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Fabrication of magnetic carbohydrate-modified iron oxide nanoparticles (Fe₃O₄/pectin) decorated with bimetallic Co/Cu-MOF as an effective and recoverable catalyst for the Biginelli reaction

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Due to their biocompatibility, facile recoverability, mechanical and thermal stability, high surface area, and active catalytic sites, magnetic nanocomposites, containing natural polymers and magnetic nanoparticles, have been used to produce supports for catalysts or biocatalysts. Pectin, an important polycarbohydrate, has abundant functional groups with excellent ability to coat the surface of the nanoparticles to fabricate composite and hybrid materials. A novel bimetallic cobalt(II) and copper(II)-based metal–organic framework (Co/Cu-MOF) immobilized pectin-modified Fe₃O₄ magnetic nanocomposite was designed and fabricated. Fe₃O₄ nanoparticles were modified *in situ* by pectin and, subsequently, used as a support for growing Co/Cu-MOF [Fe₃O₄/pectin/(Co/Cu)MOF]. The properties of the nanocomposite were investigated by FT-IR, XRD, SEM, EDS, VSM, STA, and BET. The nanocomposite exhibited both magnetic characteristics and a high surface area, making it a suitable candidate for catalytic applications. Then, the Fe₃O₄/pectin/(Co/Cu)MOF nanocomposite was utilized in the Biginelli reaction for the production of biologically active dihydropyrimidinones. Due to paramagnetism, Fe₃O₄/pectin/(Co/Cu)MOF was easily recovered and reused in six cycles without significant loss in reactivity. This green method comprises several benefits, such as mild reaction conditions, free-solvent media, high yields, easy workup, short reaction times and reusability of the prepared catalyst.

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1. Introduction

The Biginelli reaction represents incredibly significant multi-component reactions that generate bioactive compounds with various therapeutic and pharmacological traits, such as anti-tumor, antibacterial, and antioxidant.^{1–3} The Biginelli reaction includes the three-component, acid-catalyzed condensation of aldehydes, β-ketoester, and urea to yield dihydropyrimidones.^{4–10} So far, various compounds, including Brønsted or Lewis acids and ionic liquids, have been utilized to catalyze the Biginelli reaction.^{4–11}

Magnetic nanoparticles (MNPs) have attracted considerable interest as exceptional supports for immobilization, owing to the presence of abundant surface hydroxyl groups, highly stable, straightforward accessibility, great permeability, inexpensiveness, and easy separation using an external magnet.^{12–14} The surface of MNPs can be functionalized by a variety of polymers.^{15,16} Among them, biopolymers receive special attention because of their biocompatibility and biodegradability.^{17,18} Pectin, as a natural biopolymer, is a water-soluble gelatin-like

material and is known as one of the structurally complicated members of polycarbohydrates abundant in the cell walls of plants with key roles in primary/secondary structure of the wall and its function.¹⁹ Carbohydrate-based coatings, especially coating by pectin, on the surface of various nanoparticles, liposomes, foods and materials have recently attracted the attentions of scientists due to the ability of carbohydrates to enhance the physicochemical and/or storage stability of the materials, or adding specific functions for different purposes such as drug delivery. Pectin possesses valuable properties, such as the availability of plenteous carboxyl and hydroxyl functional groups, non-toxicity, low-cost, high flexibility, and biodegradability that has made it an outstanding biomolecule for use in surface engineering nanotechnology.²⁰

Metal–organic frameworks (MOFs) represent a class of porous crystalline materials, constructed from metal ions or metal oxide clusters bridged by organic linkers that exhibit substantial porosity, high surface area, and diverse functionalities.^{21,22} MOFs are of special interest owing to their extensive range of potential applications, such as catalysis, medicinal, gas storage, separation, and chemical sensors.^{23–31} However, the actual application of MOFs has been hampered by their poor stability, chemical instability, and difficult recovery. Recently, composites based on MOFs have been constructed by the

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combination of MOFs with materials including metal nanoparticles, quantum dots, natural enzymes, and polymers, which represent physical and chemical properties that are superior to those of the individual components, due to the combination of features from the two ingredients.^{32,33} The combination of functionalized magnetic nanoparticles and MOFs creates magnetic porous composites, which have attracted considerable attention due to a large number of advanced applications in diverse fields.³⁴ These magnetic porous composites possess a core-shell architecture containing a porous shell on the magnetic core. The resulting nanostructures presented magnetic properties and high surface area without aggregation of magnetite cores, enabling them to become desired candidates for the catalysts of organic reactions.³⁵ Bimetallic MOFs are MOFs comprising two various metal ions in their frameworks. These MOFs can be easily fabricated using the mixture of different metals *via* two synthetic methods: the one-pot synthesis method or the post-synthetic ion exchange method.^{36–38} These mixed-metal MOFs represent a superior catalytic activity compared to the corresponding single-metal MOFs, owing to the existence of two different metal active sites, which are suitable for orthogonal and multicomponent reactions as well as other synthetic routes requiring advanced multisite catalysts.^{39,40}

In the present work, Fe₃O₄/pectin/(Co/Cu)MOF nanocomposite was hydrothermally prepared and utilized as an effective and cost-effective heterogeneous catalyst for the Biginelli reaction under solvent-free conditions. The synergistic effects of Fe₃O₄, Co-MOF, and Cu-MOF are expected to enhance catalytic performance.

2. Experimental

2.1. Materials

The reagents and chemicals including iron(II) chloride tetrahydrate (FeCl₂·4H₂O), iron(III) chloride hexahydrate (FeCl₃·6H₂O), sodium hydroxide (NaOH), urea (CO(NH₂)₂), ethyl acetoacetate, cobalt(II) nitrate hexahydrate (Co(NO₃)₂·6H₂O), copper(II) nitrate trihydrate (Cu(NO₃)₂·3H₂O), 1,4-benzenedicarboxylic acid (1,4-BDC, terephthalic acid), *N,N*-dimethylformamide (DMF), ethanol, benzaldehyde, 4-nitrobenzaldehyde, 4-chlorobenzaldehyde, 2-chlorobenzaldehyde, 4-methoxybenzaldehyde, 4-methylbenzaldehyde, and 2-hydroxybenzaldehyde were purchased from Merck Company (Darmstadt, Germany) and used without further purification. Pectin was supplied by Lidoma Golden Day Ltd company (Tehran, Iran).

2.2. Apparatus

FT-IR spectra were recorded with a Jasco-680 spectrometer (Japan) in the range of 4000–400 cm⁻¹. FT-IR spectra of the compounds were collected by making their pellets in KBr, as a medium. The X-ray diffraction pattern of the prepared materials was recorded in the reflection mode using a Bruker, D8 Advance diffractometer. The ultrasonic bath (Tecno-GAZ SPA Ultrasonic system, Italy) was used at a frequency of 60 Hz and

power of 130 W. The surface morphology of the nanocomposite was examined by scanning electron microscopy (SEM; EM10C-ZEISS, 80 kV, Zeiss Co., Germany). Nitrogen adsorption/desorption isotherm was measured by Brunauer–Emmett–Teller analysis (Belsorp Mini II, Japan). The thermal stability of the nanocomposite was determined by Simultaneous Thermal Analyzer (STA, PerkinElmer STA6000, USA).

2.3. Preparation of Fe₃O₄/pectin

1.00 g of pectin was dissolved in 100 mL of distilled water under continuous stirring at room temperature. In another vessel, 5.00 g of FeCl₃·6H₂O and 2.00 g of FeCl₂·4H₂O were dissolved in 10 mL of distilled water. The solutions were mixed, and then the pH of the solution increased to 12 with NaOH. Finally, the mixture was stirred at 80 °C for 1 h and then cooled to room temperature. The resultant precipitate was collected through an external magnet, washed with deionized water and ethanol, and dried at 70 °C under vacuum.

2.4. Synthesis of Fe₃O₄/pectin/(Co–Cu)MOF

0.3 g of Fe₃O₄/pectin was dispersed in 30 mL of DMF under ultrasound irradiation for 20 min. Consequently, 0.1 g of Co(NO₃)₂·6H₂O, 0.09 g of Cu(NO₃)₂·3H₂O, and 0.13 g of 1,4-BDC were added to the above mixture and sonicated for 30 min. The mixture was placed in a 150 mL Teflon-lined autoclave and heated up to 140 °C for 48 h, then cooled back down to room temperature. The dark brown precipitate was collected using an external magnet, washed with ethanol, and dried at 70 °C under vacuum.

2.5. The Biginelli reaction using Fe₃O₄/pectin/(Co–Cu)MOF as a catalyst

General procedure: a mixture of aldehydes (1.0 mmol), ethyl acetoacetate (1.0 mmol), urea (1.5 mmol), and Fe₃O₄/pectin/(Co–Cu)MOF (25 mg) was stirred under solvent-free conditions at 85 °C. The progress of the reaction was monitored by thin-layer chromatography (TLC). After the reaction was completed, warm ethanol (10 mL) was added to the reaction mixture and Fe₃O₄/pectin/(Co–Cu)MOF was separated using a magnet and washed with ethanol. The solvent was evaporated and the solid was recrystallized from ethanol to produce the pure product. The products were identified by matching their melting points and FT-IR analysis. The recovered Fe₃O₄/pectin/(Co–Cu)MOF was reused in six runs under similar conditions as the first run to reveal the recyclability and stability of the catalyst.

5-(Ethoxycarbonyl)-6-methyl-4-phenyl-3,4-dihydropyrimidin-(1H)-one (4a): m.p.: 202–204 °C. ¹H NMR (400 MHz, DMSO-*d*₆) δ 9.27 (s, 1H), 7.89 (s, 1H), 7.47 (t, 2H), 7.58 (d, 3H), 5.32 (d, *J* = 3.4 Hz, 1H), 3.83 (q, *J* = 7.3 Hz, 2H), 2.21 (s, 3H), 1.12 (t, *J* = 7.3 Hz, 3H).

5-(Ethoxycarbonyl)-6-methyl-4-(4-nitrophenyl)-3,4-dihydropyrimidin-2(1H)-one (4b): m.p.: 202–203 °C. ¹H NMR (400 MHz, DMSO-*d*₆) δ 9.42 (s, 1H), 8.53 (d, *J* = 8.9 Hz, 2H), 7.86 (s, 1H), 7.63 (d, *J* = 8.9 Hz, 2H), 5.48 (d, *J* = 3.4 Hz, 1H), 3.76 (q, *J* = 7.4 Hz, 2H), 2.22 (s, 3H), 1.25 (t, *J* = 7.1 Hz, 3H).



5-(Ethoxycarbonyl)-6-methyl-4-(2-chlorophenyl)-3,4-dihydropyrimidin-2(1H)-one (4d): m.p.: 221–223 °C. ^1H NMR (400 MHz, $\text{DMSO-}d_6$) δ 9.47 (s, 1H), 7.68 (s, 1H), 7.13–7.38 (m, 4H), 5.71 (d, $J = 2.8$ Hz, 1H), 3.92 (q, $J = 7.4$ Hz, 2H), 2.41 (s, 3H), 0.89 (t, $J = 7.4$ Hz, 3H).

3. Results and discussion

3.1. Synthesis of $\text{Fe}_3\text{O}_4/\text{pectin}/(\text{Co-Cu})\text{MOF}$

The synthesis of $\text{Fe}_3\text{O}_4/\text{pectin}/(\text{Co-Cu})\text{MOF}$ for the Biginelli reaction was reported using commercially available chemical materials. Pectin is a natural biopolymer that is obtained from many fruits. This biopolymer is a polysaccharide containing carboxylic acid groups in each monomer. The carboxylic acid groups on the surface of pectin are covalently chelated to Fe_3O_4 nanoparticles and direct the nanostructure. Moreover, pre-treating Fe_3O_4 nanoparticles with pectin improved their compatibility with MOFs and facilitated the growth of MOFs around Fe_3O_4 nanoparticles. $\text{Fe}_3\text{O}_4/\text{pectin}$ was treated with $\text{Co}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$ and $\text{Cu}(\text{NO}_3)_2 \cdot 3\text{H}_2\text{O}$ (as inorganic constituents), 1,4-BDC, and DMF (as a solvent) *via* heating to 140 °C for 48 h to form $\text{Fe}_3\text{O}_4/\text{pectin}/(\text{Co-Cu})\text{MOF}$ nanocomposite. The

synthesis route of $\text{Fe}_3\text{O}_4/\text{pectin}/(\text{Co-Cu})\text{MOF}$ nanocomposite is displayed in Fig. 1.

3.2. Characterization of the prepared compounds

Fig. 2 displays the FT-IR spectra of pectin (a), $\text{Fe}_3\text{O}_4/\text{pectin}$ (b), $(\text{Co/Cu})\text{MOF}$ (c) and $\text{Fe}_3\text{O}_4/\text{pectin}/(\text{Co/Cu})\text{MOF}$ (d). In the FT-IR spectrum of pectin, the bands at 3389, and 2932 cm^{-1} are attributed to the stretching vibrations of O–H and C–H bonds, respectively. The peak which appeared around 1760 cm^{-1} corresponds to C=O stretching vibration. The peak at 1626 cm^{-1} is assigned to the bending vibration of O–H and the peak around 1120 cm^{-1} is related to the stretching vibration of

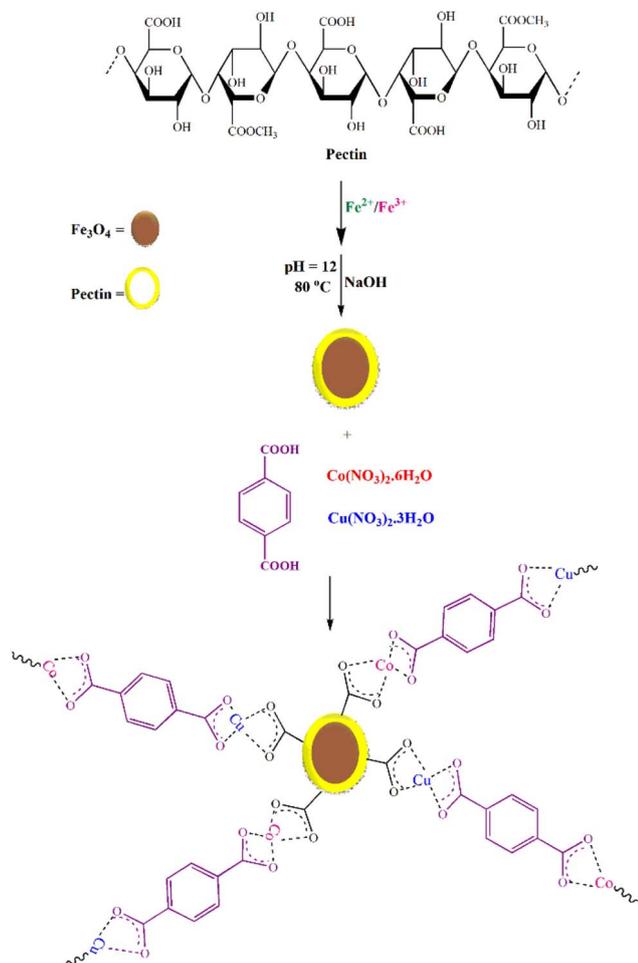


Fig. 1 Preparation of $\text{Fe}_3\text{O}_4/\text{pectin}/(\text{Co/Cu})\text{MOF}$ nanocomposite.

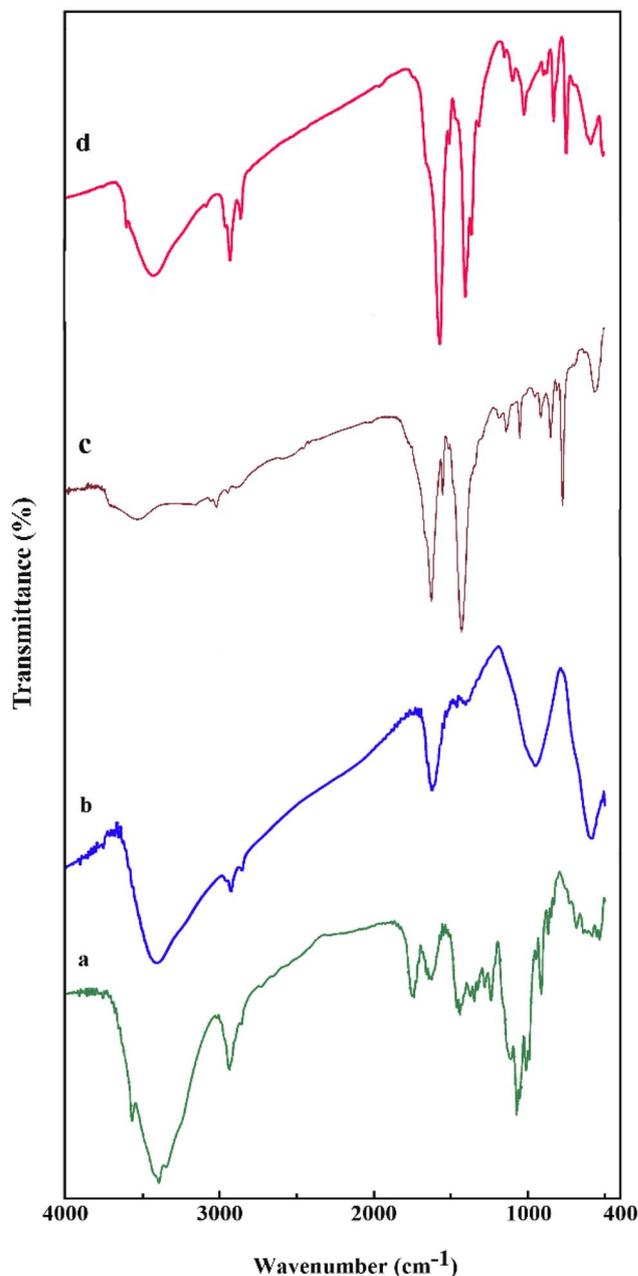


Fig. 2 FT-IR spectra of pectin (a), $\text{Fe}_3\text{O}_4/\text{pectin}$ (b), $(\text{Co/Cu})\text{MOF}$ (c) and $\text{Fe}_3\text{O}_4/\text{pectin}/(\text{Co/Cu})\text{MOF}$ (d).



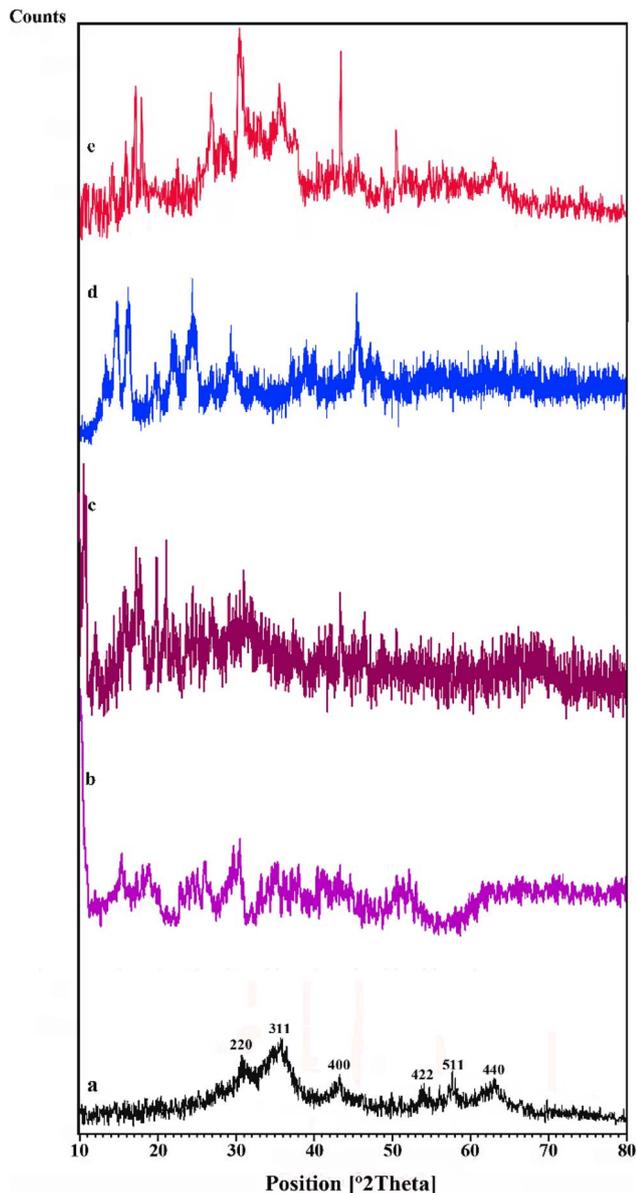


Fig. 3 XRD patterns of $\text{Fe}_3\text{O}_4/\text{pectin}$ (a), Co-MOF (b), (Co/Cu)MOF (c), Cu-MOF (d), $\text{Fe}_3\text{O}_4/\text{pectin}/(\text{Co}/\text{Cu})\text{MOF}$ (e).

C–O. In the spectrum of $\text{Fe}_3\text{O}_4/\text{pectin}$, the absorption peaks at 3401, 2932, 1620, and 584 cm^{-1} are related to the stretching vibrations of O–H, C–H, C=O, and Fe–O bonds, respectively. In the spectrum of (Co–Cu)MOF, the two absorption peaks appeared at 1620 and 1434 cm^{-1} correspond to the asymmetric and symmetric stretching vibrations of the O=C–O bonded to Cu and Co. The peak appeared at 597 cm^{-1} in the spectrum of $\text{Fe}_3\text{O}_4/\text{pectin}/(\text{Co}/\text{Cu})\text{MOF}$ nanocomposite is attributed to Fe–O stretching vibration, and the absorption peaks observed at 1563 and 1400 cm^{-1} correspond to the asymmetric and symmetric stretching of the O=C–O bonded to Co and Cu metals.

Fig. 3 represents the XRD patterns of $\text{Fe}_3\text{O}_4/\text{pectin}$ (a), Co-MOF (b), (Co/Cu)MOF (c), Cu-MOF (d), and $\text{Fe}_3\text{O}_4/\text{pectin}/(\text{Co}/\text{Cu})\text{MOF}$ (e). As can be seen, the XRD profile of $\text{Fe}_3\text{O}_4/\text{pectin}/$

(Co/Cu)MOF reflects the characteristic peaks of standard cubic spinel Fe_3O_4 nanoparticles that appeared at $2\theta = 30.4^\circ$, 35.9° , 43.2° , 54.1° , 57.2° and 63.1° . These correspond to the (220), (311), (400), (422), (511) and (440) diffraction planes (JCPDS No. 19–0629). However, the additional peaks observed at $2\theta = 14.3^\circ$, 15.0° , 17.1° , 18.0° , 22.5° , 25.2° , 27.9° , 28.7° , 35.5° , 47.7° , 48.8° and 50.5° are assigned to Co-MOF and Cu-MOF crystalline phases.

The surface morphology of $\text{Fe}_3\text{O}_4/\text{pectin}/(\text{Co}/\text{Cu})\text{MOF}$ nanocomposite was investigated by scanning electron microscopy (SEM). The SEM images showed that the $\text{Fe}_3\text{O}_4/\text{pectin}/(\text{Co}/\text{Cu})\text{MOF}$ nanocomposite possesses nanoparticles with spherical morphology with uniform shapes and smooth surface (Fig. 4). In addition, the EDS analysis of the prepared catalyst confirms the existence of Co and Cu in the framework. It displays the presence of carbon, oxygen and iron atoms (Fig. 5).

The thermal stability of $\text{Fe}_3\text{O}_4/\text{pectin}/(\text{Co}/\text{Cu})\text{MOF}$ nanocomposite was examined by simultaneous thermal analysis (STA, Fig. 6). The first weight loss that appeared in a temperature range of 210–400 °C can be attributed to the removal of immobilized pectin groups onto the Fe_3O_4 surface. The second weight loss from 420 to 550 °C is related to the degradation of pectin and the decomposition of MOF shells on the material surface. These data successfully prove the good thermal stability of $\text{Fe}_3\text{O}_4/\text{pectin}/(\text{Co}/\text{Cu})\text{MOF}$ at 200 °C, demonstrating its sufficient stability for catalytic applications.

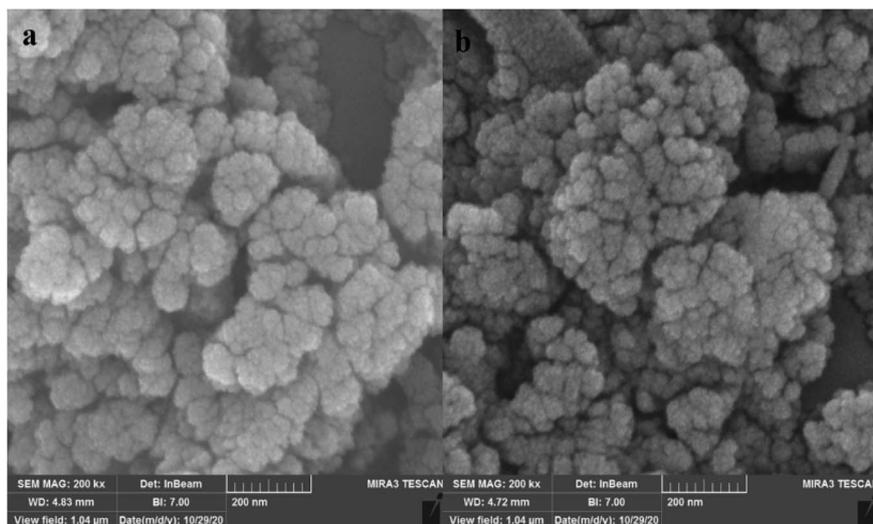
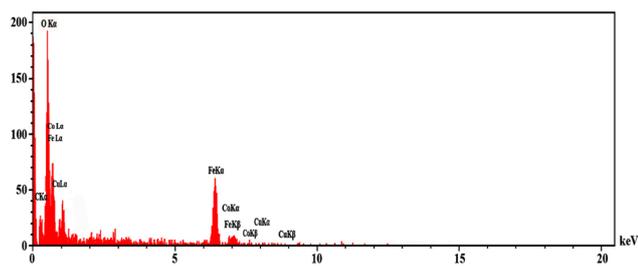
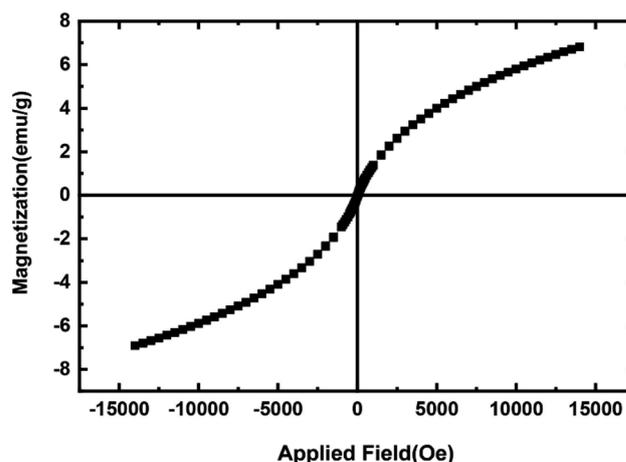
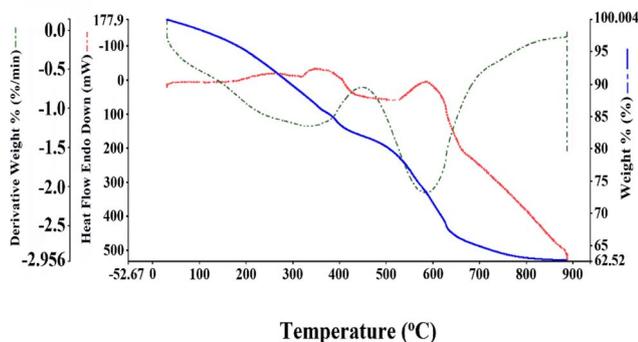
The magnetic property of $\text{Fe}_3\text{O}_4/\text{pectin}/(\text{Co}/\text{Cu})\text{MOF}$ was measured by a vibrating sample magnetometer (VSM). The saturation magnetization of $\text{Fe}_3\text{O}_4/\text{pectin}/(\text{Co}/\text{Cu})\text{MOF}$ nanocomposite is about 7 emu g^{-1} (Fig. 7), which is lower than that of Fe_3O_4 (about 65 emu g^{-1}). This decrease in saturation magnetization confirms the successful coating of (Co/Cu)MOF shell on the surface of the Fe_3O_4 nanoparticles. Also, the magnetic isolating of $\text{Fe}_3\text{O}_4/\text{pectin}/(\text{Co}/\text{Cu})\text{MOF}$ nanocomposite is tested by setting a magnet near the glass bottle of the nanocomposite.

The porosity of $\text{Fe}_3\text{O}_4/\text{pectin}/(\text{Co}/\text{Cu})\text{MOF}$ is examined by measuring N_2 adsorption–desorption isotherms at 77 K. According to this analysis, the surface area, the total pore volume, and the mean diameter of the cavities were $207\text{ m}^2\text{ g}^{-1}$, $0.1756\text{ cm}^3\text{ g}^{-1}$, and 6.591 nm, respectively (Fig. 8).

3.3. Catalytic activity test

The catalytic activity of $\text{Fe}_3\text{O}_4/\text{pectin}/(\text{Co}/\text{Cu})\text{MOF}$ was tested in the reaction of benzaldehyde, ethyl acetoacetate, and urea as a model reaction (Table 1). The effect of catalyst loading was investigated by carrying out the model reaction in the presence of different amounts of $\text{Fe}_3\text{O}_4/\text{pectin}/(\text{Co}/\text{Cu})\text{MOF}$. Upon increasing the amount of catalyst from 10 to 25 mg, a noteworthy increase in the yield of the product was observed (Table 1, entries 2–5), which proves that the progress of the reaction is affected by the catalyst loading. By increasing the catalyst loading to 30 mg, no significant change in the yield of the product is observed (Table 1, entry 6). The best amount of $\text{Fe}_3\text{O}_4/\text{pectin}/(\text{Co}/\text{Cu})\text{MOF}$ is 25 mg which created the desired product with 96% yield (Table 1). The reaction was also affected



Fig. 4 FE-SEM images of $\text{Fe}_3\text{O}_4/\text{pectin}/(\text{Co}/\text{Cu})\text{MOF}$.Fig. 5 EDS spectrum of $\text{Fe}_3\text{O}_4/\text{pectin}/(\text{Co}/\text{Cu})\text{MOF}$.Fig. 7 VSM analysis of $\text{Fe}_3\text{O}_4/\text{pectin}/(\text{Co}/\text{Cu})\text{MOF}$.Fig. 6 STA thermogram of $\text{Fe}_3\text{O}_4/\text{pectin}/(\text{Co}/\text{Cu})\text{MOF}$.

by temperature, and the best result was obtained at 85 °C (Table 1, entries 7–13). Additionally, various solvents such as THF, DMF, H_2O , ethanol, methanol, toluene (Table 1, entries 14–19), and solvent-free conditions were applied. The reaction yield under solvent-free conditions was higher in comparison with solvent conditions, and the reaction time was also shortened. Based on these studies, 25 mg of $\text{Fe}_3\text{O}_4/\text{pectin}/(\text{Co}-\text{Cu})\text{MOF}$ catalyst, 85 °C, and no solvent were chosen as optimum conditions. To further assess $\text{Fe}_3\text{O}_4/\text{pectin}/(\text{Co}-\text{Cu})\text{MOF}$, its components separately [$\text{Fe}_3\text{O}_4/\text{pectin}$, Co-MOF, Cu-MOF and

(Co-Cu)MOF] were used in the model reaction under the same conditions as the $\text{Fe}_3\text{O}_4/\text{pectin}/(\text{Co}-\text{Cu})\text{MOF}$ catalyst (Table 2). The results showed that the product with a lower yield was obtained using each of these components as a catalyst. Under these conditions, the activity of the $\text{Fe}_3\text{O}_4/\text{pectin}/(\text{Co}-\text{Cu})\text{MOF}$ was then investigated in the Biginelli reaction of several aldehyde substrates with ethyl acetoacetate and urea (Table 3). As seen, both electron-withdrawing and electron-releasing substituents on the aldehyde aryl ring were tolerated and reacted with ethyl acetoacetate and urea under optimized conditions. Strikingly, in all cases the corresponding products were isolated in high yields in pure crystal form. The results prove that the type and position of the substituent have no important effect on the activity of the catalyst. These observations confirm the high efficiency of the catalyst for the conversion of an extensive range of aldehyde substrates.

According to the recycling study, no significant reduction in the reactivity of the catalyst was observed. However, to verify the



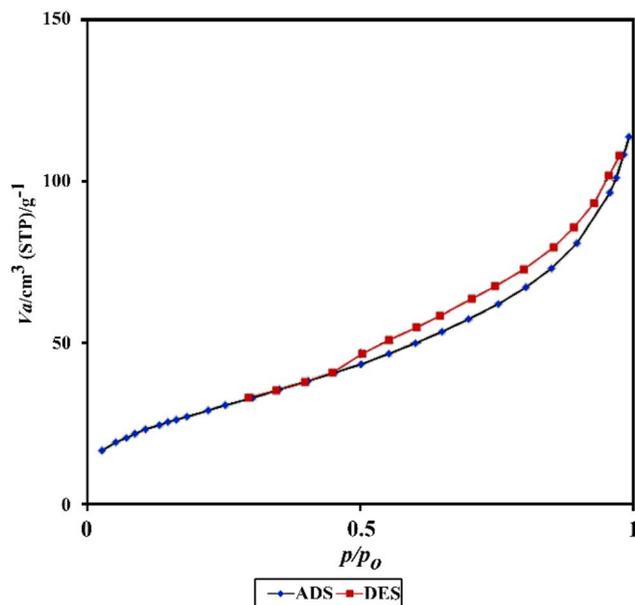


Fig. 8 N_2 adsorption-desorption isotherms of Fe_3O_4 /pectin/(Co/Cu) MOF.

Table 1 Effect of catalyst loading, temperature and solvent in the synthesis of 3,4-dihydropyrimidinone^a

Entry	Catalyst (mg)	Solvent	<i>T</i> (°C)	Time (min)	Yield (%)
1	—	—	85	60	Trace
2	10	—	85	45	72
3	15	—	85	55	76
4	20	—	85	40	84
5	25	—	85	30	96
6	30	—	85	35	95
7	25	—	35	60	21
8	25	—	45	45	41
9	25	—	55	45	53
10	25	—	65	50	67
11	25	—	75	30	89
12	25	—	85	30	96
13	25	—	95	35	91
14	25	THF	85	50	73
15	25	DMF	85	65	62
16	25	H ₂ O	85	55	65
17	25	Ethanol	85	35	89
18	25	Methanol	85	35	66
19	25	Toluene	85	60	68
20	25	Solvent-free	85	30	96

^a Reaction conditions: benzaldehyde (1 mmol), ethyl acetoacetate (1 mmol), urea (1.5 mmol), catalyst (25 mg).

extent of copper species leaching from the material during the reaction process, in the next investigation a hot filtration test was carried out for the reaction of benzaldehyde, urea, and ethyl

Table 2 Preparation of DHPM using Fe_3O_4 /pectin, Co-MOF, Cu-MOF and (Co-Cu)MOF^a

Entry	Catalyst	Time (min)	Yield (%)
1	Fe_3O_4 /pectin	60	62
2	Co-MOF	60	73
3	Cu-MOF	60	61
4	(Co-Cu)MOF	60	82

^a Reaction conditions: aldehyde (1 mmol), ethyl acetoacetate (1 mmol), urea (1.5 mmol), catalyst (25 mg). *T* = 85 °C.

acetoacetate after ~40% of the coupling reaction was completed. The filtrate was then transferred to another flask, and the reaction was continued under the same conditions as before. After 2 h, an additional conversion of only 5% was observed in the coupling reaction. Furthermore, atomic adsorption spectroscopy, performed for the filtrate, demonstrated that the amount of leached cobalt and copper is less than 1 ppm. This observation illustrates that the catalyst may operate in a heterogeneous manner.

To prove the heterogeneity of the catalyst, the filtration experiment of Fe_3O_4 /pectin/(Co-Cu)MOF was performed. The catalyst was separated from the reaction mixture after 15 min,

Table 3 The Biginelli reaction of several aromatic aldehydes, ethyl acetoacetate and urea in the presence of Fe_3O_4 /pectin/(Co/Cu)MOF catalyst^a

Entry	R	Time (min)	Yield (%)	m.p. (°C)
1 (4a)	H	30	96	202–204
2 (4b)	4-NO ₂	20	98	202–203
3 (4c)	4-Cl	30	94	218–219
4 (4d)	2-Cl	35	92	221–223
5 (4e)	4-OMe	20	93	205–207
6 (4f)	2-OMe	20	91	266–268
7 (4g)	4-Me	45	92	217–219
8 (4h)	4-Br	35	93	212–214
9 (4i)	2-OH	25	85	218–220
10 (4j)	4-CH(CH ₃) ₂	45	76	210–212

^a Reaction conditions: aldehyde (1 mmol), ethyl acetoacetate (1 mmol), urea (1.5 mmol), catalyst (25 mg). *T* = 85 °C.



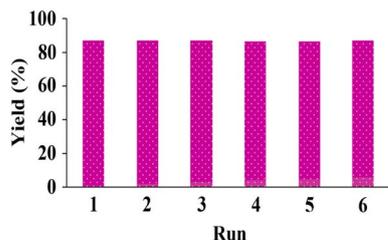


Fig. 9 Reusability of Fe₃O₄/pectin/(Co/Cu)MOF.

and the resulting filtrate was further stirred for 15 min. The conversion yield of the benzaldehyde remains unchanged for the filtrate, even at an extended time, indicating that the catalytic process is heterogeneous, and there is not any progress for the reaction in the homogeneous phase.

3.4. Reusability of Fe₃O₄/pectin/(Co–Cu)MOF

In the next study, the recycling performance of the Fe₃O₄/pectin/(Co–Cu)MOF nanocatalyst was tested in the condensation of benzaldehyde, ethyl acetoacetate, and urea under normal conditions. To this end, after the completion of the reaction, the mixture was filtered and thoroughly washed with hot ethanol. The recovered catalyst was then reused in the next run under the

same conditions as the first run. More experiments were performed than before, and the results illustrated that the catalyst could be recovered and reused at least 5 times without significant lowering activity (Fig. 9). This observation strongly confirms the high recycling efficiency of the catalyst, which is a noteworthy property from economic and environmental points of view. The FT-IR, and SEM analyses of recycled Fe₃O₄/pectin/(Co/Cu)MOF (after the sixth run) were performed and the results were compared with the FT-IR, and SEM image of the fresh catalyst. The absorptions in the spectrum of recycled Fe₃O₄/pectin/(Co/Cu)MOF exhibited that the structure of the recovered catalyst was retained during the recycling process. Moreover, the amorphous nature and morphology of the recovered catalyst are similar to fresh catalyst demonstrating high stability of Fe₃O₄/pectin/(Co/Cu)MOF during the catalytic process (Fig. 10).

3.5. Comparison of the proposed catalyst with the previously reported catalysts for the Biginelli reaction

The comparison between the performance of the Biginelli reaction based on the Fe₃O₄/pectin/(Co–Cu)MOF catalyst and some previously reported catalysts involving the Biginelli reaction is listed in Table 4. It is found that Fe₃O₄/pectin/(Co–Cu)MOF exhibits advantages in terms of cost-effectiveness, simplicity, and short reaction time. Additionally, it has a short

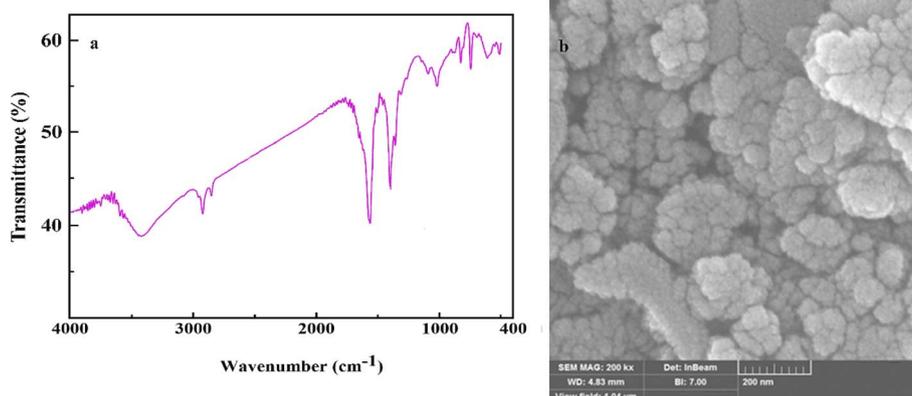
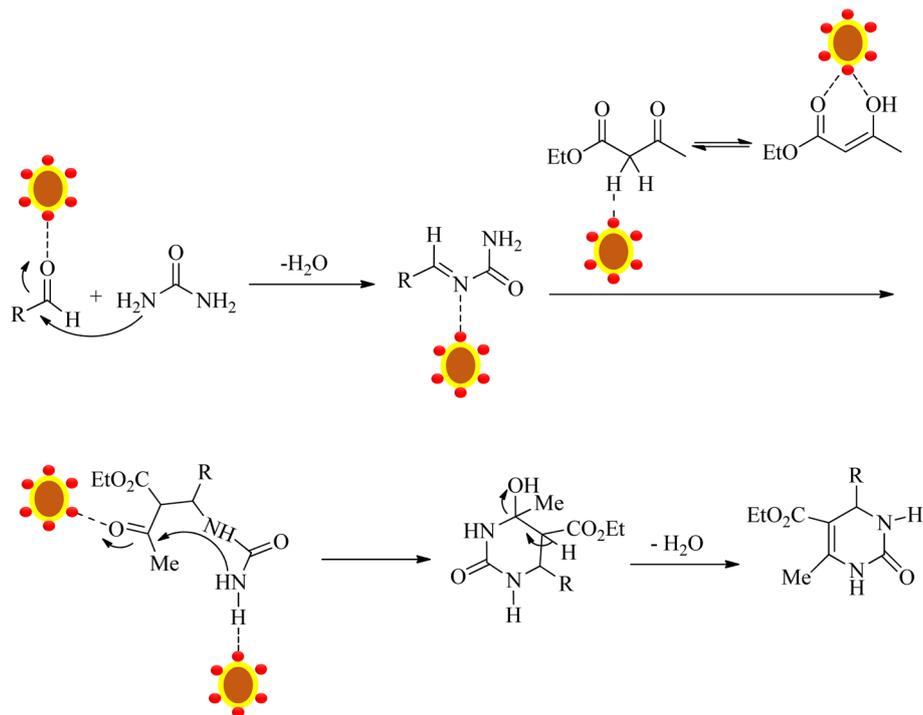


Fig. 10 FT-IR spectrum (a), and SEM image (b) of recycled Fe₃O₄/pectin/(Co–Cu)MOF.

Table 4 Comparison of the proposed catalyst with reported catalysts for the Biginelli reaction

Catalyst	Amount	Time	Solvent	<i>T</i> (°C)	Yield (%)	Ref.
Anionic Zn(II)-MOF	10 wt%	2 h	—	60	93	41
0.2NbCl ₅ -BMImCl	5 mol%	30 min	—	100	90	42
PTA@MIL-101	0.006 mmol	1 h	—	100	90	43
Cu-based MOF	0.1 mmol	2 h	—	60	86	44
Ompg-C ₃ N ₄ /SO ₃ H	30 mg	30 min	EtOH	Reflux	94	45
Ni-DDIA	0.005 mmol	30 min	—	80	84.6	7
BIL-SO ₃ H	2.5 mol%	3 h	EtOH	80	98	46
COF-IM-SO ₃ H	8.64 mg (1.0 mol%)	2.5 h	—	90	98	47
Fe ₃ O ₄ /pectin/(Co/Cu)MOF	25 mg	30 min	—	85	96	This work





Scheme 1 Proposed mechanism for the construction of DHPMs catalyzed by $\text{Fe}_3\text{O}_4/\text{pectin}/(\text{Co}-\text{Cu})\text{MOF}$.

reaction time and mild conditions in the Biginelli reaction compared with the literature.

In the proposed catalytic mechanism, *N*-acylimine intermediate was formed from the reaction of aldehyde and urea, which aldehyde activated by Lewis acidic $[\text{Co}(\text{II})$ and $\text{Cu}(\text{II})]$ and Brønsted acidic sites (the uncoordinated $-\text{COOH}$ groups on the pectin surface). Consequently, the reaction of *N*-acylimine coordinated with Lewis acidic sites was carried out with enol of ethyl acetoacetate to provide ureide. The cyclization of ureide in the presence of catalyst gave the heterocyclic intermediate, which on dehydration affords the corresponding 3,4-dihydropyrimidin-2(1H)-one (DHPM) products (Scheme 1). Here $\text{Fe}_3\text{O}_4/\text{pectin}/(\text{Co}-\text{Cu})\text{MOF}$ has a large number of dispersed Brønsted acidic $-\text{COOH}$ groups and Lewis acidic $\text{Co}(\text{II})$ and $\text{Cu}(\text{II})$ centers, which gives excellent catalytic performance to the reaction.

4. Conclusion

Bimetallic Co/Cu -MOF grown on $\text{Fe}_3\text{O}_4/\text{pectin}$ was prepared and applied as a recoverable nanocatalyst for the one-pot synthesis of DHPM derivatives by the three-component condensation of diverse aldehydes, ethyl acetoacetate, and urea under solvent-free conditions. The catalyst was easily recovered and reused without a significant decrease in activity. The advantages of this catalyst system include short reaction time, low loading of catalyst, solvent-free conditions, and easy catalyst separation.

Data availability

All data generated or analyzed during this study are included in this published article.

Conflicts of interest

There are no conflicts to declare.

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