadiazole.



The catalytic activity arises some structural changes in GO which were analyzed by FTIR, XRD, SEM, HR-TEM, and EDX analysis. The XRD spectra of fresh GO and recycled catalyst (GO after 3<sup>rd</sup> run and 5<sup>th</sup> run) are shown in Fig. 2. A comparison of spectra indicates the reduction in the intensity of the first characteristic peak of GO ( $2\theta = 10.01$ ) and the appearance of a new peak at ( $2\theta = 24.62$ ) due to the formation of partially reduced GO/reduced graphene oxide upon reuse. These results confirm the reduction of the functional groups of GO during the reaction.

The comparison of the FTIR spectra revealed that the peak at  $1720\text{ cm}^{-1}$  in fresh GO has completely disappeared after reuse. In addition to this, the peak intensity of the hydroxyl group at  $3400\text{ cm}^{-1}$  decreases after reuse. FTIR data strongly support the reduction of GO to rGO in this oxidative cyclization reaction (Fig. 3).

A morphological study of GO and GO after the 5<sup>th</sup> run was carried out using SEM and HR-TEM to investigate the disintegration of graphene oxide sheets after the reaction. In HR-TEM, the graphene oxide sheets are disintegrated into smaller sheets with slight aggregation after recycle (Fig. 4).

Moreover, the SEM images (Fig. 5) also reveal the formation of multiple small GO sheets after reuse. As GO catalyzes the reaction, its reduction to reduced graphene oxide possibly leads to its disintegration into smaller sheets.

The contribution of oxygen-containing functionalities during the reaction was further confirmed by the EDX analysis (Fig. 6). The carbon content was increased from 52.65% (fresh GO) to 71.79% (GO after 5<sup>th</sup> run) and the oxygen content was decreased from 47.35% (fresh GO) to 28.21% (GO after 5<sup>th</sup> run). The decrease in the oxygen content, therefore, indicates the role of GO in this cyclization reaction as an oxidizing agent. The universality and the dual catalytic activity of GO were established by a plausible mechanism (Scheme 1).

## Mechanism

A plausible mechanism of GO catalyzed synthesis of 3,5-disubstituted 1,2,4-oxadiazole has been proposed (Scheme 1) based on literature reports<sup>55</sup> and our controlled experiments (Table 2). Now, we propose the formation of amidoxime intermediate (I) from benzonitrile and hydroxylamine hydrochloride. However, in the first step, a base is required to neutralize hydroxylamine hydrochloride. In the 2<sup>nd</sup> step protonation of aldehyde, oxygen occurs and subsequently, a nucleophilic attack by amidoxime occurs at the electrophilic center of aldehyde. After that, the intermediate (II) undergoes an oxidative cyclization in presence of GO to produce 1,2,4-oxadiazoles. This mechanism is in good agreement with the control experiments as described in Table 2. However, in presence of only H<sub>2</sub>O<sub>2</sub> oxidant the yield of the reaction was diminished (Table 2, entry 11). The role of GO as an acid catalyst and an oxidant was confirmed as its absence did not lead to the oxadiazole product. The oxygen containing functional groups of GO are consumed during the reaction and the activity of GO gradually decreases. The activity of recycled GO is lower than that of the pristine GO. Good yield of the product was obtained even under an inert atmosphere which strongly establish (Table 2, entry 15), the prime role of GO in absence of atmospheric oxygen.

In connection to our previous work, the catalytic activity of synthesized GO was investigated in the case of 2,4,6-triarylpyridine synthesis. To find out the optimized condition of the reaction, acetophenone (2 mmol), benzaldehyde (1 mmol), and ammonium acetate (2 mmol) were selected as model substrates and the results were summarized in Table 4. As can be seen from Table 4 that neither polar nor non-polar solvents were found suitable for the reaction. The best result was obtained under neat or solvent-free conditions (Table 4, entry 11). The effect of temperature and the amount of catalyst was also examined to find out the optimized condition. Studies reveal

Table 4 Optimization of reaction condition for the reaction of 2,4,6-triarylpyridine<sup>a</sup>

Entry	Temp (°C)	Solvent	Catalyst GO (mg)	Ammonia source	Yield <sup>b</sup> (%)
1	100	H <sub>2</sub> O	15	NH <sub>4</sub> OAc	65
2	80	Ethanol	15	NH <sub>4</sub> OAc	55
3	100	DMF	15	NH <sub>4</sub> OAc	53
4	100	DMSO	15	NH <sub>4</sub> OAc	45
5	100	Toluene	15	NH <sub>4</sub> OAc	50
6	80	CH <sub>3</sub> CN	15	NH <sub>4</sub> OAc	30
7	100	Ethylene glycol	15	NH <sub>4</sub> OAc	60
8	100	Neat	15	NH <sub>4</sub> OAc	83
9	120	Neat	30	NH <sub>4</sub> OAc	90
10	150	Neat	30	NH <sub>4</sub> OAc	86
11	100	Neat	30	NH <sub>4</sub> OAc	92
12	80	Neat	30	NH <sub>4</sub> OAc	80
13	100	Neat	—	NH <sub>4</sub> OAc	Trace
14	100	Neat	30	(NH <sub>4</sub> ) <sub>2</sub> CO <sub>3</sub>	48
15	100	Neat	30	(NH <sub>4</sub> ) <sub>2</sub> SO <sub>4</sub>	Trace
16	RT	Neat	30	NH <sub>4</sub> OAc	<20

<sup>a</sup> Reaction condition: acetophenone (2 mmol), benzaldehyde (1 mmol), ammonium acetate (2 mmol), reaction time: 2 h <sup>b</sup> Isolated yields.

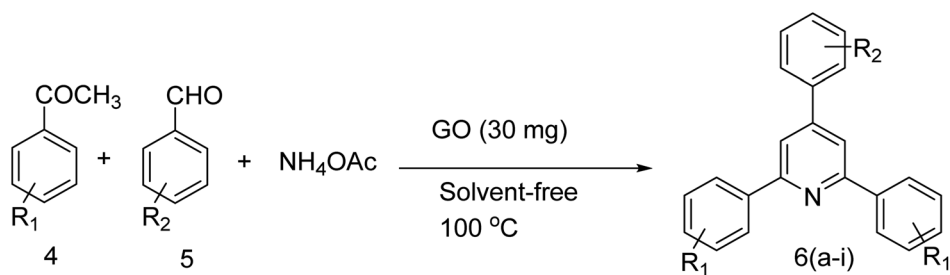


Table 5 Synthesis of 2,4,6-triarylpyridine derivatives in presence of GO<sup>a</sup>

Entry	R <sub>1</sub>	R <sub>2</sub>	Product	Time (h)	Yield <sup>b</sup> (%)
1	4-H	4-H		2 h	92
2	4-H	4-Me		2 h	86
3	4-H	4-Cl		1 h	93
4	4-H	4-NO <sub>2</sub>		1 h	88



Table 5 (Contd.)



Entry	R <sub>1</sub>	R <sub>2</sub>	Product	Time (h)	Yield <sup>b</sup> (%)
5	4-H	4-OMe		2 h	83
6	4-H	Furan-2-carbaldehyde		2 h	78
7	4-Me	4-H		2 h	87
8	4-Br	4-H		1 h	90



Table 5 (Contd.)

Entry	R <sub>1</sub>	R <sub>2</sub>	Product	Time (h)	Yield <sup>b</sup> (%)
9	4-Br	4-Cl	 6i	1 h	94

<sup>a</sup> Reaction condition: acetophenone (2 mmol), benzaldehyde (1 mmol), ammonium acetate (2 mmol) and GO (30 mg). <sup>b</sup> Isolated yields after purification through column chromatography on silica gel.

that the yield increases with increasing temperature. Room-temperature reaction afforded only 20% of the product which strongly indicates the vital role of temperature in governing the reaction (entry 16). However, after 120 °C the yield decreases with a further increase in temperature (Table 4, entry 10). To ascertain the catalytic function of GO, the reaction was performed in absence of catalyst and only a trace amount of product was obtained. The amount of the catalyst was also altered and optimum condition offered a neat reaction with 30 mg of GO at 100 °C temperature. Ammonia sources other than ammonium acetate produced the corresponding product with a low yield (Table 4, entries 14 and 15).

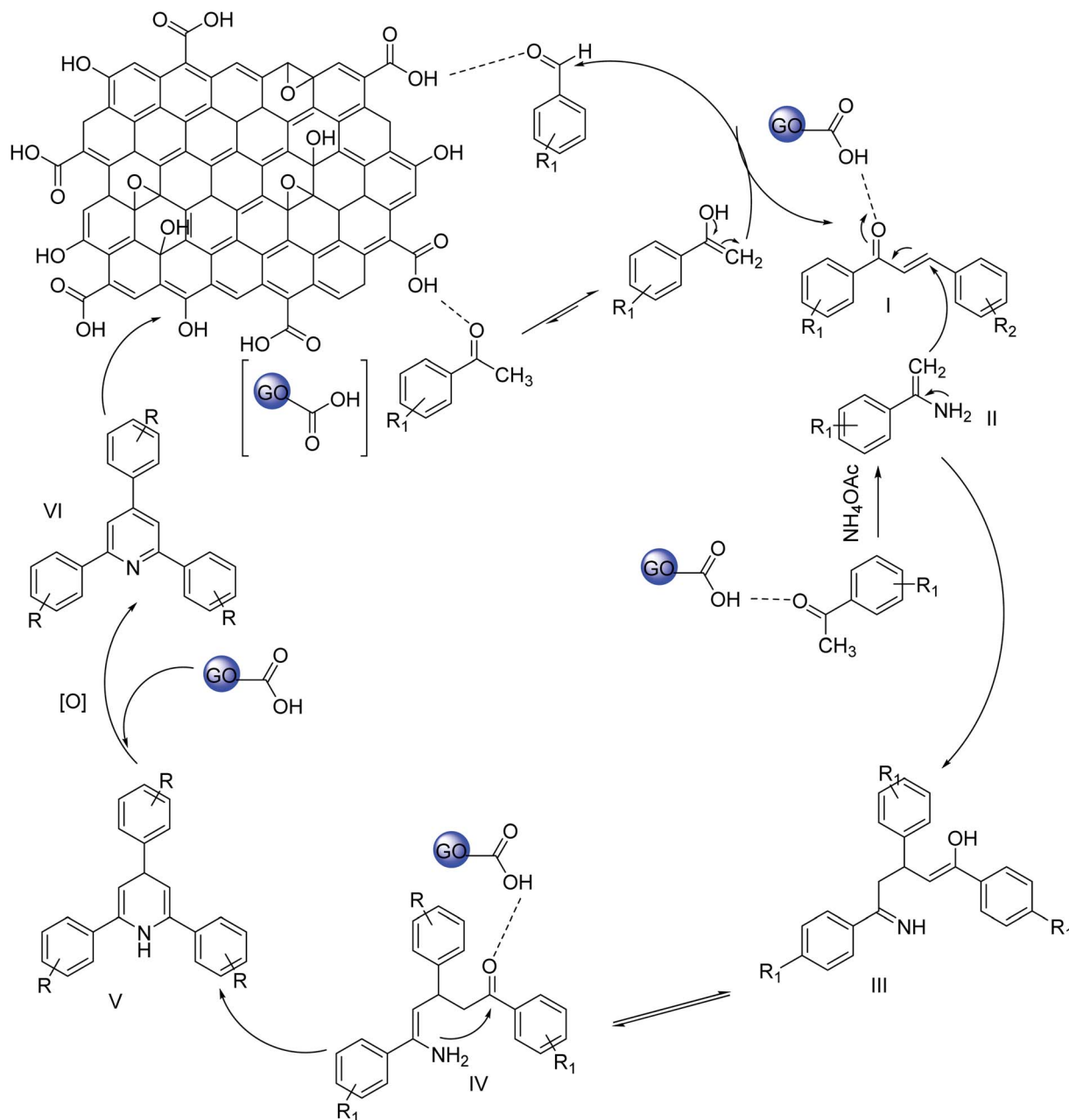
To explore the catalytic activity of GO, a wide variety of aromatic aldehydes and substituted acetophenones were subjected to synthesize 2,4,6-triarylpyridines. Based on the above-optimized results, GO catalyzed reaction was carried out at 100 °C temperature under solvent-free condition and the results are summarized in Table 5. First, the compatibility of the substituents in the phenyl ring of acetophenone and benzaldehyde was examined. All the electron-donating and electron-withdrawing substituents on the aromatic ring are equally capable of producing the corresponding product with a good yield. However, aldehydes with electron-withdrawing groups (Table 5, entries 3, 4 and 9) exerted excellent yield and reacted faster than the aromatic aldehydes with electron-donating groups (Table 5, entries 2, 5, 7). In the case of heterocyclic

aldehydes, the reaction has smoothly proceeded as can be seen from entry 6.

The probable mechanism for the synthesis of 2,4,6-triarylpyridines using GO is described in Scheme 2. At the very first step, aldol condensation occurs between acetophenone and aromatic aldehyde. Acetophenone is activated by the acidic group of GO and the nucleophilic attack occurs at the carbonyl carbon of aromatic aldehyde. After that, an acetophenone molecule is reacted with an ammonia source to form enamine (II). In the third stage, Michael's addition between enamine (II) and the aldol condensation product (I) occurs. GO protonates the condensation product (I), thereby facilitating the Michael addition by enamine (II). The intermediate (III) is formed by Michael's addition and undergoes cyclization to form dihydropyridine (V). At the last step, oxidation to dihydropyridine occurs and gives the ultimate product 2,4,6-triarylpyridine (VI).

The main advantage of heterogeneous catalysts is their reusability in organic transformation. For this purpose, acetophenone, benzaldehyde, and ammonium acetate were taken in a reaction vial in presence of 120 mg of GO. The model reaction was carried out for an adequate time and after completion of the reaction, ethyl acetate (30 mL) was added into the reaction vial and centrifuged for four times. The supernatant liquid after centrifugation was decanted off and the residual catalyst was washed repeatedly with water and acetone. The dry GO was then collected and reused for the 2<sup>nd</sup> run. It was observed that GO





Scheme 2 A possible route of GO catalyzed synthesis of 2,4,6-triarylpyridine.

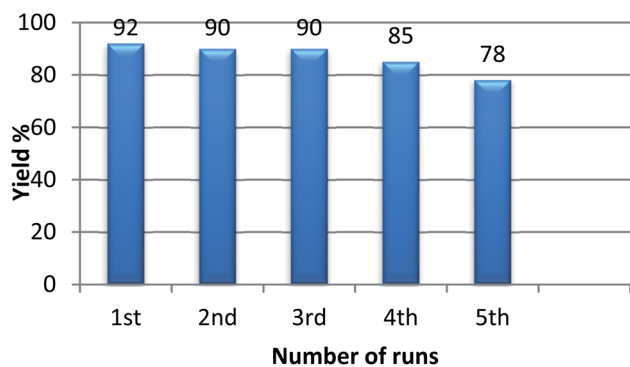


Fig. 7 Recyclability experiment of catalyst GO for the synthesis of 2,4,6-triarylpyridines.

could easily retain its acidic property without significant loss in its catalytic activity even after 5 successive runs (Fig. 7). Although there may be loss of some oxygenated groups due to subsequent runs, the recovered catalyst shows almost equal efficiency with the fresh GO.

## Conclusion

In conclusion, carbocatalyst based metal-free catalytic pathway for the synthesis of 3,5-disubstituted 1,2,4-oxadiazoles and 2,4,6-triarylpyridines has been established. The solid acid catalyst, GO facilitates the synthesis of oxadiazoles and triarylpyridines with good yield, easy recovery, and under mild



reaction conditions. The dual catalytic activity of GO has been demonstrated without any undesired by-product under benign conditions. The present protocol gives a clean strategy to provide a wide variety of substituted oxadiazoles and pyridines.

## Conflicts of interest

The authors declare no competing financial interest.

## Acknowledgements

PB is thankful to CSIR-New Delhi, India for the CSIR-Senior Research Fellowship [File no: 09/285(0072)/2016-EMR-I].

## References

- 1 M. Carbone, Y. Li, C. Irace, E. Mollo, F. Castelluccio, D. A. Pascale, G. Cimino, R. Santamaria, W. Y. Guo and M. Gavagnin, *Org. Lett.*, 2011, **13**, 2516–2519.
- 2 A. V. Gulevich, A. S. Dudnik, N. Chernyak and V. Gevorgyan, *Chem. Rev.*, 2013, **113**, 3084–3213.
- 3 R. J. Mathvink, A. M. Barritta, M. R. Candelore, M. A. Cascieri, L. Deng, L. Tota, C. D. Strader, M. J. Wyvratt, M. H. Fisher and A. E. Weber, *Bioorg. Med. Chem. Lett.*, 1999, **9**, 1869–1874.
- 4 F. I. Carroll, J. L. Gray, P. Abrahm, M. A. Kuzemko, A. H. Lewin, J. W. Boja and M. J. Kuhar, *J. Med. Chem.*, 1993, **36**, 2886–2890.
- 5 J. W. Clitherow, P. Beswick, W. J. Irving, D. I. C. Scopes, J. C. Barnes, J. Clapham, J. D. Brown, D. J. Evans and A. G. Hayes, *Bioorg. Med. Chem. Lett.*, 1996, **6**, 833–838.
- 6 C. B. Vu, E. G. Corpuz, T. J. Merry, S. G. Pradeepan, C. Bartlett, R. S. Bohacek, M. C. Botfield, C. J. Eyermann, B. A. Lynch, I. A. MacNeil and M. K. Ram, *J. Med. Chem.*, 1999, **42**, 4088–4098.
- 7 J. Matsumoto, T. Takahashi, M. Agata, H. Toyofuku and N. Sasada, *Jpn. J. Pharmacol.*, 1994, **65**, 51–57.
- 8 B. L. Mylari, T. A. Beyer, P. J. Scott, C. E. Aldinger, M. F. Dee, T. W. Siegel and W. J. Zembrowski, *J. Med. Chem.*, 1992, **35**, 457–465.
- 9 B. S. Orlek, F. E. Blaney, F. Brown, M. S. G. Clark, M. S. Hadley, J. Hatcher, G. J. Riley, H. E. Rosenberg, H. J. Wadsworth and P. Wyman, *J. Med. Chem.*, 1991, **34**, 2726–2735.
- 10 T. Nakamura, M. Asano, Y. Sekiguchi, Y. Mizuno, K. Tamaki, F. Nara, Y. Kawase, Y. Yabe, D. Nakai, E. Kamiyama, Y. Urasaki-Kaneno, T. Shimozato, H. Doi-Komuro, T. Kagari, W. Tomisato, R. Inoue, M. Nagasaki, H. Yuita, K. Oguchi-Oshima, R. Kaneko and T. Nishi, *Eur. J. Med. Chem.*, 2012, **51**, 92–98.
- 11 B. Y. Kim, J. BokAhn, H. W. Lee, S. K. Kang, J. H. Lee, J. S. Shin, S. K. Ahn, C. Hong and S. S. Yoon, *Eur. J. Med. Chem.*, 2004, **39**, 433–447.
- 12 L. Tian, J. Song, J. Wang and B. Liu, *Chin. Chem. Lett.*, 2009, **20**, 288–291.
- 13 R. D. Allen and G. A. R Johnston, *Med. Res. Rev.*, 1983, **3**, 91–118.
- 14 E. C. Constable, C. E. Housecroft, M. Neuburger, D. Phillips, P. R. Raithby, E. Schofield, E. Sparr, D. A. Tocher, M. Zehnder and Y. Zimmermann, *Dalton Trans.*, 2000, **13**, 2219–2228.
- 15 E. Figgemeier, E. C. Constable, C. E. Housecroft and Y. C. Zimmermann, *Langmuir*, 2004, **20**, 9242–9248.
- 16 (a) Y. Wang, R. L. Miller, D. R. Sauer and S. W. Djuric, *Org. Lett.*, 2005, **7**, 925–928; (b) S. Kandre, P. R. Bhagat, R. Sharma and A. Gupte, *Tetrahedron Lett.*, 2013, **54**, 3526–3529; (c) B. Kaboudin and L. Malekzadeh, *Tetrahedron Lett.*, 2011, **52**, 6424–6426; (d) B. Kaboudin and F. Saadati, *Tetrahedron Lett.*, 2007, **48**, 2829–2832.
- 17 J. K. Augustine, V. Akabote, S. G. Hegde and P. Alagarsamy, *J. Org. Chem.*, 2009, **74**, 5640–5643.
- 18 D. S. Bolotin, K. I. Kulish, N. A. Bokach, G. L. Starova, V. V. Gurzhiy and V. Y. Kukushkin, *Inorg. Chem.*, 2014, **53**, 10312–10324.
- 19 (a) M. Okimoto and Y. Takahashi, *Bull. Chem. Soc. Jpn.*, 2003, **76**, 427–428; (b) D. B. Repke, H. P. Albrecht and J. G. Moffat, *J. Org. Chem.*, 1975, **40**, 2481–2487.
- 20 W. Wang, H. Xu, Y. Xu, T. Ding, W. Zhang, Y. Ren and H. Chang, *Org. Biomol. Chem.*, 2016, **14**, 9814–9822.
- 21 F. Saadati, B. Kaboudin, R. Hasanloei, Z. Namazifar, X. Marset and G. Guillena, *Appl. Organomet. Chem.*, 2020, **34**, 5838–5849.
- 22 M. Adib, A. H. Jahromi, N. Tavoosi, M. Mahdavi and H. R. Bijanzadeh, *Tetrahedron Lett.*, 2006, **47**, 2965–2967.
- 23 N. Montazeri and S. Mahjoob, *Chin. Chem. Lett.*, 2012, **23**, 419–422.
- 24 J. Safari, S. Gandomi-Ravandi and M. Borujeni, *Chem. Sci.*, 2013, **125**, 1063–1070.
- 25 M. M. Heravi, K. Bakhtiari, Z. Daroogheha and F. F. Bamoharram, *Catal. Commun.*, 2007, **8**, 1991–1994.
- 26 L. Nagarapu, A. R. Peddiraju and S. Apuri, *Catal. Commun.*, 2007, **8**, 1973–1976.
- 27 A. Davoodnia, M. Bakavoli, R. Moloudi, N. Tavakoli-Houseini and M. Khashi, *Monatsh. Chem.*, 2010, **141**, 867–870.
- 28 A. Maleki and R. Firouzi-Haji, *Sci. Rep.*, 2018, **8**, 17303–17311.
- 29 E. Tabrizian, A. Amoozadeh, S. Rahmani, E. Imanifar, S. Azhari and M. Malmir, *Chin. Chem. Lett.*, 2015, **26**, 1278–1282.
- 30 G.-L. Wu and Q.-P. Wu, *Adv. Synth. Catal.*, 2018, **360**, 1949–1953.
- 31 R. Mishra, A. Jana, A. K. Panday and L. H. Choudhury, *Org. Biomol. Chem.*, 2018, **16**, 3289–3302.
- 32 C.-Y. Chen, W.-P. Hu, P.-C. Yan, G. C. Senadi and J.-J. Wang, *Org. Lett.*, 2013, **15**, 6116–6119.
- 33 C. C. Răzvan, R. Eelco and V. A. O. Romano, *Green Chem.*, 2014, **16**, 2958–2975.
- 34 D. S. Su, S. Perathoner and G. Centi, *Chem. Rev.*, 2013, **113**, 5782–5816.
- 35 O. Mohammadi, M. Golestanzadeh and M. Abdouss, *New J. Chem.*, 2017, **41**, 11471–11497.
- 36 N. Oger, Y. F. Lin, E. L. Grogneec, F. Rataboul and F. X. Felpin, *Green Chem.*, 2016, **18**, 1531–1537.



- 37 D. R. Dreyer and C. W. Bielawski, *Chem. Sci.*, 2011, **2**, 1233–1240.
- 38 S. Navalon, A. Dhakshinamoorthy, M. Alvaro and H. Garcia, *Chem. Rev.*, 2014, **114**, 6179–6212.
- 39 S. Zhu, J. Wang and W. Fan, *Catal.: Sci. Technol.*, 2015, **5**, 3845–3858.
- 40 C. Su and K. P. Loh, *Acc. Chem. Res.*, 2013, **46**, 2275–2285.
- 41 (a) D. R. Dreyer, H. P. Jia and C. W. Bielawski, *Angew. Chem., Int. Ed.*, 2010, **49**, 6813–6816; (b) D. R. Dreyer, H. P. Jia, A. D. Todd, G. Jeng and C. W. Bielawski, *Org. Biomol. Chem.*, 2011, **9**, 7292–7295.
- 42 T. Szabo, E. Tombacz, E. Illes and I. Dekany, *Carbon*, 2006, **44**, 537–545.
- 43 H. P. Jia, D. R. Dreyer and C. W. Bielawski, *Tetrahedron*, 2011, **67**, 4431–4434.
- 44 H. Huang, J. Huang, Y. M. Liu, H. Y. He, Y. Cao and K. N. Fan, *Green Chem.*, 2012, **14**, 930–934.
- 45 G. A. B. Gonçalves, S. M. G. Pires, M. M. Q. Simoes, M. G. P. M. S. Neves and P. A. A. P. Marques, *Chem. Commun.*, 2014, **50**, 7673–7676.
- 46 X. Chu, Q. Zhu, W.-L. Dai and K. Fan, *RSC Adv.*, 2012, **2**, 7135–7139.
- 47 G. Lv, H. Wang, Y. Yang, T. Deng, C. Chen, Y. Zhu and X. Hou, *ACS Catal.*, 2015, **5**, 5636–5646.
- 48 T. A. J. Siddiqui, B. G. Ghule, S. Shaikh, P. V. Shinde, K. C. Gunturu, P. K. Zubaidha, J. M. Yun, C. O. Dwyer, R. S. Mane and K. H. Kim, *RSC Adv.*, 2018, **8**, 17373–17379.
- 49 B. Roy, D. Sengupta and B. Basu, *Tetrahedron Lett.*, 2014, **55**, 6596–6600.
- 50 R. Wang, Z. Wu, Z. Qin, C. Chen, H. Zhu, J. Wu, G. Chen, W. Fan and J. Wang, *Catal.: Sci. Technol.*, 2016, **6**, 993–997.
- 51 S. Bhattacharya, P. Ghosh and B. Basu, *Tetrahedron Lett.*, 2018, **59**, 899–903.
- 52 Z. Chen, Y. Wen, Y. Fu, H. Chen, M. Ye and G. Luo, *Synlett*, 2017, **28**, 981–985.
- 53 J. Porwal, N. Karanwal, S. Kaul and S. L. Jain, *New J. Chem.*, 2016, **40**, 1547–1553.
- 54 H. P. Mungse, N. Bhakuni, D. Tripathi, O. P. Sharma, B. Sain and O. P. Khatri, *J. Phys. Org. Chem.*, 2014, **27**, 944–951.
- 55 K. B. Dhopte, R. S. Zambare, A. V. Patwardhan and P. R. Nemade, *RSC Adv.*, 2016, **6**, 8164–8172.

