

Cite this: *Nanoscale Adv.*, 2019, 1, 3177

# Supercritical water assisted preparation of recyclable gold nanoparticles and their catalytic utility in cross-coupling reactions under sustainable conditions†

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Preparation of gold nanoparticles (AuNPs) in environmentally friendly water without using any reducing agents under supercritical conditions is demonstrated. PXRD, XPS, FE-SEM and HR-TEM analysis confirmed the formation of phase-pure and crystalline AuNPs of the size of ~10–30 nm. The catalytic potential of AuNPs was manifested through a generalized green procedure that could accommodate both Sonogashira as well as Suzuki coupling under aqueous conditions at low catalytic loading (0.1 mol%). The AuNP catalyst was found to be recuperated after the reaction and reused for up to six catalytic cycles with no leaching out of gold species as confirmed through ICP-OES analysis. With no confinement of AuNP catalysis to cross-coupling reaction, synthetic extension to one-flask preparation of  $\pi$ -conjugated semiconductors (4 examples) and their optoelectronic properties were also investigated. Other significant features of the present work include short reaction time, site-selectivity, wide substrate scope, high conversion, good chemical yields and applicability in gram-scale synthesis. Overall, the results of this paper signify an operationally sustainable supercritical fluid processing method for the synthesis of AuNPs and their catalytic application towards cross-coupling reactions in green media.

Received 12th April 2019  
Accepted 26th June 2019

DOI: 10.1039/c9na00240e

rsc.li/nanoscale-advances

## 1. Introduction

Noble metal nanoparticles (*viz.* Ag, Au, Pt and Pd) have received considerable attention in recent years, due to their unique morphology-dependent electronic, optical and catalytic properties.<sup>1,2</sup> Because of the robust and green nature of gold catalysts, organic transformations based on gold nanoparticles (AuNPs) have received a special place in the area of heterogeneous gold catalysis.<sup>3–10</sup> The main advantage that AuNPs possess compared to their bulk form is the high surface area to volume ratio (*sa/vol*) and modifiable shapes which lead to high chemical reactivity. Their easy recyclability, reuse and environmentally friendly appeal established their catalytic potential for a myriad of organic reactions. Based on the preparation method, the size of AuNPs or their distribution on the support can be controlled, which would directly influence the reactivity as well as selectivity of the given chemical transformation.

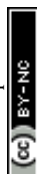
Therefore, preparation of AuNPs with controlled size and shape is of utmost significance not only to catalytic but also towards other applications such as sensors, biomedical devices and energy storage/conversion.<sup>11–13</sup> It is appropriate to mention that most of the chemical methods require citric acid, hydrazine or sodium borohydride as reducing agents for the synthesis of AuNPs<sup>14,15</sup> and thus possess significant environmental impacts.<sup>16</sup> Reportedly, this issue was addressed through the rapid reduction of  $[\text{AuCl}_4]^-$  species in sonicated water. However, it leads to the formation of triangular and hexagonal Au nanoplates coexisting with spherical AuNPs with a diameter of ~40 nm.<sup>17</sup> To circumvent this limitation, a preparative method for the controlled growth of AuNPs without using reducing agents/additional immobilizing stabilizers/rare earth oxide support in an environmentally benign route is of paramount importance. With no confinement to gold, recent years have witnessed many greener synthetic routes for the preparation of an array of transition metal nanoparticles.<sup>18</sup> Especially, the synthesis, characterization and application of different metallic nanoparticles constitute an important class of catalytic systems in cross-coupling reactions.<sup>19–24</sup> On the other hand, it is pertinent to mention that the catalytic potential of AuNPs is exemplified in oxidative esterification and hydrogen transfer reactions.<sup>25</sup> As part of our research interest in new reaction development<sup>26–30</sup> based on gold catalysis,<sup>31–37</sup> we became

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† Electronic supplementary information (ESI) available. See DOI: 10.1039/c9na00240e

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interested in the sustainable preparation of AuNPs and subsequent catalytic application in cross-coupling reactions. In particular, we desired to develop a simple and practical procedure for AuNP catalyzed Sonogashira and Suzuki reactions. Differing from previous gold catalysis disclosures on these reactions (Fig. 1), our working hypothesis comprises a generalized procedure that could accommodate both Sonogashira<sup>38–45</sup> as well as Suzuki<sup>46–49</sup> reactions. Keeping in mind the principles of green chemistry, our focus is also to execute the above-mentioned transformations in aqueous media.<sup>50,51</sup>

As far as the preparative method is concerned, huge consideration has been paid in recent years for the synthesis of nanomaterials with precisely defined morphology and dimensions using supercritical fluid (SCF) processing. In this approach, the temperature and pressure of fluids/solvents are increased to their critical points to attain a homogeneous SCF medium, wherein the SCF phase has both gas and liquid-like properties. This condition allows the rapid and facile preparation of nanomaterials in high yields in a shorter reaction time.<sup>52–54</sup> Moreover, SCF exhibits interesting physicochemical properties such as high diffusion coefficient, excellent wetting

of surfaces, low interfacial tension and low viscosity. Indeed, these are some of the critical parameters for the control of nucleation and growth of nanomaterials.<sup>55</sup> Hence, nanomaterials with the desired size and shape without using any surfactants can be accomplished by varying the fluid/temperature/pressure. Recently, AuNPs have been shown to deposit on the surface of graphene oxide nanosheets,<sup>56</sup> on supported silica<sup>57</sup> and inside the pores of MCM-48 using supercritical carbon dioxide (scCO<sub>2</sub>)<sup>58</sup> for various important applications. In pursuance of our continuing efforts in SCF promoted nanoparticle/sheet synthesis,<sup>59–64</sup> we envisaged the supercritical fluid assisted direct conversion of tetrachloroauric acid trihydrate (HAuCl<sub>4</sub>·3H<sub>2</sub>O) to AuNPs using water as the green solvent. Our idea was to employ water both as a fluid as well as a base catalyst for the preparation of AuNPs by sequential reduction and decomposition of aqueous HAuCl<sub>4</sub>·3H<sub>2</sub>O by SCF processing. This is because of the fact that supercritical water can form a homogeneous phase with inorganic/organic substances and virtually serves as a base catalyst under supercritical conditions.<sup>65</sup> Under these conditions the critical temperature, pressure and density of water molecule are 374 °C, 22.06 MPa and

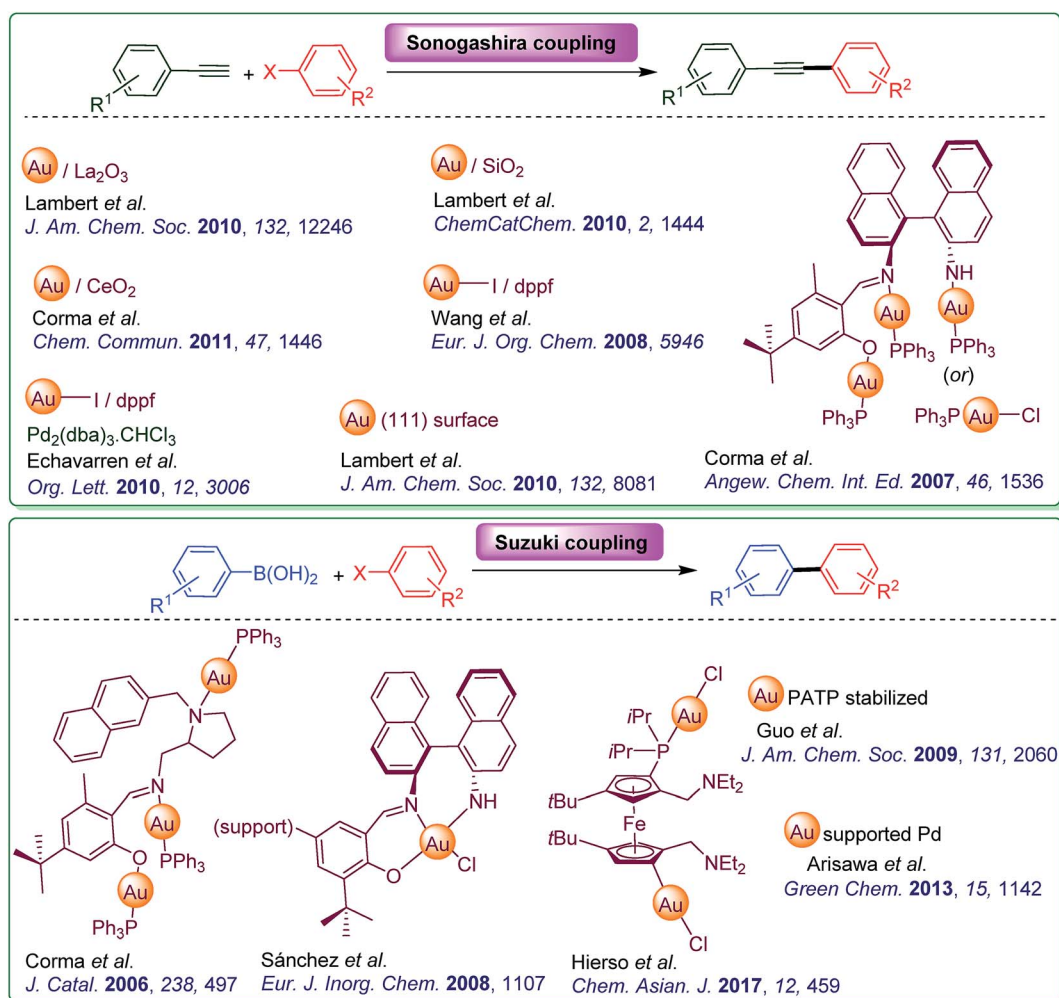


Fig. 1 Literature precedents of gold catalyzed Sonogashira and Suzuki coupling.



0.32 g mL<sup>-1</sup> respectively. Herewith, we report the sustainable preparation of AuNPs from HAuCl<sub>4</sub>·3H<sub>2</sub>O using water as a green solvent without using any reducing agents. The as-prepared AuNPs were thoroughly characterized by PXRD, XPS, UV-vis, FE-SEM and HR-TEM analysis followed by the demonstration of their catalytic potential towards Sonogashira and Suzuki cross-coupling reactions under aqueous conditions. To highlight the synthetic utility of the developed methodology, we also established the one-pot synthesis of  $\pi$ -semiconductors along with optoelectronics investigation and all results are compiled in this paper.

## 2. Experimental section

### 2.1 Chemicals, methods and instruments

HAuCl<sub>4</sub>·3H<sub>2</sub>O (99.995% trace metal basis), halobenzenes, arylboronic acids, and terminal acetylenes were purchased from Sigma-Aldrich. All chemicals and solvents were used as received without further purification. All reactions were carried out in ultra-clean glass vials and stir bars. Analytical TLC was performed on pre-coated plastic sheets of silica gel G/UV-254 of 0.2 mm thickness (Macherey-Nagel, Germany) using analytical grade solvents and visualized with iodine spray (10% w/w I<sub>2</sub> in silica gel), UV light ( $\lambda = 254$  and 365 nm) and alkaline KMnO<sub>4</sub> solution. Column chromatography was performed using thick-walled glass columns along with a mixture of petroleum ether and ethyl acetate on silica gel (100–200 mesh, SRL, India). The relative proportions of solvents in chromatography solvent mixtures refer to the volume to volume ratio (v/v). FT-IR spectra were recorded on a Bruker TENSOR 27 spectrometer in the frequency range from 400 to 4000 cm<sup>-1</sup> using the KBr pellet technique. <sup>1</sup>H and <sup>13</sup>C NMR spectra were obtained in CDCl<sub>3</sub> on a BRUKER spectrometer at 400 and 100 MHz respectively. Proton chemical shifts ( $\delta$ ) are relative to tetramethylsilane (TMS,  $\delta = 0.00$ ) as internal standard and expressed in parts per million (ppm). The number of protons ( $n$ ) for a given resonance is indicated as  $n$ H. Spin multiplicities are given as s (singlet), d (doublet), t (triplet) and m (multiplet). Coupling constants ( $J$ ) are given in hertz (Hz). High resolution mass spectra (HRMS) were obtained using a Q-TOF and a JEOL-GCMate-II mass spectrometer. GC-MS spectra were obtained using a PerkinElmer instrument. Inductively coupled plasma (ICP) measurements were done using Agilent Technologies 5110-ICP-OES equipped with an auto sampler capacity of 240 tubes. For ICP-OES analysis, 1.77 g of Et<sub>3</sub>PAuCl (based on the calculation of molecular weight of Et<sub>3</sub>PAuCl/atomic weight of gold) was made up to 1 L using 2% HNO<sub>3</sub> solution. This corresponds to 1000 ppm of gold standard and was used as the stock solution. 1 mL of filtrate obtained after removing the solid AuNP catalyst was made up to 10 mL using a standard gold solution in a Teflon vessel and subjected to microwave digestion. This was again made up to 50 mL and used for ICP-OES analysis. UV-visible spectra for all solutions and thin film phases were recorded as chloroform solutions and a thin film respectively on a SHIMADZU UV-3600 plus UV-VIS-NIR spectrometer. Photoluminescence spectra were recorded on a Jasco spectrofluorometer FP-8500. Cyclic voltammetry (CV) experiments were

performed with an Autolab potentiostat galvanostat (Model PGSTAT-30). All CV measurements were carried out at room temperature with a conventional three-electrode configuration employing glassy carbon as the working electrode, silver wire as the pseudo-reference electrode and platinum wire as the counter electrode. Tetrabutylammonium perchlorate (0.1 M) as the supporting electrolyte was used in argon-purged acetonitrile at a scan rate of 50 mV s<sup>-1</sup>. A thermal evaporator with a sputtering unit (model no. VRSU04D) and thickness monitor (model no. CTM-200) was used for chemical vapour deposition. The OPV device performance was measured by recording the current-voltage characteristics using a potentiostat (Keithley 2400) under illumination from a Solar Simulator (Science-Tech) with an AM1.5G filter set and an illumination intensity of 100 mW cm<sup>-2</sup>.

### 2.2 Catalyst characterization

The formation of AuNPs was confirmed using powder X-ray diffraction (PXRD) analysis, which was performed on a BRUKER D8 ADVANCE X-ray diffractometer with Cu K $\alpha$  radiation ( $\alpha = 1.5418$  Å) from  $2\theta = 10$  to  $80^\circ$  in steps of  $0.02^\circ$  and a count time of 0.2 s. UV-visible absorption spectra were measured using a VARIAN (Cary 500 Scan) UV-vis-NIR spectrophotometer system. The surface morphology of AuNPs was characterized by field emission scanning electron microscopy (FE-SEM) and high-resolution transmission electron microscopy (HR-TEM) analysis. SEM and HR-TEM images were captured at different locations of the AuNP coated substrate using a Carl Zeiss AG, Supra 55VP with an accelerating voltage of 0.1–30 kV and Tecnai G2 F20S-Twin working at an accelerating voltage of 200 kV, respectively. The formation of AuNPs was further confirmed by X-ray photoelectron spectroscopy (XPS) analysis using a Thermo Scientific MULTILAB 2000 base system with X-ray, Auger and ISS attachments using Mg K $\alpha$  (1253.6 eV) as the X-ray source.

## 3. Results and discussion

### 3.1 Catalyst preparation and characterization

At the outset, we intended to prepare AuNPs by reducing highly pure HAuCl<sub>4</sub>·3H<sub>2</sub>O (99.995% trace metal basis).<sup>66–68</sup> In a typical synthesis, 250 mg of HAuCl<sub>4</sub>·3H<sub>2</sub>O was dissolved in 25 mL of distilled water and sonicated for 5 min at room temperature. This homogeneous solution was filled in a stainless-steel supercritical reactor (35 mL volume capacity) and both ends of the reactor were tightly closed (Fig. 2). Then, the sealed supercritical reactor was placed in a preheated tubular furnace at 400 °C for 60 min and subsequently the hot reactor was cooled in an ice-cold water bath. The resulting AuNPs formed were collected by washing the precipitate with distilled water and ethanol using repeated centrifugation. The powder sample obtained was dried in a hot air oven at 80 °C for 12 h. It is worth mentioning that there was no product formation when the reaction time was less than 60 min. If the reaction time was allowed for >60 min at high temperature (>400 °C), doping of chromium and iron (leached out from the supercritical reactor) elements on the surface of AuNPs increases. Therefore, we



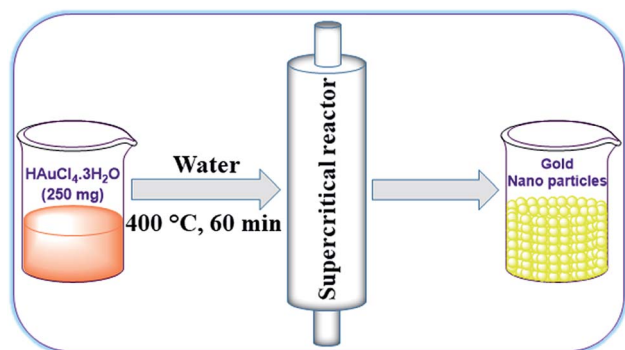


Fig. 2 Preparation of AuNPs in supercritical water.

preferred a reaction condition of 60 min at 400 °C for the synthesis of AuNPs.

Initially, powder X-ray diffraction (PXRD) analysis was used to confirm the formation of AuNPs. Fig. 3a shows the XRD pattern of as-synthesized AuNPs (bottom panel) and the standard pattern of Au (top panel). The obtained well-defined diffraction lines of the as-synthesized material are in good agreement with the reference pattern of the face centred cubic (fcc) structure of Au (JCPDS card no. 04-0784), which revealed the formation of AuNPs. The diffraction lines also demonstrate that the obtained AuNPs are highly crystalline in nature even without applying any external thermal treatments to improve crystallinity. The observed diffraction lines at  $2\theta = 38.19^\circ$ ,

$44.39^\circ$ ,  $64.58^\circ$  and  $77.55^\circ$  correspond to the (111), (200), (220) and (311) lattice planes of the cubic phase of AuNPs respectively. It is pertinent to note that no other reflection lines can be found along with the AuNP diffraction peaks, which confirms the phase purity of the prepared AuNPs. Thus, these findings emphasize the advantage of the SCF approach in producing phase pure nanomaterials with high crystallinity in a short reaction time. The well-known Scherrer formula ( $D = K\lambda/\beta \cos \theta$ ) has been used to calculate the crystallite size of AuNPs. Here,  $D$  is the crystallite size of the synthesized sample,  $K$  is the dimensionless shape factor,  $\lambda$  is the wavelength of the X-ray beam,  $\beta$  is the full width half maximum (FWHM) and  $\theta$  is the diffraction angle. The calculated crystallite size at  $2\theta = 38.19^\circ$ ,  $44.39^\circ$ ,  $64.58^\circ$  and  $77.55^\circ$  was found to be 29.15, 23.08, 39.04 and 35.27 nm, respectively. Thus, the average crystallite size of the synthesized AuNPs is 31.6 nm.

The chemical state of the synthesized AuNPs was explicitly confirmed by X-ray photoelectron spectroscopic (XPS) analysis. Fig. 3b represents the deconvoluted Au 4f spectra of AuNPs which show their characteristic doublets at 83.7 and 87.4 eV which are in good agreement with the binding energies of the Au 4f<sub>7/2</sub> and Au 4f<sub>5/2</sub> spin orbit pair respectively.<sup>69–73</sup> These values clearly correspond to the metallic Au(0) state. It also ascertains that supercritical water serves the purpose of both fluid as well as base to effectively reduce the Au(III) ions to Au(0) metal. XPS results unambiguously confirmed that under SCF conditions, the formation of AuNPs can be accomplished without using any reducing agents. Hence, the SCF approach

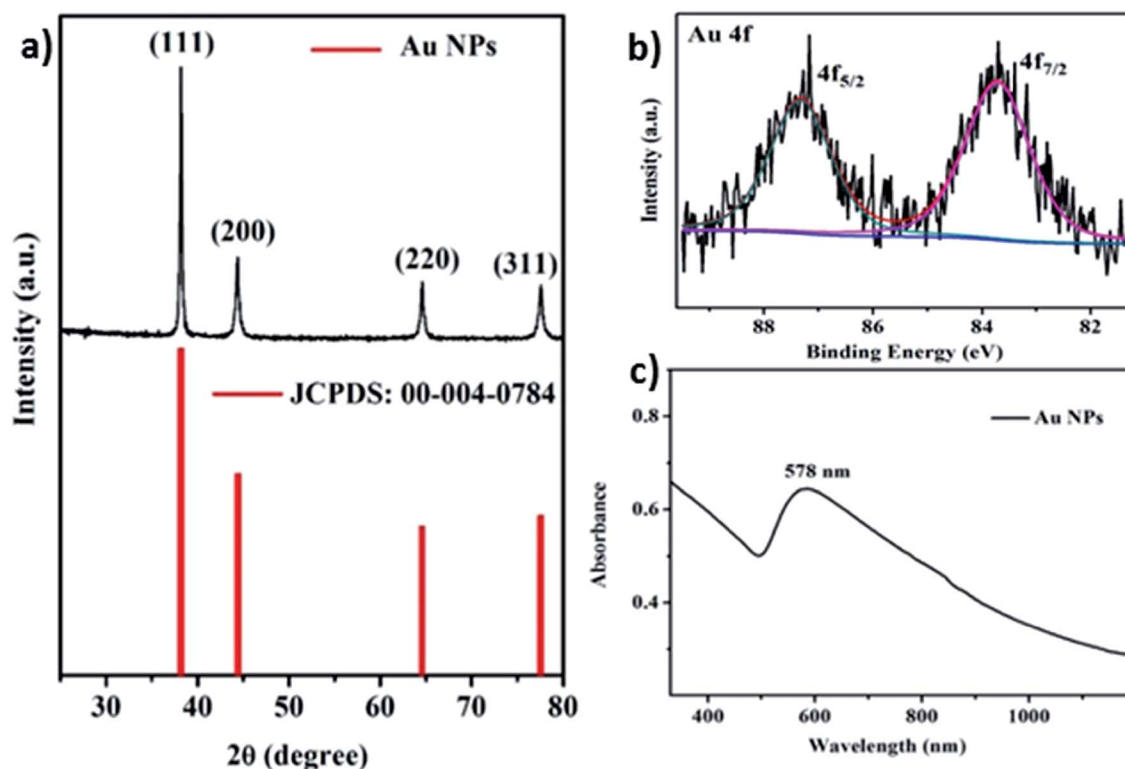


Fig. 3 (a) PXRD pattern of synthesized AuNPs (bottom) and its reference pattern (top); (b) Au 4f high resolution XPS of AuNPs; (c) UV-vis absorption spectrum of AuNPs dispersed in aqueous medium.



can be considered as an alternative energy-efficient and greener method to reduce other noble metal ions for sustainable energy and environmental related applications.

In general, the absorption peak position for the surface plasmon resonance (SPR) band depends on the size and shape of the AuNPs, *i.e.* the band can shift either bathochromically or hypsochromically.<sup>74</sup> Therefore, electronic absorption spectroscopy was used to further elucidate the formation of AuNPs. The as-prepared AuNP powder was dispersed in water by ultrasonication for 15 min prior to recording UV-vis spectra (Fig. 3c). Interestingly, AuNPs exhibited a broad SPR band around 578 nm, which matches well with the literature reports on AuNPs with particle size of 3–100 nm.<sup>75,76</sup> In addition, the broadness of the SPR band suggests the large particle size distribution, which was later confirmed by SEM and HR-TEM analysis.

The surface morphology of AuNPs was characterized using electron microscopic analysis. Fig. 4 shows the FE-SEM images of AuNPs coated over a silica wafer substrate at different magnifications. It clearly indicates that AuNPs are relentlessly stacked together, in which agglomeration of ultra-small individual AuNPs can be clearly seen in high magnification. It also suggests that the assembling of AuNPs may not be controlled by SCF processing, since no reducing agents or surfactants have been employed during the synthetic stage. It is worth mentioning that the surface hydroxyl groups are mainly accountable for the aggregation of any nanoparticles, wherein the rate of aggregation depends on the nature of the fluid medium. In other words, the aggregation will be more in the aqueous compared to the non-aqueous medium.<sup>77</sup> However, the nucleation and growth of AuNPs can be significantly hampered by the rapid reaction time and cooling of the supercritical reactor in an ice-cold water bath after SCF processing. Hence, ultra-small AuNPs with controlled size can be achieved which

may be suitable as a heterogeneous catalyst for organic reactions.

To identify the presence of elements in the AuNPs, SEM-EDX analysis was performed (Fig. 5). A high intense elemental peak corresponding to gold atoms can be seen along with the absence of elemental chloride peak, which confirms the complete reduction of  $\text{HAuCl}_4 \cdot 3\text{H}_2\text{O}$  to AuNPs. However, elemental peaks corresponding to chromium and iron were observed along with the Au peak although in very less quantity. This could be attributed to the leaching out of alloys of the supercritical reactor by the highly acidic  $\text{HAuCl}_4 \cdot 3\text{H}_2\text{O}$  and also by *in situ* generated HCl. As a consequence, the leached-out Cr and Fe possibly deposit on the surface of AuNPs under low pH at high temperature/pressure SCF conditions. It is appropriate to mention that this unwanted reaction can be excluded by choosing suitable Au precursors. Nevertheless, these elemental impurities were not detected in the XRD analysis due to their much lower concentration and also because they are spread

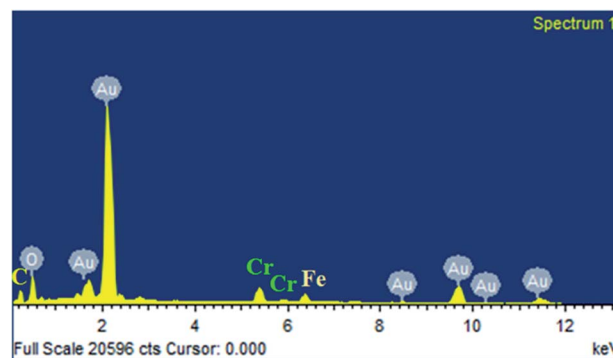


Fig. 5 SEM-EDX analysis of the synthesized AuNPs.

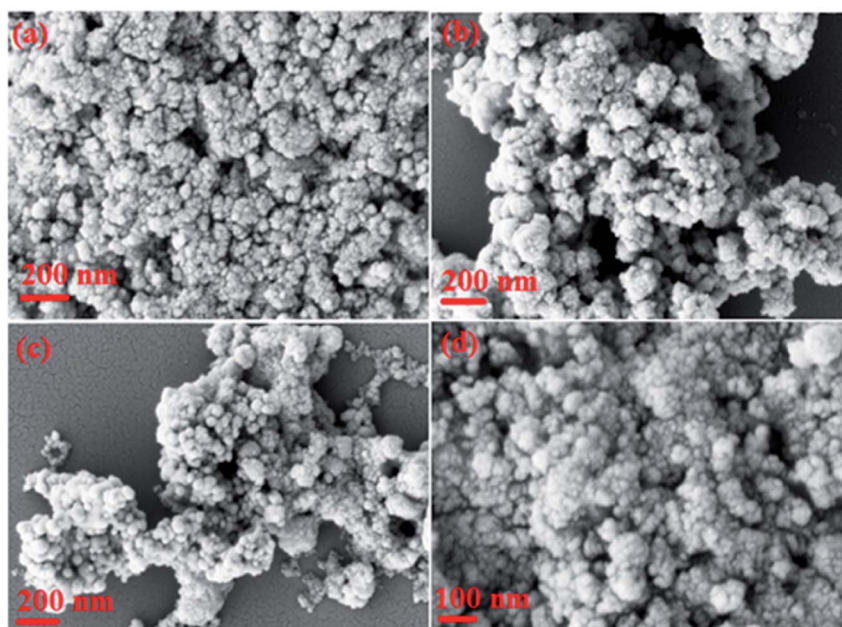


Fig. 4 (a–d) FE-SEM images of AuNPs at different magnifications.



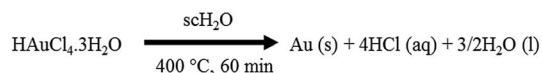
over AuNPs only as a very thin layer. Yet, these elements are not expected to contribute much in the subsequent reactions because of their catalytic inertness in cross-coupling chemistry.

On the basis of the above results, the formation of AuNPs under SCF conditions can be explained as follows: AuNPs are formed by high temperature/pressure assisted decomposition of  $\text{HAuCl}_4 \cdot 3\text{H}_2\text{O}$ , in which supercritical water ( $\text{scH}_2\text{O}$ ) acts both as a fluid as well as a base catalyst. It is pertinent to mention that, Montero *et al.* recently disclosed the direct thermal decomposition of  $\text{HAuCl}_4 \cdot n\text{H}_2\text{O}$  under atmospheric conditions to prepare AuNPs along with the formation of  $\text{H}_2\text{O}$ ,  $\text{HCl}$  and  $\text{Cl}_2$  as by-products.<sup>78</sup> Based on this report, we propose a reasonable chemical reaction for the synthesis of AuNPs under SCF conditions (Scheme 1).

The formation of size-reduced AuNPs with insignificant agglomeration without using any capping agents has been our expectation based on the unique physico-chemical properties of SCFs. These properties include diffusion coefficient, low viscosity, thermal conductivity, molecular density, excellent wetting of surfaces, low surface tension, high degree of super saturation and higher nucleation rate. Indeed, these are some of the critical parameters to control the size and morphology of the metal nanoparticles. However, we obtained AuNPs only with large particle-size distribution which may be because of the low density of the supercritical water ( $0.32 \text{ g mL}^{-1}$ ). Under this condition, the formed nanoparticles settle down easily and might have longer residence time which facilitates the formation of large particles by crystal growth. To further confirm the surface morphology and size of AuNPs, HR-TEM analysis at different magnifications was conducted (Fig. 6). It can be clearly seen that AuNPs are accumulated at different regions, in which the lateral size of individual AuNPs ranges from 10 to 30 nm and are found to be in parallel agreement with FE-SEM analysis. It is worth mentioning that the well distributed AuNPs with 10–30 nm in size are observed throughout the HR-TEM analysis. The lattice fringe value for the AuNPs was calculated to be 0.24 nm which corresponds to the (111) crystal plane of the cubic phase of AuNPs.<sup>79</sup> The electron diffraction pattern of a selected area reveals the crystalline nature of AuNPs which were in agreement with the PXRD results.

### 3.2 Catalytic tests towards cross-coupling reactions

Having prepared and characterized the requisite AuNPs, we next set out to investigate their scope towards Suzuki and Sonogashira coupling reactions. At the outset, it is appropriate to mention that all reactions were performed in ultraclean glass vials and stir bars not previously used for Pd-catalyzed reactions. The purpose of this precautionary measure is to exclude the possibility of participation of Pd species, which otherwise would catalyze the reaction even in ppm levels.<sup>40,41</sup> It is



Scheme 1 Reduction of  $\text{HAuCl}_4 \cdot 3\text{H}_2\text{O}$  under supercritical conditions.

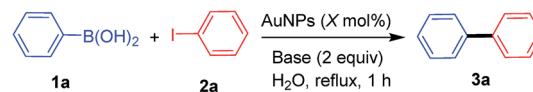
important to note that there are several gold catalyzed methodologies disclosed in the literature for cross-coupling reactions and are either limited to Sonogashira<sup>38–45</sup> or Suzuki.<sup>46–49</sup> Our approach differs from these protocols with an objective of establishing a generalized protocol that can accommodate an array of substrates suitable for both Suzuki as well as Sonogashira coupling. Towards this end, we initiated our studies by refluxing a mixture of iodobenzene (0.01 mol), phenylboronic acid (0.01 mol), NaOH (0.02 mol) and AuNPs (0.5 mol%) in water for 1 h (Table 1). To our delight, we obtained the expected biphenyl **3a** in 61% yield (entry 1). Switching over to  $\text{K}_3\text{PO}_4$  as the base in place of NaOH, the yield improvised to 83% (entry 2). Instead of  $\text{K}_3\text{PO}_4$ , a mild base like  $\text{K}_2\text{CO}_3$  afforded the cross-coupled product in an excellent 98% yield (entry 3). This is quite contradictory to some of the related reports, wherein  $\text{K}_3\text{PO}_4$  was reported to be an effective base compared to  $\text{K}_2\text{CO}_3$ .<sup>80,81</sup> Nevertheless, it is to be remembered that these two reports deal with heterogenized Au-NHC complexes and not bare gold nanoparticles. Reducing the catalytic amount of AuNPs from 0.5 to 0.1 mol% still affords more or less similar yield (entry 4). Performing a blank reaction devoid of AuNPs resulted in no product formation at all (entry 5), which evidently indicates the catalytic role of AuNPs in promoting this transformation. As a final screen, a room temperature reaction yielded only a trace amount of product suggesting the need for high temperature (entry 6).

As per entry 4, the best conditions for the prototype reaction are as follows: heating a reaction mixture comprising 0.01 mol each of iodobenzene and phenylboronic acid,  $\text{K}_2\text{CO}_3$  (0.02 mol), AuNPs (0.1 mol%) in demineralized water for 1 h. To this end, we utilized the same reaction conditions for other structurally and electronically different substrates (Table 2). At first, the influence of electronically different substituents such as 4-Me and 4-Ac was assessed, which revealed that our protocol can accommodate both electron releasing as well as electron withdrawing substituents equally (entries 1 and 2). Modifying the steric environment from the C4 to the C2-position also led to identical reactivity with a yield of 90% (entry 3). At this point of our investigation, we also conducted the reaction by varying the

Table 1 Optimization of reaction conditions

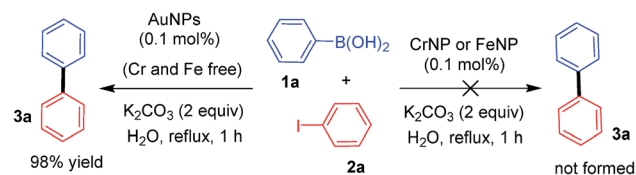
Entry	X	Base	Isolated yield <sup>b</sup> (g)	Yield <sup>c</sup> (%)
1	0.5	NaOH	0.94	61
2	0.5	$\text{K}_3\text{PO}_4$	1.28	83
3	0.5	$\text{K}_2\text{CO}_3$	1.48	96
4	0.1	$\text{K}_2\text{CO}_3$	1.51	98
5	—	$\text{K}_2\text{CO}_3$	—	—
6 <sup>a</sup>	0.1	$\text{K}_2\text{CO}_3$	0.31	20

<sup>a</sup> Reaction performed at rt for 1 h. <sup>b</sup> 0.01 mol each of **1a** and **2a** were used for the reaction. <sup>c</sup> Isolated yield by flash chromatography.



substituents at the boronic acid reacting partner. Hence, testing with 4-MeO and 4-F phenylboronic acid afforded the coupled products with only a slight diminution in yield (entries 4 and 5). As exemplified in entries 6 and 7, the catalytic activity of AuNPs is not only limited to the iodo precursor, but also can be extended to less reactive bromides and chlorides with the same efficacy (entries 6 and 7). The catalytic behaviour towards these halo derivatives can be attributed to the precisely controlled size of AuNPs in conjunction with the absence of any stabilizer, which otherwise could act as a catalyst inhibitor for cross-coupling reactions.<sup>82</sup> It is to be remembered that all the tested reactions resulted only in cross-coupling with no formation of undesired homocoupling products, which highlights the excellent site-selectivity of the present methodology. This is indeed advantageous from a synthetic chemist perspective, since nanostructured gold was also reported to catalyze the self-coupling of aryl iodides (Ullman coupling)<sup>83,84</sup> and aryl boronic acids (homocoupling)<sup>49,71,84–88</sup> to form different products. Thus, the observed hetero-selectivity is possibly due to the absence of the immobilizer in AuNPs which leads to sterically unhindered facets that allow easy access of both reactants towards the catalyst surface.

It is to be remembered that trace amounts of Cr and Fe contents were present in the AuNPs as indicated by the SEM-EDX analysis. Therefore, to exclude the possibility of Cr or Fe in catalyzing the cross-coupling reaction, we conducted two control experiments for the reaction between **1a** and **2a**. In these experiments, we used commercially available Cr (~70 nm) and Fe (~25 nm) nanopowder instead of AuNPs, which resulted in no reaction at all with quantitative recovery of **2a** (Scheme 2). In order to exclude any potential effect of Cr or Fe, a control reaction with AuNPs completely devoid of these metal impurities would be a confirmatory indication. Therefore, we prepared AuNPs (~40 nm) with no metal contamination by the sonolysis of an aqueous solution of HAuCl<sub>4</sub>.<sup>17</sup> AuNPs prepared in this way afforded **3a** in almost quantitative yields similar to our catalytic system. These experimental findings clearly revealed the

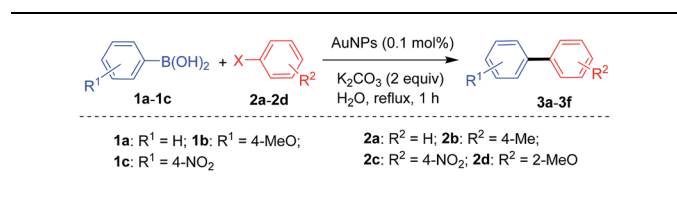


Scheme 2 Control reactions using CrNPs and FeNPs.

catalytic inertness of Cr and Fe towards cross-coupling and confirm AuNPs as the actual catalytic species.

As per our objective, we also look into the possibility of exploring our AuNP catalysis for Sonogashira coupling by including terminal alkynes as coupling partners instead of boronic acids. Gratifyingly, the desired cross-coupling proceeds well for the tested acetylenes (**4a–4c**) and haloarenes (**2a**, **2b** and **2e**) of different electronic natures (Table 3). However, a slight diminution in yield was observed for these couplings compared to Suzuki. Nevertheless, no formation of homo-coupling side products is suggestive of the good site-selectivity of the AuNP catalysis. This could be attributed to the influence of the large particle size of AuNPs (~10–30 nm) in controlling the selectivity of the heterogeneous cross-coupling chemistry and is in sharp agreement with the findings of Lambert *et al.*<sup>42</sup> As evidenced by crude NMR analysis, some of the iodoarene and alkyne remains unreacted even after 1 h of the reaction, which accounts for the lack of complete conversion. Interestingly, replacing the alkyne with an olefin such as styrene **4a'** to effect Mizoroki–Heck coupling with iodobenzene **2a** under our established conditions resulted only in 22% yield of stilbene **5a'**. We attribute this poor reactivity of alkenes to the low acidity compared to alkynes. As far as the selectivity is concerned, both Suzuki as well as Sonogashira resulted exclusively in cross-coupling and no homocoupling was observed.

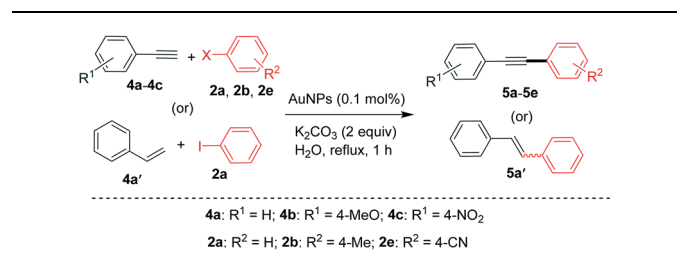
Table 2 Suzuki coupling under AuNP catalysis



Entry	R <sup>1</sup>	R <sup>2</sup>	X	Product	Isolated yield <sup>a</sup> (g)	Yield <sup>b</sup> (%)
1	H	4-Me	I	<b>3b</b>	1.61	96
2	H	4-Ac	I	<b>3c</b>	1.72	88
3	H	2-MeO	I	<b>3d</b>	1.65	90
4	4-MeO	H	I	<b>3e</b>	1.49	81
5	4-F	H	I	<b>3f</b>	1.53	89
6	H	H	Br	<b>3a</b>	1.41	92
7	H	H	Cl	<b>3a</b>	1.29	84

<sup>a</sup> 0.01 mol each of aryl boronic acid and haloarene were used for the reaction. <sup>b</sup> Isolated yield after flash chromatography.

Table 3 Sonogashira and Heck coupling under AuNP catalysis



Entry	R <sup>1</sup>	R <sup>2</sup>	X	Product	Isolated yield <sup>a</sup> (g)	Yield <sup>a,b</sup> (%)
1	H	H	I	<b>5a</b>	1.41	79
2	H	H	Br	<b>5a</b>	1.21	68
3	4-MeO	H	I	<b>5b</b>	1.68	81
4	4-NO <sub>2</sub>	H	I	<b>5c</b>	1.36	61
5	H	4-Me	I	<b>5d</b>	1.44	75
6	H	4-CN	I	<b>5e</b>	1.40	69
7	H	H	I	<b>5a'</b>	0.39	22

<sup>a</sup> Isolated yield after flash chromatography. <sup>b</sup> Trace amount of the homo-coupled product was detected in all cases by crude NMR analysis.



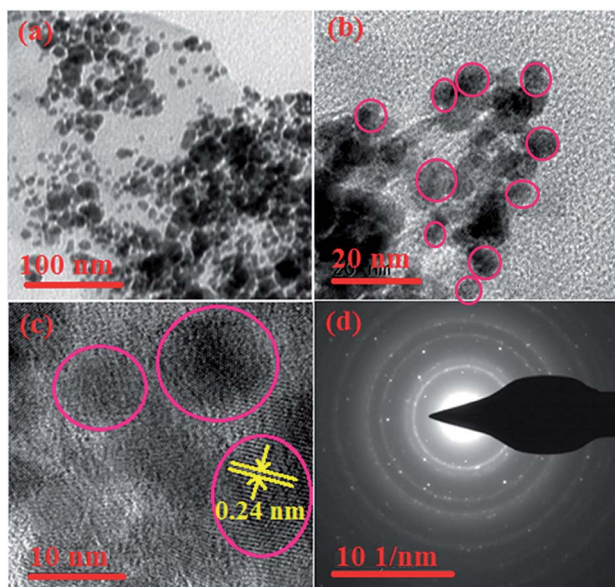


Fig. 6 (a–c) HR-TEM images and (d) SAED pattern of AuNPs.

In order to evaluate the conversion of different substrates under the optimized conditions, individual reaction solutions were subjected to HPLC analysis in a time interval of every 15 minutes (Fig. 7a and b). As depicted in Fig. 7a, time-dependent behaviour in the conversion for Suzuki coupling was observed, where complete or nearly complete conversion was possible only after allowing the reaction for 1 h. Similar conversion with respect to time was observed in the case of Sonogashira coupling, but not all substrates went onto complete conversion (Fig. 7b). In these reactions, complete conversion was hampered by the formation of trace quantities of homo-coupled by-products through a competition of hetero-coupling vs. homo-coupling. Consequently, the Sonogashira reaction of all substrates led to cross-coupled products with slightly lower turnover numbers (TONs =  $\sim$ 3 to 4) than that of Suzuki

coupling (TONs =  $\sim$ 4 to 5). Nevertheless, the findings of high conversion efficiency are consistent with a heterogeneous catalysis pathway by AuNPs.<sup>42,45</sup>

### 3.3 Catalyst recycling

Another significant advantage of our methodology is the ease with which the catalyst can be recovered and reused after the coupling reaction. The recyclability test of the catalyst was proved through a gram-scale synthesis of **3a** in a reaction between **1a** (0.1 mol, 12.2 g), **2a** (0.1 mol, 20.4 g) and AuNPs (0.1 mmol, 19.7 mg) (Fig. 8a). After completion of the reaction, the catalyst was recovered by simple centrifugation followed by filtration from an ethanol suspension. This was followed by sequential washing with ethanol, water, acetone and finally dried at 40 °C for 1 h. Thus, the recovered gold after each reaction cycle was found to be 18.71 mg (95%), 18.51 mg (94%), 18.12 mg (92%), 17.73 mg (90%), 17.33 mg (88%) and 16.55 mg (84%). It was found that up to six cycles, the recovered AuNPs can be reused, after which the catalytic activity reduced gradually owing to the lower mass recovery in each step. This means, the amount of recycled catalyst available for subsequent reactions is lower compared to the previous cycle. This gradual loss of catalyst after each cycle could account for the slight decrease in the yield after each cycle. To quantify the amount of any leached-out gold in the filtrate, ICP-OES analysis was performed. The process involves the introduction of product solution into the plasma using a nebulizer and a spray chamber. To this end, freshly prepared standard gold solutions at concentrations ranging between 0.05 and 10 ppm were initially analyzed, in which gold was detected at an emission wavelength of 242.794 nm (Fig. 8b). The standard linear curve exhibited a positive correlation between the concentration and intensity with a regression coefficient of 0.9995 (Fig. 8c). The gold catalyst recovered after four catalytic cycles was used for the trace element analysis, the result of which indicates no significant peak at 242.794 nm, clearly revealing the absence of gold content (Fig. 8d). As depicted in Fig. 8e, spiking (standard

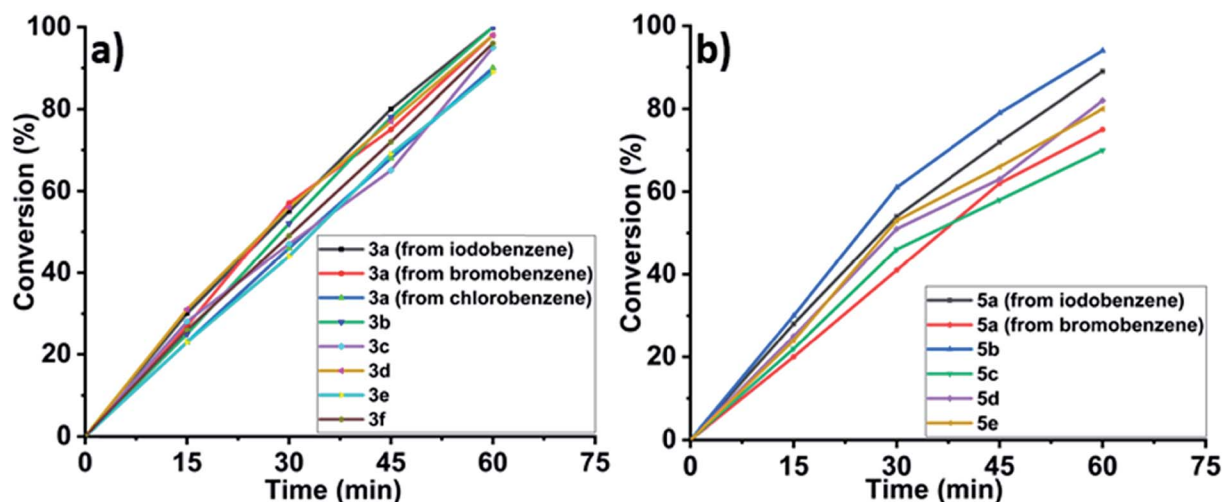


Fig. 7 Conversion of substrates in (a) Suzuki coupling and (b) Sonogashira coupling.





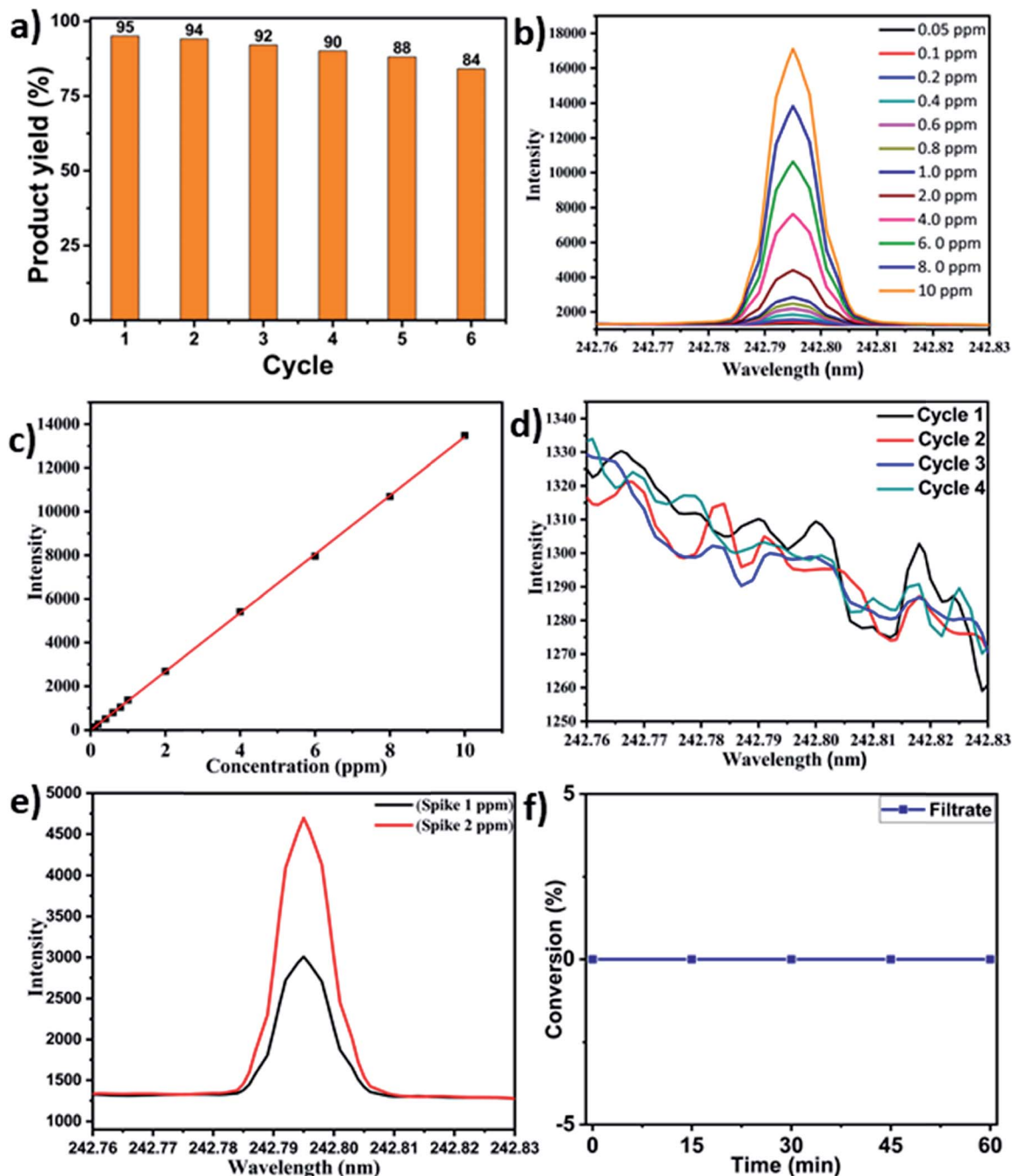


Fig. 8 (a) Recyclability of AuNPs tested in the synthesis of **3a**; (b) ICP-OES curve for the gold standard at different concentrations; (c) linearity curve for the gold standard; (d) ICP-OES curve of first four recycled samples; (e) spiked curve; (f) filtrate test of Suzuki coupling.

addition method) was carried out to confirm the above results. Two levels of concentrations such as 1.02 ppm and 2.01 ppm were spiked in the first recycled sample and the content of gold was found to be  $1.02 \mu\text{g mL}^{-1}$  and  $2.06 \mu\text{g mL}^{-1}$  respectively with 100.0% and 101.9% recoveries, which confirmed that the leaching of gold is negligibly small. Nevertheless, a filtrate test was performed by adding **1a** (0.01 mmol), **2a** (0.01 mmol) and  $\text{K}_2\text{CO}_3$  (0.02 mmol) to the obtained filtrate and refluxed. After several intervals, it was found that no conversion to product **3a** was detected which confirms that the reaction does not

proceed in the absence of the solid AuNP catalyst (Fig. 8f). This could possibly be because of the absence of AuNPs in the filtrate and thus validates the heterogeneity of the catalytic process. Additionally, a hot filtration test was carried out to prove the heterogeneity of AuNPs in a reaction between **1a** and **2a** under standard conditions.<sup>89</sup> When the reaction was stopped at 35 min, the isolated yield of **3a** was found to be 55%. Then, the catalyst was removed by centrifugation and filtered-off from the solution and the reaction was allowed to continue for another 1 h. However, no increase in product



yield was observed which testifies to the catalyst's heterogeneous nature.

In another series of experiments, the spent catalyst obtained after six catalytic cycles was subjected to PXRD, XPS, SEM and FE-SEM analysis (Fig. 9). As depicted in Fig. 9a, the XRD peak positions of the recovered catalyst at  $2\theta$  values of  $38.21^\circ$ ,  $44.41^\circ$ ,  $64.62^\circ$  and  $77.60^\circ$  [JCPDS card no. 01-089-3697] correspond to the (111), (200), (220) and (311) lattice planes respectively. Since the  $2\theta$  values match well with the fcc phase of the as-prepared sample, it clearly indicates no change in phase purity after recycling. The calculated crystallite sizes corresponding to the different  $2\theta$  values were found to be 29.40, 30.11, 40.22 and 43.94 nm respectively. Thus, the average crystallite size of the recycled catalyst is calculated to be 35.9 nm, which is 4.3 nm greater than that of the as-prepared sample (31.6 nm). This increase in crystallite size after recycling is attributed to the agglomeration during catalytic tests which is in parallel agreement with the gradual decrease in catalytic activity after every recycling. It is to be remembered at this juncture that during recycling studies, a gradual loss in catalyst amount was observed after each catalytic run, which was considered to be the reason for the slight decrease in yield after each cycle. We believe that probably both the increase in crystallite size as well as loss in catalyst amount after each cycle could contribute for the gradual loss of yield. To exclude the possibility of oxidation

of Au(0) to Au(I) and/or Au(III) during the course of catalytic tests or recycling, XPS analysis was carried out on the recovered catalyst (Fig. 9b). The spectra featured a large 4f peak as a doublet with  $4f_{7/2}$  and  $4f_{5/2}$  spin orbit split peaks centred at 84.1 and 87.7 eV respectively. These binding energy values are in good agreement with gold in its zerovalent state, thus revealing no change in the oxidation state during catalysis. This in turn confirms that Au(0) is indeed the catalytic species. Moreover, the surface morphology of the spent catalyst was assessed by SEM and FE-SEM analysis (Fig. 9c). Notably, the recovered AuNP catalyst maintains its unique morphology even after six catalytic cycles, which is a clear indication of the stability of the catalyst.

Based on the above results, a mechanistic basis representing the probable sequence of events is provided in Fig. 10. For Suzuki coupling, the reaction is initiated by the interaction of aryl boronic acids **1** and haloarenes **2** on the catalyst surface to form transient species **I**, which further reorganize to form organometallic intermediates **II**. The latter in the presence of  $K_2CO_3$  and water deliver the biaryl products **3** along with  $KHCO_3$ ,  $KX$  and  $B(OH)_3$  as byproducts.<sup>90</sup> In the case of Sonogashira coupling, the reaction is likely to proceed through the chemisorption of terminal alkynes **4** and haloarenes **2** on the catalyst surface to form adsorbed species **III**. The adsorbed molecules then undergo reorganization on the surface to form intermediate **IV**. The latter intermediate under the action of

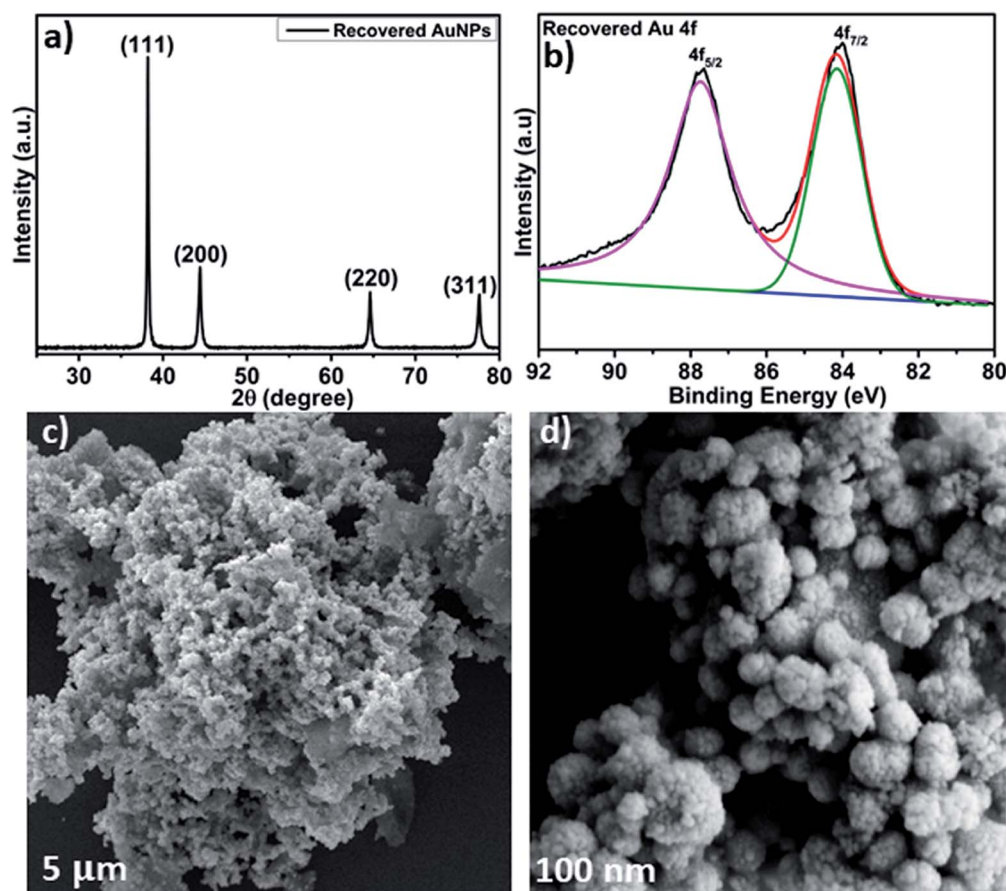


Fig. 9 Recovered AuNPs after six catalytic cycles: (a) PXRD pattern, (b) Au 4f high resolution XPS, (c) SEM image, (d) FE-SEM image.



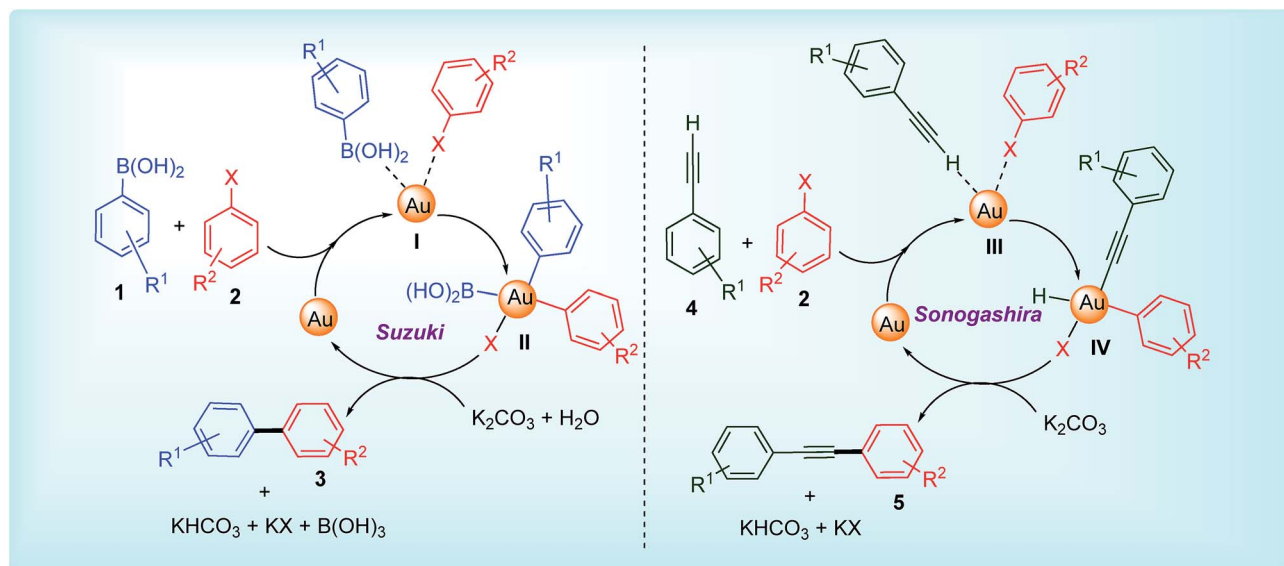


Fig. 10 Plausible mechanism for the AuNP catalyzed Suzuki and Sonogashira coupling.

$K_2CO_3$  affords the desired coupling products **5** accompanied by  $KX$  and  $KHCO_3$  with the regeneration of the catalyst.

### 3.4 Applications in the one-pot synthesis of optoelectronic chromophores

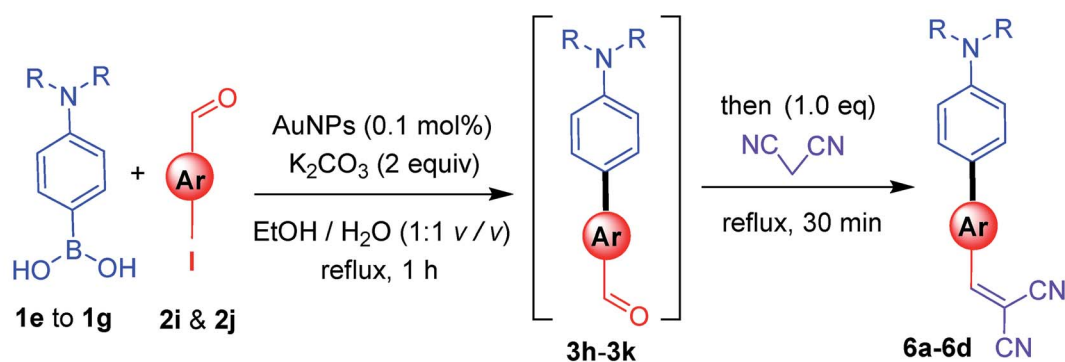
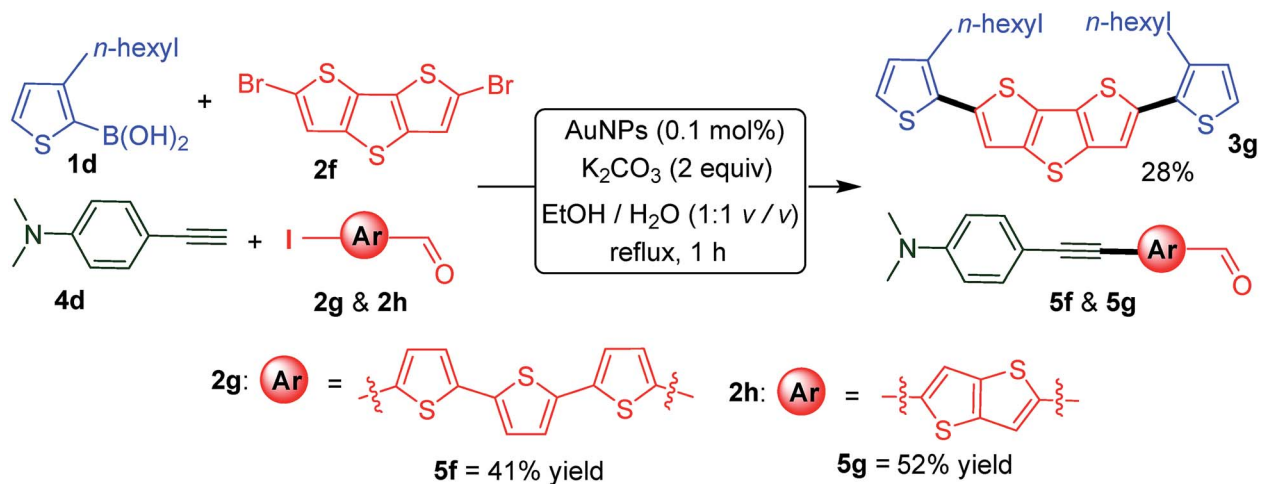
As an application of AuNP catalysis coupled with our pursuit on photoactive functional chromophores,<sup>28–30</sup> we envisaged the synthesis of optoelectronic  $\pi$ -semiconductors by taking advantage of our sustainable conditions (Scheme 3). In this connection, **3g**,<sup>91,92</sup> an important thiophene oligomer intermediate frequently encountered in dye-sensitized solar cells (DSSCs) has been prepared by the Suzuki reaction between **1d** and **2f**, albeit in low yield (28%). Sonogashira coupling of **4d** with **2g** and **2h** respectively affords the photoactive alkynylated adducts **5f** and **5g** in moderate yields (41 and 52%). At this point of our investigation, the feasibility of preparing D- $\pi$ -A (push-pull type) chromophores under one-flask conditions by applying our standard conditions was also investigated (Scheme 3, middle panel). For this study, we chose 4-aminoarylboronic acid (**1e–1f**), iodo-arylcarboxaldehyde (**2i** and **2j**) and malononitrile as the donor (D),  $\pi$ -spacer ( $\pi$ ) and acceptor (A) respectively. All the screened reactions were carried out initially by coupling the boronic acid with iodo precursors for 1 h. Without isolating the coupled products (**3h–3k**), malononitrile was subsequently added to the same flask (*in situ*) and the reaction was continued for the next 30 min under reflux, which allowed the one-flask preparation of the targeted D- $\pi$ -A chromophores **6a–6d** in good to excellent yields (77–91%). Moreover, separate experiments were performed to check the conversion of the precursors (**1e–1g**, **2i** and **2j**) to the intermediates **3h–3k** by discontinuing the reaction at the coupling stage, which afforded an isolated yield of 84 to 97%. These intermediates were characterized by  $^1H$  and  $^{13}C$  NMR, which clearly revealed that the cross-coupling step proceeds well before the ultimate condensation step. It is to be remembered that all these reactions were carried out in EtOH/

$H_2O$  medium (1 : 1 v/v) to enhance the solubility of the reactants and products. Finally, to investigate the catalytic activity of AuNPs in the preparation of bithiophene (**3k**), 2-halothiophenes (**2k–2m**) were subjected to Suzuki coupling with thiophene 2-boronic acid (**1f**). Under identical conditions, the yields gradually reduced from iodo (78%) to bromo (59%) to chloro (22%) precursors, which is in accordance with the electronic difference of the halogens (Scheme 3, bottom panel).

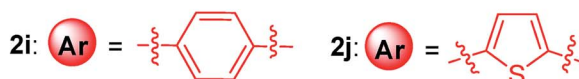
### 3.5 Application of synthesized chromophores

Having prepared the desired chromophores **6a–6d** under one-pot conditions, we next set out to investigate their photo-physical, electrochemical and photovoltaic properties in view of exploring their possible application in organic photovoltaics (Table 4). UV-visible electronic spectra recorded for these chromophores in solution and solid state are depicted in Fig. 11a and S1† respectively. All chromophores in solution phase exhibited an intense  $\pi$ - $\pi^*$  absorption band in the blue region of the visible spectrum with  $\lambda_{max}$  ranging from 462 to 506 nm. The molar absorptivity ( $\epsilon$ ) of all chromophores was found to be in between 2600 and 27 400  $M^{-1} cm^{-1}$  which is indicative of their good light harvesting properties. Thin-film spectra recorded on a quartz plate revealed an extended absorption edge ( $\lambda_{onset} = 595$  to 611 nm) compared to the solution state spectra ( $\lambda_{onset} = 532$  to 571 nm). This bathochromic shift can be attributed to the solid-state  $J$ -aggregation of the chromophores, thus leading to a decrease in the optical band gap ( $E_g^{opt} = 2.08$  to 2.02 eV), which in turn is essential to enhance the short circuit current ( $J_{sc}$ ) in organic solar cells.<sup>93</sup> Prompted by the change in colour with respect to solvent polarity, we also evaluated the absorption behavior of all chromophores in different solvents (*viz.* hexane, THF and MeOH), which led to the finding of a hypsochromic shift in their respective absorption spectra compared to  $CHCl_3$  (Fig. S2 and S3†). Intrigued by the intense colour of all chromophores under UV-vis illumination, we also

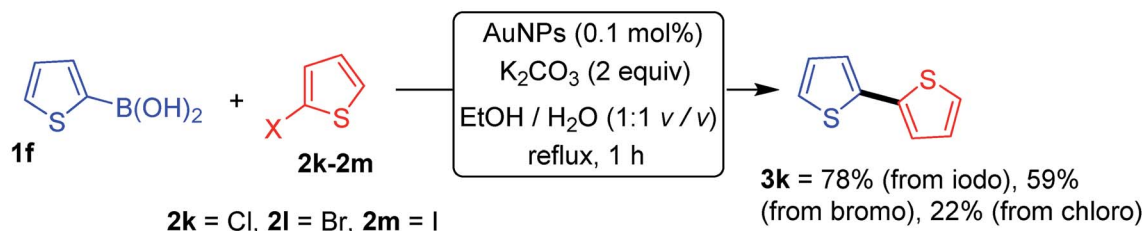




1e: R = Ph; 1f: R = Me; 1g: R = Et



1e & 1g	2i & 2j	3h-3k, yield <sup>a</sup>	6a-6d, yield <sup>a</sup>
1e	2i	3h, 88%	6a, 83%
1e	2j	3i, 84%	6b, 77%
1f	2i	3j, 97%	6c, 91%
1g	2i	3k, 90%	6d, 86%



Scheme 3 AuNP catalyzed synthesis of functional chromophores.

recorded their emission fluorescence (Fig. S4†). All chromophores displayed an orange to red emission maxima ( $\lambda_{em} = 616$  to  $667 \text{ nm}$ ) with Stokes shift ranging between  $60 \text{ nm}$  and  $89 \text{ nm}$

$\text{cm}^{-1}$ . This is an indication of the high molecular flexibility of these  $\pi$ -semiconductors which allows the excited molecules to rapidly relax to the lower frequency vibrational modes of the



Table 4 Optical, electrochemical and photovoltaic data of **6a–6d**

	Abs. (solution) $\lambda_{\max}/\lambda_{\text{onset}}^a$ (nm)	$\epsilon$ ( $\text{M}^{-1} \text{cm}^{-1}$ )	Abs. (film) $\lambda_{\max}/\lambda_{\text{onset}}^b$ (nm)	$E_{\text{g}}^{\text{optc}}$ (eV)	$E^{\text{oxd}}$ (V)	$E^{\text{rede}}$ (V)	$E_{\text{HOMO}}^f$ (eV)	$E_{\text{LUMO}}^g$ (eV)	$E_{\text{g}}^{\text{cvh}}$ (eV)	$V_{\text{oc}}^i$ (V)	$J_{\text{sc}}^i$ ( $\text{mA cm}^{-2}$ )	FF	$\eta^j$ (%)
<b>6a</b>	462/532	21 300	506/595	2.08	1.00	−1.03	−5.35	−3.32	2.03	0.52	6.84	0.48	1.70
<b>6b</b>	506/571	27 400	507/605	2.04	0.90	−1.07	−5.25	−3.28	1.97	0.53	9.20	0.45	2.19
<b>6c</b>	472/563	2600	468/607	2.04	0.84	−0.91	−5.19	−3.44	1.75	0.47	5.85	0.4	1.09
<b>6d</b>	479/546	12 000	497/611	2.02	0.59	−1.02	−4.94	−3.33	1.61	0.48	4.83	0.6	1.39

<sup>a</sup> Absorption in chloroform solution ( $2 \times 10^{-5}$  M). <sup>b</sup> Absorption in thin film. <sup>c</sup> Optical band gap was derived from  $\lambda_{\text{onset}}$  of thin film absorption using the formula,  $E_{\text{g}}^{\text{optc}}$  (eV) =  $1240/\lambda_{\text{onset}}$ . <sup>d</sup> Anodic potential onset. <sup>e</sup> Cathodic potential onset. <sup>f</sup>  $E_{\text{HOMO}}$  was calculated using the relationship,  $E_{\text{HOMO}} = -[(E^{\text{ox}} - E_{1/2(\text{ferrocene})}) + 4.8]$  eV. <sup>g</sup>  $E_{\text{LUMO}}$  was calculated using the relationship,  $E_{\text{LUMO}} = -[(E^{\text{red}} - E_{1/2(\text{ferrocene})}) + 4.8]$  eV, given that  $E_{1/2(\text{ferrocene})} = 0.45$  V. <sup>h</sup>  $E_{\text{g}}^{\text{cvh}}$  (eV) calculated from  $E_{\text{LUMO}} - E_{\text{HOMO}}$ . <sup>i</sup> Current characteristics of photovoltaic devices fabricated from a solution of **6a–6d**:PCBM (1 : 1 blend) in *o*-dichlorobenzene and thermally annealed at 70 °C for 5 min. <sup>j</sup> Efficiencies of inverted OPVs under AM1.5G illumination ( $100 \text{ mW cm}^{-2}$ ).

first excited state ( $S_1$ ) by dissipating excess energy. The cyclic voltammetry of all chromophores was carried out to evaluate the formal redox potential and HOMO–LUMO energy levels of **6a–6d** (Fig. 11b). With reference to the  $\text{Fc}^+/\text{Fc}$  couple, all chromophores showed one oxidation wave with the onset anodic potential ( $E^{\text{ox}}$ ) between 0.59 and 1.0 V and one reduction potential with the onset cathodic potential ( $E^{\text{red}}$ ) between −0.91 and −1.07 V. The anodic potential values are characteristic of electron rich aryl/alkyl amine groups, while the cathodic potential corresponds to the nitrile reduction.<sup>94,95</sup> By applying the Pommerehne empirical relationship,<sup>85</sup> the Frontier molecular orbital energies ( $E_{\text{HOMO}}$  and  $E_{\text{LUMO}}$ ) were calculated from the respective  $E^{\text{ox}}$  and  $E^{\text{red}}$  values. The results revealed that the HOMOs (*ca.* −4.94 to −5.35 eV) and LUMOs (*ca.* −3.28 to −3.44 eV) are appropriately located to enable the chromophores to act as hole-transporting materials (p-type) against a standard n-type material like PCBM (Fig S5†).<sup>96–98</sup> Therefore, a preliminary bulk heterojunction (BHJ) solar cell was fabricated using these

chromophores in conjunction with a standard n-type material (PCBM) in a cell configuration of ITO/ZnO/BHJ/ $\text{V}_2\text{O}_5/\text{Ag}$  (Fig. 11c). The photovoltaic performance revealed that all chromophores displayed similar open circuit voltage ( $V_{\text{oc}} = 0.47$  to 0.53 V) and fill-factor (FF = 0.40 to 0.60). However, there is a notable difference in the short circuit-current density ( $J_{\text{sc}} = 4.83$  to  $9.20 \text{ mAcm}^{-2}$ ), which in turn is influential in enhancing the power conversion efficiency ( $\eta$ ) of the overall device. Among the tested chromophores, **6b** emerged as the best in terms of its high  $\eta$  (2.19%). The efficiencies reported are merely from unoptimized device structures and can be improved further by varying the cell configuration, optimizing the surface morphology and understanding the charge-carrier mobility which falls outside the realm of the present work.<sup>99,100</sup> Nevertheless, these initial results highlight the catalytic potential of AuNPs in preparing small molecule organic solar cells (SMOSCs) essentially under one-pot conditions.

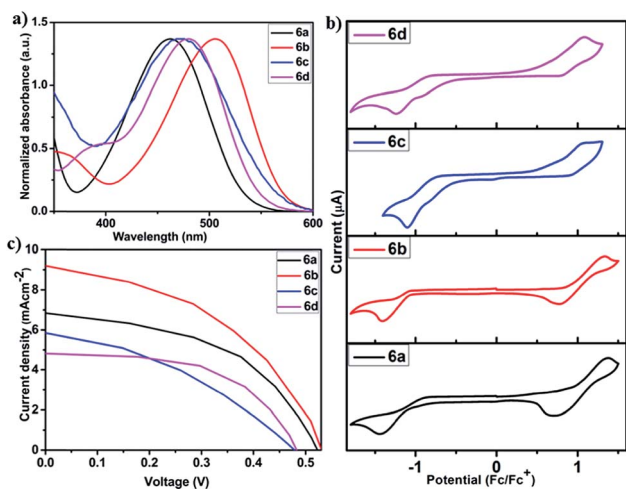


Fig. 11 (a) Normalized UV-vis absorption spectra of **6a–6d** in  $\text{CHCl}_3$  ( $5 \times 10^{-5}$  M); (b) CV of **6a–6d** in dichloroethane ( $2 \times 10^{-3}$  M) using  $n\text{-Bu}_4\text{ClO}_4$  (0.1 M) as the supporting electrolyte at a scan rate of  $100 \text{ mV s}^{-1}$ ; (c) current–voltage ( $J$ – $V$ ) characteristics of fabricated devices of **6a–6d**.

## 4. Conclusions

To summarize, we have developed a supercritical processing method for the preparation of AuNPs using environmentally benign water as the solvent. The formation of phase pure and crystalline AuNPs in a facet of 10–30 nm was characterized by PXRD, XPS, UV-VIS, FE-SEM and HR-TEM analysis. Unlike chemical methods, the current protocol neither requires any reducing agents for AuNP preparation nor a surfactant or rare earth metal oxide as a stabilizing agent. Application of the as-prepared AuNPs as a heterogeneous recyclable catalyst was exemplified in Suzuki as well as Sonogashira cross-coupling reactions under sustainable conditions. To delineate the synthetic scope, one-pot preparation of  $\pi$ -conjugated small molecular chromophores was also highlighted. Following their optoelectronic assessment by UV-vis and CV, a preliminary organic photovoltaic device was constructed which resulted in efficiencies ranging between 1.09 and 2.19%. We are currently engaged in unravelling the mechanistic insights of the AuNP catalyzed coupling processes, which will be disclosed shortly.



## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

One of the authors P. T. thanks UGC-CSIR, India for the Senior Research Fellowship (SRF). C. P. acknowledges the Science and Engineering Research Board (SERB) and Department of Science and Technology (DST), India for providing the fast-track grant (SB/FT/CS-001/2015) and INSPIRE faculty award (IFA14-MS27) respectively.

## References

- 1 Y. Lua and W. Chen, *Chem. Soc. Rev.*, 2012, **41**, 3594–3623.
- 2 J. M. Campelo, D. Luna, R. Luque, J. M. Marinas and A. A. Romero, *ChemSusChem*, 2009, **2**, 18–45.
- 3 A. Corma and H. Garcia, *Chem. Soc. Rev.*, 2008, **37**, 2096–2126.
- 4 M. Stratakis and H. Garcia, *Chem. Rev.*, 2012, **112**, 4469–4506.
- 5 T. Mitsudome and K. Kaneda, *Green Chem.*, 2013, **15**, 2636–2654.
- 6 X. Liu, L. He, Y.-M. Liu and Y. Cao, *Acc. Chem. Res.*, 2014, **47**, 793–804.
- 7 R. Ye, A. V. Zhukhovitskiy, C. V. Deraedt, F. D. Toste and G. A. Somorjai, *Acc. Chem. Res.*, 2017, **50**, 1894–1901.
- 8 G. Li and R. Jin, *Nanotechnol. Rev.*, 2013, **2**, 529–545.
- 9 B. S. Takale, M. Bao and Y. Yamamoto, *Org. Biomol. Chem.*, 2014, **12**, 2005–2027.
- 10 Y. Zhang, X. Cui, F. Shi and Y. Deng, *Chem. Rev.*, 2012, **112**, 2467–2505.
- 11 M. Padilla, F. Peccati, J. L. Bourdelande, X. S. Monfort, G. Guirado, M. Sodupe and J. Hernando, *Chem. Commun.*, 2017, **53**, 2126–2129.
- 12 G. Bharath, A. Naldoni, K. H. Ramsait, A. A. Wahab, R. Madhu, E. Alsharaeh and N. Ponpandian, *J. Mater. Chem. A*, 2016, **4**, 6385–6394.
- 13 Y. C. Yeh, B. Creran and V. M. Rotello, *Nanoscale*, 2012, **4**, 1871–1880.
- 14 I. O. Jiménez, F. M. Romero, N. G. Bastús and V. Puntes, *J. Phys. Chem. C*, 2010, **114**, 1800–1804.
- 15 C. Deraedt, L. Salmon, S. Gatard, R. Ciganda, R. Hernandez, J. Ruiz and D. Astruc, *Chem. Commun.*, 2014, **50**, 14194–14196.
- 16 P. Pati, S. McGinnis and P. J. Vikesland, *Environ. Eng. Sci.*, 2014, **31**, 1–11.
- 17 T. Sakai, S. Miwa, T. Okada and S. Mishima, *AIP Conf. Proc.*, 2012, **1474**, 167–170.
- 18 R. S. Varma, *Curr. Opin. Chem. Eng.*, 2012, **1**, 123–128.
- 19 T. Baran, *Catal. Lett.*, 2019, **149**, 1496–1503.
- 20 T. Baran, *Carbohydr. Polym.*, 2018, **195**, 45–52.
- 21 T. Baran, *Ultrason. Sonochem.*, 2018, **45**, 231–237.
- 22 T. Baran, *Catal. Lett.*, 2019, **149**, 1721–1729.
- 23 N. Y. Baran, T. Baran and A. Menteş, *Carbohydr. Polym.*, 2018, **181**, 596–604.
- 24 T. Baran, N. Y. Baran and A. Menteş, *Appl. Organomet. Chem.*, 2018, **32**, e4075.
- 25 M. B. Gawande, A. K. Rathi, J. Tucek, K. Safarova, N. Bundaleski, O. M. N. D. Teodoro, L. Kvitek, R. S. Varma and R. Zboril, *Green Chem.*, 2014, **16**, 4137–4143.
- 26 M. Vadivelu, S. Sugirdha, P. Dheenkumar, Y. Arun, K. Karthikeyan and C. Praveen, *Green Chem.*, 2017, **19**, 3601–3610.
- 27 K. Parthasarathy, C. Praveen, K. Saranraj, C. Balachandran and P. S. Kumar, *Med. Chem. Res.*, 2016, **25**, 2155–2170.
- 28 A. A. Raheem, S. Kamaraj, V. Sannasi and C. Praveen, *Org. Chem. Front.*, 2018, **5**, 777–787.
- 29 A. A. Raheem, S. Gopi, M. Kathiresan and C. Praveen, *RSC Adv.*, 2019, **9**, 1895–1902.
- 30 C. Kumar, A. A. Raheem, K. Pandian, V. Nandakumar, R. Shanmugam and C. Praveen, *New J. Chem.*, 2019, **43**, 7015–7027.
- 31 C. Praveen and D. B. Babu, *Bioorg. Med. Chem. Lett.*, 2016, **26**, 2507–2512.
- 32 C. Praveen and P. T. Perumal, *Synthesis*, 2016, **48**, 855–864.
- 33 K. Parthasarathy, C. Praveen, J. C. Jeyaveeran and A. A. M. Prince, *Bioorg. Med. Chem. Lett.*, 2016, **26**, 4310–4317.
- 34 J. C. Jeyaveeran, C. Praveen, Y. Arun, A. A. M. Prince and P. T. Perumal, *J. Chem. Sci.*, 2016, **128**, 787–802.
- 35 J. C. Jeyaveeran, C. Praveen, Y. Arun, A. A. M. Prince and P. T. Perumal, *J. Chem. Sci.*, 2016, **128**, 73–83.
- 36 C. Praveen, *Coord. Chem. Rev.*, 2019, **392**, 1–34.
- 37 C. Praveen, *Catal. Rev.: Sci. Eng.*, 2019, **61**, 1–41.
- 38 C. González-Arellano, A. Abad, A. Corma, H. García, M. Iglesias and F. Sánchez, *Angew. Chem., Int. Ed.*, 2007, **46**, 1536–1538.
- 39 S. K. Beaumont, G. Kyriakou and R. M. Lambert, *J. Am. Chem. Soc.*, 2010, **132**, 12246–12248.
- 40 T. Lauterbach, M. Livendahl, A. Rosellón, P. Espinet and A. M. Echavarren, *Org. Lett.*, 2010, **12**, 3006–3009.
- 41 A. Corma, R. Juárez, M. Boronat, F. Sánchez, M. Iglesias and H. García, *Chem. Commun.*, 2011, **47**, 1446–1448.
- 42 G. Kyriakou, S. K. Beaumont, S. M. Humphrey, C. Antonetti and R. M. Lambert, *ChemCatChem*, 2010, **2**, 1444–1449.
- 43 V. K. Kanuru, G. Kyriakou, S. K. Beaumont, A. C. Papageorgiou, D. J. Watson and R. M. Lambert, *J. Am. Chem. Soc.*, 2010, **132**, 8081–8086.
- 44 P. Li, L. Wang, M. Wang and F. You, *Eur. J. Org. Chem.*, 2008, 5946–5951.
- 45 C. González-Arellano, A. Corma, H. García, M. Iglesias and F. Sánchez, *Eur. J. Inorg. Chem.*, 2008, 1107–1115.
- 46 J. Han, Y. Liu and R. Guo, *J. Am. Chem. Soc.*, 2009, **131**, 2060–2061.
- 47 M. Al-Amin, M. Akimoto, T. Tameno, Y. Ohki, N. Takahashi, N. Hoshiya, S. Shuto and M. Arisawa, *Green Chem.*, 2013, **15**, 1142–1145.
- 48 N. Dwadina, J. Roger, N. Pirio, H. Cätte, R. B. Salem and J.-C. Hierso, *Chem.-Asian J.*, 2017, **12**, 459–464.
- 49 C. González-Arellano, A. Corma, M. Iglesias and F. Sanchez, *J. Catal.*, 2006, **238**, 497–501.



- 50 M.-O. Simon and C.-J. Li, *Chem. Soc. Rev.*, 2012, **41**, 1415–1427.
- 51 C.-J. Li, *Chem. Rev.*, 2005, **105**, 3095–3166.
- 52 T. Adschiri, Y. Hakuta and K. Arai, *Ind. Eng. Chem. Res.*, 2000, **39**, 4901–4907.
- 53 R. S. Oakes, A. A. Clifford and C. M. Rayner, *J. Chem. Soc., Perkin Trans. 1*, 2001, **0**, 917–941.
- 54 K. P. Johnston and P. S. Shah, *Science*, 2004, **303**, 482–483.
- 55 T. Adschiri, Y.-W. Lee, M. Goto and S. Takami, *Green Chem.*, 2011, **13**, 1380–1390.
- 56 Y. Meng, F. Su and Y. Chen, *ACS Appl. Mater. Interfaces*, 2017, **9**, 39549–39559.
- 57 B. Wong, S. Yoda and S. M. Howdle, *J. Supercrit. Fluids*, 2007, **42**, 282–287.
- 58 M. Chatterjee, Y. Ikushima, Y. Hakuta and H. Kawanami, *Adv. Synth. Catal.*, 2006, **348**, 1580–1590.
- 59 P. Thangasamy and M. Sathish, *J. Mater. Chem. C*, 2016, **4**, 1165–1169.
- 60 M. Sathish, S. Mitani, T. Tomai and I. Honma, *J. Mater. Chem. A*, 2014, **2**, 4731–4738.
- 61 P. Thangasamy, K. Selvakumar, M. Sathish, S. M. Senthil Kumar and P. Thangamuthu, *Catal. Sci. Technol.*, 2017, **7**, 1227–1234.
- 62 P. Thangasamy, T. Partheeban, S. Sudanthiramoorthy and M. Sathish, *Langmuir*, 2017, **33**, 6159–6166.
- 63 P. Thangasamy, V. Maruthapandian, V. Saraswathy and M. Sathish, *Catal. Sci. Technol.*, 2017, **7**, 3591–3597.
- 64 P. Thangasamy, M. Santhanam and M. Sathish, *ACS Appl. Mater. Interfaces*, 2016, **8**, 18647–18651.
- 65 D. Rangappa, K. Sone, M. Wang, U. K. Gautam, D. Golberg, H. Itoh, M. Ichihara and I. Honma, *Chem.–Eur. J.*, 2010, **16**, 6488–6494.
- 66 C. Gunanathan, Y. Ben-David and D. Milstein, *Science*, 2007, **317**, 790–792.
- 67 S. S. Zalesskiy, A. E. Sedykh, A. S. Kashin and V. P. Ananikov, *J. Am. Chem. Soc.*, 2013, **135**, 3550–3559.
- 68 A. Moragues, F. Neațu, V. I. Pârvulescu, M. D. Marcos, P. Amoros and V. Michelet, *ACS Catal.*, 2015, **5**, 5060–5067.
- 69 K. Nobusawa, N. Okamoto, K. S. L. Chong, X. Lin, K. Iwahori and I. Yamashita, *ACS Omega*, 2017, **2**, 1416–1424.
- 70 X. Zhang, H. Shi and B.-Q. Xu, *Angew. Chem., Int. Ed.*, 2005, **44**, 7132–7135.
- 71 H. Tsunoyama, H. Sakurai, N. Ichikuni, Y. Negishi and T. Tsukuda, *Langmuir*, 2004, **20**, 11293–11296.
- 72 Z. Zhong, J. Lin, S.-P. Teh, J. Teo and F. M. Dautzenberg, *Adv. Funct. Mater.*, 2007, **17**, 1402–1408.
- 73 B. Vilhanová, J. Václavík, L. Artiglia, M. Ronocchiaro, A. Togni and J. A. van Bokhoven, *ACS Catal.*, 2017, **7**, 3414–3418.
- 74 V. Amendola, R. Pilot, M. Frascioni, O. M. Maragò and M. A. Iatì, *J. Phys.: Condens. Matter*, 2017, **29**, 203002.
- 75 D. Liu, C. Li, F. Zhou, T. Zhang, H. Zhang, X. Li, G. Duan, W. Cai and Y. Li, *Sci. Rep.*, 2015, **5**, 7686.
- 76 C. Li, K. L. Shuford, M. Chen, E. J. Lee and S. O. Cho, *ACS Nano*, 2008, **2**, 1760–1769.
- 77 T. He, D. Chen and X. Jiao, *Chem. Mater.*, 2004, **16**, 737–743.
- 78 M. A. Montero, M. R. Gennero de Chialvo and A. C. Chialvo, *J. Mater. Chem.*, 2009, **19**, 3276–3280.
- 79 Y. Tan, Y. Liu, L. Kong, L. Kang and F. Ran, *J. Power Sources*, 2017, **363**, 1–8.
- 80 A. Corma, C. González-Arellano, M. Iglesias, S. Pérez-Ferreras and F. Sánchez, *Synlett*, 2007, 1771–1774.
- 81 A. Corma, E. Gutiérrez-Puebla, M. Iglesias, A. Monge, S. Pérez-Ferreras and F. Sánchez, *Adv. Synth. Catal.*, 2006, **348**, 1899–1907.
- 82 A. Biffis, M. Zecca and M. Basato, *J. Mol. Catal. A: Chem.*, 2001, **173**, 249–274.
- 83 K. Layek, H. Maheswaran and M. L. Kantam, *Catal. Sci. Technol.*, 2013, **3**, 1147–1150.
- 84 A. Monopoli, P. Cotugno, G. Palazzo, N. Ditaranto, B. Mariano, N. Cioffi, F. Ciminale and A. Nacci, *Adv. Synth. Catal.*, 2012, **354**, 2777–2788.
- 85 L. Wang, H. Wang, W. Zhang, J. Zhang, J. P. Lewis, X. Meng and F.-S. Xiao, *J. Catal.*, 2013, **298**, 186–197.
- 86 A. Primo and F. Quignard, *Chem. Commun.*, 2010, **46**, 5593–5595.
- 87 C. González-Arellano, A. Corma, M. Iglesias and F. Sánchez, *Chem. Commun.*, 2005, 1990–1992.
- 88 S. Carretin, J. Guzman and A. Corma, *Angew. Chem., Int. Ed.*, 2005, **44**, 2242–2245.
- 89 S. Hemmati, L. Mehrazin, M. Pirhayati and H. Veisi, *Polyhedron*, 2019, **158**, 414–422.
- 90 J. Sun, Y. Fu, G. He, X. Sun and X. Wang, *Appl. Catal., B*, 2015, **165**, 661–667.
- 91 T.-H. Kwon, V. Armel, A. Nattestad, D. R. MacFarlane, U. Bach, S. J. Lind, K. C. Gordon, W. Tang, D. J. Jones and A. B. Holmes, *J. Org. Chem.*, 2011, **76**, 4088–4093.
- 92 J. Frey, A. D. Bond and A. B. Holmes, *Chem. Commun.*, 2002, 2424–2425.
- 93 H. Yao, L. Ye, H. Zhang, S. Li, S. Zhang and J. Hou, *Chem. Rev.*, 2016, **116**, 7397–7457.
- 94 H. G. Roth, N. A. Romero and D. A. Nicewicz, *Synlett*, 2016, **27**, 714–723.
- 95 O. R. Luca, J. L. Gustafson, S. M. Maddox, A. Q. Fenwick and D. C. Smith, *Org. Chem. Front.*, 2015, **2**, 823–848.
- 96 J. Pommerehne, H. Vestweber, W. Guss, R. F. Mahrt, H. Bässler, M. Porsch and J. Daub, *Adv. Mater.*, 1995, **7**, 551–554.
- 97 L.-L. Chua, J. Zaumseil, J.-F. Chang, P. E. C.-W. Ou, K.-H. Ho, H. Sirringhaus and R. H. Friend, *Nature*, 2005, **434**, 194–199.
- 98 B. C. Thompson and J. M. J. Fréchet, *Angew. Chem., Int. Ed.*, 2008, **47**, 58–77.
- 99 H. Spanggaard and F. C. Krebs, *Sol. Energy Mater. Sol. Cells*, 2004, **83**, 125.
- 100 W. J. Potscavage, A. Sharma and B. Kippelen, *Acc. Chem. Res.*, 2009, **42**, 1758–1767.

