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# Nanomagnetic nickel complex based on salicylamide and L-proline ligands as an efficient heterogeneous catalyst for synthesis of tetrazoles

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A novel salicylamide–L-proline–nickel(II) complex, supported on magnetic iron oxide [Fe<sub>3</sub>O<sub>4</sub>@salicylamide–L-proline–Ni(II)], was synthesized through a three-step procedure. This included the functionalization of Fe<sub>3</sub>O<sub>4</sub> with amine groups using glycine as a linker, followed by direct amidation of salicylic acid and its subsequent coordination with Ni(II) and L-proline as a co-ligand to form the nanomagnetic Ni(II) complex. The resulting catalyst was comprehensively characterized by several techniques. The catalyst exhibited outstanding catalytic performance in the homoselective synthesis of 5-substituted-1*H*-tetrazoles from benzonitriles. Notably, it demonstrated excellent recyclability, maintaining high efficiency over eight reaction cycles. The use of a low-cost linker, ligand, and complex catalyst, combined with easy magnetic separation, minimal leaching, and scalability, renders this approach both environmentally sustainable and economically advantageous compared to traditional Ni-based methods.

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## 1. Introduction

Azoles, a class of five-membered, nitrogen-containing heterocyclic compounds, represent a significant area of study within heterocyclic organic chemistry.<sup>1</sup> These aromatic parent structures, characterized by two double bonds, encompass a range of derivatives, including reduced analogs such as azolines and azolidines, which possess fewer nitrogen atoms. The number of nitrogen atoms within these heterocycles profoundly influences their chemical properties and diverse applications.<sup>2–4</sup> Ranging from one to five nitrogen atoms, these structures give rise to

a variety of scaffolds, including pyrrole,<sup>5</sup> imidazole,<sup>6</sup> pyrazole,<sup>7</sup> triazole,<sup>8</sup> tetrazole, and pentazole.<sup>9</sup> Although pyrrole, imidazole, and pyrazole (containing one or two nitrogen atoms, respectively) occur naturally, azoles with more than two nitrogen atoms are predominantly synthetic and require laboratory synthesis.<sup>10</sup>

Tetrazole stands out among these as the aromatic five-membered organoheterocyclic compound with the highest nitrogen content (four nitrogen atoms and one carbon atom), making it a particularly noteworthy moiety in organic chemistry.<sup>10</sup> Pentazole, while possessing a higher nitrogen content, is not considered an organic molecule in the same way due to its lack of a carbon atom within the ring system.<sup>9</sup> Tetrazoles are versatile building blocks with diverse applications, notably in medicinal chemistry, but also in other fields like materials science and high-energy applications.<sup>11–13</sup> Due to the importance of these structures several synthetic procedures were provided for preparation of these materials from various synthons including nitriles, isocyanides, aldehydes, amines, and aryldiazonium salts.<sup>10,14–19</sup>

The synthesis of substituted tetrazoles from nitriles *via* [3 + 2] cycloaddition mechanism has garnered significant attention, evolving from traditional, metal-free approaches to more modern, metal-catalyzed methodologies.<sup>20–26</sup> While older methods often relied on elevated temperatures and solvents like DMF or toluene, sometimes incorporating catalysts such as triethylamine hydrochloride, iodine, or sulfamic acid, current research increasingly focuses on metal-catalyzed reactions. These modern methods, employing catalysts like palladium,<sup>27</sup>

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copper, indium,<sup>28</sup> silver,<sup>29</sup> neodymium,<sup>30</sup> erbium,<sup>31</sup> gadolinium,<sup>32</sup> samarium<sup>33</sup> and zinc, offer advantages such as milder reaction conditions including lower temperatures and shorter times. However, most of the reported methodologies use toxic solvents, expensive catalysts, high temperature and pressure, and harsh reaction conditions. As a result, these processes generate significant toxic waste, harming the environment and human health. This has led to stricter regulations and a push for green chemistry, which reduces waste and improves efficiency, but further catalyst development is still needed.

In accordance with the green chemistry principles, heterogeneous supported catalysts offer a compelling strategy for enhancing catalytic activity, promoting recyclability, and minimizing waste generation.<sup>34–36</sup> The integration of such catalysts with magnetic nanoparticles (MNPs) presents a particularly attractive avenue, affording not only improved catalytic performance but also facile separation and recovery.<sup>37–41</sup> MNPs have attracted considerable interest due to their inherent stability and the ease of manipulation *via* external magnetic fields.<sup>37,42–44</sup> The judicious combination of MNPs with catalytically active complexes, particularly those derived from sustainable and cost-effective precursors, results in a robust heterogeneous catalytic system.<sup>45–47</sup> This synergistic approach confers several distinct advantages, including a high surface area, excellent catalytic activity, simplified separation, and demonstrable reusability.<sup>44,48</sup> Consequently, such hybrid materials provide a promising platform for diverse applications in catalysis and chemical synthesis.<sup>49</sup>

To date, various nanomagnetic catalysts have been utilized for the synthesis of 5-substituted-1*H*-tetrazoles. However, many of these catalysts rely on toxic and expensive linkers, ligands, and metals, often requiring multi-step immobilization processes and harsh reaction conditions. Therefore, a greener approach is needed. To address this challenge, we explore a novel approach using glycine as a safe and affordable amine linker, salicylic acid and *L*-proline as green ligands, and nickel as an inexpensive and environmentally friendly metal center. The resulting nanocomposite effectively immobilizes the nickel catalytic species, and the supported Ni complex demonstrates high catalytic efficiency for the synthesis of 5-substituted-1*H*-tetrazoles under environmentally benign conditions.

## 2. Experimental

### 2.1. Materials and instruments

All chemicals and solvents were purchased from Merck Millipore and Sigma-Aldrich and used as received. Reactions were monitored by thin-layer chromatography (TLC) on commercial glass-backed plates, with spot visualization under UV light at 254 nm. Fourier transform infrared (FT-IR) spectra were recorded using a PerkinElmer 597 spectrophotometer with KBr plates. X-ray powder diffraction (XRD) analysis was carried out using a PHILIPS PW1730 powder diffractometer with Cu K $\alpha$  radiation ( $\lambda = 1.54056 \text{ \AA}$ ). Inductively Coupled Plasma Optical Emission Spectroscopy (ICP-OES) was performed using a Varian Vista Pro Analyzer. Energy-dispersive X-

ray (EDX) and EDX mapping analyses were performed using a TESCAN VEGA 3 scanning electron microscope (SEM). Transmission electron microscopy (TEM) images were obtained using a CM120 microscope. The magnetic properties of the sample were analyzed using Vibrating Sample Magnetometry (VSM) analysis *via* a VSM-250 (YP, Jilin, China). Brunauer–Emmett–Teller (BET) analysis was performed using a TriStar 3020 analyzer.

### 2.2. Typical procedure for synthesis of [Fe<sub>3</sub>O<sub>4</sub>@salicylamide-*L*-proline-Ni(II)] nanocomposite

Fe<sub>3</sub>O<sub>4</sub> MNPs were prepared according to a previously reported procedure.<sup>50</sup> Subsequently, 10 g of Fe<sub>3</sub>O<sub>4</sub> MNPs were dispersed in 200 mL of a 1 : 1 (v/v) ethanol/water mixture and sonicated for 60 minutes. Then 50 mL aqueous solution of 0.5 M glycine (1.88 g) was then added dropwise to the suspension under vigorous stirring. The resulting mixture was refluxed under a nitrogen atmosphere with vigorous stirring for 48 hours. After the completion of reaction, the obtained Fe<sub>3</sub>O<sub>4</sub>@Gly MNPs were magnetically separated, washed sequentially with hot water and ethanol, and dried in vacuum oven at 80 °C for 24 h. In the subsequent modification step, 5 g of the Fe<sub>3</sub>O<sub>4</sub>@Gly MNPs were dispersed in 100 mL of dry toluene through sonication for 1 h to form a uniform suspension. Then a mixture of 2.06 g dicyclohexylcarbodiimide (DCC) (10 mmol), 1.38 g salicylic acid (10 mmol), and 5 mL of pyridine was then added to this suspension. The reaction mixture was first stirred at room temperature for 2 hours, followed by reflux under a nitrogen atmosphere for 48 h. The resulting Fe<sub>3</sub>O<sub>4</sub>@salicylamide product was then isolated magnetically, washed sequentially with hot ethanol and water, and dried at 80 °C for 4 h. Finally, 5 g of Fe<sub>3</sub>O<sub>4</sub>@salicylamide was dispersed in a 1 : 1 water–ethanol mixture (150 mL) for 30 minutes. To this suspension, 2.488 g Ni(OAc)<sub>2</sub>·4H<sub>2</sub>O (10 mmol), 1.15 g *L*-proline (10 mmol) and five drops of triethylamine were introduced. The reaction mixture was refluxed for 24 hours. The [Fe<sub>3</sub>O<sub>4</sub>@salicylamide-*L*-proline-Ni(II)] nanocomposite was then magnetically separated, washed with hot water and ethanol, and dried at 80 °C for 6 hours.

### 2.3. General procedure for synthesis of 5-substituted-1*H*-tetrazoles catalyzed by [Fe<sub>3</sub>O<sub>4</sub>@salicylamide-*L*-proline-Ni(II)] nanocomposite

A mixture of aryl nitrile (1.0 mmol), sodium azide (1.3 mmol), and the [Fe<sub>3</sub>O<sub>4</sub>@salicylamide-*L*-proline-Ni(II)] nanocomposite catalyst (5 mg) was added to a round-bottom flask containing 3 mL of PEG-400. The reaction mixture was stirred at 120 °C until completion, as determined by thin-layer chromatography (TLC). Upon completion, the reaction mixture was diluted with hot water, and the catalyst was magnetically separated. The resulting aqueous solution was acidified to pH  $\sim$  1 with 2 N hydrochloric acid and subsequently extracted with ethyl acetate. The combined organic extracts were dried over anhydrous sodium sulfate, filtered, and concentrated under reduced pressure. The crude product was purified by column chromatography on silica gel using a petroleum ether/ethyl acetate mixture as the eluent.



### 3. Results and discussions

In this study, a novel nickel complex was successfully immobilized on  $\text{Fe}_3\text{O}_4$  MNPs through an initial amine functionalization step using glycine as an inexpensive and environmentally friendly linker. Salicylic acid, a cost-effective ligand compared to aldehydes and ketones, was then grafted onto the MNPs surface through direct amidation with the surface amine groups. This heterostructure was subsequently used to coordinate with nickel(II) acetate tetrahydrate and L-proline as an efficient co-ligand, resulting in the formation of a six-coordinated nickel(II) complex. This strategy provides an effective nanomagnetic electron donor environment for nickel complexation (Scheme 1). The synthesis and immobilization of the targeted complex were thoroughly monitored and confirmed through various techniques, including FT-IR, XRD, TGA, EDX, ICP-OES, elemental mapping, FE-SEM, TEM, and VSM, to assess their structural and magnetic properties.

#### 3.1. Catalyst characterization

The FTIR spectra of pristine  $\text{Fe}_3\text{O}_4$ ,  $\text{Fe}_3\text{O}_4@\text{Gly}$ ,  $\text{Fe}_3\text{O}_4@\text{salicylamide}$ , and the  $[\text{Fe}_3\text{O}_4@\text{salicylamide-L-proline-Ni(II)}]$  nanocomposite are presented in Fig. 1. The FTIR spectrum of pristine  $\text{Fe}_3\text{O}_4$  displays an Fe–O stretching band at  $564\text{ cm}^{-1}$ , while the functionalized samples show this band shifted to the range of  $555\text{--}558\text{ cm}^{-1}$ . The broad peaks observed between  $3200$  and  $3600\text{ cm}^{-1}$  and at  $1624\text{ cm}^{-1}$  correspond to the O–H bond vibrations of hydroxyl groups on the  $\text{Fe}_3\text{O}_4$  surface. In the FTIR spectrum of  $\text{Fe}_3\text{O}_4@\text{Gly}$ , additional peaks are observed at  $3488$ ,  $3434$ ,  $2973$ ,  $2920$ ,  $1629$ ,  $1303$ , and  $1100\text{ cm}^{-1}$ . These peaks are attributed to the stretching vibrations of  $\text{NH}_2$ , aliphatic C–H, and carboxylate groups, respectively, providing clear evidence of successful functionalization with glycine. The FTIR spectrum of  $\text{Fe}_3\text{O}_4@\text{salicylamide}$  displays new peaks at  $3132$  and  $3044\text{ cm}^{-1}$ , which are assigned to the N–H stretching vibration of the amide group and the aromatic C–H stretching

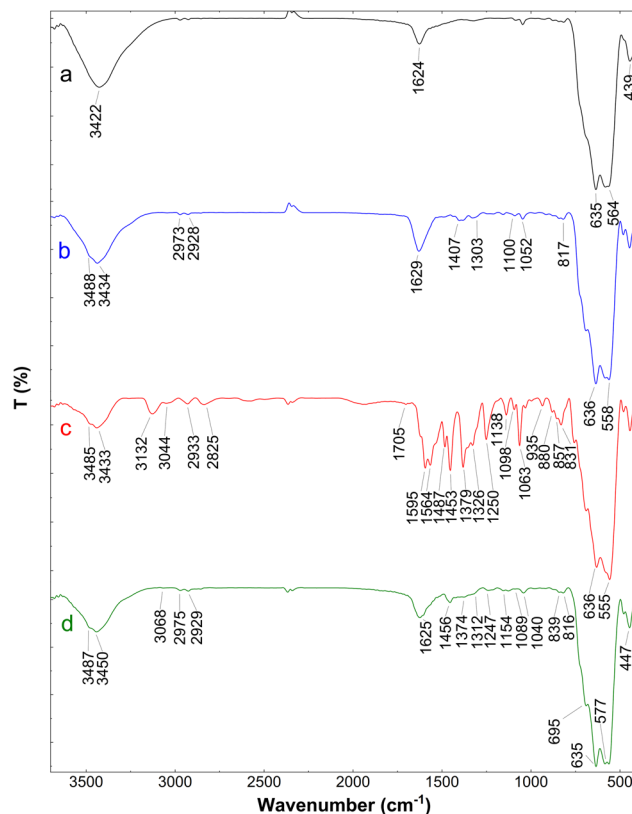
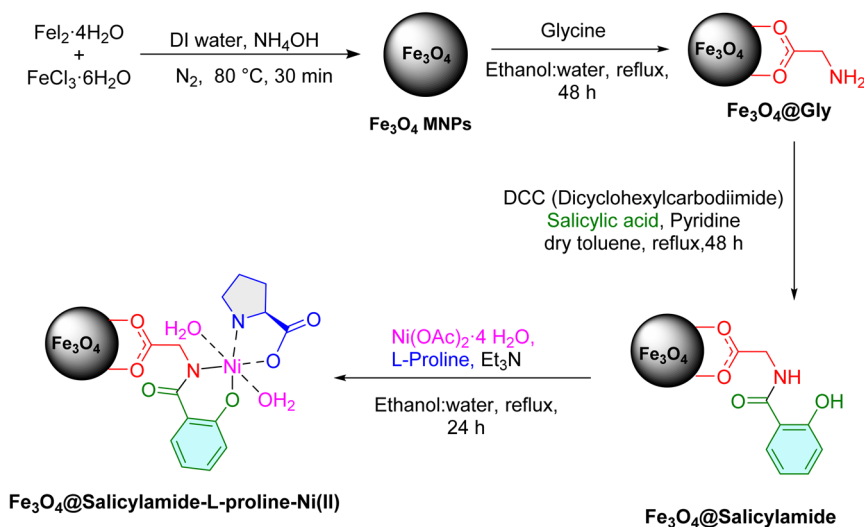


Fig. 1 FT-IR analysis of (a)  $\text{Fe}_3\text{O}_4$ , (b)  $\text{Fe}_3\text{O}_4@\text{Gly}$  (c)  $\text{Fe}_3\text{O}_4@\text{salicylamide}$  and (d)  $[\text{Fe}_3\text{O}_4@\text{salicylamide-L-proline-Ni(II)}]$  nanocomposite.

vibration. The band at  $1705\text{ cm}^{-1}$  corresponds to the carbonyl stretching vibration, while aromatic C–C stretching vibrations are observed at  $1595$  and  $1487\text{ cm}^{-1}$ . An Ar–O stretching vibration is observed at  $1250\text{ cm}^{-1}$ , further supporting successful modification of the  $\text{Fe}_3\text{O}_4$  surface with salicylamide. The FTIR spectrum of the  $[\text{Fe}_3\text{O}_4@\text{salicylamide-L-proline-Ni(II)}]$



Scheme 1 Stepwise synthesis of  $[\text{Fe}_3\text{O}_4@\text{salicylamide-L-proline-Ni(II)}]$  nanocomposite.



nanocomposite reveals a peak at  $1154\text{ cm}^{-1}$ , indicating the presence of L-proline in the complex.<sup>51</sup> The incorporation of Ni(II) was confirmed through EDX, ICP, and elemental mapping analyses, providing additional validation of the successful synthesis of the complex and the presence of Ni.

Powder X-ray diffraction (P-XRD) analysis was used to examine the crystalline structure of the  $[\text{Fe}_3\text{O}_4@\text{salicylamide-L-proline-Ni(II)}]$  nanocomposite. The resulting diffraction pattern displayed sharp peaks at  $2\theta$  values of  $30.34^\circ$ ,  $35.64^\circ$ ,  $43.24^\circ$ ,  $53.79^\circ$ ,  $57.14^\circ$ , and  $62.94^\circ$ , corresponding to the (220), (311), (400), (422), (511) and (440) lattice planes of  $\text{Fe}_3\text{O}_4$ . The observed diffraction pattern closely matched the characteristic peaks of  $\text{Fe}_3\text{O}_4$  MNPs (JCPDS 88-0866),<sup>52</sup> further verifying that the  $\text{Fe}_3\text{O}_4$  crystalline phase was maintained throughout the functionalization process Fig. 2.

The thermal stability of the  $[\text{Fe}_3\text{O}_4@\text{salicylamide-L-proline-Ni(II)}]$  nanocomposite was assessed using TGA analysis (Fig. 3). A weight loss of approximately 4.2% observed below  $200^\circ\text{C}$  is attributed to the removal of physisorbed moisture and solvents. A second, more substantial weight loss of about 14% occurs between  $200$  and  $600^\circ\text{C}$ , corresponding to the degradation of the salicylamide and L-proline organic components on the catalyst surface through the pyrolysis reaction. These findings confirm the presence of organic moieties on the catalyst surface and indicate its thermal stability up to  $200^\circ\text{C}$ , suggesting its suitability for reactions conducted at temperatures below this limit.

The chemical composition of the synthesized  $[\text{Fe}_3\text{O}_4@\text{salicylamide-L-proline-Ni(II)}]$  nanocomposite was examined using energy dispersive spectroscopy (EDS) (Fig. 4). The analysis revealed peaks corresponding to iron (55.21 wt%, 26.23 at%) and oxygen (22.63 wt%, 37.51 at%), confirming their presence as integral components of the  $\text{Fe}_3\text{O}_4$  structure. Additionally, the detection of carbon (11.97 wt%, 26.47 at%) and nitrogen (3.57 wt%, 6.77 at%) peaks indicates the successful incorporation of organic linkers and ligands on the surface of  $\text{Fe}_3\text{O}_4$ . The observed nickel (6.62 wt%, 2.98 at%) peak further confirms the presence of nickel complexed with the ligands on the surface of

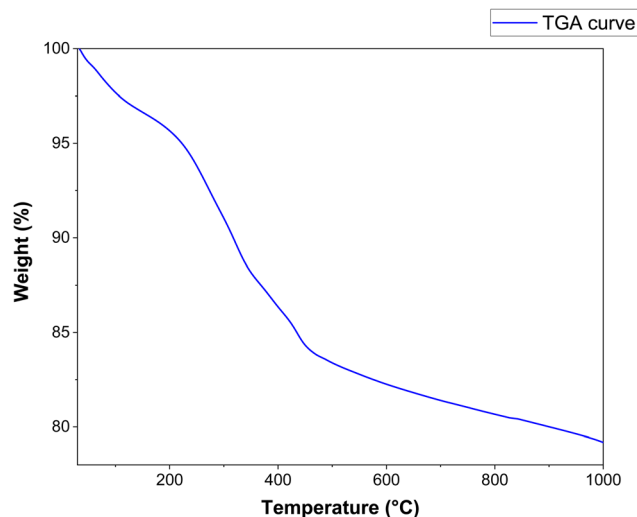


Fig. 3 TGA analysis of  $[\text{Fe}_3\text{O}_4@\text{salicylamide-L-proline-Ni(II)}]$  nanocomposite.

the magnetic nanoparticles (MNPs). Additionally, ICP-OES analysis revealed that the nickel content in the  $[\text{Fe}_3\text{O}_4@\text{salicylamide-L-proline-Ni(II)}]$  nanocomposite was approximately  $1.17 \times 10^{-3}\text{ mol g}^{-1}$ .

The elemental mapping images (Fig. 5) show that the surface of the  $[\text{Fe}_3\text{O}_4@\text{salicylamide-L-proline-Ni(II)}]$  nanocomposite is predominantly made up of iron and oxygen, which originate from the magnetic support. Additionally, carbon, nitrogen, and nickel are present in lower amounts but are evenly distributed across the surface. This indicates that the ligands and complexes are securely attached to the surface, ensuring that guest reactants can easily access the active sites.

The morphology of the synthesized  $[\text{Fe}_3\text{O}_4@\text{salicylamide-L-proline-Ni(II)}]$  nanocomposite was investigated using scanning electron microscopy (SEM) analysis (Fig. 6). The SEM images show spherical shape crystallites arranged in a well-dispersed

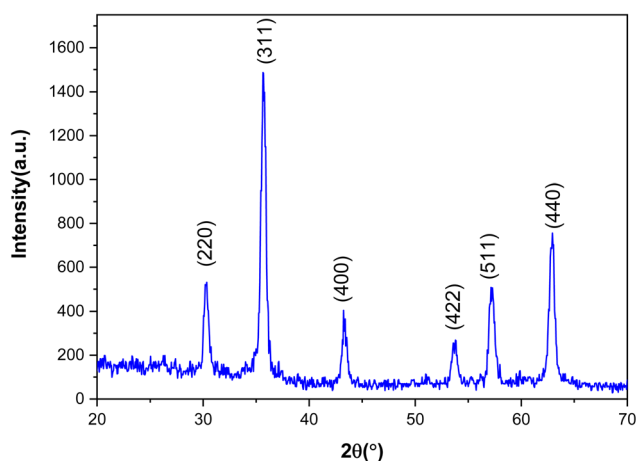


Fig. 2 XRD analysis of  $[\text{Fe}_3\text{O}_4@\text{salicylamide-L-proline-Ni(II)}]$  nanocomposite.

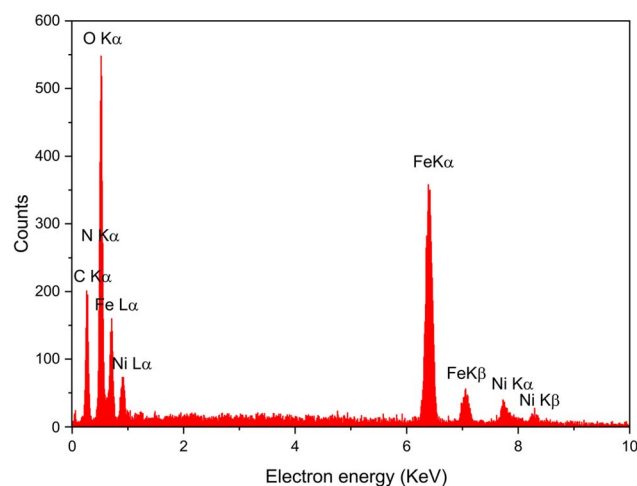


Fig. 4 EDAX analysis of  $[\text{Fe}_3\text{O}_4@\text{salicylamide-L-proline-Ni(II)}]$  nanocomposite.



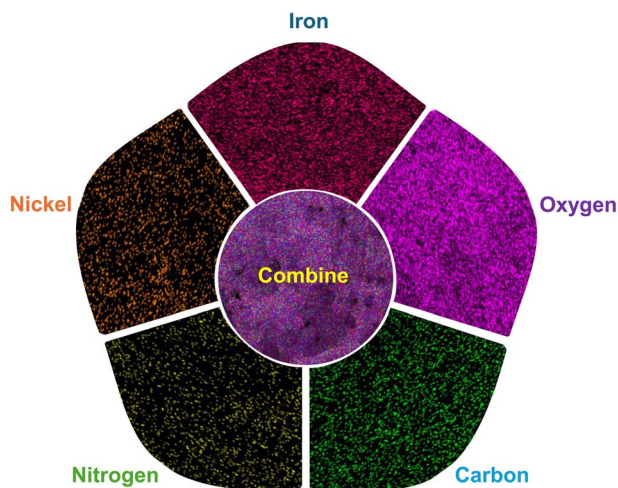


Fig. 5 EDX elemental mapping images of  $[\text{Fe}_3\text{O}_4@\text{salicylamide-L-proline-Ni(II)}]$  nanocomposite.

manner. The overall surface morphology exhibits the characteristic features of iron oxide nanoparticles, confirming their stability throughout the post-synthetic modification process.<sup>53</sup>

While some degree of agglomeration is observed, it is likely attributed to the surface modification, which further supports the successful synthesis of the target catalyst.

The TEM micrographs of the  $[\text{Fe}_3\text{O}_4@\text{salicylamide-L-proline-Ni(II)}]$  nanocomposite clearly depict a spherical morphology, with particle sizes ranging from 14 to 25 nm and a uniform dispersion. These images reveal distinct surface characteristics, further confirming the successful immobilization of the targeted complex onto the MNPs support (Fig. 7).

Fig. 8 displays the magnetization curves of the prepared samples at room temperature. The saturation magnetization ( $M_s$ ) values for bare  $\text{Fe}_3\text{O}_4$ , as well as for the synthesized  $\text{Fe}_3\text{O}_4@\text{Gly}$ ,  $\text{Fe}_3\text{O}_4@\text{salicylamide}$ , and  $[\text{Fe}_3\text{O}_4@\text{salicylamide-L-proline-Ni(II)}]$  nanocomposite, were approximately 60.97, 47.31, 41.02, and 29.96  $\text{emu g}^{-1}$ , respectively. The reduction in  $M_s$  for each functionalized MNPs can be attributed to the increased sample mass caused by the incorporation of diamagnetic organic ligands, which weaken the magnetic properties of the  $\text{Fe}_3\text{O}_4$  core. Nevertheless, the complex still retains notable magnetic properties, making it well-suited for various magnetically guided applications.

To analyze the porous structure of the  $[\text{Fe}_3\text{O}_4@\text{salicylamide-L-proline-Ni(II)}]$  nanocomposite,  $\text{N}_2$  physisorption was used.

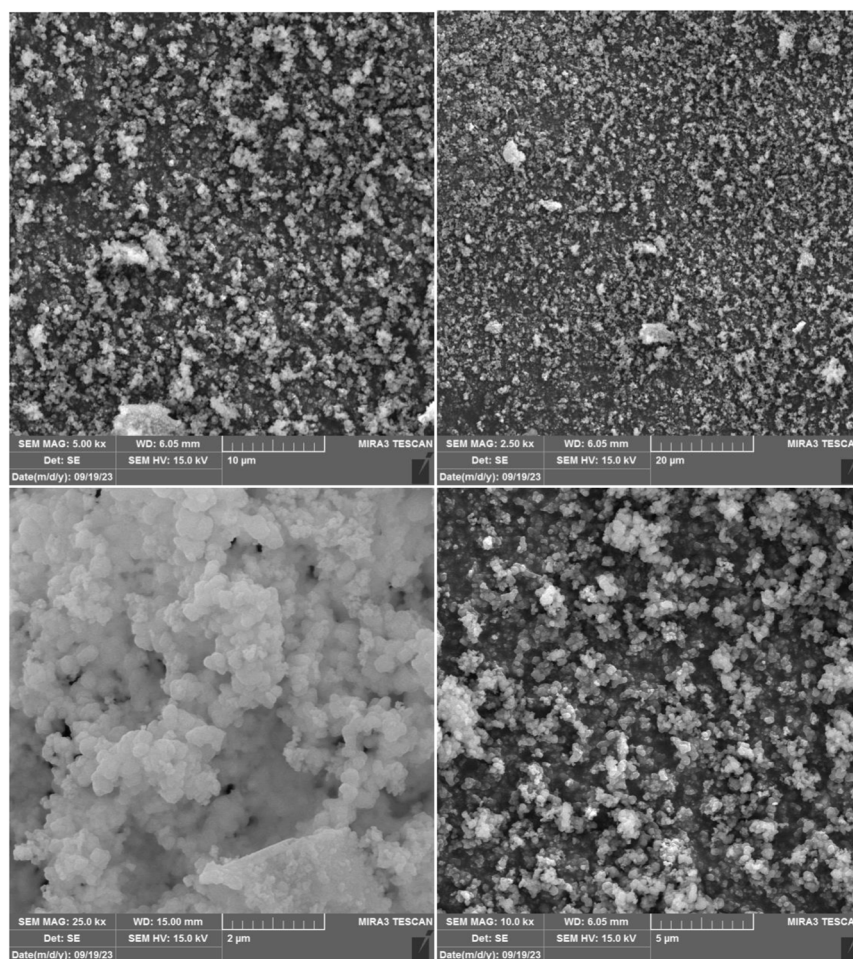


Fig. 6 SEM images of  $[\text{Fe}_3\text{O}_4@\text{salicylamide-L-proline-Ni(II)}]$  nanocomposite.



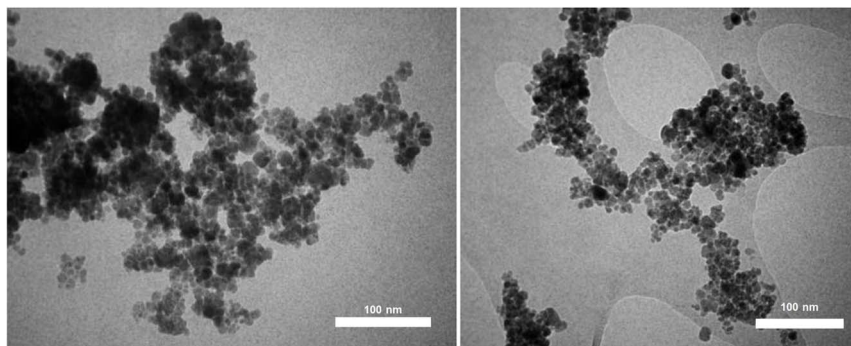


Fig. 7 TEM images of  $[\text{Fe}_3\text{O}_4@\text{salicylamide-L-proline-Ni(II)}]$  nanocomposite.

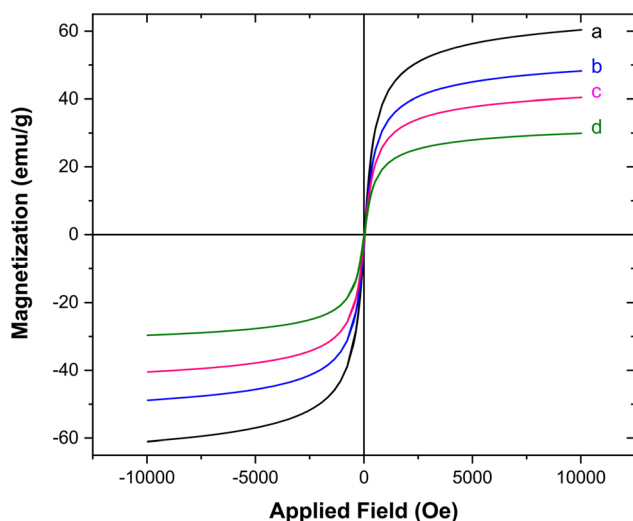


Fig. 8 VSM analysis of (a)  $\text{Fe}_3\text{O}_4$ , (b)  $\text{Fe}_3\text{O}_4@\text{Gly}$ , (c)  $\text{Fe}_3\text{O}_4@\text{salicylamide}$  and (d)  $[\text{Fe}_3\text{O}_4@\text{salicylamide-L-proline-Ni(II)}]$  nanocomposite.

The resulting type IV isotherms with a noticeable hysteresis loop suggest a mesoporous character, which was further supported by a pore diameter of about 42.168 nm and a total pore volume of  $0.098 \text{ cm}^3 \text{ g}^{-1}$ . The calculated specific surface area of  $47.55 \text{ m}^2 \text{ g}^{-1}$  is lower than that of pure  $\text{Fe}_3\text{O}_4$  ( $72.148 \text{ m}^2 \text{ g}^{-1}$ ),<sup>54</sup> likely due to pore obstruction and cavity filling during the functionalization process (Fig. 9).

### 3.2. Catalytic study

After thoroughly characterizing the  $[\text{Fe}_3\text{O}_4@\text{salicylamide-L-proline-Ni(II)}]$  nanocomposite, we investigated its catalytic activity in various organic reactions, including the synthesis of tetrazoles. In the first stage, we optimized key reaction parameters such as catalyst loading, solvent, and temperature (Table 1). The optimization was carried out using the  $[\text{Fe}_3\text{O}_4@\text{salicylamide-L-proline-Ni(II)}]$  nanocomposite as the catalyst in a model reaction involving 1 mmol of benzonitrile and 1.3 mmol of sodium azide. Initially, the reaction was conducted without a catalyst, yielding no conversion (Table 1, entry 1). When the reaction was performed in PEG-400 under reflux,

varying amounts of catalyst were tested, and a significant improvement in yield was observed (Table 1, entries 2–5). A catalyst loading of 1 mg resulted in a low yield (Table 1, entry 2), while increasing the catalyst amount to 5 mg led to an excellent product yield (Table 1, entry 5). However, further increasing the catalyst loading to 7 mg did not enhance the yield or reduce the reaction time (Table 1, entry 6). Subsequently, various green solvents, including water, ethanol and their mixture (1 : 1), were also tested, but none produced satisfactory results (Table 1, entries 7 and 8). Furthermore, reducing the reaction temperature to  $80^\circ\text{C}$  or room temperature resulted in only moderate or negligible product formation, respectively (Table 1, entries 10 and 11). The influence of sodium azide stoichiometry on reaction efficacy was subsequently examined (Table 1, entries 12 and 13). Quantification of product yields revealed that an

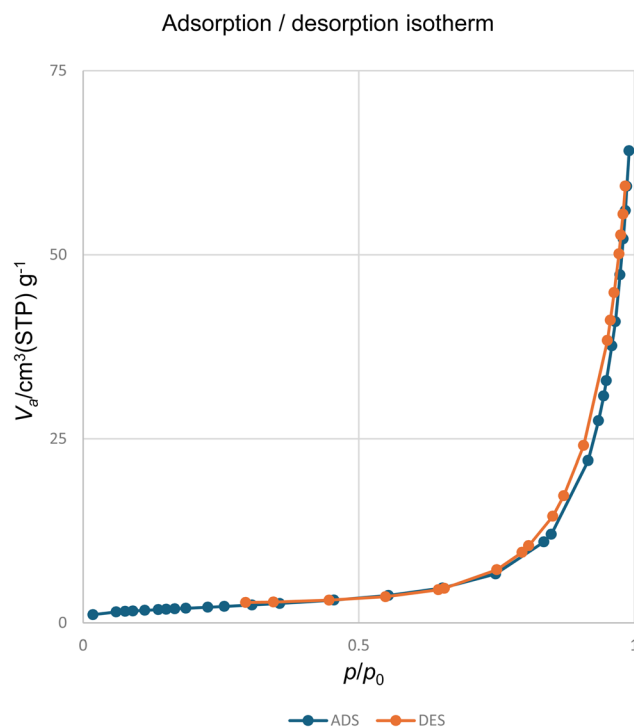
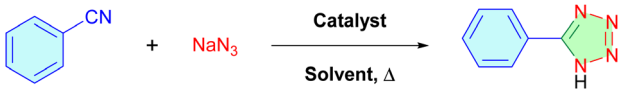


Fig. 9  $\text{N}_2$  adsorption–desorption isotherm of  $[\text{Fe}_3\text{O}_4@\text{salicylamide-L-proline-Ni(II)}]$  nanocomposite.



Table 1 Optimization of synthesis of 5-phenyl-1*H*-tetrazol in the presence of [Fe<sub>3</sub>O<sub>4</sub>@salicylamide-*L*-proline-Ni(II)] complex


| Entry | Catalyst   | Catalyst amount (mol%) | Solvent                 | Temperature (°C) | Time (min) | Yield <sup>a,b</sup> (%) |
|-------|--|------------------------|-------------------------|------------------|------------|--------------------------|
| 1     | [Fe <sub>3</sub> O <sub>4</sub> @salicylamide- <i>L</i> -proline-Ni(II)] | —                      | PEG-400                 | 120              | 1 day      | NR                       |
| 2     | [Fe <sub>3</sub> O <sub>4</sub> @salicylamide- <i>L</i> -proline-Ni(II)] | 1                      | PEG-400                 | 120              | 90         | 36                       |
| 3     | [Fe <sub>3</sub> O <sub>4</sub> @salicylamide- <i>L</i> -proline-Ni(II)] | 2                      | PEG-400                 | 120              | 50         | 69                       |
| 4     | [Fe <sub>3</sub> O <sub>4</sub> @salicylamide- <i>L</i> -proline-Ni(II)] | 4                      | PEG-400                 | 120              | 30         | 92                       |
| 5     | [Fe <sub>3</sub> O <sub>4</sub> @salicylamide- <i>L</i> -proline-Ni(II)] | 5                      | PEG-400                 | 120              | 25         | 97                       |
| 6     | [Fe <sub>3</sub> O <sub>4</sub> @salicylamide- <i>L</i> -proline-Ni(II)] | 7                      | PEG-400                 | 120              | 25         | 97                       |
| 7     | [Fe <sub>3</sub> O <sub>4</sub> @salicylamide- <i>L</i> -proline-Ni(II)] | 5                      | Water                   | Reflux           | 180        | 45                       |
| 8     | [Fe <sub>3</sub> O <sub>4</sub> @salicylamide- <i>L</i> -proline-Ni(II)] | 5                      | Ethanol                 | Reflux           | 135        | 73                       |
| 9     | [Fe <sub>3</sub> O <sub>4</sub> @salicylamide- <i>L</i> -proline-Ni(II)] | 5                      | Water : ethanol (1 : 1) | Reflux           | 30         | 64                       |
| 10    | [Fe <sub>3</sub> O <sub>4</sub> @salicylamide- <i>L</i> -proline-Ni(II)] | 5                      | PEG-400                 | 80               | 2 h        | 51                       |
| 11    | [Fe <sub>3</sub> O <sub>4</sub> @salicylamide- <i>L</i> -proline-Ni(II)] | 5                      | PEG-400                 | r.t.             | 6 h        | Trace                    |
| 12    | [Fe <sub>3</sub> O <sub>4</sub> @salicylamide- <i>L</i> -proline-Ni(II)] | 5                      | PEG-400                 | 120              | 25         | 97 <sup>c</sup>          |
| 13    | [Fe <sub>3</sub> O <sub>4</sub> @salicylamide- <i>L</i> -proline-Ni(II)] | 5                      | PEG-400                 | 120              | 25         | 94 <sup>d</sup>          |
| 14    | Fe <sub>3</sub> O <sub>4</sub>   | 5                      | PEG-400                 | 120              | 120        | Trace                    |
| 15    | Fe <sub>3</sub> O <sub>4</sub> @Gly                                      | 5                      | PEG-400                 | 120              | 120        | Trace                    |
| 16    | Fe <sub>3</sub> O <sub>4</sub> @salicylamide                             | 5                      | PEG-400                 | 120              | 120        | Trace                    |

<sup>a</sup> Isolated yield. <sup>b</sup> Conditions: benzonitrile (1 mmol), sodium azide (1.2 mmol), [Fe<sub>3</sub>O<sub>4</sub>@salicylamide-*L*-proline-Ni(II)] catalyst (mg) and solvent (3 mL). <sup>c</sup> The reaction was carried out with 1.4 mmol of NaN<sub>3</sub>. <sup>d</sup> The reaction was carried out with 1 mmol of NaN<sub>3</sub>.

increase in NaN<sub>3</sub> from 1.2 mmol to 1.4 mmol did not significantly enhance reaction performance. Conversely, a reduction to 1.0 mmol resulted in a demonstrable decrease in yield. Furthermore, to elucidate the catalytic role of the Ni species, a comparative study was conducted utilizing various catalyst intermediates: bare Fe<sub>3</sub>O<sub>4</sub>, Fe<sub>3</sub>O<sub>4</sub>@Gly, and Fe<sub>3</sub>O<sub>4</sub>@salicylamide (Table 1, entries 14–16). These intermediates yielded only trace amounts of the desired product, underscoring the indispensable contribution of the nickel(II) component in achieving optimal catalytic activity for the [Fe<sub>3</sub>O<sub>4</sub>@salicylamide-*L*-proline-Ni(II)] catalyst.

Overall, the optimal conditions were found to be 5 mg of the [Fe<sub>3</sub>O<sub>4</sub>@salicylamide-*L*-proline-Ni(II)] catalyst in PEG-400 at 120 °C, which yielded the best results for the synthesis of 5-substituted 1*H*-tetrazoles.

The scope and efficiency of the optimized methodology was evaluated in various substrates including substituted benzonitriles, as well as different aliphatic cyano compounds (Table 2). The results revealed that all substrates reacted effectively, yielding the desired tetrazoles in good to excellent amounts. The electron-withdrawing groups on the aryl ring increased reactivity, while electron-donating groups had the opposite effect. Moreover, the reaction was successfully applied to aliphatic nitriles, leading to the formation of the desired tetrazoles with yields reaching up to 90%. Additionally, the reaction of dicyanoarenes including; phthalonitrile, isophthalonitrile, and terephthalonitrile derivatives (Table 2 entries 5–7), with a stoichiometric equivalent of azide reagent afforded the desired 2-(1*H*-tetrazol-5-yl)benzonitrile derivative as the sole product. The observed retention of the second cyano

substituent demonstrates the remarkable homoselectivity of this catalytic methodology for the synthesis of 5-substituted tetrazoles. Finally, a 10-fold scale-up of the reaction, using benzonitrile as a model reaction maintaining stoichiometric ratios, yielded 94% of the desired product in 35 min, demonstrating the method's feasibility for gram-scale production. This efficient and rapid transformation confirms the method's potential for industrial application in 5-substituted 1*H*-tetrazole synthesis.

The plausible mechanism for the catalytic cycle in the synthesis of 5-aryl-1*H*-tetrazoles, facilitated by the [Fe<sub>3</sub>O<sub>4</sub>@salicylamide-*L*-proline-Ni(II)] nanocomposite, is outlined in Scheme 2. In the first step, the Ni(II) center interacts with the sodium azide salt (NaN<sub>3</sub>), generating a reactive nitrogen species. This intermediate then reacts with an organic nitrile (R-CN) on the catalyst surface. The interaction between the intermediate and the Ni complex induces the subsequent ring closure, resulting in the formation of the tetrazole structure. The cycle concludes with the regeneration of the catalyst, making it ready for the next catalytic iteration.<sup>65</sup>

### 3.3. Reusability of catalyst

The ability to recycle a catalyst from the reaction medium is a powerful characteristic that makes it particularly valuable for catalysis applications, especially from an environmental perspective.<sup>66</sup> In this regard, we investigated the model reaction under optimized conditions for the synthesis of 5-phenyl-1*H*-tetrazole derivatives. After the reaction reached completion, as indicated in Table 2, the reaction was stopped, and the catalyst was separated following the procedure outlined in the



Table 2 The scope of synthesis of 5-substituted 1*H*-tetrazoles catalyzed by [Fe<sub>3</sub>O<sub>4</sub>@Salicylamide-L-proline-Ni(II)] nanocomposite

| Entry | Aryl nitrile | Product | Time (min) | Yield <sup>a,b</sup> (%) | TON    | TOF (min <sup>-1</sup> ) | Melting point [ref.] |
|-------|--------------|---------|------------|--------------------------|--------|--------------------------|----------------------|
| 1     |              |         | 25         | 97                       | 16 440 | 657                      | 218–220 (ref. 55)    |
| 2     |              |         | 20         | 95                       | 16 101 | 805                      | 159–161 (ref. 56)    |
| 3     |              |         | 25         | 97                       | 16 440 | 657                      | 155–156 (ref. 55)    |
| 4     |              |         | 15         | 99                       | 16 779 | 1118                     | 219–221 (ref. 55)    |
| 5     |              |         | 20         | 92                       | 15 593 | 779                      | 209–212 (ref. 57)    |
| 6     |              |         | 27         | 96                       | 16 271 | 602                      | 210–212 (ref. 57)    |
| 7     |              |         | 18         | 98                       | 16 610 | 922                      | 251–254 (ref. 57)    |
| 8     |              |         | 30         | 88                       | 14 915 | 497                      | 295–296 (ref. 58)    |
| 9     |              |         | 40         | 93                       | 15 762 | 394                      | 152–154 (ref. 59)    |
| 10    |              |         | 28         | 96                       | 16 271 | 581                      | 249–251 (ref. 55)    |



Table 2 (Contd.)

| Entry | Aryl nitrile | Product | Time (min) | Yield <sup>a,b</sup> (%) | TON    | TOF (min <sup>-1</sup> ) | Melting point [ref.] |
|-------|--------------|---------|------------|--------------------------|--------|--------------------------|----------------------|
|       |              |         |            |                          |        |                          |                      |
| 11    |              |         | 85         | 88                       | 14 915 | 175                      | 222–224 (ref. 60)    |
| 12    |              |         | 45         | 95                       | 16 101 | 537                      | 200–204 (ref. 61)    |
| 13    |              |         | 80         | 91                       | 15 423 | 192                      | 266–269 (ref. 55)    |
| 14    |              |         | 80         | 89                       | 15 084 | 188                      | 225–227 (ref. 62)    |
| 15    |              |         | 37         | 95                       | 16 101 | 435                      | 244–246 (ref. 63)    |
| 16    |              |         | 80         | 92                       | 15 593 | 194                      | 231–234 (ref. 55)    |
| 17    |              |         | 30         | 97                       | 16 440 | 548                      | 140–143 (ref. 64)    |
| 18    |              |         | 25         | 97                       | 16 440 | 657                      | 154–155 (ref. 55)    |

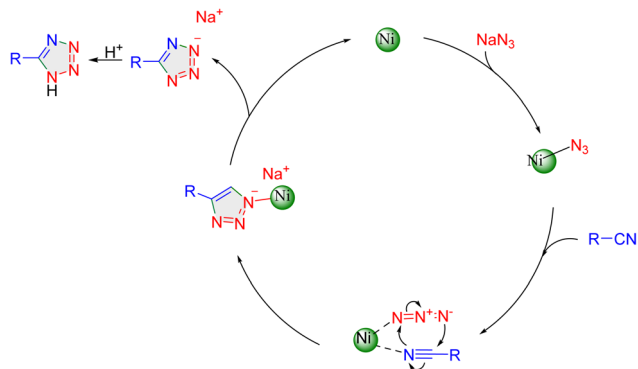
<sup>a</sup> Isolated yield. <sup>b</sup> Conditions: aryl nitrile (1.0 mmol), sodium azide (1.2 mmol) and [Fe<sub>3</sub>O<sub>4</sub>@salicylamide-L-proline-Ni(II)] complex (5 mol%) in PEG-400 (2 mL) at 120 °C.

Experimental section. It was then washed several times with ethyl acetate (EtOAc), water, and ethanol, before being dried in an oven at 80 °C. The recovered catalyst was subsequently reused in the next reaction cycle. The results demonstrate excellent reusability, with only a negligible loss in efficiency over eight consecutive reaction cycles (Fig. 10). These findings

confirm the reusability and recyclability of the prepared catalyst under the optimized conditions.

To evaluate the stability of the [Fe<sub>3</sub>O<sub>4</sub>@salicylamide-L-proline-Ni(II)] nanocomposite following recyclability experiments, FT-IR, VSM and ICP-OES analyses were performed. The consistent FT-IR pattern observed after eight cycles (Fig. 11)





Scheme 2 Possible mechanism for synthesis of 5-substituted-1H-tetrazoles over the catalysis of  $[\text{Fe}_3\text{O}_4@\text{salicylamide-L-proline-Ni(II)}]$  nanocomposite.

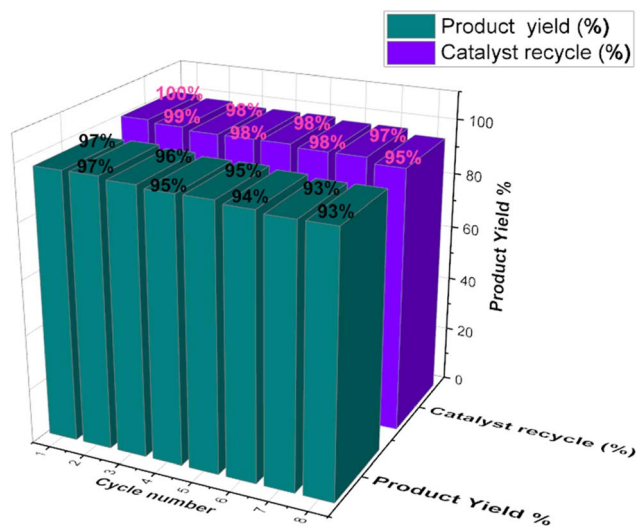


Fig. 10 The reusability of  $[\text{Fe}_3\text{O}_4@\text{salicylamide-L-proline-Ni(II)}]$  nanocomposite in the model reaction of 5-phenyl-1H-tetrazole synthesis.

demonstrates the exceptional chemostability and recyclability of the catalyst. Additionally, the  $M_s$  value of the recycled  $[\text{Fe}_3\text{O}_4@\text{salicylamide-L-proline-Ni(II)}]$  catalyst was determined to be  $26.37 \text{ emu g}^{-1}$  (Fig. 12). A comparative analysis with the fresh catalyst demonstrated only a modest reduction of  $3.59 \text{ emu g}^{-1}$  in the  $M_s$  value following eight recycling cycles. This observation indicates a high degree of magnetic stability, confirming that the material maintains sufficient magnetic responsiveness for effective separation *via* neodymium magnets. Finally, the ICP-OES results show that the Ni content in the recycled catalyst is nearly identical to that of the fresh catalyst ( $1.16 \times 10^{-3} \text{ mol g}^{-1}$ ), indicating no significant metal leaching from the catalyst surface during the reaction cycles.

#### 3.4. Leaching and hot filtration test

The nature of the catalytic species and their potential leaching into the reaction medium was examined using a hot filtration test and instrumental analysis. For this investigation, the model

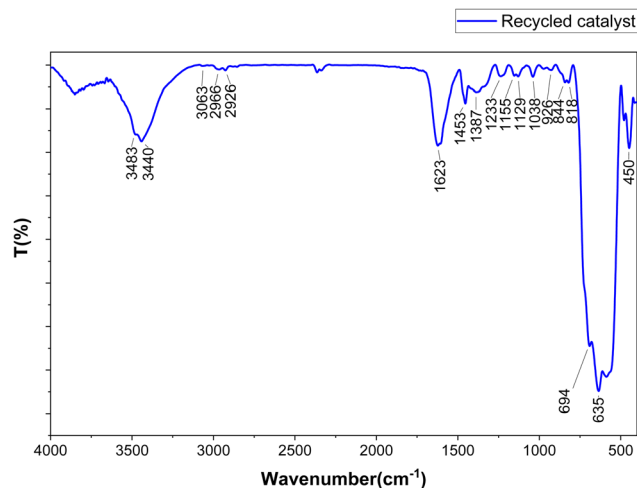


Fig. 11 FT-IR analysis of recovered  $[\text{Fe}_3\text{O}_4@\text{salicylamide-L-proline-Ni(II)}]$  nanocomposite.

reaction for synthesizing 5-phenyl-1H-tetrazole was selected. At the halfway point of the reaction, the  $[\text{Fe}_3\text{O}_4@\text{salicylamide-L-proline-Ni(II)}]$  nanocomposite was magnetically separated from the reaction mixture (with a product yield of 67% at this time), and the reaction was allowed to proceed for an additional half of the reaction time. The results revealed that the product yield increased by only a negligible 2% after catalyst removal, confirming the catalyst's predominantly heterogeneous nature. In addition, the ICP-OES analysis of the filtrate showed no trace of nickel, suggesting that there was very little metal leaching throughout the reaction.

#### 3.5. Comparison study of catalytic activity

In recent years, several catalytic methods have been developed for synthesizing 5-aryl-1H-tetrazoles, given the significant importance of these compounds. However, a comparison between the catalytic performance of the  $[\text{Fe}_3\text{O}_4@\text{salicylamide-}$

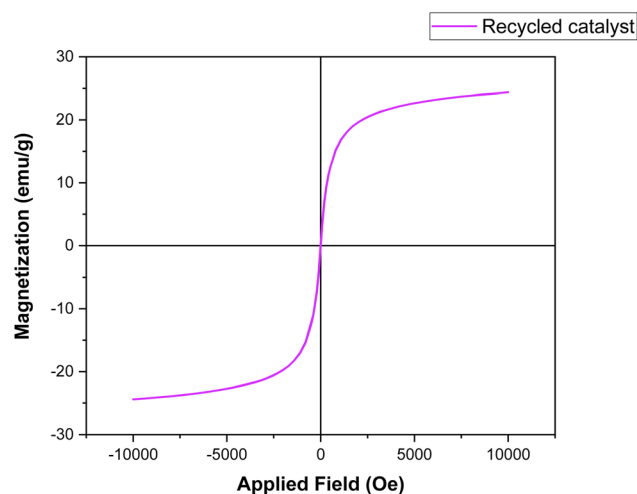


Fig. 12 VSM analysis of recovered  $[\text{Fe}_3\text{O}_4@\text{salicylamide-L-proline-Ni(II)}]$  nanocomposite.



**Table 3** Comparison the  $[\text{Fe}_3\text{O}_4@\text{salicylamide-L-proline-Ni(II)}]$  nanocomposite catalytic activity in synthesis of 5-phenyl-1*H*-tetrazole model reaction

| Entry | Catalyst (active site)   | Time (min) | Isolated yield (%) | Reference |
|-------|--|------------|--------------------|-----------|
| 1     | $\text{Fe}_3\text{O}_4@\text{L-lysine-Pd(0)}$ (palladium)                  | 60         | 99                 | 27        |
| 2     | $[\text{Fe}_3\text{O}_4@\text{TAM-Schiff-base-Cu}^{\text{II}}]$ (copper)   | 100        | 98                 | 68        |
| 3     | $\text{Fe}_3\text{O}_4@\text{SiO}_2@\text{SBA-3}@2\text{-ATP-Cu}$ (copper) | 65         | 94                 | 69        |
| 4     | $\text{Co-(PYT)}_2@\text{BNPs}$ (cobalt)                                   | 120        | 98                 | 70        |
| 5     | CF/MC/HA/A (silver)  | 10         | 93                 | 29        |
| 6     | Ag NPs (silver)  | 3 h        | 88                 | 71        |
| 7     | $\text{CoFe}_2\text{O}_4@\text{amino glycol/Gd}$ (gold)                    | 10         | 97                 | 32        |
| 8     | AuNPs (gold)   | 30 min     | 95                 | 72        |
| 9     | MCM-41@serine/Er (erbium)  | 10         | 96                 | 31        |
| 10    | MCM-41/3,4,5-tri-hydroxyphenyl acetic acid/Sm (samarium)                   | 20         | 92                 | 33        |
| 11    | $\text{InCl}_3$ (indium)   | 15 h       | 92                 | 28        |
| 12    | Nd-Schiff-base@BMNPs (neodymium)   | 3 h        | 98                 | 30        |
| 13    | Polymer-bound $\text{AlCl}_3$ (aluminium)                                  | 6 h        | 79                 | 20        |
| 14    | SSA (sulfuric acid)  | 4 h        | 88                 | 67        |
| 15    | $[\text{Fe}_3\text{O}_4@\text{salicylamide-L-proline-Ni(II)}]$ (nickel)    | 25         | 97                 | This work |

L-proline-Ni(II)] nanocomposite and the most efficient green methods in the literature reveals several limitations of the latter. While previous catalysts often suffer from issues like high costs (especially for precious metals such as palladium, indium,<sup>28</sup> silver,<sup>29</sup> neodymium,<sup>30</sup> erbium,<sup>31</sup> gadolinium,<sup>32</sup> and samarium<sup>33</sup>), the need for harsh acidic conditions,<sup>67</sup> long reaction times, and toxic reagents or solvents, the  $[\text{Fe}_3\text{O}_4@\text{salicylamide-L-proline-Ni(II)}]$  nanocomposite stands out as a more sustainable alternative. It performs excellently under environmentally friendly conditions, achieving high yields in short reaction times. Moreover, it was synthesized using low-cost, green reagents through a simple and multi-gram scale strategy and shows minimal metal leaching. This makes it a promising, efficient, and eco-friendly option compared to conventional methods (Table 3).

## 4. Conclusions

In conclusion, a novel nanomagnetic nickel complex was successfully synthesized by a straightforward three-step post-synthesis method. This involved functionalizing  $\text{Fe}_3\text{O}_4$  magnetic nanoparticles with glycine, converting it to salicylamide, and subsequently complexing with nickel and proline. The resulting material exhibits excellent thermal stability, a high surface area, and a spherical morphology with particle sizes ranging from 14 to 25 nm. This catalyst proved highly effective in the homoselective synthesis of 5-substituted-1*H*-tetrazoles *via* click condensation of various organic nitriles and sodium azide in PEG-400 at 120 °C, affording good to excellent yields (88–99 yield% in 15–85 min). Furthermore, the catalyst demonstrated remarkable reusability, enhanced stability, and minimal leaching. These findings highlight the potential of this catalyst for advancing catalysis, particularly in surface amination using glycine and the immobilization of carboxyl-containing ligands *via* direct amidation. These heterogeneous catalysts hold considerable promise for diverse catalytic applications. Future studies could explore the applicability of this

direct amidation method on the catalyst surface for immobilizing carboxylic acid-containing ligands, further enhancing its potential for catalytic applications.

## Data availability

The authors declare that all the data are available the data within the paper.

## Author contributions

Chou-Yi Hsu: characterization of catalyst. Ghusoon Faidhi Hameed: writing original draft, laboratory works. Abhinav Kumar: conceptualization, analysis, review draft, acquiring research funding and supervision. Subbulakshmi Ganesan and S. Sunitha: laboratory works, catalysis studies and analysis. Irfan Ahmad and Aman Shankhyan: conceptualization, software and review/editing. Rajashree Panigrahi: laboratory works, analysis, review/editing.

## Conflicts of interest

The authors declare that they have no competing interests.

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