


 Cite this: *RSC Adv.*, 2023, 13, 10367

Facile synthesis of pyrazolopyridine pharmaceuticals under mild conditions using an algin-functionalized silica-based magnetic nanocatalyst (Alg@SBA-15/Fe₃O₄)[†]

 Fereshte Hassanzadeh-Afruzi,^{ID} Zeinab Amiri-Khamakani, Mahdi Saeidirad, Mohammad Mehdi Salehi, Reza Taheri-Ledari^{ID} and Ali Maleki^{ID*}

Pyrazolopyridines are common scaffolds in various bioactive compounds, which have several therapeutic effects and unique pharmacological properties. In this study, we fabricated a novel environmentally friendly silica-based nanocomposite as a multifunctional catalytic system for the synthesis of pyrazolopyridine derivatives. This novel heterogeneous nanocomposite named Alg@SBA-15/Fe₃O₄ (Alg stands for alginic acid), was prepared in several steps. In this regard, SBA-15 was synthesized by the hydrothermal method. Next, it was magnetized by Fe₃O₄ nanoparticles *via* an *in situ* co-precipitation process. Then, SBA-15/Fe₃O₄ particles were functionalized with 3-minopropyltriethoxysilane (APTES). Afterward, Alg@SBA-15/Fe₃O₄ was obtained by a nucleophilic substitution reaction between SBA-15/Fe₃O₄-NH₂ and an as-synthesized methyl-esterified alginic. Different analyses such as Fourier-transform infrared (FTIR), energy-dispersive X-ray (EDX) spectroscopy, field-emission scanning-electron microscopy (FESEM), vibrating-sample magnetometer (VSM), X-ray diffraction (XRD), thermogravimetric analysis (TGA), and BET (Brunauer–Emmett–Teller) have been used to confirm the structure of the fabricated catalyst. The magnetic properties of the Alg@SBA-15/Fe₃O₄ catalytic system imparted by Fe₃O₄ MNPs enable it to be conveniently isolated from the reaction mixture by using an external magnet. According to the obtained results, the prepared nanocatalyst has high thermal stability and it lost approximately 26% of its weight up to 800 °C. Interestingly, a small amount of prepared nanocatalyst (0.02 g) has shown excellent catalytic performance in the synthesis of pyrazolopyridine derivatives (90–97%) in a short reaction time (20–30 min) at room temperature which can be attributed to its porous structure and large surface area, and the presence of many acidic and basic functional groups. In general, it can be argued that the Alg@SBA-15/Fe₃O₄ nanocomposite deserves more attention due to its non-toxicity, ease of preparation, good recyclability, and its high catalytic efficiency.

Received 14th November 2022

Accepted 20th March 2023

DOI: 10.1039/d2ra07228a

rsc.li/rsc-advances

1. Introduction

It has been stated that nitrogen-containing heterocycles account for more than 90% of substances examined by pharmaceutical companies.¹ Pyrazolopyridine as a kind of nitrogen-rich heterocycle has attracted a lot of interest due to having valuable pharmacological and biological activities, including anti-virus² and anti-leishmania,³ as well as anti-malarial⁴ properties. Tracazolate, cartazolate, and etazolate are some of pyrazolopyridine derivatives that are used as anxiolytic drugs. Other pyrazolopyridine-containing biologically-active compounds involve a GSK-3 inhibitor, and BAY 41-2272 can

be utilized as cardiovascular therapeutic agents.⁵ Although various multi-step procedures for the construction of pyrazolopyridines have been developed in recent years, some of them suffer from one or more challenges for instance long reaction time, low yields, laborious work-up procedure, as well as generation of by-products. It seems that the development of efficient methods for the synthesis of pyrazolopyridines is still an important topic in chemistry because of their extensive biological activities. Multicomponent reactions (MCRs) is a practical approach to addressing the challenges of the pyrazolopyridine synthesis. MCRs are one-pot processes in which three or more reactants are reacted to form a particular product containing all of the reactants' atoms.^{6,7} These reactions have been broadly utilized for the synthesis of complicated bio-active organic compounds, particularly heterocyclic materials for instance the pyrazolopyridine, owing to their outstanding advantages such as atom economy, experimental simplicity,

Catalysts and Organic Synthesis Research Laboratory, Department of Chemistry, Iran University of Science and Technology, Tehran 16846-13114, Iran. E-mail: maleki@iust.ac.ir; Fax: +98-21-73021584; Tel: +98-21-73228313

[†] Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d2ra07228a>



and the formation of several bonds in a single operation.⁸ The use of magnetic nanocatalysts in MCRs is an attractive idea for the development of sustainable methods in green synthetic chemistry.^{9–11}

Porous materials with outstanding properties such as, uniform pore size, adjustable porous structure, and high surface area have piqued the curiosity of scientists.¹² According to the IUPAC classification, porous materials can be categorized as microporous materials, whose pore width is less than 2 nm, mesoporous materials, whose pore width is 2–50 nm, and macroporous materials, whose pore width is more than 50 nm.¹³ Mesoporous materials were first discovered by Mobil Research and Development Corporation in 1992, and their synthesis and application have received great attention since then.¹⁴ Zhao and coworkers at the University of California, Santa Barbara reported the SBA-15 mesoporous silica for the first time in 1998.^{13,15–17} Well-defined pore size between 4 to 30 nm, thick pore walls, pore architecture,^{11,18} wide surface area (up to 1000 m² g⁻¹), high thermal and hydrothermal stability, and excellent capability to be modified/functionalized are among the features that make materials based on SBA-15 have different applications, including adsorption, drug delivery, imaging, sensor, and catalysis.^{13,19–22} One-pot synthesis or co-condensation, and grafting technique or post-synthesis are the two main methods used to modify/functionalize silica mesoporous materials.¹³ Incorporation of Fe₃O₄ magnetic nanoparticles as a superparamagnetic substance into silica-based mesoporous supports is a very practical step towards green chemistry, as it can lead to the easy separation of these materials from the reaction medium.^{23–27} Chemical modification of SBA-15 can lead to the formation of versatile highly-efficient porous catalysts by additional active sites for interaction with the reactants.²⁸ A catalytic application on SBA-15-based materials have been extensively studied in the literature.^{29,30} Safaei Ghomi *et al.* reported several of supported SBA-15 catalysts. Generally, in these reported works, the SBA-15 mesoporous silica was employed as a support metal species, and a highly dispersed metal catalyst was utilized in different multicomponent reactions,^{31–33} for example, they prepared functionalized-SBA-15 using a tetradentate ligand to construct a new catalyst. They investigated the efficiency of this catalyst in the synthesis of 2-azapyrrolizidine alkaloids, which have different biological properties.³⁴ In 2019, by grafting the nickel Schiff base complex on the SBA-15 surface, Noroozi *et al.* prepared a nano-catalyst called Ni(II)-Schiff base/SBA-15. It was an eco-friendly, and effective catalyst for the synthesis of 2-amino-4-aryl-5-oxo-4H,5H-pyrano[3,2-*c*]chromene-3-carbonitriles derivatives through a tandem reaction.³⁵ In another example, in 2021, research on the synthesis of dihydropyranopyrazole compounds by using a guanidine functionalized SBA-15/Fe₃O₄ mesoporous nanocomposite was performed. In this research, synthesis efficiency of dihydropyranopyrazole derivatives by employing a guanidine functionalized SBA-15/Fe₃O₄, SBA-15/Fe₃O₄, and Fe₃O₄ reported 95%, 71%, and negligible, respectively, which is a good indication of the ability of SBA-15 in catalytic applications.¹³

Alginic acid, also called algin, is generally extracted from cell-wall of brown seaweeds such as macroalgae, ascophyllum,

and kelp has attracted so much attention due to its natural availability and non-toxicity.^{36,37} Since this natural linear polysaccharide is obtained from natural resources, it can be utilized for construction of green catalysis of organic reactions.³⁸ Brilliant properties such as chemical versatility, renewability, hydrophilicity, cost-effectiveness, and biodegradability make alginic acid as an environmentally friendly organocatalyst.^{39,40} However, the use of this natural polymer as a catalyst may be associated with weaknesses such as low chemical and physical stability, relatively low chemical variability, and undesirable catalytic efficiency that limit its catalytic application. Modification of alginic acid and adding it to heterogeneous systems seems to be a suitable solution to overcome these limitations and obtain a heterogeneous catalyst with high catalytic capacity.³⁸

Herein, we introduce a new multifunctional catalytic system for the synthesis of pyrazolopyridine derivatives which mainly composed of SBA-15 as a mesoporous component, iron oxide MNPs as magnetic part, and modified alginic acid as natural polymer. In the fabrication of this novel mesoporous nanocomposite, formulated as “Alg@SBA-15/Fe₃O₄”, first SBA-15 was magnetized with Fe₃O₄ MNPs, then magnetized SBA-15 was modified with APTES to obtain SBA-15/Fe₃O₄-NH₂. An important step in the preparation of the final composite is the conversion of alginic acid to methyl-esterified alginic acid *via* chemical esterification reaction. After the preparation of methyl-esterified alginic acid, in the final stage, the SBA-15/Fe₃O₄-NH₂ formed a covalent bond with methyl-esterified alginic acid through nucleophilic substitution reaction. This inorganic–organic mesoporous nanocomposite has exhibited unique physicochemical characteristics which originate from the combination of its components. The outstanding characteristic properties of the fabricated heterogeneous catalytic system, for instance its porosity, average size, and thermal stability were carefully examined utilizing various analyses. Afterward, the ability of this catalyst in the organic synthesis was carefully investigated. In summary, it was found that using the Alg@SBA-15/Fe₃O₄ catalytic system at room temperature can lead to the synthesis of various high-yield pyrazolopyridine derivatives in short times. Also, the catalytic reusability of Alg@SBA-15/Fe₃O₄ was investigated for five successive runs, which showed that less than 10% of its catalytic capacity decreases after consecutive times compared to the first time. The easy separation of Alg@SBA-15/Fe₃O₄ catalytic system from the reaction medium stemmed from its magnetic property, is an effective step toward green chemistry. Overall, because of its cost-effectiveness and efficiency, the presented nanocatalyst is suggested for large-scale synthesis.

2. Results and discussion

2.1. Preparation of Alg@SBA-15/Fe₃O₄ nanocomposites

To prepare Alg@SBA-15/Fe₃O₄ nanocomposite, several steps were performed which are briefly described in this section. After preparing SBA-15, the *in situ* co-precipitation method was used to magnetize the SBA-15, resulting in SBA-15/Fe₃O₄. Then, the modification of the magnetic SBA-15 surface was performed



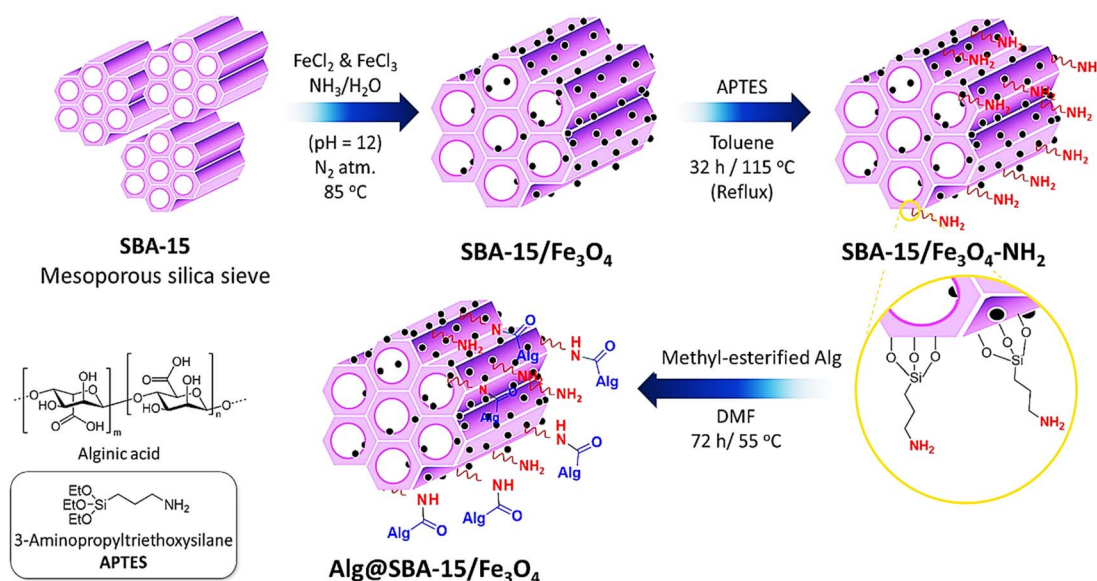


Fig. 1 Schematic preparation route of Alg@SBA-15/Fe₃O₄ catalytic system.

with APTES. The reason for using APTES in this study was that the surface of SBA-15 obtained the appropriate functional groups to bind to methyl-esterified alginic acid. Eventually, to prepare the Alg@SBA-15/Fe₃O₄ nanocomposites, the methyl-esterified alginic acid was attached to SBA-15/Fe₃O₄-NH₂ through an amide bond (Fig. 1).

2.2. Characterization of Alg@SBA-15/Fe₃O₄ nanocomposites

2.2.1. FTIR spectroscopy. To investigate the existence of functional groups in prepared samples during the preparation of the final nanocomposite, FTIR analysis was utilized. The silica network has characteristic absorption bands at about 463 cm⁻¹ (rocking bending vibration of Si-O-Si), about 804-960 cm⁻¹ (stretching vibration of Si-OH), about 1084 cm⁻¹ (symmetric stretching vibration of Si-O-Si), and about 3445 cm⁻¹ (stretching vibration of OH groups) (Fig. 2a).^{26,41,42} Almost in the same areas of the SBA-15 peaks, there are peaks in the FTIR spectrum of SBA-15/Fe₃O₄, SBA-15/Fe₃O₄-NH₂, and Alg@SBA-15/Fe₃O₄ (Fig. 2b, c and f), it can be a step towards proving the claim that these compounds contain SBA-15 in their structure. In the spectrum of SBA-15/Fe₃O₄ and Alg@SBA-15/Fe₃O₄, an absorption appeared at about 577 cm⁻¹ associated to Fe-O vibration (Fig. 2b and f).^{43,44} After the prepared SBA-15/Fe₃O₄ was functionalized with APTES, two peaks emerged in its FTIR curve. One of them appeared in 1558 cm⁻¹ related to the bending vibration of NH₂, and the other one appeared in 2926 cm⁻¹, which is related to the stretching vibrations of the C-H bond, which originate from the propyl chain of APTES (Fig. 2c).¹³ As can be seen in the FTIR curve of alginic acid (Fig. 2d), the peak of the carbonyl of the carboxylic acid functional group of alginic acid has observed at 1624 cm⁻¹, while the carbonyl peak of the ester of methyl-esterified alginic acid appeared at 1720 cm⁻¹ (Fig. 2e),⁴⁵ this can confirm successful chemical esterification on alginic acid. The FTIR curve of the

Alg@SBA-15/Fe₃O₄ (Fig. 2f) shows other peaks, the most important of them is the C=O peak at about 1624 cm⁻¹, which probably belongs to the amide functional group. This peak is important in that it can be a reason for the creation of amide bonds between SBA-15/Fe₃O₄-NH₂ and methyl-esterified alginic acid.

2.2.2. XRD analysis. To characterize the crystalline structure of the samples, X-ray diffraction (XRD) analysis was carried out. The diffractograms of prepared Fe₃O₄ (Fig. 3b) shows characteristic diffraction peaks at $2\theta = 30.43^\circ, 35.86^\circ, 43.47^\circ, 57.38^\circ, 63.19^\circ$ were indexed to the (2 2 0), (3 1 1), (4 0 0), (3 3 3), (4 4 0) miller indices of cubic Fe₃O₄ with Card No. JCPDS 01-075-1609.⁴⁶ According to Fig. 3a, the diffractograms of SBA-15 shows a broad distinctive peak in the range of $2\theta: 20^\circ-30^\circ$ attributed to the amorphous SiO₂ character.^{47,48} The XRD pattern of SBA-15/Fe₃O₄ shows peaks at $2\theta = 30.35^\circ(2\ 2\ 0), 35.98^\circ(3\ 1\ 1), 43.56^\circ(4\ 0\ 0), 57.70^\circ(3\ 3\ 3)$ (Fig. 3b) that are also present in the diffractograms of the prepared Fe₃O₄ nanoparticles. This can be a reason to claim that SBA-15 has been magnetized with Fe₃O₄ nanoparticles. Similar to the XRD pattern of SBA-15, the XRD patterns of SBA-15/Fe₃O₄ and Alg@SBA-15/Fe₃O₄ (Fig. 3d) show a relatively broad peak at $2\theta: 20^\circ-30^\circ$, with less intensity in compared to unmodified SBA-15. It is worth mentioning that the lower intensity of the broad peak at $2\theta: 20^\circ-30^\circ$ in Alg@SBA-15/Fe₃O₄ than its counterpart in SBA-15 may be due to the fact that SBA-15 was modified three times. Moreover, characteristic peaks of Fe₃O₄ MNPs were also observed at $2\theta: 30.38^\circ, 35.74^\circ, 43.40^\circ, 58^\circ, 63.22^\circ$. It can be concluded that fabrication of Fe₃O₄ nanoparticles on SBA-15 substrate even with modifications has improved its crystallinity.

2.2.3. TGA analysis. The thermogravimetric analysis (TGA) curve of the SBA-15/Fe₃O₄ shows a steady weight loss in the range of 50 °C–800 °C with a mild slope (Fig. 4). The remarkable thermal stability of SBA-15/Fe₃O₄ is proven by the fact that only about 9% of its weight percentage is reduced up to 800 °C. This



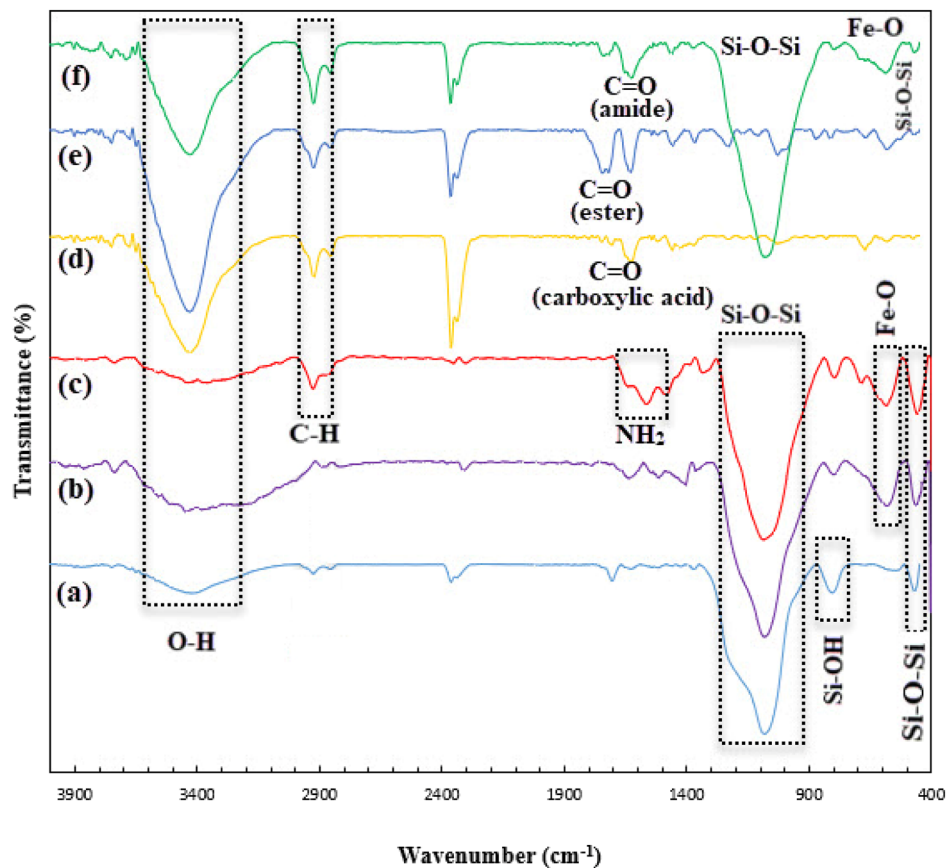


Fig. 2 FTIR spectra of (a) SBA-15, (b) SBA-15/ Fe_3O_4 , (c) SBA-15/ Fe_3O_4 -NH₂, (d) alginic acid, (e) methyl-esterified alginic acid, and (f) Alg@SBA-15/ Fe_3O_4 .

reduction is attributed to of vaporization of adsorbed water and moisture in the sample's cavities, as well as dehydrogenation or dehydroxylation of its surface.¹³ According to the TGA curve of Alg@SBA-15/ Fe_3O_4 , which is shown in Fig. 4, with increasing

the sample temperature up to 800 °C, the mass percentage of Alg@SBA-15/ Fe_3O_4 decreases with a gentle slope. More precisely, as the temperature increases from 50 °C to 160 °C, about 3% of the weight percentage of Alg@SBA-15/ Fe_3O_4

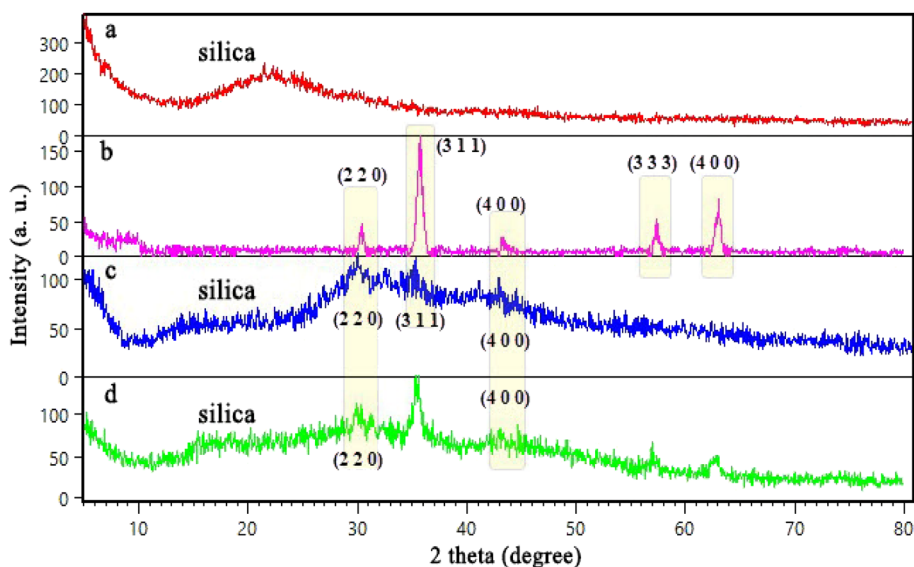


Fig. 3 XRD patterns of (a) SBA-15, (b) Fe_3O_4 MNPs, (c) SBA-15/ Fe_3O_4 , (d) Alg@SBA-15/ Fe_3O_4 .



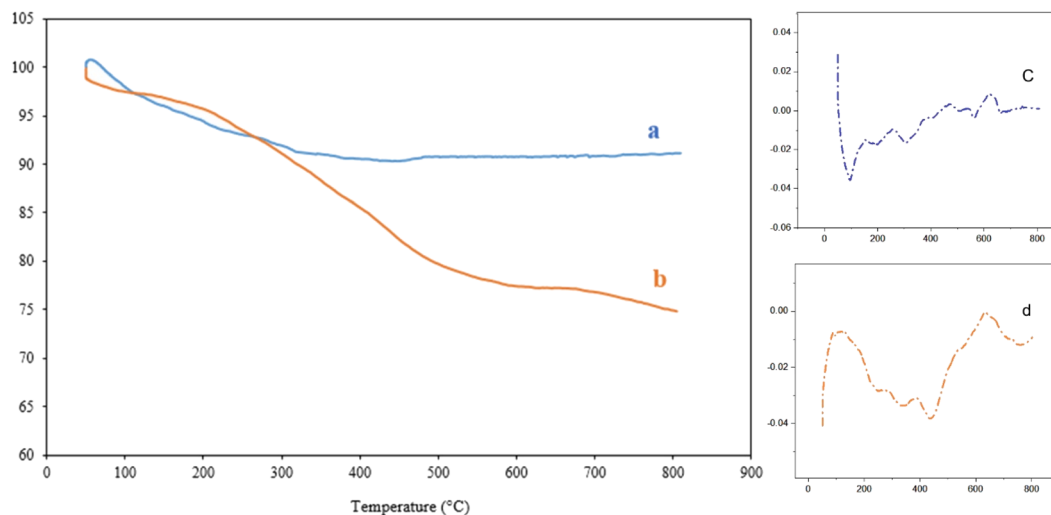


Fig. 4 TGA/DTG curves of (a and c) SBA-15/Fe₃O₄, and (b and d) Alg@SBA-15/Fe₃O₄ nanocomposite systems.

decreases. This reduction is due to the vaporization of water and moisture absorbed inside the cavities of the nanocomposite and on its surface. As the temperature increases to 550 °C, the weight percentage of the sample decreases by about 18.5% in comparison with the weight percentage of the sample at 160 °C. This reduction may be because of the separation and thermal decomposition of the organic constituents of Alg@SBA-15/Fe₃O₄ (alkyl chain and alginate) attached to SBA-15 with covalent bonding. By increasing the temperature to 800 °C, the weight percentage of Alg@SBA-15/Fe₃O₄ decreases by more than 3% compared to the mass percentage of the same substance at 550 °C. This reduction might be related to the condensation of the silane groups of Alg@SBA-15/Fe₃O₄. After heating the nanocomposite to a temperature of 800 °C, its remaining weight was about 74.8%.¹³

2.2.4. VSM analysis. Vibrating-sample magnetometer (VSM) was employed as a common technique for assessing magnetic properties of the materials at ambient temperature. As is observed in the magnetization curve of the Alg@SBA-15/Fe₃O₄, the absence of hysteresis loop, coercivity (H_c) and remanent magnetization (M_r) at room temperature indicate that the fabricated nanocomposite exhibited superparamagnetic (SPM) behavior. The magnetization saturation (M_s) of Alg@SBA-15/Fe₃O₄ is about 10 emu g⁻¹ is less than the M_s value of SBA-15/Fe₃O₄ (Fig. 5), which seems to be due to the fact that a large part of Alg@SBA-15/Fe₃O₄ was composed of non-magnetic components.¹³ One of the significant properties of the fabricated catalyst in this study is its paramagnetic property. This feature allows Alg@SBA-15/Fe₃O₄ to be recycled magnetically and reused while maintaining performance.

2.2.5. Electron microscopy and EDX. FESEM was employed to study morphologies and structure of materials and prepared samples. Fig. 6a–e showed the FESEM images of the Alg, SBA-15, SBA-15/Fe₃O₄ and SBA-15/Fe₃O₄-NH₂, and Alg@SBA-15/Fe₃O₄ in same scales (1 μm). Fig. 6a shows the morphology of alginate, and cylindrical shape of SBA-15 obviously can be seen in Fig. 6b. In Fig. 6c which belongs to SBA-15/Fe₃O₄, Fe₃O₄ NPs

with a spherical shape are found on the cylindrical mesopore of the SBA-15. By further functionalization of SBA-15/Fe₃O₄ with APTES, the mesoporous structure of SBA-15 support with cylindrical shape has been retained. But in preparing the final composite structure, *i.e.* Alg@SBA-15/Fe₃O₄, SBA-15 underwent changes that resulted in changes in the morphology of SBA-15 (Fig. 6e). The chemical composition of the prepared samples were determined using EDX spectroscopy (Fig. 6). The EDX spectrum of each sample was presented next to its FESEM image. In the EDX spectrum of alginate, the C, N, O and Na peaks were detected (Fig. 6a). As is shown in the EDX spectrum of SBA-15, O and Si are the elemental constituent of SBA-15. The presence of O, Si and Fe elements in the EDX spectrum of SBA-15/Fe₃O₄ confirmed its chemical composition, observing two Fe peaks are connected to the existence of Fe₃O₄ MNPs in the SBA-15/Fe₃O₄ composition (Fig. 6b and c). The emerging the N and C peaks in the SBA-15/Fe₃O₄-NH₂'s EDX spectrum, along with O, Si, and Fe peaks verified the successful modification of the SBA-15/Fe₃O₄ by the N-contained organosilane reagent, *i.e.*, APTES

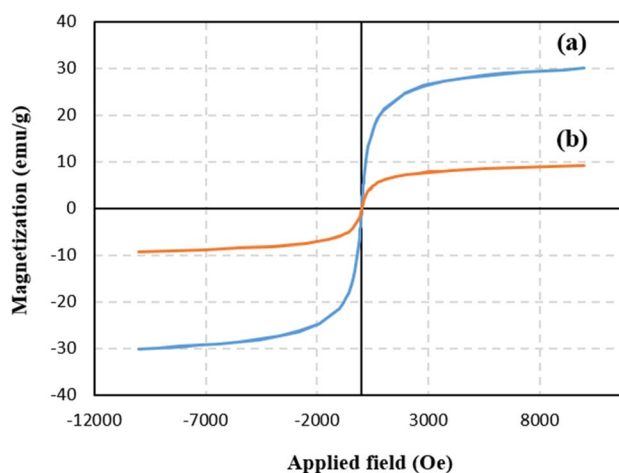


Fig. 5 The magnetization curves of (a) SBA-15/Fe₃O₄, (b) Alg@SBA-15/Fe₃O₄.

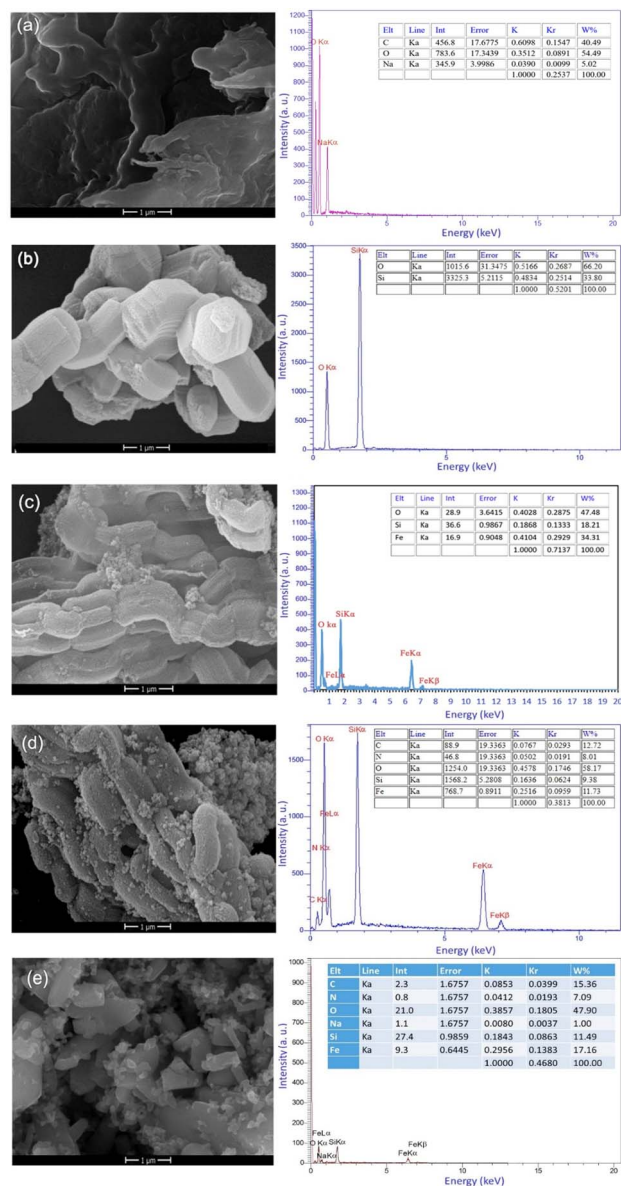


Fig. 6 FESEM images and EDX spectra and tabulated data of (a) alginate, (b) SBA-15, (c) SBA-15/Fe₃O₄, (d) SBA-15/Fe₃O₄-NH₂, and (e) Alg@SBA-15/Fe₃O₄.

(Si-(CH₂)₃-NH₂) (Fig. 6d). Finally, the EDX result of the final nanocomposite, the Alg@SBA-15/Fe₃O₄, proves the presence of all predicted elements including O, Si, Fe, C, N, and Na (Fig. 6e). The elemental composition of O, Si, Fe, C, N, Na is 47.90, 11.49, 17.16, 15.36, 7.09, and 1.00%, respectively.

2.2.6. The N₂ adsorption-desorption isotherm. The N₂ adsorption-desorption isotherms of the SBA-15, SBA-15/Fe₃O₄, and Alg@SBA-15/Fe₃O₄ are shown in Fig. 7a. The type-IV curves that are typical of mesoporous compounds are seen in all three samples. The results of the surface area determined by the BET technique, the pore volume, and pore size (width) calculated by the BJH (Barrett-Joyner-Halenda) method are presented in Table 1. The surface area of the neat SBA-15 (687 m² g⁻¹) is more than the surface area of its magnetized compound, *i.e.*, SBA-15/

Fe₃O₄ (384 m² g⁻¹) and Alg@SBA-15/Fe₃O₄ (about 113.4 m² g⁻¹). This result can be attributed to the blockage of a part of SBA-15 pores by Fe₃O₄ NPs incorporating on the pores of SBA-15 and its consequent modifications, which led to a reduction in accessible surface area for gas adsorption.^{13,49} In Table 1, the BJH pore size distributions of the prepared sample and compounds are shown. It can be seen that the SBA-15/Fe₃O₄ sample has a slightly larger pore size (diameter) and also has relatively broader pore size range than its precursor matrix.

2.3. Application of Alg@SBA-15/Fe₃O₄ composite in the synthesis of pyrazolopyridine derivatives

2.3.1. Optimization of the catalyzed reaction conditions in the synthesis of pyrazolopyridine. To obtain the best catalytic performance of Alg@SBA-15/Fe₃O₄ nanocomposite for the synthesis of pyrazolopyridine derivatives, various experimental conditions were investigated¹⁵ in the four components reaction of ethyl acetoacetate (2.0 mmol), hydrazine hydrate (2.0 mmol), 3-nitrobenzaldehyde (1.0 mmol) and ammonium acetate (1.0 mmol), the details of which are in Table 2. Firstly, the model multicomponent reaction was performed without Alg@SBA-15/Fe₃O₄ catalyst and solvent in two various temperatures, in which the product yield was inconsiderable (Table 2, entries 1 and 2). When the Alg@SBA-15/Fe₃O₄ nanocatalyst was added in the solvent free conditions, the reaction efficiency of the model reached approximately 65% (Table 2, entry 3). The further step, which can be seen in Table 2, entry 4, to examine the effect of the solvent, water was used as the solvent with 0.02 g of the Alg@SBA-15/Fe₃O₄ catalytic system at room temperature. In this case, the yield percentage of the reaction increased compared to the condition investigated in Table 2, entry 3. Among the obtained yields, the best was 97% obtained in EtOH media at room temperature (Table 2, entry 5). Even though the 97% efficiency was also obtained in Table 2, entry 9, but we prefer Table 2, entry 5 because less catalyst was used. Besides, the reaction condition and solvent, different amounts of Alg@SBA-15/Fe₃O₄ were investigated shown in Table 2, entries 7 to 9. The yield of the model multicomponent reaction was also examined under other different conditions which are shown in Table 2, entries 10 and 11. Also, the catalytic ability of a mixture of Alg and SBA-15/Fe₃O₄ and a mixture of methyl-esterified Alg and SBA-15/Fe₃O₄ were investigated, which showed yields of 91% and 90%, respectively (Table 2, entries 13 and 14). Although these efficiencies are high and close to the catalytic efficiency shown by the Alg@SBA-15/Fe₃O₄ catalytic system (Table 2, entry 5), an important issue in the performance of catalysts is that catalytic systems must be stable in the reaction medium. The Alg@SBA-15/Fe₃O₄ catalytic system, in addition to having high catalytic efficiency, has significant stability in the reaction medium.

2.3.2. Catalyzed synthesis of the pyrazolopyridine derivatives. To assess the catalytic efficacy of the fabricated Alg@SBA-15/Fe₃O₄ composite, the organic synthesis of pyrazolopyridine derivatives was investigated (Scheme 1), and the results are summarized in Table 3. Interestingly, a wide range of high-yield pyrazolopyridine derivatives were obtained with using the Alg@SBA-15/Fe₃O₄ catalyst in 20 to 30 minutes.



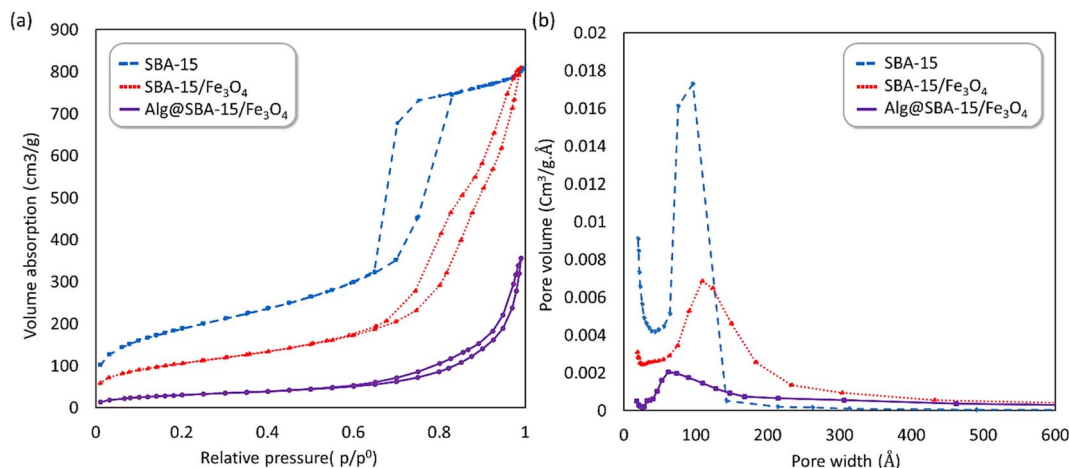


Fig. 7 (a) N_2 adsorption–desorption isotherms and (b) the pore size distribution curve of SBA-15 (blue), SBA-15/ Fe_3O_4 (red) and Alg@SBA-15/ Fe_3O_4 (purple).

Table 1 BET Surface area, pore volume, and pore size (diameter) of SBA-15, SBA-15/ Fe_3O_4 and Alg@SBA-15/ Fe_3O_4

Sample	Surface area ^a ($m^2 g^{-1}$)	Pore volume ^b ($cm^3 g^{-1}$)	Pore size ^b (nm)
SBA-15	686.90	1.28	7.35
SBA-15/ Fe_3O_4	383.91	1.26	12.64
Alg@SBA-15/ Fe_3O_4	113.14	0.55	17.7

^a The surface area obtained by BET technique. ^b Pore volume and pore diameter measured by BJH technique.

2.3.3. Suggested mechanism. The most likely assisted mechanism to synthesis of the different pyrazolopyridine derivatives with Alg@SBA-15/ Fe_3O_4 catalyst is a four-steps is shown in Fig. 8. The Alg@SBA-15/ Fe_3O_4 mesoporous nanocomposite by having abundant hydroxyl groups, amides and to some extent carboxylates, Lewis acid site (Fe^{3+} in Fe_3O_4), and also because of its physical properties such as high porous structure as well as great surface area, can play a critical role in activation of reactants in all steps of this multicomponent reaction as demonstrated in Fig. 8.¹³ Initially, Alg@SBA-15/ Fe_3O_4 activates the carbonyl ethyl acetoacetate group by forming a hydrogen bond and also by interacting with Lewis acid

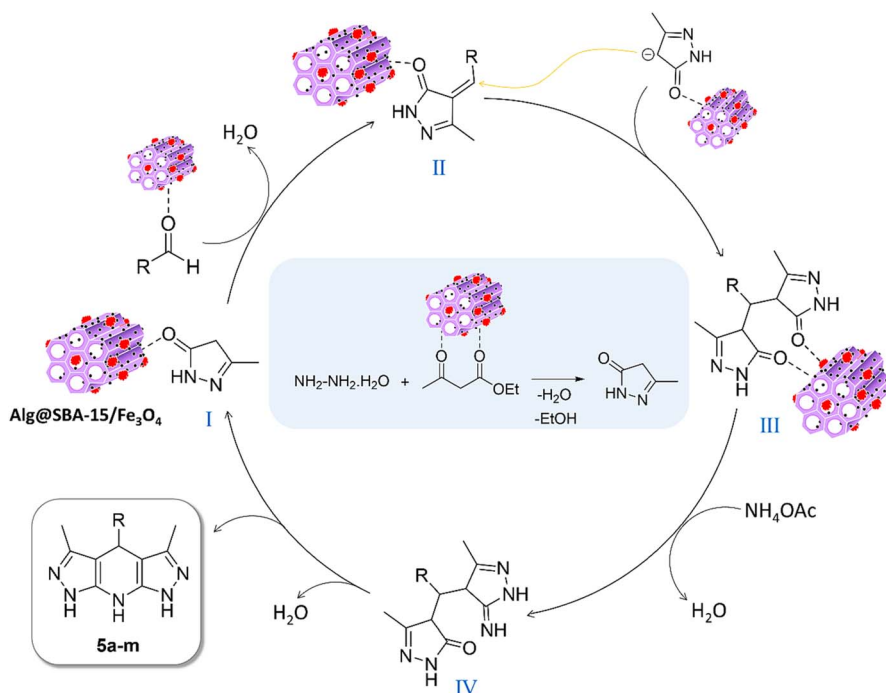


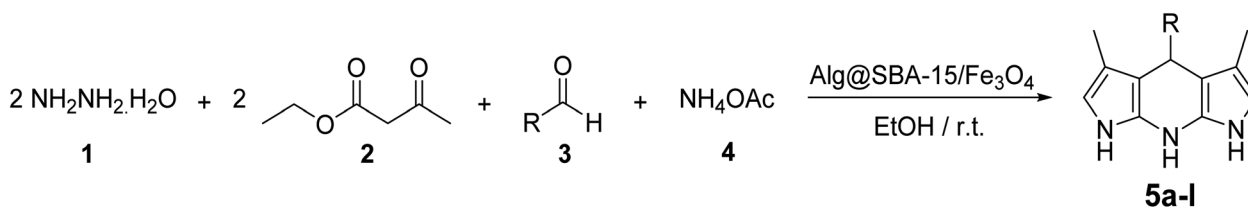
Fig. 8 A plausible mechanism for catalyzing the synthesis reaction of pyrazolopyridine derivatives by Alg@SBA-15/ Fe_3O_4 .



Table 2 Optimizing the multicomponent reaction conditions in the synthesis of pyrazolopyridine derivatives

Entry	Catalyst	Catalyst loading (g)	Solvent	Temp. (°C)	Yield ^a (%)
1	—	—	—	r.t.	Trace
2	—	—	—	80	Trace
3	Alg@SBA-15/Fe ₃ O ₄	0.02	—	r.t.	<65
4	Alg@SBA-15/Fe ₃ O ₄	0.02	H ₂ O	r.t.	78
5	Alg@SBA-15/Fe ₃ O ₄	0.02	EtOH	r.t.	97
6	Alg@SBA-15/Fe ₃ O ₄	0.02	EtOH/H ₂ O (1 : 1)	r.t.	89
7	Alg@SBA-15/Fe ₃ O ₄	0.015	EtOH	r.t.	94
8	Alg@SBA-15/Fe ₃ O ₄	0.010	EtOH	r.t.	92
9	Alg@SBA-15/Fe ₃ O ₄	0.025	EtOH	r.t.	97
10	Alg	0.020	EtOH	r.t.	57
11	SBA-15/Fe ₃ O ₄	0.020	EtOH	r.t.	58
12	Methyl-esterified Alg	0.020	EtOH	r.t.	48
13	Mixture of Alg and SBA-15/Fe ₃ O ₄	0.020	EtOH	r.t.	91
14	Mixture of methyl-esterified Alg and SBA-15/Fe ₃ O ₄	0.020	EtOH	r.t.	90

^a Multicomponent reaction conditions: ethyl acetoacetate (2 mmol), hydrazine hydrate (2 mmol), 3-nitrobenzaldehyde (1 mmol) and ammonium acetate (1 mmol), catalyst (10–25 mg). The yields relate to the isolated product.

Scheme 1 Synthesis of pyrazolopyridine derivatives (5a–l) in the presence of Alg@SBA-15/Fe₃O₄ nanocatalyst.

sites followed by nucleophilic attack of hydrazine on them, which is accompanied by a ring-forming reaction, intermediate **I**, a pyrazolone ring, is formed. Next, *via* the Knoevenagel condensation reaction between the activated aldehyde and the pyrazolone ring, intermediate **II** is formed. The further step, by

performing the Michael addition reaction between the pyrazolone ring and intermediate **II**, intermediate **III** is formed. Then, intermediate **IV** is constructed by the nucleophilic attack of ammonia (from ammonium acetate) on intermediate **III** with the aid of the Alg@SBA-15/Fe₃O₄ catalyst. In the last step,

Table 3 Synthesis yields of different pyrazolopyridine derivatives by using Alg@SBA-15/Fe₃O₄ catalyst under the optimized conditions

Entry	Aldehyde	Product	Time (min)	Yield ^a (%)	Mp (°C)	
					Found	Reported
1	Benzaldehyde	5a	25	95	240–243	240–241 (ref. 8)
2	3-NO ₂ benzaldehyde	5b	20	97	287–290	285–287 (ref. 8)
3	2,4-Cl ₂ benzaldehyde	5c	20	97	307–310	308 (ref. 8)
4	4-Cl benzaldehyde	5d	20	93	257–259	255–257 (ref. 8)
5	4-Me benzaldehyde	5e	25	97	246–248	244–246 (ref. 8)
6	4-OMe benzaldehyde	5f	30	92	184–186	185–186 (ref. 50)
7	3-Me benzaldehyde	5g	30	90	279–280	280–282 (ref. 51)
8	2-Cl benzaldehyde	5h	30	93	220–223	219–221 (ref. 52)
9	2-Me benzaldehyde	5i	25	91	290–293	290–292 (ref. 53)
10	4-F benzaldehyde	5j	20	92	258–260	259–261 (ref. 8)
11	2-OH, 3-OMe benzaldehyde	5k	30	91	202–204	200–202 (ref. 54)
12	3,4,5-(OMe) ₃ benzaldehyde	5l	25	93	249–251	248–250 (ref. 51)
13	4-CH(Me) ₂ benzaldehyde	5m	30	91	281–284	280–282 (ref. 51)

^a The yields relate to the isolated product.



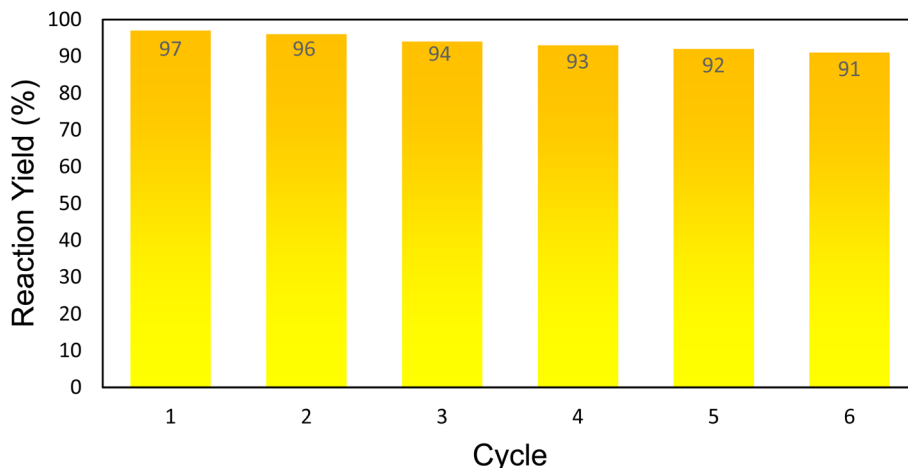


Fig. 9 Recycling diagram of the Alg@SBA-15/Fe₃O₄ catalyst in the synthesis of 5b.

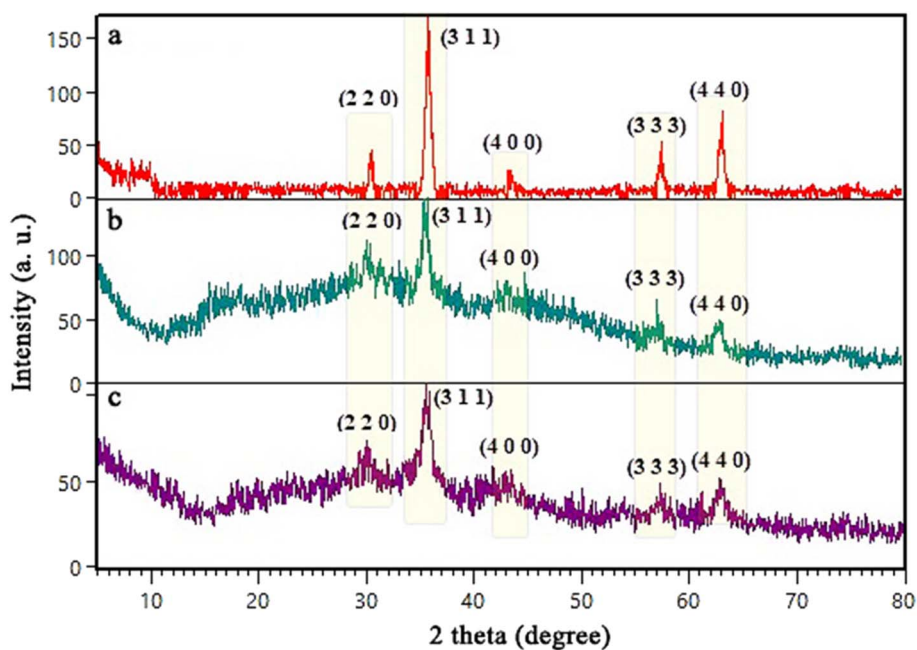


Fig. 10 The XRD pattern of (a) Fe₃O₄ MNPs, (b) the fresh Alg@SBA-15/Fe₃O₄ catalyst (c) after five times reusing in the reaction.

through intramolecular ring formation of intermediate **IV**, which leads to the removal of water from it, as well as product tautomerism, the final product of pyrazolopyridine is synthesized.

2.3.4. Catalyst recyclability and stability. In this study, the reusability of the prepared Alg@SBA-15/Fe₃O₄ catalytic system in the synthesis of pyrazolopyridine derivatives was evaluated, since it is an important issue in green chemistry.^{55–57} After the reaction was completed, the Alg@SBA-15/Fe₃O₄ catalyst was isolated from the reaction medium by using an external magnet and then diluted with ethanol and dried in an oven at 55 °C for 6 h to prepare for the subsequent catalytic run. For following runs, a fixed amount of the recovered Alg@SBA-15/Fe₃O₄ catalyst was utilized. According to Fig. 9, after five

consecutive uses of Alg@SBA-15/Fe₃O₄ catalyst in the synthesis of pyrazolopyridine derivatives, less than 9% of its catalytic power was reduced compared to the first use in the same reaction. As a result, the Alg@SBA-15/Fe₃O₄ catalyst can be used for five runs without considerable reduction in its catalytic activity. Furthermore, the XRD analysis of the recycled catalyst (Fig. 10) was also performed and compared with diffractograms of the Fe₃O₄ MNPs and the fresh Alg@SBA-15/Fe₃O₄ catalyst to determine the structural and mechanical stability of the fabricated catalyst. The results indicate that the structure of the Alg@SBA-15/Fe₃O₄ has remained unchanged after reuses in the reaction compared to fresh catalysts and all of the indicative peaks of the nanocomposite appeared in XRD pattern of recycled one.

3. Experimental section

3.1. Materials and equipment

All used chemicals in present work, were bought from Merck and Sigma-Aldrich companies. In order to show the correct construction of the Alg@SBA-15/Fe₃O₄ catalyst and its characteristics, several analyses were used. A Shimadzu IR-470 spectrometer was used to get Fourier-transform infrared (FT-IR) spectra of the samples. Energy dispersive X-ray analysis (EDX) of the neat materials and prepared samples was performed employing a Numerix DXP-X10P and TESCAN, MIRA II. The morphology and surface property of neat materials and prepared compounds were examined with the help of images obtained from FESEM (model KYKY-EM8000). By employing a Lakeshore 7407, the magnetic behavior of the samples was studied at ambient temperature. By utilizing an STA504 device, thermal stability of compounds was assessed by TGA. The produced samples' X-ray diffraction (XRD) pattern was achieved utilizing an X-ray diffractometer (JEOL JDX-8030 (20 mA, 30 kV)). The N₂ adsorption-desorption isotherm of this study was also obtained using Micromeritics ASAP 2010. Also, a, KQ-250 DE (40 kHz) device was utilized as an ultrasound cleaning bath.

3.2. Preparation method

3.2.1. Synthesis of SBA-15. Well-ordered SBA-15 was synthesized by hydrothermal method. For this, in a 250 mL flask, 3.5 mL of pluronic P123, 120.0 mL of distilled water and 20.0 mL of 37% hydrochloric acid were poured and then stirred at room temperature to mix thoroughly. Then, after the temperature of the reaction mixture fixed at 40 °C, the flask containing the components was stirred at 40 °C for 24 h. After this time, 8.5 mL of TEOS (tetraethyl orthosilicate) was dropped to the above solution while stirring vigorously. After the addition of TEOS, the resulting mixture was spun at 120 rpm for a day. Next, the reaction mixture was transferred to a teflon-lined autoclave reactor and heated at 150 °C for 24 h in an oven. After the synthesis process, the resulting mixture cooled to room temperature and the resulting sediment was separated by filtration. The resulting solid was rinsed with distilled water many times and dried overnight at 100 °C. The dried powder was calcination was performed by heating for 5 h at 500 °C to remove of surfactant and constructing the ordered channel of mesoporous SBA-15.

3.2.2. Synthesis of SBA-15/Fe₃O₄. 0.43 g of FeCl₂·4H₂O and 0.54 g of FeCl₃·6H₂O with 100.0 mL of distilled water were poured into a 250 mL flask and stirred for 40 minutes at room temperature. Then 0.5 g of white powder of SBA-15 which was synthesized in the previous step was added to the reaction mixture under N₂ atmosphere and stirred for 30 min. After raising the reaction mixture temperature to 80 °C, 10.0 mL of 25% ammonia solution was dropped to the reaction mixture. The resulting brown mixture was stirred at 80 °C for one hour. Using an external magnet, SBA-15/Fe₃O₄ was isolated, and it was washed many times with distilled water. The resulting product was placed in an oven at 100 °C for several hours to dry.

3.2.3. Synthesis of SBA-15/Fe₃O₄-NH₂. 1.0 g of SBA-15/Fe₃O₄ synthesized in the previous step was poured into 40.0 mL of toluene in a 100 mL flask. After that, an ultrasonic bath was used to thoroughly disperse the reaction mixture. After that, 2.0 mL of APTES was added and the temperature was increased to 120 °C, then it was stirred under reflux for 32 h. After these 32 h, the precipitate was washed several times with water and ethanol and placed in an oven at 60 °C for 12 h to dry.

3.2.4. Preparation of methyl-esterified alginic acid. Methyl-esterified alginic acid was prepared *via* chemical esterification of alginic acid by using methanol in mildly acidic aqueous solution.⁵⁸ 2.0 g of alginic acid was poured in a 500 mL flask containing 200.0 mL of methanol and 2.0 mL of sulfuric acid 98%. Then, the temperature of the reaction was increased to 60 °C and it was stirred for 72 h at the same temperature. After 72 h, the resulting powder was diluted with distilled water and acetone, then placed in an oven at 55 °C to dry.

3.2.5. Synthesis of Alg@SBA-15/Fe₃O₄. 0.7 g of synthesized SBA-15/Fe₃O₄-NH₂ with 0.05 g of methyl-esterified alginic acid were poured in 10 mL of DMF solvent and was stirred for 72 h at 55 °C. The product was then gathered using a magnet and rinsed three times with distilled water and ethanol. Afterwards it was dried in an oven at 55 °C.

3.3. Procedure for the synthesis of pyrazolopyridine derivatives with the Alg@SBA-15/Fe₃O₄ catalytic system

Four reactants including; ethyl acetoacetate (2.0 mmol), hydrazine hydrate (2.0 mmol), aromatic aldehydes (1.0 mmol) and ammonium acetate (1.0 mmol) were mixed and reacted in the presence of 0.02 g of the Alg@SBA-15/Fe₃O₄ catalytic system in the presence of 2.0 mL EtOH at room temperature. TLC (Thin layer chromatography) technique was employed to investigate the reaction completion process. After the reaction was finished, the magnetic nanocatalyst was effortlessly removed from the reaction medium by an external magnet. In the final step, by adding water to the reaction mixture, pure products were got, filtered and washed with further water. Most of the time, additional purification was not required.

3.3.1. Spectral data of two products

3.3.1.1 *3,5-Dimethyl-4-phenyl-1,4,7,8-tetrahydrodipyrzolo [3,4-*b*:4',3'-*e*] pyridine (5a).* FT-IR (KBr) ν_{\max} : 3540, 3301, 2925, 1614, 1515, 1375 cm⁻¹; ¹H-NMR (300 MHz, DMSO-*d*₆): δ_{H} (ppm) = 2.06 (6H, 2-CH₃, singlet), 4.81 (1H, -CH, singlet), 7.14–7.20 (5H, H-aromatic, multi), 11.24–11.32 (3H, -NH, broad singlet); ¹³C NMR (75 MHz, DMSO-*d*₆): δ (ppm) = 10.8, 33.1, 104.6, 125.8, 127.9, 128.1, 140.2, 143.7, 161.5. CHN Anal. Calcd for C₁₅H₁₅N₅: C: 67.90, H: 5.70, N: 26.40 found C: 67.86, H: 5.67, N: 26.37, mass for C₁₅H₁₅N₅: [M-H] 265.133 found 265.122.

3.3.1.2 *4-(4-Chlorophenyl)-3,5-dimethyl-1,4,7,8-tetrahydrodipyrzolo[3,4-*b*:4',3'-*e*] pyridine (5d).* FT-IR (KBr) ν_{\max} : 3100, 2854, 1612, 1526, 1368 cm⁻¹; ¹H-NMR (300 MHz, DMSO-*d*₆): δ_{H} (ppm) = 2.06 (6H, 2-CH₃, singlet), 4.8 (1H, -CH, singlet), 7.12 (5H, H-aromatic, multi), 7.24 (2H, H-aromatic, broad singlet), 11.35 (3H, -NH, broad singlet). ¹³C NMR (75.47 MHz, DMSO-*d*₆): δ (ppm) = 10.7, 32.6, 104.3, 128.0, 129.8, 130.4, 140.1, 142.7, 161.4. CHN Anal.



Calcd for C₁₅H₁₄ClN₅: C: 60.10, H: 4.71, N: 23.36 found C: 60.08, H: 4.69, N: 23.35.

3.3.1.3 4-(4-Fluorophenyl)-3,5-dimethyl-1,4,7,8-tetrahydrodipyrazolo[3,4-b:4',3'-e]pyridine (**5j**). FT-IR (KBr) ν_{\max} : 3165, 3067, 1602, 1511; ¹H NMR (300.13 MHz, DMSO-d₆): δ_{H} (ppm) = 2.07 (6H, 2-CH₃, singlet), 4.81 (1H, -CH, singlet), 7.01–7.05 (2H, H-aromatic, multi), 7.08–7.13 (m, 2H, H-Aromatic, multi), 11.34 (3H, -NH, broad singlet); ¹³C NMR (75.47 MHz, DMSO-d₆): δ = 10.8, 32.5, 104.6, 114.8, 129.6, 129.7, 139.8, 161.4, 162.4; Anal. Calcd for C₁₅H₁₄FN₅: C: 63.59, H: 4.98, N: 24.72 found C: 63.57, H: 5.01, N: 24.69; mass for C₁₅H₁₄FN₅ [M-H] 283.12 found 283.11.

4. Conclusion

In this study, a novel mesoporous nanocomposite based on SBA-15 was fabricated in which SBA-15 was modified with Fe₃O₄ nanoparticles, APTES and methyl-esterified alginate. For the synthesis of pyrazolopyridine derivatives, the prepared nanocomposite demonstrated excellent catalytic performance. The corresponding products were synthesized in high yields without using a complicated work-up process. TGA result revealed that this nanocomposite is thermally stable, losing only around 25% of its weight up to 800 °C. The XRD analysis also confirmed that Alg@SBA-15/Fe₃O₄ has properly been prepared. Also, the results of other analyzes, including VSM, FESEM, BET and EDX, were consistent with the confirmation of the correct synthesis of Alg@SBA-15/Fe₃O₄. The catalytic system synthesized in this study can be considered efficient catalyst for the synthesis of pyrazolopyridine derivatives. One of the reasons for this is its superparamagnetic property, which was also confirmed by VSM analysis, which allows this catalyst to be easily separated from the medium and reused for several times. In the model reaction of this study for the synthesis of various derivatives of pyrazolopyridine, while a small amount of Alg@SBA-15/Fe₃O₄ (0.02 g) was used, high yield products were obtained in a short time.

Conflicts of interest

The authors declare no conflict of interest.

Acknowledgements

The authors are grateful of Iran University of Science & Technology (IUST) for the partial support.

References

- 1 A. P. Marjani, J. Khalafy and S. Akbarzadeh, *Green Process. Synth.*, 2019, **8**, 533–541.
- 2 K. S. Gudmundsson, S. D. Boggs, P. R. Sebahar, L. D. A. Richardson, A. Spaltenstein, P. Golden, P. B. Sethna, K. W. Brown, K. Moniri, R. Harvey and K. R. Romines, *Bioorg. Med. Chem. Lett.*, 2009, **19**, 4110–4114.
- 3 H. De Mello, A. Echevarria, A. M. Bernardino, M. Canto-Cavaleiro and L. L. Leon, *J. Med. Chem.*, 2004, **47**, 5427–5432.
- 4 R. G. Stein, J. H. Biel and T. Singh, *J. Med. Chem.*, 1970, **13**, 153–155.
- 5 R. Navari, S. Balalaie, S. Mehrparvar, F. Darvish, F. Rominger, F. Hamdan and S. Mirzaie, *Beilstein J. Org. Chem.*, 2019, **15**(85), 874–880.
- 6 R. Taheri-Ledari, S. M. Hashemi and A. Maleki, *RSC Adv.*, 2019, **9**, 40348–40356.
- 7 F. Hassanzadeh-Afruzi, M. M. Salehi, G. Heidari, A. Maleki and E. N. Zare, *J. Mol. Struct.*, 2023, **1274**, 134490.
- 8 R. Taheri-Ledari, J. Rahimi and A. Maleki, *Ultrason. Sonochem.*, 2019, **59**, 104737.
- 9 R. Taheri-Ledari and A. Maleki, *J. Pept. Sci.*, 2020, **26**, e3277.
- 10 F. Hassanzadeh-Afruzi, Z. Amiri-Khamakani, S. Bahrami, M. R. Ahghari and A. Maleki, *Sci. Rep.*, 2022, **12**(1), 1–16.
- 11 F. Hassanzadeh-Afruzi, A. Maleki and E. N. Zare, *J. Nanostruct. Chem.*, 2022, **2022**, 1–12.
- 12 R. Taheri-Ledari, W. Zhang, M. Radmanesh, N. Cathcart, A. Maleki and V. Kitaev, *J. Nanobiotechnol.*, 2021, **19**, 1–21.
- 13 F. Hassanzadeh-Afruzi, S. Asgharnasl, S. Mehraeen, Z. Amiri-Khamakani and A. Maleki, *Sci. Rep.*, 2021, **11**(1), 1–15.
- 14 J. Safaei-Ghomi and R. Teymuri, *Appl. Organomet. Chem.*, 2019, **33**, e5150.
- 15 P. Verma, Y. Kuwahara, K. Mori, R. Raja and H. Yamashita, *Nanoscale*, 2020, **12**, 11333–11363.
- 16 D. Zhao, Q. Huo, J. Feng, B. F. Chmelka and G. D. Stucky, *J. Am. Chem. Soc.*, 1998, **120**, 6024–6036.
- 17 I. M. El-Nahal, J. K. Salem, N. S. Tabasi, R. Hempelmann and F. S. Kodeh, *Chem. Phys. Lett.*, 2018, **691**, 211–218.
- 18 S. Singh, R. Kumar, H. D. Setiabudi, S. Nanda and D. V. N. Vo, *Appl. Catal., A*, 2018, **559**, 57–74.
- 19 B. N. Mahato and T. Krithiga, *Can. J. Chem.*, 2022, **100**, 9–17.
- 20 J. Wang, H. Ge and W. Bao, *Mater. Lett.*, 2015, **145**, 312–315.
- 21 A. Ghorbani-Choghamarani, M. Mohammadi, T. Tamoradi and M. Ghadermazi, *Polyhedron*, 2019, **158**, 25–35.
- 22 F. Hassanzadeh-Afruzi, F. Esmailzadeh, S. Asgharnasl, F. Ganjali, R. Taheri-Ledari and A. Maleki, *Sep. Purif. Technol.*, 2022, **291**, 120956.
- 23 K. Valadi, S. Gharibi, R. Taheri-Ledari and A. Maleki, *Solid State Sci.*, 2020, **101**, 106141.
- 24 F. Ganjali, A. Kashtiaray, S. Zarei-Shokat, R. Taheri-Ledari and A. Maleki, *Nanoscale Adv.*, 2022, **4**, 1263–1307.
- 25 A. Maleki, R. Taheri-Ledari and M. Soroushnejad, *ChemistrySelect*, 2018, **3**, 13057–13062.
- 26 Z. Hajizadeh, F. Hassanzadeh-Afruzi, D. F. Jelodar, M. R. Ahghari and A. Maleki, *RSC Adv.*, 2020, **10**, 26467–26478.
- 27 F. Hassanzadeh-Afruzi, H. Dogari, F. Esmailzadeh and A. Maleki, *Appl. Organomet. Chem.*, 2021, **35**, e6363.
- 28 A. Hakiki, B. Boukoussa, H. Habib Zahmani, R. Hamacha, N. H. Hadj Abdelkader, F. Bekkar, F. Bettahar, A. P. Nunes-Beltrao, S. Hacini, A. Bengueddach and A. Azzouz, *Mater. Chem. Phys.*, 2018, **212**, 415–425.
- 29 S. Yuan, M. Wang, J. Liu and B. Guo, *J. Environ. Manage.*, 2020, **254**, 109787.
- 30 M. Mureddu, F. Ferrara and A. Pettinau, *Appl. Catal., B*, 2019, **258**, 117941.
- 31 J. Safaei Ghomi, Z. Akbarzadeh and A. Bakhtiari, *J. Coord. Chem.*, 2019, **72**, 826–840.
- 32 J. S. Ghomi and A. Bakhtiari, *ChemistrySelect*, 2018, **3**, 12704–12711.



- 33 J. Safaei-Ghomi, R. Teymuri and A. Bakhtiari, *BMC Chem.*, 2019, **13**, 1–12.
- 34 J. Safaei-Ghomi and A. Bakhtiari, *RSC Adv.*, 2019, **9**, 19662–19674.
- 35 N. Noroozi Pesyan, G. Rezanejade Bardajee, E. Kashani, M. Mohammadi and H. Batmani, *Res. Chem. Intermed.*, 2019, **46**(1), 347–367.
- 36 A. Srivastava, A. Yadav and S. Samanta, *Tetrahedron Lett.*, 2015, **56**, 6003–6007.
- 37 I. P. S. Fernando, D. Kim, J. W. Nah and Y. J. Jeon, *Chem. Eng. J.*, 2019, **355**, 33–48.
- 38 A. Maleki, Z. Varzi and F. Hassanzadeh-Afruzi, *Polyhedron*, 2019, **171**, 193–202.
- 39 Z. Durmus, H. Sözeri, B. Unal, A. Baykal, R. Topkaya, S. Kazan and M. S. Toprak, *Polyhedron*, 2011, **30**, 322–328.
- 40 X. Guo, Y. Wang, Y. Qin, P. Shen and Q. Peng, *Int. J. Biol. Macromol.*, 2020, **162**, 618–628.
- 41 R. Taheri-Ledari, W. Zhang, M. Radmanesh, *et al.*, *Small*, 2020, **16**, 2002733.
- 42 A. Maleki, R. Taheri-Ledari, J. Rahimi, *et al.*, *ACS Omega*, 2019, **4**, 10629–10639.
- 43 R. Taheri-Ledari, A. Maleki, E. Zolfaghari, M. Radmanesh, H. Rabbani, A. Salimi and R. Fazel, *Ultrason. Sonochem.*, 2020, **61**, 104824.
- 44 R. Taheri-Ledari, K. Valadi, S. Gharibi and A. Maleki, *Mater. Res. Bull.*, 2020, **130**, 110946.
- 45 M. Munir, M. Ahmad, M. Mubashir, S. Asif, A. Waseem, A. Mukhtar, S. Saqib, H. Siti Halimatul Munawaroh, M. K. Lam, K. Shiong Khoo, A. Bokhari and P. Loke Show, *Bioresour. Technol.*, 2021, **328**, 124859.
- 46 A. Maleki, R. Taheri-Ledari, R. Ghalavand and R. Firouzi-Haji, *J. Phys. Chem. Solids*, 2020, **136**, 109200.
- 47 Z. P. Zanele, F. M. Mtunzi, S. M. Nelana, A. N. Ebelegi, N. Ayawei, E. D. Dikio, D. Wankasi and P. N. Diagboya, *Langmuir*, 2021, **37**, 9764–9773.
- 48 E. F. Aboelfetoh, M. E. Zain Elabedien and E. Z. M. Ebeid, *J. Environ. Chem. Eng.*, 2021, **9**, 104817.
- 49 K. Dashtian, R. Zare-Dorabei, R. Jafarinia and M. Saghanejhad Tehrani, *J. Environ. Chem. Eng.*, 2017, **5**, 5233–5240.
- 50 Kavita, P. Verma, D. K. Verma, B. Kumar, A. K. Singh, N. Shukla, V. Srivastava and R. B. Rastogi, *RSC Adv.*, 2020, **10**, 10188–10196.
- 51 R. R. Chinthaparthi, V. L. Chittiboena, S. Jorepalli and C. S. R. Gangireddy, *J. Heterocycl. Chem.*, 2021, **58**, 1104–1116.
- 52 F. Tamaddon and D. Arab, *RSC Adv.*, 2019, **9**, 41893–41902.
- 53 M. Dashteh, M. Yarie, M. A. Zolfigol, A. Khazaei and S. Makhdoomi, *Appl. Organomet. Chem.*, 2021, **35**, e6222.
- 54 J. Safaei-Ghomi, R. Sadeghzadeh and H. Shahbazi-Alavi, *RSC Adv.*, 2016, **6**, 33676–33685.
- 55 H. Zhao, M. Cheng, J. Zhang and M. Cai, *Green Chem.*, 2014, **16**, 2515–2522.
- 56 Z. Abbasi, S. Rezayati, M. Bagheri and R. Hajinasiri, *Chin. Chem. Lett.*, 2017, **28**, 75–82.
- 57 S. Sadjadi, *J. Mol. Liq.*, 2021, **323**, 114994.
- 58 T. Taubner, M. Marounek and A. Synytsya, *Int. J. Biol. Macromol.*, 2017, **103**, 202–207.

