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Application of Nanoparticle Mediated N-Arylation of Amines for the Synthesis of Pharmaceutical Entities using Vit-E analogues as Amphiphile in Water

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First CuI-nanoparticle catalyzed inter and intramolecular N-arylation of amines using vitamin E analogues (TPGS) as amphiphile have been developed in water. Application of this transition metal-amphiphile C-N bond formation methodology is further extended for the synthesis substituted indole, bioactive natural product tryptanthrin and intermediates of pharmaceutical entities such as Imatinib, Nilotinib, selective D3 agonist/antagonist ligand, and Oxcarbazpine.

Introduction

In twenty first century the chemical transformations are not only evaluated by yield of synthesis /process, but a complete art of synthesis is required for the evaluation of synthesis /process. The formula for evaluation of synthesis becomes YES (Yield, Economics - eco friendliness and Stereochemistry). To address economics and environmental sustainability emphasis is direct towards development of green medium, and recyclable reagents. Development of green medium is fascinating area since the ionic liquid has been introduced but later on IL biodegradability and greenness in under question mark. Therefore water is consider as the best available choice in nature. Water a solvent of life, as almost all the biochemical reaction under goes in water. However, poor aqueous solubility of organic compounds or reactivity of many transition-metal/organo catalysts are the main limitation, which can be overcome by the introducing organic microenvironment in aqueous phase.

A smart strategy to solve these issues is micellar catalysis. Recently developed the transition metal coupling reaction catalyzed in water with TPGS-750-M / PTS. TPGS-750-M is a biodegradable and water-soluble derivative of natural vitamin E, which is formed by esterification of vitamin E succinate with polyethylene glycol (PEG) 1000. It is consisting of a tocopherol (vitamin E) hydrophobic group and a PEG hydrophilic group. Micellar catalysis greenness methodology which allows the E factor approach to zero. Micelles are mainly simple spherical supramolecules, which are formed by amphiphiles in water and have ability to solubilization of both nonpolar and polar substrates, reagents, and catalysts.

Previous report:

\[
\begin{align*}
\text{R}_1 \text{Br} & + \text{H}_2 \text{Ar} \rightarrow \text{R}_1 \text{Ar} \\
\text{R}_2 = \text{H}, \text{Ph}, \text{Me} \\
\text{CuI-NP, NaOH} & \\
\text{TPGS-750-M/water} & \\
\text{cat} & \text{[(allyl)PdCl\textsubscript{2}]} \\
\text{ligand 2, KOH} & \\
\text{2 % PTS/H\textsubscript{2}O, rt} & \\
\end{align*}
\]

This work:

![Figure 1. N-arylation of amines reaction using Vit-E Amphiphile in Water and their position on HLB Scale](image)

Lipshutz et al. reported N-arylation of amines reaction using PTS as nonionic amphiphile with [(p-allyl)PdCl\textsubscript{2}] as catalyst, Takasago’s as ligand and base in presence of water. Inspired by
this report we wish to report here first CuI-nanoparticle catalyzed inter and intramolecular N-arylation of amines using vitamin E analogues (TPGS) as amphiphile in water. To the best of our knowledge this is first report on Cu nano and Vitamin E analogues as amphiphile for catalyzing C-N bond formation. We utilised this methodology for the synthesis of medicinally important indoles, anticancer and antitubercular natural product tryptanthrin. We further extended the protocol for the synthesis of intermediates of pharmaceutical entities such as clinically used anticancer drug Imatinib, Nilotinib, selective D3 agonist/antagonist ligand for the treatment of alzimeres and, drug addiction therapy and anticonvulsant Oxacarbazpine. Our initial efforts were focused on the actual effectiveness of an efficient nano catalyst for the N3arylation of amines reaction in water. For this investigation the reaction of 3-bromobenzene with morpholine, sodium hydroxide and CuI nanoparticles in water without surfactant was carried out. After that we was explore nonionic amphiphile such as oil/water emulsifier and detergent. However, the use of amphiphile (Brij 30) and nano CuI resulted in the formation of the desired product in 65% yield at room temperature. In search of an efficient amphiphile, we further screened various amphiphiles in the model N-arylation of amines reaction in water such as Brij 30, Triton X-100 and PTS. Therefore, TPGS was found to be the best amphiphile with Nano CuI for the synthesis of N-arylation of amines in water.

The copper(I) salts such as CuBr, CuCl, CuI, and CuI were found to be inferior to CuI nanoparticles (Table 1, entries 11–15). Among the bases studied, Et₂N, K₂CO₃, KOtBu, K₂PO₃, provided lower yields than NaOH (Table 1, entries 1–5). Of the copper catalysts investigated, CuI nanoparticles were significantly superior to other catalyst (Table 1, entries 1). In order to evaluate the catalyst loading, the reaction was carried out using 1.5, 3.5, and 5.0 % mole nano CuI at room temperature in water. It was found that 3.5 % mole NP-Cul is sufficient to give the desired product in excellent yield with an enhanced rate (Table 1, entries 8–10).

A control reaction carried out “on water” (i.e., in the absence of amphiphile) confirmed the importance of micellar catalysis in facilitating N-arylation of amines coupling reaction in aqueous media (Table 2).

**Table 1.** Optimization of reaction conditions for synthesis of N-Aryl amines

<table>
<thead>
<tr>
<th>Entry</th>
<th>Conc. (wt %)</th>
<th>Amphiphile</th>
<th>Yield (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>2%</td>
<td>TPGS</td>
<td>92</td>
</tr>
<tr>
<td>2</td>
<td>1%</td>
<td>TPGS</td>
<td>70</td>
</tr>
<tr>
<td>3</td>
<td>5%</td>
<td>TPGS</td>
<td>82</td>
</tr>
<tr>
<td>4</td>
<td>0%</td>
<td>TPGS</td>
<td>10</td>
</tr>
</tbody>
</table>

**Table 2: Optimisation of Amphiphiles Concentration**

<table>
<thead>
<tr>
<th>Entry</th>
<th>Conc. (wt %)</th>
<th>Amphiphile</th>
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<td>0 %</td>
<td>TPGS</td>
<td>10</td>
</tr>
</tbody>
</table>

A lower concentration of amphiphile was therefore deemed favorable, although too low a concentration or its complete absence was detrimental. Consequently, TPGS (2 wt %) was selected for further studies. A wide array of aromatic amines, azoles, secondary amines were under goes N-arylation with aryl halides in excellent yields (Scheme 1).

**Scheme 1: N-arylation of amines via intermolecular amination**

We further carried out the reaction with nano CuI catalyst at room temperature in water for 10 h but found very less yield. After that we was explore nonionic amphiphile such as oil/water emulsifier and detergent. However, the use of amphiphile (Brij 30) and nano CuI resulted in the formation of the desired product in 65% yield at room temperature. In search of an efficient amphiphile, we further screened various amphiphiles in the model N-arylation of amines reaction in water such as Brij 30,
To explore this methodology, we have economically synthesized the intermediate of high profile anticancer drugs such as Imatinib, Nilotinib. The process exhibits a broad scope for the synthesis of intermediates of imatinib, nilotinib, selective D3 receptor ligand, bioactive natural product tryptanthrin, substituted indoles, and oxacarbazepine respectively (Schemes 2-7). After completion of reaction, the catalyst was recovered from the reaction mixture by centrifugation and reused for the next fresh reaction. It is noteworthy that the catalyst could be reused at least five times without any significantly loss of efficiency. Therefore, we suspected that because of large surface area, nanoparticles have very high catalytic properties as compared to other catalyst.

**Application of intermolecular N-arylation**:

**Scheme 2**: Synthesis of Intermediate of Imatinib:

**Scheme 3**: Synthesis of N1-Selective Intermediate of Nilotinib

**Scheme 4**: Synthesis of Intermediate of Selective D3 Receptor ligand

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The surface property and the composition of the catalyst were characterized from Scanning electron microscope (SEM), Transmission electron microscope (TEM) and energy dispersive X-ray analysis (EDX). The EDX spectrum (shown in Figure 3) further authenticates the presence of Cu in the nanocomposite. In addition, in the SEM, TEM analysis of Cu nanoparticles, interestingly, the shape and size of the nanoparticles remained unchanged before and after the reaction.
Figure 2. (a) SEM images of catalyst before the reaction (b) After the 5th run (c) TEM images before the reaction (d) After the 5th run, (e) EDX image of fresh catalyst, and (f) EDX image of catalyst after 5th run

Table 3. Recyclability of CuI Nanoparticles

<table>
<thead>
<tr>
<th>Run</th>
<th>Catalyst recovery (%)</th>
<th>Product Yield (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1^a</td>
<td>85</td>
<td>92</td>
</tr>
<tr>
<td>2^b</td>
<td>81</td>
<td>86</td>
</tr>
<tr>
<td>3^b</td>
<td>74</td>
<td>82</td>
</tr>
<tr>
<td>4^b</td>
<td>72</td>
<td>80</td>
</tr>
<tr>
<td>5^b</td>
<td>70</td>
<td>75</td>
</tr>
</tbody>
</table>

^aCu nanoparticles (3.5 mol%), bromobenzene (1.0 mmol), morpholine (1.5 mmol) base (1.5 equiv.), aqueous TPGS (2 wt %) amphiphile solution (5 ml) as solvent for 4.5 hr at rt (room temperature). ^bThe recovered catalyst was used under identical reaction conditions to those for the first run.

Conclusions

In summary, We have developed first CuI-nanoparticle catalyzed inter and intramolecular N-arylation of amines using vitamin E analogues (TPGS) as amphiphile in aqueous environment. This methodology is further extended to bioactive natural product tryptanthrin, intermediates of pharmaceutical entity such as life saving anticancer drugs Imitinib, Nilotinib and anticonvulsant drug Oxcarbazepine. The amphiphile used is biodegradable and catalyst have have good recyclability. The nano CuI mediated organic synthesis (NAMO-Synthesis) in aqueous media via micellar catalysis has immense future in application in the area of medicinal chemistry and material science.

Experimental Section

General procedure for preparation of CuI nanoparticles

0.464 g (4 mmol) of dimethylglyoxime (dmgH) and 0.400 g (2 mmol) of Cu(OAc)₂.H₂O were added into 50 ml of absolute ethanol in sequence, which was stirred at 0°C for 30 min to get brown precipitates Cu(dmg)₂. Then the collected precipitates dispersed in 50 ml of absolute ethanol again, 0.664 g (4 mmol) KI was added and stirred vigorously for 2 h. After that, the mixture was transferred into 60 mL Teflon-lined stainless steel autoclave. The autoclave was sealed and heated at 180°C for 6 h, and then the reactor bomb is allowed to cool to room temperature. Black precipitates were obtained, then centrifuged and washed with ethanol and deionized water for three times to ensure the removal of the impurities. The final product was then dried in a vacuum oven at room temperature for 12 h.

General procedure for the N-Arylation of Amine / Azoles:

The N-arylation of amines was carried out in a round bottomed flask. In a typical experiment, a mixture of chlorobenzene (1 mmol), morpholine (1.5 mmol), CuI NPs (3.5 mol%) and NaOH (3 equiv.) were dissolved in aqueous TPGS (2 wt %) amphiphile solution (5 ml) as solvent and stirred for the 4.5 hours at room temperature. The reaction was monitored to completion using TLC. At the end of reaction, the mixture was then cooled to room temperature and poured into distilled water. The products were extracted using EtOAc and the organic layer was dried over anhydrous sodium sulphate (Na₂SO₄). The solvent was evaporated in vacuo, the crude products were purified by column chromatography using EtOAc / hexane solvent system.

General procedure for the 2-BromoIndole derivatives in one-pot.

The intramolecular amination reaction was carried out in a round bottomed flask. In a typical experiment, a mixture of 2-(2,2-dibromovinyl)aniline (1 mmol), CuI NPs (3.5 mol%) and NaOH (3.0 equiv.) were dissolved in aqueous TPGS (2 wt %) amphiphile solution (5 ml) as solvent and stirred for the 15 hours at 90°C temperature. The reaction was monitored to completion using TLC. At the end of reaction, the mixture was then cooled to room temperature and poured into distilled water. The products were extracted using EtOAc and the organic layer was dried over anhydrous sodium sulphate (Na₂SO₄). The solvent was evaporated in vacuo, the crude products were purified by silica column chromatography using EtOAc / hexane solvent system.

Note and References:


