RSC Advances



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PAPER



Cite this: RSC Adv., 2024, 14, 27092

Received 8th May 2024 Accepted 20th August 2024

DOI: 10.1039/d4ra03401e

rsc.li/rsc-advances

Introduction

In catalysis, chemists use ligands as tools because minute changes in their steric and electronic properties have a significant impact on the performance, properties and functioning of the catalyst. Such changes can also be used to influence the catalytic reactions. The impact of ligand's structure on catalytic processes has been investigated both experimentally1-9 and theoretically.¹⁰⁻¹² In the case of coupling reactions, which are heavily applied tools in chemical transformations, bondmaking and bond-breaking processes are very important. A variety of ligands, including Schiff bases, pincer ligands, and heterocyclic compounds such as triazoles and pyridines, have become well-known for synthesizing transition metal catalytic systems for such reactions.13-16 Additionally, over the last decade, a variety of organosulphur and organoselenium ligands have also been explored for designing homogeneous, heterogeneous, and nanocatalytic systems for such reactions.17-25 However, the use of organotellurium ligands is very much

Catalytic system having an organotellurium ligand on graphene oxide: immobilization of Pd(0) nanoparticles and application in heterogeneous catalysis of cross-coupling reactions⁺

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First heterogeneous catalytic system, having a covalently linked hybrid bidentate organotellurium ligand *[i.e.,* PhTe-CH₂-CH₂-NH₂] on the surface of graphene oxide, has been synthesized with immobilized and stabilized Pd(0) nanoparticles. To the best of our knowledge, it is the first such catalytic system in which a heterogenized organotellurium ligand has been used. It has been well-characterized using different physicochemical characterization techniques *viz.* P-XRD, XPS, HR-TEM, EELS, FE-SEM, EDX, TGA, BET surface area analysis, FT-IR spectroscopy, and Raman spectroscopy. The Pd content of the final system has been quantified using ICP-OES. Its applications have been explored in Suzuki–Miyaura C-C cross coupling and C-O cross coupling reactions. Hot filtration experiments corroborate the heterogeneous nature of the catalysis. It is recyclable for up to five reaction cycles in Suzuki–Miyaura and C-O cross coupling with marginal loss in performance. It also catalyzes the reactions of chloroarenes such as chlorobenzene, 4-chloroaniline, 1-chloro-4-nitrobenzene, 4-chloroacetophenone, 4-chlorobenzophenone for Suzuki coupling, and 1-chloro-4-nitrobenzene, 4-chlorobenzonitrile, chlorobenzene, and 4-chlorotoluene for C-O coupling. P-XRD, FE-SEM, and EDX study reveals that the catalytic system retains its structural originality and functionality after recycling.

limited, and that too only to the designing of homogeneous complexes.²⁶⁻³³ Unlike organosulphur and organoselenium ligands, they have been rarely used in designing heterogeneous or nanocatalytic systems.

In heterogeneous catalytic systems, an appropriate inert support (oxide of metals,^{32,33} silica,³⁴ double hydroxides of layered structure³⁵ and graphene/graphene oxide^{36,37}) is used to immobilize the metal nanoparticles38 or coordination complex of the metal.³⁹ Such catalytic systems are known for their specific advantages including efficiency, selectivity, high conversion rate, surface area, and reusability under mild conditions.⁴⁰⁻⁴⁷ Stabilizing the noble metal nanoparticles with organic capping agents (i.e. stabilizers) is common, but these stabilizers can impact catalytic activity due to their binding to the metal surface.48 However, if small nanoparticles of uniform size are immobilized over the surface of a solid support to obtain a stable catalytic system, high activity may be achieved due to the high surface area to volume ratio and high population of catalytically active surface Pd atoms. Such modification and functionalization of the surface of the solid support with nanoparticles are very advantageous from the point of view of desired catalytic properties for specific applications.⁴⁹ Among various solid supports used for this purpose, graphene oxide is very attractive because of its unique electronic structure,

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[†] Electronic supplementary information (ESI) available. See DOI: https://doi.org/10.1039/d4ra03401e

mechanical strength, and chemical stability.⁵⁰ It has epoxy and hydroxyl groups at the basal planes and carboxylic groups at the edges of the sheets.⁵¹ Materials of tremendous potential have been synthesized by grafting nanomaterials on graphene oxide (GO).⁵² Graphene/GO embedded with palladium nanoparticles are known as potent species for catalyzing multiple reactions *e.g.*, C–C cross coupling, and C–O coupling reactions.^{52–59}

Tellurium, a metalloid, is distinct from other elements in its group, and its chemistry has been relatively understudied despite its unique properties. Organotellurium compounds have distinct characteristics compared to similar sulphur and selenium derivatives. Due to the presence of Te, such compounds are known to possess antioxidant,60 anti-inflammatory,61 neuroprotective,62 chemopreventive,63 and glutathione peroxidase⁶⁴ like activities. In some cases, they also possess toxic effects to the central nervous systems of various rodents and mammals, which can lead to mutagenicity and carcinogenicity.65 They may also have a dual biological functioning. Depending on the biological redox environment, they may have both harmful and helpful effects.⁶⁰⁻⁶⁶ When Te is present in the framework of the organic compounds, they may behave as ligands. Hence, they can also be used as potentially valuable candidates in catalyst development.27,28,67,68 The combination of tellurium and palladium demonstrates synergistic effects and potential for enhanced catalytic performance.69,70 It has, therefore, been thought worthwhile to modify the surface of GO with immobilization of hybrid organotellurium ligand, i.e., Ph-Te-CH₂-CH₂-NH₂, through covalent interactions so that a novel system may be obtained having a combination of chelating donor (Te and N) groups on the surface of GO. Presence of such donor groups allows the system to offer strong binding sites to the Pd NPs and hold them efficiently without any significant agglomeration and leaching problems. It may also lead to the development of Pd NPs based stable and recyclable catalytic system. A pair of Te and N donor groups, i.e., a combination of soft and hard donor sites capable of chelating, is also worth exploring in the context of heterogenized form. Thus, the GO surface modified with covalently linked Te donor site containing primary amine is reported herein for the development of heterogeneous catalytic system having Pd NPs. It has the potential to efficiently catalyze Suzuki coupling and C-O coupling reactions at low catalyst loading. Pd NPs immobilized on heterogenized organotellurium compound (i.e., tellurated ethyl amine) on the surface of GO and their application in the catalysis of the coupling reactions is reported for the first time to the best of our knowledge. The catalytic system [i.e., nanosized GO-TeN-Pd(0)] is highly stable, efficient, heterogeneous and reusable. These reactions, catalyzed with it, could be beneficial, as these reactions are considered as important tools in the synthesis of agrochemical, pharmaceutical, and perfume industry.71-74

Results and discussion

Synthesis of GO-TeN-Pd(0) nanocatalyst

Hummers' method⁷⁵ has been adopted to produce graphene oxide (Scheme 1) which is required for the development of the



Scheme 1 Protocol for surface modification of graphene oxide with organotellurium ligand and immobilization of Pd(0) nanoparticles onto Te, N ligand-functionalized graphene oxide.

heterogeneous catalytic system. It is subjected to wet chemical treatment in the next step to functionalize it with hydroxyl, epoxide, and carboxyl groups. 2-(Phenyltelluro)ethylamine has been synthesized using reported procedure.76 The GO structure has COOH groups which have the ability to form chemical bonds with 2-(phenyltellanyl)ethanamine (Ph-Te-CH2-CH2-NH₂) through an amide linkage. However, it is preferred to increase the abundance of carboxyl groups on GO before immobilization of the ligand. Hence, the reaction of GO has been carried out with chloroacetic acid. It constituted a part of the wet chemical treatment process. During this reaction, the epoxy and hydroxyl groups of GO are targeted and converted into COOH groups. In the next step, the treatment of the 2 was done with thionyl chloride to convert COOH groups into -COCl groups (*i.e.*, the species 3). In the next step, $Ph-Te-CH_2-CH_2-$ NH₂ was immobilized covalently over the 3 (*i.e.*, functionalized GO) using a reaction in which it has been treated (Scheme 1) with the 3 (GO-COCl) in DMF. This system includes Te and N bidentate donor sites. In the last step, the GO-TeN is treated with Na₂PdCl₄ and NaOH solution to achieve the catalytic system [i.e., GO-TeN-Pd(0)].

Stepwise characterization of GO, GO-COOH, GO-TeN and GO-TeN-Pd(0)

Various spectroscopic and microscopic characterization techniques, including powder-XRD (Fig. 1), XPS (Fig. 2), Raman (Fig. 3), HR-TEM (Fig. 4), HAADF-STEM (Fig. 6), FE-SEM (Fig. 7), EELS (Fig. 8–13 and ESI: Fig. S9–S11†), TGA (Fig. 14), BET surface area studies (Fig. 15), EDS (ESI: Fig. S1–S4†), FT-IR (ESI:



Fig. 1 P-XRD patterns of (a) GO, (b) GO–COOH, (c) GO–TeN and (d) GO–TeN–Pd(0).

Fig. S5–S8†), and ICP-OES have been used to study the structural and morphological properties of graphene oxide and its functionalized surface at every step. All these studies corroborate the formation of the nanocatalytic system [*i.e.*, GO–TeN–Pd(0)].

Powder X-ray diffraction (P-XRD)

The study of the powder X-ray diffraction (P-XRD) patterns (Fig. 1) provide substantial evidence of the formation of a heterogeneous catalyst. In the initial state, graphene oxide (GO) exhibits a well-defined peak [Fig. 1(a)] at $2\theta = 10.5^{\circ}$, corresponding to the (001) plane, affirming the presence of GO. Upon carboxylation of GO, this peak at $2\theta = 10.5^{\circ}$ vanishes, giving way to a new broad peak [Fig. 1(b)] at $2\theta = 24.4^{\circ}$ (111). The emergence of this (111) peak signifies the conversion of hydroxyl (-OH) and epoxy groups in GO into carboxylic groups. This transformation is consistent with literature reports.⁵³ The alterations in the intensities of P-XRD patterns of GO-TeN in comparison to GO-COOH [Fig. 1(c)] suggest the formation of amide linkages with the ligand (Ph-Te-CH₂-CH₂-NH₂). Additionally, peaks at $2\theta = 37.9^{\circ}$ (102) and 48.8° (201) are attributed to the presence of Te. Finally, the P-XRD pattern of GO-TeN-Pd(0) [Fig. 1(d)] gives strong indication of the presence of a cubic crystalline Pd(0) phase. Peaks at (hkl) 111, 200, 220, and 311, corresponding to Pd(0) nanoparticles, appear at $2\theta = 39.9^{\circ}$, 46.2°, 67.1°, and 81.5°, unequivocally confirming the existence of the Pd(0) phase.

X-ray photoelectron spectroscopy (XPS)

X-ray photoelectron spectroscopic (XPS) studies (Fig. 2) have been conducted for GO–COOH, GO–TeN, and GO–TeN–Pd(0) to gain insights into the chemical entities incorporated during the functionalization procedure. Gaussian functions were



Fig. 2 XPS spectra of (i) GO-COOH; (A1): C 1s (A2): O 1s (A3): full, (ii) GO-TeN; (B1): C 1s (B2): O 1s (B3): Te 3d (B4): full, (iii) GO-TeN-Pd(0); (C1): O 1s (C2): Te 3d (C3): Pd 3d (C4): full.

employed to fit the XPS spectra (Fig. 2). In the XPS data for GO-COOH, three distinct C 1s peaks [Fig. 2(A1)] were observed at binding energies of 284.7, 286.7, and 288.9 eV. These peaks attributed to sp² carbon bonds within the graphene sheet, carbon atoms attached to oxygen atoms *via* single bonds, and the presence of carboxylic acid groups (–COOH) on the sheet's surface. Subsequently, the XPS survey exhibits 2 discernible O 1s peaks [Fig. 2(A2)], representing C–O and C(=O)OH groups.⁷⁷

Species 4 (*i.e.*, GO–TeN) contains organotellurium compound, covalently linked on the surface of GO. Its XPS spectrum has a single peak for N 1s [Fig. 2(B1)] and Te 3d [Fig. 2(B3)], along with 2 prominent C 1s peaks [Fig. 2(B4)]. These C 1s peaks are attributed to the formation and existence of the C–N bond between the ligand and sp^2 carbons of the carbonyl group of GO (*i.e.*, the graphene sheet).

Subsequently, GO–TeN is transformed (Scheme 1) into the final nanocatalytic system, *i.e.*, species 5 [GO–TeN–Pd(0)]. The XPS data [Fig. 2] validates the existence of all anticipated elements (*i.e.*, C, N, O, Te, and Pd) in this system. The peaks, corresponding to Pd $3d_{5/2}$ & Pd $3d_{3/2}$ [Fig. 2(C4)], corroborate the presence of Pd NPs.^{78–81} These 3d peaks of Pd correspond to spin–orbit splitting, resulting in $3d_{5/2}$ and $3d_{3/2}$ states. The binding energies of the $3d_{5/2}$ & $3d_{3/2}$ states of Pd have been determined and found to be 335.1 eV and 340.4 eV, respectively. The values indicate the presence of Pd (0),^{79–81} in which the order of $3d_{5/2}$ and $3d_{3/2}$ states follows the trend *i.e.*, $3d_{5/2} < 3d_{3/2}$.

Raman spectroscopy

Raman spectroscopy is a non-invasive tool, used for unveiling the intricate structural and compositional traits inherent in carbon-based materials. Raman spectroscopic studies (Fig. 3) have been conducted systematically on graphene oxide (GO), GO-COOH, GO-TeN, and GO-TeN-Pd(0) to understand the changes experienced by GO during its transformation (Scheme 1) from **1** to **5** [*i.e.*, GO-TeN-Pd(0)]. The Raman spectrum of GO



Fig. 3 Raman spectra: (a) GO, (b) GO–COOH, (c) GO–TeN and (d) GO–TeN–Pd(0).

[Fig. 3(a)] has the characteristic D and G bands, located at 1358 and 1599 cm⁻¹, respectively. However, the D band in GO-COOH (*i.e.*, 2), GO-TeN (*i.e.*, 4), and 5 [*i.e.*, GO-TeN-Pd(0)] is seen [Fig. 3(b)-(d)] at 1352, 1347, and 1341 cm⁻¹, respectively. On the other hand, the G band resonates at 1591, 1584, and 1578 cm⁻¹ in these species (*i.e.*, 2, 4 and 5). These changes in the spectrum suggest that the wet chemical treatment damages the carbon-carbon conjugation in the graphene sheets.

The quantified intensity ratio of the D- and G-bands, represented as I_D/I_G , is considered as an indicator of the quality of the graphitic structure. The reliability of this indicator is known to be high. When pyrolyzed graphite is highly ordered, its value asymptotically approaches zero.⁸² The values of I_D/I_G ratios for GO, GO-COOH, GO-TeN, and GO-TeN-Pd(0) have been unequivocally determined and found to be 1.18, 1.43, 1.75, and 1.82 respectively. This increase (*i.e.*, the progressive change) in the values of the ratio from GO to GO-TeN-Pd(0) signifies a concurrent diminishment in the dimensions of the sp² domains and an amplification in the degree of structural disorder and surface irregularities. This increase during the conversion of the 4 into the 5 may be used to infer the presence of Pd(0) nanoparticles in the 5.

High resolution-transmission electron microscopy (HR-TEM)

High resolution-transmission electron microscopic (HR-TEM) studies (Fig. 4) provide interesting information about the surface morphology, structure, and configuration of both GO and the nanocatalytic system 5 [*i.e.*, GO–TeN–Pd(0)]. The HR-TEM images reveal that GO [Fig. 4(a)] is made up of translucent sheets exhibiting wrinkles and folds. These features are indications of the successful exfoliation of graphene-stacked layers during the oxidation process. Such a process is



Fig. 4 TEM images: (a) GO at 100 nm, (b)–(d) GO–Te–Pd(0) at 100 nm, 10 nm and 5 nm respectively.



Fig. 5 Size distribution graph of Pd(0) nanoparticles present in the catalytic system.

a prerequisite for chemical functionalization of the sheets. Spherical Pd(0) NPs have size distribution with a mean particle size of \sim 9 nm (Fig. 5) are uniformly dispersed [Fig. 4(b)–(d)] over the surface of Te and N functionalized GO sheets. HR-TEM image Fig. 4(d) showing the lattice fringes have a distance of 0.233 nm, assigned to the Pd(0) (111) crystal plane.

High angle annular dark field scanning TEM (HAADF-STEM) images [Fig. 6(a) and (b)] of GO–TeN–Pd(0) also reveal the dispersion of palladium nanoparticles over the functionalized graphene oxide sheet. The intensity of the HAADF-STEM images depends on the thickness when the objects have homogeneous composition. This image shows constant intensity, indicating the presence of palladium nanoparticles.

Field emission-scanning electron microscopy (FE-SEM)

The FE-SEM images [Fig. 7(a) and (b)] of GO and GO–TeN–Pd(0) give intriguing insights. In the case of GO, a typical layer structure characteristic of graphene is observed [Fig. 7(a)]. Fig. 7(b) showcases the surface of telluroamine-functionalized GO immobilized with Pd(0) nanoparticles. The SEM-EDX spectrum (ESI: Fig. S1–S4†), unequivocally confirms the presence of palladium, tellurium, nitrogen, carbon and oxygen in the nanocatalyst [*i.e.*, GO–TeN–Pd(0)]. The atomic composition of



Fig. 6 HAADF-STEM image of GO–TeN–Pd(0) at (a) 1 μm; (b) 0.2 μm.



Fig. 7 FE-SEM images: (a) GO and (b) GO-Te-Pd(0).

the nanocatalyst, as quantified by EDX analysis, includes carbon (57.84 wt%), nitrogen (5.23 wt%), oxygen (22.47 wt%), palladium (9.86 wt%), and tellurium (4.61 wt%).

Electron energy loss spectroscopy (EELS)

Electron energy loss spectroscopic studies have been carried out (Fig. 8–13 and ESI: Fig. S9–S11[†]) to study the elemental composition of graphene oxide (GO) and the nanocatalytic system [*i.e.*, GO–TeN–Pd(0)]. For GO, major peaks at 284 eV and 532 eV, due to the presence of carbon and oxygen in the structure, are prominent. In GO–TeN–Pd(0), additional peaks at 401 eV for nitrogen and 572 eV for tellurium are also observed. Peak at 335 eV corresponding to the palladium is also visible in the EELS analysis of GO–TeN–Pd(0). These results support the formation of the solid support (GO) and the heterogeneous catalytic system [*i.e.*, GO–TeN–Pd(0)].

Fourier transform-infrared (FT-IR) spectroscopy

FT-IR spectroscopic investigations have been carried out. The results are helpful in the validation of the functionalization and chemical modification of GO and immobilization of Pd(0) nanoparticles onto the surface of graphene oxide (GO). The FT-IR spectrum of GO (ESI: Fig. S5†) exhibits distinct bands at 3430, 1724, 1619, and 1225 cm⁻¹. These bands correspond to the stretching vibrations of O–H, C=O, C=C, and C–O (epoxy) bonds, respectively. The values of these bands are in agreement with literature reports,⁸³ and hence corroborate the formation of graphene oxide. FT-IR spectrum of GO–COOH (ESI: Fig. S6†) shows the decrease in the intensity of the band for C–O bond of



Fig. 8 Low-loss EELS spectra of GO-TeN-Pd(0).



epoxy groups, and increase in the intensity of the band for C=O bond. It confirms the carboxylation and functionalization on the GO surface. When organotellurium ligand gets immobilized and the 4 is formed (Scheme 1), novel peaks emerge at 3400, 1695 and ~1125 cm⁻¹ in IR spectrum (ESI: Fig. S7†). They may be due to N-H, C=O and C-N stretching vibrations in the amide linkage. When GO-TeN-Pd(0) is formed (Scheme 1), clear reduction in the intensity of the band at 3430 cm⁻¹ (for O-H stretching) can be noticed in the IR spectrum (ESI: Fig. S5–S8†).⁸¹ The successful development of the GO-Te-Pd(0) nanocatalyst is validated through the emergence of novel bands at 3430, 1724, 1619, and 1225 cm⁻¹ and 3400, 1695 and ~1125 cm⁻¹ for carboxyl group, and amide group linkage. These bands have significant differences with respect to those observed in the IR spectrum of GO (ESI: Fig. S5 and S8†). Such



Fig. 10 Oxygen K-edge (a and b) EELS spectra of GO-TeN-Pd(0).



Fig. 11 Nitrogen K-edge (a and b) EELS spectra of GO-TeN-Pd(0).

differences are easily noticeable on matching spectrum of GO with that of GO–TeN–Pd(0).

Thermogravimetric analysis (TGA)

The TGA curve (Fig. 14) of GO–TeN–Pd(0) NPs exhibits a minor weight loss (approximately 9%) up to 100 °C. This loss may be attributed to the removal of the absorbed/adsorbed molecules of water/moisture. With the increase in the temperature beyond 250 °C, gradual weight loss occurs until complete degradation at 800 °C. Notably, GO–TeN–Pd(0) experiences a loss of 15% in weight up to 350 °C. The magnitude of this loss is significantly lesser than that observed in bare GO.⁸⁴ This comparative analysis indicates that GO attains higher stability after immobilization of organotellurium (Te, N ligand), and Pd(0)



Fig. 12 Tellurium M-edge (a and b) EELS spectra of GO-TeN-Pd(0).



Fig. 13 Palladium M-edge (a and b) EELS spectra of GO-TeN-Pd(0).

nanoparticles on its surface. A sharp decline in catalyst weight at around 400 °C indicates carbon skeleton pyrolysis. These findings imply that GO-TeN-Pd(0) is more thermally-stable than GO. This stability is suitable in context of its catalytic applications for C-C and C-O coupling reactions at 100 °C and 110 °C respectively.

Brunauer-Emmett-Teller (BET) studies

BET (Brunauer-Emmett-Teller) analysis has been done using N_2 gas within a range of 0.05 to 1.0 P/P_0 at a very low temperature (*i.e.*, 77 K) achieved with the help of liquid N_2 . The N_2 adsorption isotherms (Fig. 15) obtained at 150 °C provide indications about the existence of the structural differences between GO and GO-TeN-Pd(0). The N2 adsorption/desorption



Fig. 14 TGA curve of GO-TeN-Pd(0) NPs.



Volume adsorbed (cm³/gm STP)

0

TeN-Pd(0)

0.2 0.6 0.8 1.0 0.0 0.4 Relative Pressure (P/P₀) Fig. 15 N₂ adsorption-desorption isotherm of (a) GO and (b) GO-

isotherms [Fig. 15(a) and (b)] of GO and GO-TeN-Pd(0) exhibit characteristics aligning with type II and IV adsorption isotherms according to the International Union of Pure and Applied Chemistry (IUPAC) classification, along with a H3 hysteresis loop spanning from $P/P_{o} = 0.25$ to $P/P_{o} = 1.0$. The presence of this H3 hysteresis loop is attributed to the presence of plate-like particle aggregates, contributing to the formation of slit-shaped pores.85 This phenomenon arises due to the coexistence of macroporous and mesoporous structures within the materials. Such structures have relatively fewer micropores in the functionalized graphene material. Presence of microporous structures, which show type I adsorption isotherm, can be ruled out easily.

Specific BET surface area of GO and GO-TeN-Pd(0) have been determined experimentally. The value of such an area of the GO is 72.399 m² g⁻¹. It matches with the earlier reports.⁸⁶ Interestingly, this value for GO-TeN-Pd(0) is very low i.e., 27.024 $m^2 g^{-1}$ in comparison to that of GO. It can be attributed solely to the alteration of the textural surface due to ligand immobilization. Experimentally determined value $(0.0476 \text{ cm}^3 \text{ g}^{-1})$ of the pore volume of GO is smaller than that $(0.1089 \text{ cm}^3 \text{ g}^{-1})$ of GO-

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TeN-Pd(0). It may be the consequence of wet-chemical treatment and functional group modification on the graphene oxide surface.

Applications of GO–TeN–Pd(0) as heterogeneous catalyst in C–C coupling reaction

Suzuki–Miyaura cross coupling reaction is the most powerful synthetic methods for the synthesis of carbon–carbon bond formation of high value chemicals such as natural products, agrochemical derivatives, pharmaceutical drugs, polymers, and biologically active compounds.⁷¹⁻⁷⁴ The catalytic potential of GO–TeN–Pd(0) has been explored for the first time for the catalysis of Suzuki–Miyaura cross coupling reaction.

Reaction conditions, which include solvent, base, reaction temperature, time and catalyst quantity, have been optimized for the reaction of 4-bromobenzonitrile with phenylboronic acid. It has been considered as a model reaction for conducting the study of catalysis.

The results indicated that the best reaction conditions leading to high % conversions were use of potassium carbonate (K₂CO₃) as a base, aqueous DMF as solvent, 20 mg as the quantity of the catalyst (i.e., 0.108 mol% Pd), 12 hours as the reaction time and 100 °C temperature (Table 1: entry 1). Maximum % conversion i.e., 94% was achieved under these reaction conditions (Table 1: entry 1). However, when GO or GO-TeN (without Pd loading) was used as a catalyst, only trace quantities of the coupled products have been obtained (Table 1: entries 3 and 4). When this model reaction was evaluated using Na₂PdCl₄ and PdCl₂ as a catalyst (without stabilizing ligand and solid support), the % conversion of products was relatively low i.e., 61% and 38% respectively (Table 1: entries 5 and 6). When NaOH was used as the base, only 47% conversion has been observed (Table 1: entry 8). When pure DMF was used as a solvent without mixing it with water, suppression in catalytic performance has been observed (Table 1: entry 7). Use of

Under the optimized reaction conditions, a variety of structurally divergent bromoarenes and chloroarenes can be converted (Table 2) into their biphenyl products. Such products have been authenticated by matching their proton NMR spectra with those reported for the known compound.87,88 The efficiency of catalytic processes varies with the nature of haloarenes for the coupling reactions. Easy removal of halide group is usually facilitated by the presence of electron-withdrawing group on phenyl ring. This enhances the positive character at para position and hence favours the nucleophilic attack in the coupling reaction. The outcomes for the C-C coupling reactions also demonstrate the same. Excellent % conversions (i.e., >94%) are achieved (Table 2: entries 1, 2, 3 and 6) for the substrates (aryl bromides) having electron withdrawing substituents (-CHO, -COMe, -CN and -NO₂) at para-position. Unlike these substrates, when 4-bromotoluene and 4-bromoanisole (bromoarenes with electron donating substituents) are used as substrates, the % conversions are significantly low (Table 2: entries 4 and 8), i.e., 73% and 61% respectively. The catalyst also performs well when aryl chlorides, the least reactive substrates, are used (Table 1: entries 9-13).

Applications of GO–TeN–Pd(0) as heterogeneous catalyst for C–O coupling

The O-arylation reaction of phenol plays a crucial role in synthetic organic chemistry and finds application in the

Table 1 Results of the optimization of the reaction conditions for C–C coupling reaction between phenylboronic acid and 4-bromobenzonitrile^a



S. no.	Catalyst	Base	Time	Temp.	Solvent	% conversion
1	GO-TeN-Pd(0)	K ₂ CO ₃	12 h	100 °C	DMF:water	$\sim 94\%$
2	GO-TeN-Pd(0)	K ₂ CO ₃	5 h	100 °C	DMF: water	${\sim}57\%$
3	GO	K ₂ CO ₃	12 h	100 °C	DMF: water	Trace
4	GO-TeN	K ₂ CO ₃	12 h	100 °C	DMF: water	Trace
5	Na_2PdCl_4	K ₂ CO ₃	12 h	100 °C	DMF: water	${\sim}61\%$
6	PdCl ₂	K ₂ CO ₃	12 h	100 °C	DMF: water	$\sim 38\%$
7	GO-TeN-Pd(0)	K ₂ CO ₃	12 h	100 °C	DMF	\sim 71%
8	GO-TeN-Pd(0)	NaOH	12 h	100 °C	DMF: water	${\sim}47\%$
9	GO-TeN-Pd(0)	K_2CO_3	12 h	100 °C	Water	Trace
10	GO-TeN-Pd(0)	K ₂ CO ₃	12 h	Room temp.	DMF: water	${\sim}42\%$
11	GO-TeN-Pd(0)	K_2CO_3	12 h	100 °C	DMF: water	${\sim}85\%^b$

^{*a*} Reaction conditions: 4-bromobenzonitrile (1.0 mmol); phenylboronic acid (1.2 mmol); catalyst (20 mg, 0.108 mol% Pd); DMF : water (4 : 1) 5 mL. ^{*b*} Catalyst – 10 mg.

Table 2 Results of C–C coupling reaction of different aryl halides with phenylboronic acid under optimized reaction conditions^a

	, ↓ ↓ ↓ ↓ ↓		
S. no.	Aryl halide	Product	% conversion
1	4-Bromobenzaldehyde		~99%
2	4-Bromoacetophenone	$H_{3C} = \frac{1}{7}$	~97%
3	4-Bromobenzonitrile	1-([1,1'-Biphenyl]-4-yl)ethanone	~94%
4	4-Bromotoluene	[1,1'-Biphenyl]-4-carbonitrile $H_3C \longrightarrow g$	~73%
5	4-Bromobenzoic acid		$\sim 85\%$
6	1-Bromo-4-nitrobenzene	[1,1'-Biphenyi]-4-carboxylic acid	~99%
7	Bromobenzene	4-Nitro-1,1'-biphenyl	~83%
8	4-Bromoanisole		~61%
9	Chlorobenzene	4-Methoxy-1,1'-biphenyl	~78%
10	4-Chloroaniline	Biphenyl H ₂ N 14	~76%
11	1-Chloro-4-nitrobenzene	[1,1'-Biphenyl]-4-amine O_2N	~87%
12	4-Chloroacetophenone	4-Nitro-1,1'-biphenyl H_{3C}	~83%
13	4-Chlorobenzophenone	1-([1,1'-Biphenyl]-4-yl)ethanone Ph [1,1'-Biphenyl]-4-yl(phenyl)methanone	~75%

^{*a*} Reaction conditions: aryl halide (1.0 mmol); phenylboronic acid (1.2 mmol); K_2CO_3 (2.0 mmol); DMF + water (5.0 mL); catalyst (20 mg, 0.108 mol% Pd); time, 12 h; bath temperature, 100 °C.

Table 3 Results of optimization of the O-arylation reaction between phenol and 1-bromo-4-nitrobenzene^a



S. no	Catalyst	Base	Time	Temp.	Solvent	% conversion
1	GO-TeN-Pd(0)	K_2CO_3	16 h	110 °C	DMSO	\sim 99%
2	GO	K_2CO_3	16 h	110 °C	DMSO	$\sim 13\%$
3	GO-TeN	K_2CO_3	16 h	110 °C	DMSO	${\sim}10\%$
4	Na_2PdCl_4	K_2CO_3	16 h	110 °C	DMSO	${\sim}58\%$
5	GO-TeN-Pd(0)	NaOH	16 h	110 °C	DMSO	\sim 77%
6	GO-TeN-Pd(0)	K_2CO_3	16 h	110 °C	EtOH	${\sim}48\%$
7	GO-TeN-Pd(0)	K_2CO_3	16 h	110 °C	H_2O	Trace
8	GO-TeN-Pd(0)	K_2CO_3	16 h	Room temp.	DMSO	${\sim}51\%$
9	GO-TeN-Pd(0)	K_2CO_3	6 h	110 °C	DMSO	\sim 57%

^a Reaction conditions: 1-bromo-4-nitrobenzene (1 mmol); phenol (1.1 mmol), solvent (5.0 mL); catalyst 20 mg (0.108 mol% Pd).

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synthesis of numerous industrially significant products.^{89–91} The GO–TeN–Pd(0) nanocatalyst also has the potential to catalyze C–O coupling reaction between phenol and a diverse range of substituted chloroarenes and bromoarenes. Preliminary investigations were conducted on the reaction between 1-bromo-4-nitrobenzene and phenol to optimize the reaction conditions (Table 3). The results (Table 3: entry 1) indicate that the best reaction conditions include the use of dimethyl sulfoxide (the solvent), presence of K_2CO_3 (the base), 110 °C temperature, 20 mg quantity of GO–TeN–Pd(0) nanocatalyst and reaction time of 16 hours.

The results of catalysis of C–O coupling reactions (Table 4) of different aryl halides (substrates) are highly interesting. The conversions achieved are remarkably high. Under the optimized reaction conditions, a variety of structurally divergent bromoarenes and chloroarenes can be converted (Table 4) into their diphenyl ether products. Such products have been authenticated by matching their proton NMR spectra with those reported for the known compounds.⁸⁷ The efficiency of the catalytic processes varies depending on the nature of the bromoarenes and chloroarenes.

The nanocatalytic system [*i.e.*, GO-TeN-Pd(0)] shows the ability to arylate the phenol even with some chloroarenes (e.g., 1-chloro-4-nitrobenzene, 4-chlorobenzonitrile, chlorobenzene, and chlorotoluene) with significant conversions ranging from 49 to 97% (Table 4: entries 2, 4, 6 and 8). It is worth noting that only a few heterogeneous nanocatalytic systems have been reported to exhibit such ability.41,42,45,87,92-95 The coupling reactions involving 1-bromo-4-nitrobenzene with phenol exhibited the highest conversion of approximately 99% (Table 4: entry 1). Good conversion rates (57-99%) of the desired products were achieved in the coupling reactions of 1-bromo-4-nitrobenzene, 4-bromobenzonitrile, bromobenzene, and bromotoluene with phenol (Table 4: entries 1, 3, 5 and 7). The presence of electronwithdrawing substituents on the aromatic ring of the aryl halides led to higher reactivity (Table 4: entries 1-4). However, the presence of an electron-donating group (*i.e.*, CH₃) on the

aryl halide (Table 4: entries 7 and 8) has a deactivating effect, resulting in lower product yields in the reaction.

To conclude the nature of catalysis, hot filtration tests were carried out for both Suzuki coupling and C-O coupling reactions. 4-Bromobenzonitrile was used as the substrate for Suzuki coupling and 1-chloro-4-nitrobenzene was used to arylate phenol in C-O coupling reaction. After 4 hours of reaction, the catalyst was extracted from the reaction mixtures via centrifugation. At this stage, the conversion of the reaction mixture, as assessed through ¹H-NMR spectroscopy, was 72% for Suzuki coupling and 84% for C-O coupling. Subsequently, the reaction mixtures were divided into two equal portions. In one portion, the catalyst [GO-TeN-Pd(0)] was reintroduced, and the reaction was allowed to continue for additional 8 hours. The other portion was subjected to heating and stirring without adding any catalyst. Interestingly, the conversion of the reactant into the product remained almost unchanged in the reaction mixtures (for both Suzuki coupling and C-O coupling) of the portion which did not contain the catalyst. On the other hand, % conversions in other sets of experiments (in which the catalyst was reintroduced in the reaction mixture) increased from 72% to 94% (for Suzuki coupling) and from 84% to 97% (for C-O coupling reaction). These observations and results provide compelling and unequivocal evidence for the heterogeneous nature of the catalysis and rule out any ambiguity regarding the leaching of Pd(0) NPs or other Pd(0) species from the heterogeneous catalyst, i.e., GO-TeN-Pd(0), into the reaction mixture.

Recyclability of GO-TeN-Pd(0) heterogeneous catalyst

From both environmental and economic perspectives, the ability to recover and reuse a catalyst offers distinct advantages. This particular aspect of the present catalyst was meticulously examined in the context of the Suzuki coupling as well as C–O coupling reactions. In Suzuki coupling, the recyclability was studied for the reaction between phenylboronic acid and 1-bromo-4-nitrobenzene under optimized conditions. In the case of C–O coupling reaction, the experiments were conducted for

Table 4 O-Arylation reaction of different aryl halides with phenol under optimized reaction conditions⁴

X

_OH

0.

	R ₁ +		
S. no.	Aryl halide	Product	% conversion
1	1-Bromo-4-nitrobenzene	0 ₂ N 16	~99%
2	1-Chloro-4-nitrobenzene	1-Nitro-4-phenoxy benzene	~97%
3	4-Bromobenzonitrile	1-Nitro-4-phenoxy benzene	~79%
4	4-Chlorobenzonitrile	4-Phenoxybenzonitrile	~71%
5	Bromobenzene	4-Phenoxybenzonitrile 18	~83%
6	Chlorobenzene	Diphenyl ether	~76%
7	4-Bromotoluene		\sim 57%
8	4-Chlorotoluene	1-Methyl-4-phenoxybenzene H ₃ C 19 1-Methyl-4-phenoxybenzene	$\sim 49\%$

^{*a*} Reaction conditions: aryl halide (1.0 mmol); phenol (1.2 mmol); K₂CO₃ (2.0 mmol); catalyst (20 mg, 0.108 mol% Pd); time, 16 h; bath temperature, 110 °C; DMSO, 5.0 mL.

the reaction between 1-chloro-4-nitrobenzene and phenol to understand the reuse of the catalyst. The results (Fig. 16) unequivocally demonstrate that the catalyst GO–TeN–Pd(0) shows almost similar catalytic activity up to four reaction cycles. Thereafter, a slight decrease in fifth cycle has been observed.

After successfully completing five reaction cycles in the C–O coupling reaction, comprehensive examination and characterization of the catalyst have been done using P-XRD (Fig. 17), FE-SEM (Fig. 18), and EDX (ESI: Fig. S12–S14, Table S3†) techniques. The results of these analyses convincingly demonstrate the presence of active catalytic sites in the catalytic system even after the completion of five reaction cycles. It gives strong indication about the remarkable robustness and stability of the catalyst.

The 20 mg quantity of the catalytic system, GO–TeN–Pd(0), is applied to catalyze the reactions. Analysis, done using ICP-OES technique, indicates that it is equivalent to catalyst loading of 0.108 mol%. In the reaction of bromobenzene (PhBr) with phenylboronic acid in the solvent mixture of DMF and H₂O at 100 °C in the presence of K₂CO₃ (base) for 12 hours, 83% conversion has been achieved. This system is quite efficient in comparison to some other heterogeneous catalysts reported for the Suzuki–Miyaura cross-coupling reaction.^{87,96–110} For instance, Pd–Fe₃O₄ heterodimer nanocrystals (Table 5: entry 1)



Fig. 16 Recyclability of the GO–TeN–Pd(0) in the C–C and C–O coupling reaction under optimized conditions.

work to catalyse the reaction [between bromobenzene and phenylboronic acid] in DME/H2O in presence of Na2CO3 and give the product in 70% yield after 24 hours of reaction under reflux conditions when the concentration of such crystals in the reaction mixture is 1.0 mol%.96 Similarly, for the catalysis of the reaction [between 4-bromoaniline and phenylboronic acid] in ethanol in the presence of sodium carbonate (Na_2CO_2) at 70 °C. 1.0 mol% quantity of the HMMS-NH₂-Pd (Table 5: entry 2) is required to give the product in 74% yield after 16 hours.97 Fe₃O₄@CS-SB@Pd is a very interesting catalyst from the point of view of reaction time, catalyst loading and temperature.¹⁰³ It allows the formation of the product in the reaction [between bromobenzene and phenylboronic acid] in very high (99%) yield in extremely short (30 minutes) reaction time (Table 5: entry 8). It works at 0.2 mol% concentration at 50 °C in PEG-200 i.e., the solvent in the presence of potassium carbonate (K₂CO₃).¹⁰³ In terms of catalyst loading, the present heterogeneous catalyst *i.e.*, GO-TeN-Pd(0) system shows remarkable performance. For the reaction [between bromobenzene and phenylboronic acid], it performs the catalysis at low catalyst loading and gives the



Fig. 17 P-XRD of recycled GO-TeN-Pd(0) nanocatalyst.



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Fig. 18 FE-SEM image (at 500 nm) of recycled GO-TeN-Pd(0) nanocatalyst.

product in high yield. Pd@graphene oxide is a known heterogeneous catalyst¹⁰² which works at 6.4 mol% concentration (Table 5: entry 7) to lead the formation of the product in 80% yield. The Pd-adenine@boehmite is an attractive catalyst because it allows the catalysis to occur in water.¹⁰⁴ It is also better than the present catalyst in terms of other parameters such as reaction time and temperature (Table 5: entry 9). It can give the product in appreciable yield *i.e.*, 90% [for the reaction between bromobenzene and phenylboronic acid] at 80 °C within 1.4 hours As far as the concentration of the catalyst is concerned, its performance is not at par with the present catalyst (Table 5: entry 9).104 It works when it is used in 0.65 mol% concentration in the presence of Na2CO3. Overall, the GO-TeN-Pd(0) catalytic system shows good performance. Its ability to perform at low catalyst loading is worth highlighting. Moreover, it also has the ability to convert aryl chlorides into the coupled products.

Ability to function at low concentration (*i.e.*, loading) is an attractive feature of the present catalyst i.e., GO-TeN-Pd(0) for O-arylation of phenol (i.e., C-O coupling reaction) too. Its ability to perform the catalysis of the arylation reaction of phenol even with chloroarene makes it an attractive system. Such ability is found only with a few heterogeneous nanocatalytic systems (Table 6: entries 1-8). When it is used in 0.108 mol% concentration, 76% conversion has been achieved for the reaction [between chlorobenzene and phenol] in DMSO in the presence of potassium carbonate (i.e., K₂CO₃) at 110 °C in 16 hours. Its comparison with other known catalysts is quite interesting.41,42,45,87,92-96 For instance, Fe3O4@SiO2@SePh@Pd(0) works at 1.0 mol% concentration and gives the product in 70% yield (Table 6: entry 1) in aqueous medium in the presence of NaOH at 80 °C.87 Fe₃O₄@SiO₂-BT-Cu is attractive because it has an inexpensive metal i.e., copper. However, it works at nearly 18 times higher concentration (i.e., equal to 2.0 mol%) in comparison to the present catalyst to produce the desired compound in 71% yield (Table 6: entry 2) for the reaction [between chlorobenzene and phenol] in eutectic solvent (i.e., PhCl/urea) in presence of K₂CO₃ at 100 °C in 24 hours.⁹² The

Table 5 Comparison with known heterogeneous catalyst for Suzuki–Miyaura cross coupling reaction between bromoarene/derivatives and phenylboronic acid

Entry	Catalytic system	Catalyst loading	Reaction conditions	Yield (%)	Reference
1	Pd–Fe ₃ O ₄ heterodimer nanocrystals	1.0 mol%	Na ₂ CO ₃ , DME/water, reflux 24 h	70	96
2	HMMS–NH ₂ –Pd	1.0 mol%	K ₂ CO ₃ , ethanol, 70 °C, 16 h	74^a	97
3	Pd(II)–NHC complex	1.0 mol%	K ₂ CO ₃ , DMA, 100 °C, 24 h	67	98
4	NHC–Pd(II) complex	1.0 mol%	Cs ₂ CO ₃ , toluene, 80 °C, 12 h	54	99
5	Pd NPs@CMC/AG	1.0 mol%	K ₂ CO ₃ , H ₂ O, 60 °C, 0.5 h	80^a	100
6	Pd NPs@APC	1.8 mol%	K_2CO_3 , MW, 5 min	88 ^{<i>a</i>}	101
7	Pd@graphene oxide framework	6.4 mol%	K ₂ CO ₃ , DMF, 80 °C, 8 h	80	102
8	Fe ₃ O ₄ @CS-SB@Pd	0.2 mol%	K ₂ CO ₃ , PEG-200, 50 °C, 30 min	99	103
9	Pd-adenine@boehmite	0.65 mol%	Na ₂ CO ₃ , H ₂ O, 80 °C, 1.4 h	90	104
10	OCMCS-SB-Pd(II)	0.4 mol%	K ₂ CO ₃ , EtOH/H ₂ O, 50 °C, 2 h	96	105
11	NHC–Pd(II) complex	1.0 mol%	K ₂ CO ₃ , DMA, 100 °C, 24 h	67	106
12	OCMCS-3a-Pd	0.04 mol%	K ₂ CO ₃ , toluene, 100 °C, 48 h	70^a	107
13	Pd-graphene hybrids (~4 nm Pd NPs)	1.1 mol%	K ₃ PO ₄ , H ₂ O, SDS, 100 °C, 5 min	29	108
14	GO-Pd ₄ S	0.5 mol%	K ₂ CO ₃ , ethanol/water, 80 °C, 6 h	78	109
15	GO–Pd ₄ S nanospheres	0.2 mol%	K ₂ CO ₃ , ethanol/water, TBAB, 80 °C, 5 h	19	110
16	GO-TeN-Pd(0)	0.108 mol%	DMF/H ₂ O, K ₂ CO ₃ , 100 °C, 12 h	83 ^b	Present system

GO-Pd₁₇Se₁₅ system is one of those rare and interesting catalytic system which have the ability to perform the reaction at room temperature.⁴¹ It works at 1.0 mol% loading, and gives the product in slightly higher (*i.e.*, 73%) yield in a short reaction of 3 hours in DMSO/K₂CO₃ (Table 6: entry 3).⁴¹ Cu NPs-reduced graphene oxide is another catalyst which has low-cost metal *i.e.*, copper. Its low quantity (50 mg) is sufficient to catalyse the reaction [between 1-chloro-4-nitrobenzene and phenol] and give the product in 66% yield (Table 6: entry 5) at 120 °C in DMSO in 12 hours in the presence of Cs₂CO₃ *i.e.*, the base which is more expensive than K₂CO₃.⁴⁵ There are some other catalytic systems (Table 6: entries 6–9) which are known for this reaction. They work at higher catalyst loading in comparison to the present system.^{93-95,111}

Experimental section

Materials and chemicals

Graphite fine powder, bromobenzene and its derivatives was supplied by M/S Alfa Aesar. Phenylboronic acid, ethyl alcohol,

N,N-dimethylformamide (DMF), chloroacetic acid, potassium carbonate, sodium borohydride, 2-chloroethylamine hydrochloride, diphenylditelluride, sodium tetrachloropalladate, potassium permanganate, sodium nitrate, hydrogen peroxide, and thionyl chloride were procure from Merck Life Science Private Limited (India). Other chemicals and solvents were purchased from local suppliers and utilized in the same form as received without any purification or drying. To assess the conversions attained in the Suzuki–Miyaura and C–O cross coupling processes, ¹H-NMR spectroscopic technique was used.

Instrumentation and analysis

The ¹H-NMR spectra were acquired using a JNM ECX-500 NMR spectrophotometer operating at 500 MHz. The JEM 3200FS electron microscope, which runs at 300 kV, was used to perform TEM investigations at the Department of Metallurgical and Materials Engineering in Indian Institute of Technology, Roorkee. In order to prepare the sample for analysis using transmission electron microscopy (TEM), the powder was dispersed properly and uniformly in methanol utilizing a probe sonicator.

Table 6 Comparison table with heterogeneous catalysts reported in literature for C–O coupling reaction between phenol and chloroarenes					
Entry	Catalytic system	Catalyst loading	Reaction conditions	Yield (%)	Reference
1	Fe ₃ O ₄ @SiO ₂ @SePh@Pd(0)	1.0 mol%	H ₂ O, NaOH, 80 °C, 6 h	70	87
2	Fe ₃ O ₄ @SiO ₂ -BT-Cu	2.0 mol%	PhCl/urea, K ₂ CO ₃ , 100 °C, 24 h	71	92
3	$GO-Pd_{17}Se_{15}$	1.0 mol%	DMSO, K ₂ CO ₃ , r.t., 3 h	73	41
4	GO-Cu _{1.8} S	1.25 mol%	DMSO, K ₂ CO ₃ , 120 °C, 8 h	58	42
5	Cu NPs-reduced graphene oxide	50 mg	DMSO, Cs ₂ CO ₃ , 120 °C, 12 h	66 ^{<i>a</i>}	45
6	CuI–zeolite (USY)	10.0 mol%	DMF, Cs ₂ CO ₃ , 120–140 °C, 24 h	65	93
7	CuO-UiO-66-NH2-MLm	50 mg	DMSO, KOH, 150 °C, 24 h	30	94
8	$Fe_3O_4@SiO_2@PPh_2@Pd(0)$	1.5 mol%	H ₂ O, NaOH, 80 °C, 3.5 h	83	95
9	Pd ₁₆ S ₇ NPs	0.5 mol%	DMSO, K ₂ CO ₃ , 100 °C, 3 h	64^b	111
10	GO-TeN-Pd(0)	0.108 mol%	DMSO, K2CO2, 110 °C, 16 h	76^{c}	Present system

^{*a*} 1-Chloro-4-nitrobenezene. ^{*b*} Bromobenzene. ^{*c*} % conversion.

The slurry was deposited onto a copper grid with a porous carbon film and allowed to dry in the air. In addition, electron energy loss spectroscopy (EELS) was performed using a JEM 3200FS electron microscope. The analysis of surface morphology and elemental composition was performed using FEI-Quanta FEG 200F model of scanning electron microscope with energy-dispersive X-ray spectroscopic (SEM-EDX) equipment. Powder X-ray diffraction (P-XRD) experiments were conducted using a SmartLab 9 kW rotating anode X-ray diffractometer. X-ray photoelectron spectroscopic (XPS) studies were conducted using the Nexsa base (Thermo Fisher) spectrometer. The Raman spectra were acquired using a Horiba LAB RAM HR evolution confocal microdispersive Raman spectrometer, with a 532 nm solid-state laser operating in the standard mode. BET studies were performed using the BEL-SORP Mini X instrument. TGA analysis was carried out using Shimadzu DTG-60 instrument to understand the thermal stability of the catalyst. FT-IR spectra were recorded on Shimadzu IRSpirit to study the change in the functional groups during the process of the formation of the catalytic system. Palladium content of the GO-TeN-Pd(0) catalyst was determined by ICP-OES analysis using Thermo Fisher Scientific iCAP PRO instrument. When it was required, an inert atmosphere using nitrogen gas was created and maintained using Schlenk techniques. Most of the reactions were conducted under ambient conditions, utilizing oven-dried glassware.

Synthesis of graphene oxide (GO)

Hummers' method was utilized to convert graphite powder into graphene oxide.75 In this method, 1.0 g of graphite powder was mixed with 0.75 g of sodium nitrate (NaNO₃) in 34 mL of concentrated H₂SO₄, and allowed to stir for one hour at 5 °C. Thereafter, 4.5 grams of potassium permanganate $(KMnO_4)$ were introduced slowly into the reaction mixture. During this, the temperature was maintained at around 5 °C. This reaction mixture was agitated for five days at 25 °C. Thereafter, it was heated at 90 °C for 2 hours and mixed gently with 50 mL of diluted H₂SO₄ (5 wt%) along with continuous stirring. Thereafter, 2.7 mL of hydrogen peroxide (30 wt%) was added to the mixture and stirring was allowed to continue for another 3 hours. The bright yellow precipitate of graphene oxide (GO) appeared. It was filtered, and subjected to washing. It was initially rinsed with 30 mL of H₂SO₄, followed by 10 mL of H₂O₂, and ultimately with 300 mL of HCl (3 wt%). This procedure was repeated until the pH of the filtrate reached a neutral state. In the final step, it was subjected to vacuum drying before its subsequent utilization.

Functionalization of graphene oxide *i.e.* carboxylation of graphene oxide (GO-COOH)

The wet chemical treatment process was employed to functionalize graphene oxide (GO) sheets (*i.e.*, **1**) with hydroxyl, epoxide, and carboxyl groups. The process begins with mixing graphene oxide (*i.e.*, **1**) (200 mg) in distilled water (100 mL) followed by ultrasonication for an hour to obtain exfoliated graphene oxide nanosheets. Subsequently, a slurry of graphene oxide is prepared by adding chloroacetic acid (1.0 g) and NaOH (1.2 g). It is subjected to sonication for 2 hours, for obtaining the 2. The pH of the resulting suspension of GO–COOH (*i.e.*, 2) is adjusted to neutral using dilute HCl. Thereafter, distilled water is used for washing to remove any traces of HCl. The wet material is dried under vacuum conditions.

Preparation of GO-TeN: immobilization of (Te, N) bidentate ligand (Ph-Te-CH₂-CH₂-NH₂) on graphene oxide sheets

In order to obtain the acyl chloride derivative of the 2 (i.e., GO-COOH), a suspension of its 200 mg quantity was subjected to sonication in 2.0 mL of N,N-dimethylformamide (DMF) for an hour. Thereafter, the suspension was stirred with an excess amount of thionyl chloride at moderate temperature (*i.e.*, 65 °C) for a duration of 12 hours. It resulted in the formation of GO-COCl (i.e., 3). The excess of thionyl chloride was eliminated using a vacuum distillation method. The GO-COCl was subjected to centrifugation, followed by washing with tetrahydrofuran (THF). Subsequently, it was dried under vacuum conditions. Thereafter, 100 mg of GO-COCl was mixed with N,N-dimethylformamide (DMF) and 108 mg of 2-(phenyltellanyl)ethanamine. The mixture was refluxed under inert atmosphere of nitrogen. In the final step, the material (i.e., 4) was separated with the help of centrifugation using ethanol, and ultimately dried under vacuum conditions.

Synthesis of the heterogeneous nanocatalytic system [GO-TeN-Pd(0)]

In the first step, a suspension was made by sonicating the mixture of 50 mg of GO–TeN (*i.e.*, the 4) with distilled water for one hour. In the next step, a solution of 30 mg of Na_2PdCl_4 in water (10 mL) was poured into the suspension along with constant stirring. The resulting mixture was left as such for 12 hours. The pH of the mixture was raised to approximately 13 by introducing a solution of NaOH (1.0 M) in distilled water. The alkaline mixture was agitated for 6 hours at moderate temperature (*i.e.*, 50 °C). In the final step, the 5 [*i.e.*, GO–TeN–Pd(0)] was washed many times with distilled water to eliminate any residual alkali. Similar washing was also done with ethanol. The material was dried under vacuum conditions.

Procedure for C-C coupling reaction

An oven-dried round bottom flask was employed to combine haloarene (1.0 mmol), phenylboronic acid (1.2 mmol), K_2CO_3 (2.0 mmol), and 20 mg (0.108 mol% Pd) of the 5 [*i.e.*, GO–TeN– Pd(0)] in a solvent mixture (5.0 mL) prepared by mixing *N*,*N*dimethylformamide (DMF) and water in 4 : 1 ratio. The mixture was allowed to reflux for a duration of twelve (12) hours. Thereafter, the mixture was filtered to extract the nanocatalytic species. The filtrate was cooled to room temperature and subjected to extraction with diethyl ether. Diethyl ether component was washed twice with distilled water. It was dried using anhydrous Na₂SO₄. The dried solution was subjected to evaporation under vacuum to remove the solvent and obtain the residue. The residue was analyzed using ¹H-NMR spectroscopy to ascertain the percentage conversion.

Procedure for C-O coupling reaction

For the C–O coupling reaction, a properly dried round-bottom flask was employed to combine 1.0 mmol of haloarene, 1.2 mmol of phenol, 2.0 mmol of K_2CO_3 , and 20 mg (0.108 mol% Pd) of the 5 [*i.e.*, GO–TeN–Pd(0) catalyst]. The mixture was refluxed in 5.0 mL of dimethyl sulfoxide (DMSO) for 16 hours. Thereafter, the nanocatalyst was filtered out and separated. The filtrate was allowed to cool to room temperature. It was mixed and shaken with ethyl acetate to extract the organic compounds. This solution of ethyl acetate was washed twice with distilled water. It was dried using anhydrous Na₂SO₄. In order to obtain the residue, the solvent was removed under vacuum. ¹H-NMR spectroscopic technique was used to analyze the residue and ascertain the percentage conversion.

Hot filtration test for Suzuki coupling

Hot filtration tests were conducted for both C–O coupling and Suzuki reactions. In the C–C coupling reaction setup, the GO– TeN–Pd(0) nanocatalyst was mixed with 4-bromobenzonitrile (1.0 mmol), phenylboronic acid (1.1 mmol), and K_2CO_3 (2.0 mmol) in 5.0 mL of the solvent mixture, prepared using DMF and water in 4:1 ratio in a round bottom flask. The reaction mixture was stirred at 100 °C for 4 hours. Thereafter, the catalyst was removed using centrifugation. The percentage of conversion was determined in the mixture at this juncture. The centrifugate was split into two portions. In one part, the catalyst was reintroduced in the same concentration. The second part did not contain the catalyst. Both portions were subjected to heating and stirring further for 8 hours. Thereafter, the reactions were quenched, and the conversions were estimated using ¹H-NMR analysis.

Hot filtration test for C-O coupling reaction

The GO–TeN–Pd(0) nanocatalyst was mixed with 1-chloro-4nitrobenzene (1.0 mmol), phenol (1.1 mmol), and K_2CO_3 (2.0 mmol) in a properly dried round bottom flask in 5.0 mL of the solvent *i.e.*, DMSO. The reaction mixture was stirred at 110 °C for 2.5 hours. Thereafter, the catalyst was separated through centrifugation, and the conversion percentage was determined. The centrifugate was evenly divided into two portions. In one portion, the catalyst was reintroduced in the same concentration. The second part remained devoid of the catalyst. Both portions were heated along with stirring for 8 hours. The reactions were worked-up. The conversions were evaluated through ¹H-NMR analysis.

Procedure for testing the recyclability for Suzuki coupling

1-Bromo-4-nitrobenzene (1.0 mmol), phenylboronic acid (1.2 mmol), K_2CO_3 (2.0 mmol) were mixed with the GO–TeN–Pd(0) nanocatalyst (50 mg) in a properly dried round bottom flask in 5.0 mL of solvent mixture prepared using DMF and water in 4 : 1 ratio. This mixture was stirred at 100 °C. Thereafter, the cross-coupled product was extracted using diethyl ether, and the 5 [*i.e.*, the nanocatalyst GO–TeN–Pd(0)] was isolated from the solution *via* centrifugation. It was subjected to drying for

subsequent utilization in a fresh C–C cross-coupling reaction. Such reactions were repeated five times.

Procedure for testing the recyclability for C-O coupling

1-Chloro-4-nitrobenzene (1.0 mmol), phenol (1.1 mmol), K_2CO_3 (2.0 mmol), and 50 mg of the 5 [*i.e.*, GO–TeN–Pd(0)] were mixed in 5.0 mL of the solvent *i.e.*, DMSO. The mixture was stirred at 110 °C for 16 hours. Thereafter, the cross-coupled product was extracted using ethyl acetate. The 5 [*i.e.*, GO–TeN–Pd(0) nanocatalyst] was separated *via* centrifugation and subjected to drying. The catalyst was used for a fresh C–O cross-coupling reaction. This process was repeated five times, and the percentage of the conversions were estimated in every cycle using proton NMR spectroscopy.

Conclusion

In conclusion, an organotellurium ligand has been used for the first time to the best of our knowledge for the development of a heterogeneous catalyst. Ph-Te-CH₂-CH₂-NH₂ has been used for the functionalization of the solid support *i.e.*, graphene oxide. The immobilization of the bidentate organotellurium ligand [i.e. (2-phenyltelluro)ethylamine] having Te (i.e., metalloid), and N donor sites was achieved through formation of covalent linkages. Due to the presence of such donor sites, the organotellurium ligand not only holds the nanoparticles of palladium(0) but also stabilizes them in a narrow size range. The use of such ligands as a stabilizer in heterogenized form will open a new field of research for developing applied nanomaterials. The novel and robust heterogeneous catalyst [GO-TeN-Pd(0)], obtained by covalently grafting hybrid (Te, N) ligand over graphene oxide sheets, holds and stabilizes the catalytically active Pd(0) nanoparticles. The role of the Te and N donor sites is likely to stabilize Pd(0) and facilitate the homogeneous distribution of Pd(0) across the graphene oxide sheets. The sites hold the nanoparticles strongly and prevent leaching of catalytically active sites. This system shows remarkable properties and potential as a heterogeneous catalyst for Suzuki-Miyaura cross coupling and O-arylation of phenols for a wide range of electronically divergent aryl halides including aryl chlorides. It catalyzes the reactions at 0.108 mol% loading. It is reusable and recyclable. Recyclability studies, carried out up to five reaction cycles, show that some loss in the performance of the catalyst occurs when it is recycled. The recycled catalyst has also been characterized using P-XRD (Fig. 17), FE-SEM (Fig. 18), and EDX (ESI: Fig. S12-S14, Table S3[†]) techniques, which corroborate the retention of active catalytic sites in its recycled form.

Data availability

Relevant data are within the paper and its ESI files.[†] Other data and parameters generated or analyzed during the study are available from the corresponding author upon reasonable request.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

S. P. acknowledge the Department of Science and Technology (DST) for INSPIRE Fellowship [DST/INSPIRE Fellowship/2021/ IF210150]. A. T. acknowledge the Department of Science and Technology (DST) for INSPIRE Fellowship [DST/INSPIRE Fellowship/2021/IF210531]. A. K. acknowledge the support of Department of Science and Technology (DST) for funding under the PURSE scheme [File Number: SR/PURSE/2023/199].

References

- 1 K. M. Engle and J.-Q. Yu, J. Org. Chem., 2013, 78, 8927-8955.
- 2 S. T. Kim, S. Kim and M. H. Baik, *Chem. Sci.*, 2020, **11**, 1017–1025.
- 3 E. Ocansey, J. Darkwa and B. C. E. Makhubela, *RSC Adv.*, 2018, **18**, 13826–13834.
- 4 H. V. Huynh, Chem. Rev., 2018, 118, 9457-9492.
- 5 G. K. Rao, A. Kumar, M. P. Singh, A. Kumar, A. M. Biradar and A. K. Singh, *J. Organomet. Chem.*, 2014, **753**, 42–47.
- 6 K. N. Sharma, H. Joshi, A. K. Sharma, O. Prakash and A. K. Singh, *Organometallics*, 2013, **32**, 2443–2451.
- 7 A. Kumar, G. K. Rao, S. Kumar and A. K. Singh, *Organometallics*, 2014, 33, 2921–2943.
- 8 D. B. Eremin and V. P. Ananikov, *Coord. Chem. Rev.*, 2017, **346**, 2–19.
- 9 R. Martin and S. L. Buchwald, Acc. Chem. Res., 2008, 41, 1461-1473.
- 10 M. Garcia-Melchor, A. A. Braga, A. Lledós, G. Ujaque and F. Maseras, *Acc. Chem. Res.*, 2013, **46**, 2626–2634.
- 11 D. J. Durand and N. Fey, Chem. Rev., 2019, 119, 6561-6594.
- 12 C. L. McMullin, N. Fey and J. N. Harvey, *Dalton Trans.*, 2014, 43, 13545–13556.
- 13 P. Oswal, A. Arora, S. Singh, G. K. Rao, S. Kumar, A. K. Singh and A. Kumar, *Catal. Commun.*, 2021, **149**, 106242.
- 14 S. Kumar, G. K. Rao, A. Kumar, M. P. Singh, F. Saleem and A. K. Singh, *RSC Adv.*, 2015, 5, 20081–20089.
- 15 G. K. Rao, A. Kumar, M. Bhunia, M. P. Singh and A. K. Singh, J. Hazard. Mater., 2014, 269, 18–23.
- 16 P. Oswal, A. Arora, S. Singh, D. Nautiyal, S. Kumar, G. K. Rao and A. Kumar, *Dalton Trans.*, 2020, **49**, 12503.
- 17 A. Arora, P. Oswal, D. Sharma, A. Tyagi, S. Purohit, P. Sharma and A. Kumar, *ChemistrySelect*, 2022, 7, e202201704.
- 18 A. Arora, P. Oswal, D. Sharma, S. Purohit, A. Tyagi, P. Sharma and A. Kumar, *Dalton Trans.*, 2022, **51**, 17114– 17144.
- 19 S. Purohit, R. Rana, A. Tyagi, A. Bahuguna, P. Oswal and A. Kumar, *Org. Biomol. Chem.*, 2024, **22**, 6215–6245.
- 20 P. Sharma, A. Arora, P. Oswal, G. K. Rao, J. Kaushal, S. Kumar and S. Kumar, *Polyhedron*, 2019, **171**, 120–127.

- 21 A. Arora, P. Oswal, G. K. Rao, J. Kaushal, S. Kumar, A. K. Singh and A. Kumar, *ChemistrySelect*, 2019, 4, 10765–10769.
- 22 P. Oswal, A. Arora, S. Gairola, A. Datta and A. Kumar, *New J. Chem.*, 2021, **45**, 21449–21487.
- 23 A. Arora, P. Oswal, S. Singh, D. Nautiyal, G. K. Rao, S. Kumar, A. K. Singh and A. Kumar, *Inorg. Chim. Acta*, 2021, 522, 120267.
- 24 J. Kaushal, S. Singh, D. Nautiyal, G. K Rao, A. K. Singh and A. Kumar, *New J. Chem.*, 2022, **46**, 14757–14781.
- 25 A. Tyagi, S. Purohit, P. Oswal, S. Rawat, V. Negi, A. K. Singh and A. Kumar, *New J. Chem.*, 2023, **47**, 12511–12535.
- 26 A. Arora, P. Oswal, G. K. Rao, S. Kumar, A. K. Singh and A. Kumar, *Catal. Lett.*, 2022, 1–13.
- 27 A. Arora, P. Oswal, A. Datta and A. Kumar, *Coord. Chem. Rev.*, 2022, **459**, 214406.
- 28 P. Oswal, A. Arora, S. Purohit, A. Bahuguna, P. Sharma, J. Roy and A. Kumar, *New J. Chem.*, 2023, **47**, 4346–4354.
- 29 P. Singh, D. Das, A. Kumar and A. K. Singh, *Inorg. Chem. Commun.*, 2012, **15**, 163–166.
- 30 A. Bahuguna, P. Oswal, S. Purohit, A. Tyagi and A. Kumar, New J. Chem., 2024, 48, 14140–14144.
- 31 G. K. Rao, A. Kumar, M. P. Singh and A. K. Singh, J. Organomet. Chem., 2014, 749, 1-6.
- 32 A. S. Burange, and M. B. Gawande, *Encyclopedia of Inorganic* and *Bioinorganic Chemistry*, 2011, pp. 1–19.
- 33 J. C. Védrine, ChemSusChem, 2019, 12, 577-588.
- 34 M. Opanasenko, P. Štěpnička and J. Čejka, RSC Adv., 2014, 4, 65137–65162.
- 35 G. Fan, F. Li, D. G. Evans and X. Duan, *Chem. Soc. Rev.*, 2014, **43**, 7040–7066.
- 36 N. M. Julkapli and S. Bagheri, *Int. J. Hydrogen Energy*, 2015, 40, 948–979.
- 37 X. Fan, G. Zhang and F. Zhang, *Chem. Soc. Rev.*, 2015, 44, 3023-3035.
- 38 S. Navalon, A. Dhakshinamoorthy, M. Alvaro and H. Garcia, *Coord. Chem. Rev.*, 2016, **312**, 99–148.
- 39 F. R. Hartley and P. N. Vezey, Adv. Organomet. Chem., 1977, 15, 189–234.
- 40 K. Nejati, S. Ahmadi, M. Nikpassand, P. D. K. Nezhad and E. Vessally, *RSC Adv.*, 2018, **8**, 19125–19143.
- 41 H. Joshi, K. N Sharma, A. K. Sharma and O. P. A. K. Singh, *Chem. Commun.*, 2013, **49**, 7483–7485.
- 42 V. V. Singh and A. K. Singh, *ACS Appl. Nano Mater.*, 2018, 1, 2164–2174.
- 43 R. Zahedi, Z. Asadi and F. D. Firuzabadi, *Colloids Surf., A*, 2019, **580**, 123728.
- 44 M. Hosseini-Sarvari and Z. Razmi, *RSC Adv.*, 2014, 4, 44105–44116.
- 45 M. Nasrollahzadeh, M. Maham, A. Rostami-Vartooni, M. Bagherzadeh and S. M. Sajadi, *RSC Adv.*, 2015, 5, 64769–64780.
- 46 P. Puthiaraj and W. S Ahn, *Catal. Sci. Technol.*, 2016, 6, 1701–1709.
- 47 A. Heuer-Jungemann, N. Feliu, I. Bakaimi, M. Hamaly,A. Alkilany, I. Chakraborty, A. Masood, M. F. Casula,A. Kostopoulou, E. Oh, K. Susumu, M. H. Stewart,

I. L. Medintz, E. Stratakis, W. J. Parak and A. G. Kanaras, *Chem. Rev.*, 2019, **119**, 4819–4880.

- 48 L. M. Rossi, J. L. Fiorio, M. A. Garcia and C. P. Ferraz, *Dalton Trans.*, 2018, 47, 5889–5915.
- 49 M. Nasrollahzadeh, Z. Issaabadi, M. M. Tohidi and M. S. Sajadi, *Chem. Rec.*, 2018, 18, 165–229.
- 50 S. Eigler and A. Hirsch, Angew. Chem., Int. Ed., 2014, 53, 7720-7738.
- 51 V. Georgakilas, M. Otyepka, A. B. Bourlinos, V. Chandra, N. Kim, K. C. Kemp, P. Hobza, R. Zboril and K. S. Kim, *Chem. Rev.*, 2012, **112**, 6156–6214.
- 52 P. Oswal, K. Sood, S. Singh, A. Arora, A. Bahuguna, S. Purohit and A. Kumar, *Dalton Trans.*, 2022, **51**, 6537– 6542.
- 53 P. Oswal, A. Arora, S. Singh, D. Nautiyal, S. Kumar and A. Kumar, *New J. Chem.*, 2022, 46, 3578–3587.
- 54 V. S. Sapner, P. P. Chavan and B. R. Sathe, *ACS Sustainable Chem. Eng.*, 2020, **8**, 5524–5533.
- 55 S. Naghdi, M. Sajjadi, M. Nasrollahzadeh, K. Y. Rhee,
 S. M. Sajadi and B. Jaleh, *J. Taiwan Inst. Chem. Eng.*, 2018,
 86, 158–173.
- 56 S. S. Narwade, S. M. Mali, V. S. Sapner and B. R. Sathe, *ACS Appl. Nano Mater.*, 2020, **3**, 12288–12296.
- 57 M. Nasrollahzadeh, Z. Nezafat, M. G. Gorab and M. Sajjadi, Mol. Catal., 2020, 484, 110758.
- 58 A. V. Munde, B. B. Mulik, R. P. Dighole and B. R. Sathe, New J. Chem., 2020, 44, 15776–15784.
- 59 V. S. Sapner, B. B. Mulik, R. V. Digraskar, S. S. Narwade and B. R. Sathe, *RSC Adv.*, 2019, 9, 6444–6451.
- 60 D. S. Avila, P. Gubert, A. Palma, D. Colle, D. Alves, C. W. Nogueira, J. B. T. Rocha and F. A. A. Soares, *Brain Res. Bull.*, 2008, **76**, 114–123.
- 61 D. S. Avila, A. Soares, and W. G. Salgueiro, Toxicology and pharmacology of organotellurium compounds, *Tellurium: Properties, Uses and Research*, Nova Science Publishers, Uruguaiana, Brazil, 2017, pp. 137–169.
- 62 D. S. Avila, D. Colle, P. Gubert, A. S. Palma, G. Puntel, F. Manarin, S. Noremberg, P. C. Nascimento, M. Aschner, J. B. T. Rocha and F. A. A. Soares, *Toxicol. Sci.*, 2010, 115, 194–201.
- 63 V. Jamier, L. A. Ba and C. Jacob, *Chem.-Eur. J.*, 2010, **16**, 10920–10928.
- 64 A. L. Braga, E. E. Alberto, L. C. Soares, J. B. T. Rocha, J. H. Sudati and D. H. Roos, *Org. Biomol. Chem.*, 2009, 7, 43-45.
- 65 T. H. Degrandi, I. M. d'Oliveira, G. S. d'Almeida, C. R. L. Garcia, I. V. Villela, T. N. Guecheva, R. M. Rosa and J. A. P. Henriques, *Mutagenesis*, 2010, **25**, 257–269.
- 66 C. W. Nogueira, G. Zeni and J. B. T. Rocha, *Chem. Rev.*, 2004, 104, 6255–6286.
- 67 A. K. Singh and S. Sharma, *Coord. Chem. Rev.*, 2000, 209, 49– 98.
- 68 V. K. Jain and R. S. Chauhan, *Coord. Chem. Rev.*, 2016, **306**, 270–301.
- 69 P. Tian, F. Xuan, D. Ding, Y. Sun, X. Xu, W. Li, R. Si, J. Xu and Yi-F. Han, *J. Catal.*, 2020, **385**, 21–29.

- 70 A. Arora, P. Oswal, G. K. Rao, S. Kumar, A. K. Singh and A. Kumar, *RSC Adv.*, 2021, **11**, 7214–7224.
- 71 S. Arshadi, A. Banaei, A. Monfared, S. Ebrahimiasl and A. Hosseinian, *RSC Adv.*, 2019, 9, 17101–17118.
- 72 C. Sambiagio, S. P. Marsden, A. J. Blacker and P. C. McGowan, *Chem. Soc. Rev.*, 2014, **43**, 3525–3550.
- 73 E. Voutyritsa, A. Theodorou, M. G. Kokotou and C. G. Kokotos, *Green Chem.*, 2017, **19**, 1291–1298.
- 74 Y. S. Won, Y. H. Ji, L. E. Mi and L. S. Young, Adv. Synth. Catal., 2018, 360, 278–283.
- 75 W. S. Hummers and R. E. Offeman, *J. Am. Chem. Soc.*, 1958, **80**, 1339.
- 76 J. Zakrzewski, B. Huras, A. Kielczewska and M. Krawczyk, *RSC Adv.*, 2016, 6, 98829–98834.
- 77 S. Guo, J. Raya, D. Ji, Y. Nishina, C. Ménard-Moyon and A. Bianco, *Nanoscale Adv.*, 2020, 2, 4085–4092.
- 78 N. Hussain, A. Borah, G. Darabdhara, P. Gogoi,
 V. K. Azhagan, M. V. Shelke and M. R. Das, *New J. Chem.*, 2015, 39, 6631–6641.
- 79 J. Liu, X. Peng, W. Sun, Y. Zhao and C. Xia, *Org. Lett.*, 2008, 10, 3933–3936.
- 80 Y. H. Ng, M. Wang, H. Han and C. C. L. Chai, *Chem. Commun.*, 2009, 37, 5530–5532.
- 81 G. Pang, M. Sun, P. Liu, H. Liu, I. Hou and F. Gao, RSC Adv., 2016, 6, 73469–73474.
- 82 A. Ferrari, Solid State Commun., 2007, 143, 47-57.
- 83 R. Bhaskar, H. Joshi, A. K. Sharma and A. K. Singh, ACS Appl. Mater. Interfaces, 2017, 9, 2223–2231.
- 84 M. E. Assal, M. R. Shaik, M. Kuniyil, M. Khan, A. Al-Warthan, M. R. H. Siddiqui, S. M. A. Khan, W. Tremel, M. N. Tahir and S. F. Adil, *RSC Adv.*, 2017, 7, 55336–55349.
- 85 Heterogeneous Photocatalysis: Relationships with Heterogeneous Catalysis and Perspectives, ed. G. Marcì, and L. Palmisano, Elsevier, 2019.
- 86 A. Ariharan, B. Viswanathan and V. Nandhakumar, *Graphene*, 2017, **6**, 41–60.
- 87 A. K. Sharma, H. Joshi and A. K. Singh, *RSC Adv.*, 2020, 10, 6452–6459.
- 88 P. Oswal, A. Arora, J. Kaushal, G. K. Rao, S. Kumar, A. K. Singh and A. Kumar, *RSC Adv.*, 2019, 9, 22313–22319.
- 89 J. Zhang, Z. Zhang, Y. Wang, X. Zheng and Z. Wang, *Eur. J.* Org Chem., 2008, 2008, 5112.
- 90 E. Sperotto, G. P. M. van Klink, J. G. de Vries and G. van Koten, *Tetrahedron*, 2010, **66**, 9009.
- 91 G. Evano, N. Blanchard and M. Toumi, *Chem. Rev.*, 2008, 108, 3054–3131.
- 92 M. Niakan, S. Karimi, M. Masteri-Farahani and H. Shekaari, *Colloids Surf.*, A, 2021, **620**, 126603.
- 93 V. Magné, T. Garnier, M. Danel, P. Pale and S. Chassaing, Org. Lett., 2015, 17, 4494–4497.
- 94 S. Sadeghi, M. Jafarzadeh, A. R. Abbasi and K. Daasbjerg, New J. Chem., 2017, 41, 12014–12027.
- 95 M. A. Zolfigol, V. Khakyzadeh, A. R. Moosavi-Zare, A. Rostami, A. Zare, N. Iranpoor, M. H. Beyzavi and R. Luque, *Green Chem.*, 2013, 15, 2132–2140.

- 96 Y. Jang, J. Chung, S. Kim, S. W. Jun, B. H. Kim, D. W. Lee,
 B. M. Kim and T. Hyeon, *Phys. Chem. Chem. Phys.*, 2011, 13, 2512–2516.
- 97 P. Wang, F. Zhang, Y. Long, M. Xe, R. Li and J. Ma, *Catal. Sci. Technol.*, 2013, **3**, 1618–1624.
- 98 Q. Xu, W. L. Duan, Z. Y. Lei, Z. B. Zhu and M. Shi, *Tetrahedron*, 2005, **61**, 11225.
- 99 T. Chen, J. Gao and M. Shi, Tetrahedron, 2006, 62, 6289.
- 100 T. Baran, Ultrason. Sonochem., 2018, 45, 231.
- 101 T. Baran, Carbohydr. Polym., 2018, 195, 45.
- 102 T. P. N. Tran, A. Thakur, D. X. Trinh, A. T. N. Dao and T. Taniike, *Appl. Catal., A*, 2018, **549**, 60–67.
- 103 A. S. Kumari, S. Layek and D. D. Pathak, *New J. Chem.*, 2017, 41, 5595–5604.
- 104 B. Tahmasbi, A. Ghorbani-Choghamarani and P. Moradi, New J. Chem., 2020, 44, 3717–3727.

- 105 Y. Dong, J. Bi, S. Ming, S. Zhang, D. Zhu, D. Meng and T. Li, *Carbohydr. Polym.*, 2021, **260**, 117815.
- 106 Q. Xu, W. Duan, Z. Lei, Z. Zhu and M. Shi, *Tetrahedron*, 2005, **61**, 11225–11229.
- 107 T. Baran, E. Açıksöz and A. Menteş, *J. Mol. Catal. A: Chem.*, 2015, **407**, 47–52.
- 108 Y. Li, X. Fan, J. Qi, J. Ji, S. Wang, G. Zhang and F. Zhang, *Nano Res.*, 2010, **3**, 429–437.
- 109 H. Joshi, K. N. Sharma, A. K. Sharma and A. K. Singh, *Nanoscale*, 2014, **6**, 4588–4597.
- 110 V. V. Singh, U. Kumar, S. N. Tripathi and A. K. Singh, *Dalton Trans.*, 2014, **43**, 12555–12563.
- 111 P. Singh and A. K. Singh, *Dalton Trans.*, 2017, **46**, 10037–10049.