Asparagine-EDTA MNPs: A Highly Efficient And Recyclable Magnetic Multifunctional Core-Shell Nanocatalyst For Green Synthesis of Biologically-Active 3,4-Dihydropyrimidin-2(1H)-One Compounds

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Research Article

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Abstract

In this study, the new asparagine grafted on the EDTA-modified Fe$_3$O$_4$@SiO$_2$ core-shell (Fe$_3$O$_4$@SiO$_2$-APTS-EDTA-asparagine) magnetic nanoparticles were prepared and their structures were properly confirmed using different spectroscopic, microscopic and magnetic methods or techniques such as FT-IR, EDX, XRD, FESEM, TEM, TGA and VSM. The Fe$_3$O$_4$@SiO$_2$-APTS-EDTA-asparagine core-shell nanomaterial was examined, as a highly efficient multifunctional and recoverable nanocatalyst, for the synthesis of a wide range of nitrogen-containing heterocycles and biologically-active 3,4-dihydropyrimidin-2(1H)-one derivatives under solvent-free conditions. It was proved that Fe$_3$O$_4$@SiO$_2$-APTS-EDTA-asparagine MNPs, as a catalyst having excellent thermally and magnetic stability, specific morphology and acidic sites, can activate the Biginelli reaction components. Moreover, environmental-friendliness and nontoxic nature properties of the catalyst, cost effectiveness, low catalyst loading, easy separation of the catalyst from products and short time of reaction are some of the remarkable advantages of this green protocol.

Introduction

Green chemistry has played a key role in the development of human civilization$^{1-4}$. In this regard, magnetic nanoparticles (MNPs) have received considerable interest because of their unique properties$^5$. The outstanding properties of MNPs have made them superior and indispensable in many areas of industry and academia including information storage$^6$, medicine$^7$, drug delivery$^8$, magnetic resonance imaging (MRI)$^9,10$, biomedical applications$^5,11,12$, and environmental remediation$^{13}$ as well as heterogeneous catalysis$^{14-17}$. In academia, MNPs represent a promising new technology for performing chemical reactions because they are separated from the reaction medium, comfortably$^{18,19}$. In industry, due to the importance of the cost of chemical processes and the reuse of catalysts, special attention is paid to these nanoparticles$^{20,21}$. MNPs tend to agglomerate under a magnetic field that reduces their surface to volume ratio and consequently decreases catalytic activity$^{22}$. Therefore, MNPs must be stabilized to improve their properties and prevent undesirable agglomeration$^{23}$. In fact, they are coated with a protective layer such as carbon layers$^{24,25}$, organic polymer$^26$ or silica$^{27,28}$.

Moreover, multi-component reactions (MCRs) are the most desirable powerful synthetic route in which three or more reactants come together in a single reaction vessel to form a wide range of acyclic or heterocyclic compounds by the one-pot processes$^{29,30}$. MCRs afford extended molecular complexity and diversity from simple starting materials with high atom economy, which have found application in medicinal and natural products chemistry$^{31,32}$. Indeed, the most significant feature of MCRs is generating almost no by-products or simple molecules such as H$_2$O or EtOH$^{33-37}$. Hence, in agreement of the green and sustainable chemistry process, development and the advancement of catalysts to promote MCRs are very important in synthetic and medicinal chemistry$^{38-40}$.

Among the various types of nitrogen-containing heterocycles, derivatives of 3,4-dihydropyrimidin-2(1H)-one, as biologically-active compounds, have found versatile applications such as anti-bacterial, anti-
inflammatory, antihypertensive agents, calcium channel blockers, antitumor compounds. A simple general protocol for access to 3,4-dihydropyrimidin-2(1H)-ones involves a three-component one-pot Biginelli cyclocondensation of ethyl acetoacetate, urea and various aldehydes accelerated by different types of catalytic systems such as polymer-supported catalysts, ionic liquids, ionic liquid/silica, metal–organic framework (MOF), montmorillonite clay, magnetic nanoparticles, Lewis acidic zirconium (IV)-salophen perfluorooctanesulfonate or sulfated polyborate, nanocrystalline CdS thin film, graphene oxide, mesoporous materials, sulfuric acid, and environmental friendly energy inputs such as ultrasound or microwave irradiation. Most of the reported methods in this regard have the role of heterogeneous catalysts and high value. However, these have problems such as complicated and tedious separation of products and catalysts, toxic reaction conditions, long reaction times and low yields. Therefore, there is still room to develop more environmentally-benign protocols to promote the Biginelli MCR condensation.

In many previous reports, ethylenediaminetetraacetic acid (EDTA) has been used as an ion exchange and chelating agent for various metal ions, but this compound has a good ability as an inexpensive and non-toxic cross-linker to make strong bonds with organic materials having nucleophilic centers. On the other hand, asparagine is one of the 20 amino acids found in the cells of the human body and is essential for maintaining balance in the central nervous system. Asparagine can act as a biocompatible precursor and bifunctional organocatalyst due to its high natural abundance and cost-effectiveness with acidic and basic sites.

In this research, we herein report the synthesis and characterizations of new asparagine grafted on the EDTA-modified Fe$_3$O$_4$@SiO$_2$ core-shell magnetic nanoparticles (Fe$_3$O$_4$@SiO$_2$-APTS-EDTA-asparagine), as a magnetically recoverable nanocatalyst, to promote the Biginelli reaction efficiently at 60°C under solvent-free conditions (Fig. 1).

**Results And Discussion**

**Characterization of the Fe$_3$O$_4$@SiO$_2$-APTS-EDTA-asparagine nanocatalyst (1)**

The overall procedure for the synthesis of Fe$_3$O$_4$@SiO$_2$-APTS-EDTA-asparagine (1) has been summarized in Fig. 1. At first, the obtained magnetic nanoparticles were characterized using different physicochemical techniques such as Fourier transform infrared (FT-IR) spectroscopy, energy-dispersive X-ray (EDX) spectroscopy, field emission scanning electron microscopy (FESEM), transmission electron microscopy (TEM), X-ray diffraction spectroscopy (XRD), vibrating sample magnetometer (VSM), and thermogravimetric analysis (TGA).

The FT-IR spectroscopy was employed to determine the functional groups and structure of Fe$_3$O$_4$ (a), Fe$_3$O$_4$@SiO$_2$ (b), Fe$_3$O$_4$@SiO$_2$-APTS (c), Fe$_3$O$_4$@SiO$_2$-APTS-EDTA (d) and Fe$_3$O$_4$@SiO$_2$-APTS-EDTA-
asparagine (e). The results are presented in Fig. 2. In the spectra of Fe$_3$O$_4$ nanoparticles (Fig. 2a) the bands displayed at 620 cm$^{-1}$ and about 3410 cm$^{-1}$ are attributed to stretching vibration of Fe−O bond and surface hydroxyl groups, respectively. These peaks were observed in all five samples isolated at the different synthetic stages. In the FT-IR spectrum of Fe$_3$O$_4$@SiO$_2$ (Fig. 2b), the absorption bands at 881 and 1036 cm$^{-1}$ can be ascribed to the presence of Si−O−Si symmetric and Si−O−Si asymmetric stretching modes, reflecting the coating of silica layer on the magnetite nanoparticles $^{71}$. SP$^3$ C−H stretching vibrations about 2922 cm$^{-1}$ confirmed the presence of the anchored (3-aminopropyl) triethoxysilane (APTS) group and the band about 1400 cm$^{-1}$ is assigned to the bending of −NH groups of Fe$_3$O$_4$@SiO$_2$-APTS MNPs (Fig. 2c) $^{72}$. In the FT-IR spectrum of Fe$_3$O$_4$@SiO$_2$-APTS-EDTA (Fig. 2d), the peaks at 1635 cm$^{-1}$, 1707 cm$^{-1}$ and 1760 cm$^{-1}$ corresponding to the C=O vibration of amide, acid and anhydride groups, respectively. In the last step, the peak at 1760 cm$^{-1}$, which belongs to the anhydride group has been removed and new peaks at 1651 cm$^{-1}$ and 1737 cm$^{-1}$ are attributed to the amide and acid groups in the surface of Fe$_3$O$_4$@SiO$_2$-APTS-EDTA-asparagine (Fig. 2e). These results from the FT-IR spectrum confirm that the silica coating and subsequent steps have been successfully performed on the surface of Fe$_3$O$_4$.

Compositional analysis of the Fe$_3$O$_4$@SiO$_2$-APTS-EDTA-asparagine magnetic nanocatalyst (1) was carried out using energy-dispersive X-ray spectroscopy (EDX). The EDX spectra of the Fe$_3$O$_4$@SiO$_2$-APTS-EDTA-asparagine nanomaterial (1) are depicted in Fig. 3. In addition, the EDX analysis showed the well-defined peaks related to C, O, N, Si and Fe in the structure of Fe$_3$O$_4$@SiO$_2$-APTS-EDTA-asparagine (1) with the percentages of 40.32, 36.57, 11.69, 6.34 and 5.08, respectively.

The morphology and texture of Fe$_3$O$_4$@SiO$_2$-APTS-EDTA-asparagine MNPs (1) were indicated by FESEM analysis and their photographs were presented in Fig. 4. According to these FESEM photographs, the size and surface shape of nanoparticles are well observed, which proves that the particles are spherical and without agglomeration. The FESEM photographs supported the formation of spherically shaped MNPs, which is in accordance with TEM analysis.

The TEM analysis of the Fe$_3$O$_4$@SiO$_2$-APTS-EDTA-asparagine (1) MNPs in two scales is shown in Fig. 5. The TEM images demonstrated structural order and the morphology suggested that the magnetite nanoparticles have an average diameter size of 41 nm.

The XRD pattern of Fe$_3$O$_4$@SiO$_2$-APTS-EDTA-asparagine (1) was shown in Fig. 6. The reflection peaks were compared with the reference standard patterns related to EDTA (card no. JCPDS, 00-033-1672), Fe$_3$O$_4$ (card no. JCPDS, 01-088-0315) and asparagine (card no. JCPDS, 00-031-1542). The sharp peaks in this pattern are generated by combining several peaks. These new sharp peaks are ascribed to the produced Fe$_3$O$_4$@SiO$_2$-APTS-EDTA-asparagine MNPs structures after modification reactions by EDTA and asparagine, respectively.
The magnetic properties of MNPs were measured via vibrating sample magnetometry (VSM). The magnetic attributes of Fe$_3$O$_4$ and Fe$_3$O$_4$@SiO$_2$-APTS-EDTA-asparagine MNPs (1) were measured out at room temperature by applied magnetic field -1000 to +1000 Oersted. According to data presented in Fig. 7, The values of the magnetization saturation (Ms) for Fe$_3$O$_4$ and Fe$_3$O$_4$@SiO$_2$-APTS-EDTA-asparagine MNPs (1) are 73.12 and 20.84 emu/g, respectively. Moreover, the VSM curves of both samples exhibit no hysteresis loops and this property demonstrated that no aggregation occurred in the presence of magnetic fields. A decrease in the magnetic saturation of the Fe$_3$O$_4$@SiO$_2$-APTS-EDTA-asparagine was observed after coating with SiO$_2$ and functionalization with APTS. However, the magnetic saturation of Fe$_3$O$_4$@SiO$_2$-APTS-EDTA-asparagine (1) is sufficient to be recovered by exerting an external magnet.

Thermal stability of the Fe$_3$O$_4$@SiO$_2$-APTS-EDTA-asparagine nanomaterial (1) was investigated under the air atmosphere over the temperature range of 50 – 800°C (Fig. 8). The Fe$_3$O$_4$@SiO$_2$-APTS-EDTA-asparagine MNPs (1) display three weight loss steps over the temperature range of TGA and the total weight loss of nanocatalyst 1 is around 60%. According to obtained results, in the first step 15% weight loss in the range of 150 – 200°C is due to the evaporation of adsorbed water and organic solvents that remain in the nanocatalyst through its preparation processes. In addition, 22% weight loss in the range of 200 – 400°C corresponds to the loss of EDTA–asparagine moiety. In the last step, the sharp weight loss of 23% at 400-700°C can be assigned to the decomposition of APTS moiety in the MNPs framework. These results also indicate that APTS, EDTA and asparagine has been successfully grafted onto the surface of Fe$_3$O$_4$@SiO$_2$. Above 700°C only Fe$_3$O$_4$ was present.

**Optimization of conditions in the Biginelli reaction using Fe$_3$O$_4$@SiO$_2$-APTS-EDTA asparagine nanocatalyst (1)**

In our preliminary experiments, the catalytic activity of as prepared catalyst 1 was evaluated in the formation of dihydropyrimidin-2(1$H$)-one derivatives by the Biginelli condensation. For this purpose, reaction conditions were optimized using the equimolar mixtures of urea (2, 1 mmol), 4-chlorobenzaldehyde (3a, 1 mmol) and ethyl acetoacetate (4a, 1 mmol) as the model reaction (Eq. 1). In a systematic screening, the reaction conditions were investigated precisely by considering of several crucial variables such as catalyst loading, reaction time, solvent and reaction temperature, as given in Table 1.

Initially, in the absence of any catalyst and solvent, the progress of model reaction was slow and the yield of the 9-(4-chlorophenyl)-3,3,6,6-tetramethyl-3,4,6,7,9,10-hexahydroacridine-1,8(2$H$,5$H$)-dione (5a) was trace, even after a long time (Table 1, entry 1). Then, in the presence of very low amount of Fe$_3$O$_4$@SiO$_2$-APTS-EDTA-asparagine (1) loading, as a nanocatalyst, without any solvent at room temperature, a good yield of the desired product 5a was obtained (Table 1, entry 2). To achieve an excellent yield, the reaction temperature was increased to 60°C (Table 1, entry 3). Afterward, the model reaction was performed with lower catalyst 1 loading under solvent-free conditions as well as polar and non-polar solvents. Furthermore, the effect of temperature and different solvents was investigated (Table 1, entries 4-13). Also, the model reactions in the presence of EDTA and asparagine were separately investigated, but lower yields of the desired product 5a were isolated (Table 1, entries 14-15).
Following the steps of optimizing the reaction conditions, the effect of different solvents and amount of catalyst loadings are summarized in Fig. 9. The model reaction was investigated under solvent-free conditions and different solvents such as EtOH, MeOH, EtOH/H$_2$O (1:1), and DMF using Fe$_3$O$_4$@SiO$_2$-APTS-EDTA-asparagine nanocatalyst (1) with different loading of the catalyst 1. According to the obtained findings summarized in Table 1 and Fig. 9, the optimum reaction conditions were found to be 10 mg of Fe$_3$O$_4$@SiO$_2$-APTS-EDTA-asparagine nanocatalyst (1) loading under solvent-free conditions at 60 °C.

After the above experiments, the scope of reaction was expanded by using aromatic aldehydes having electron-withdrawing or electron-donating groups under the optimized conditions (Eq. 2). The results are summarized in Table 2. As expected, in this novel magnetic heterogeneous catalytic system the reaction rate of aldehydes with electron-donating groups was slower than electron-releasing ones and required more time to complete the reaction. An alternative variation in this reaction was accomplished by utilizing methyl acetoacetate (4b) instead of ethyl acetoacetate (4a) for the synthesis of different Biginelli products. It is worth noting that all the reactions represented very good to excellent yields under solvent-free conditions in short time.

**The proposed mechanism for the synthesis of 3,4-dihydropyrimidin-2(1H)-one derivatives in the presence of Fe$_3$O$_4$@SiO$_2$-APTS-EDTA-asparagine nanocatalyst (1)**

The proposed mechanism based on the three-component strategy for the synthesis of 3,4-dihydropyrimidin-2(1H)-one derivatives catalyzed by Fe$_3$O$_4$@SiO$_2$-APTS-EDTA-asparagine nanocatalyst (1) is presented in Fig. 10. At first, the carbonyl group of aldehyde is activated by Fe$_3$O$_4$@SiO$_2$-APTS-EDTA-asparagine (1) to form intermediate (I) through the condensation with urea (2). Afterward, iminium intermediate (III) is produced after leaving the H$_2$O in the presence of the magnetic nanocatalyst. Meanwhile, intermediate (III) reacts with the enol form of the alkyl acetoacetate (4) and the corresponding intermediate (IV) is generated. Eventually, intramolecular cyclization occurs which is followed by dehydration of intermediate (V). At the end of the catalytic cycle, 3,4-dihydropyrimidin-2(1H)-ones are produced and the catalyst is recycled$^{60}$.

**Green chemistry metrics**

In this part of our research, green chemistry metrics for the synthesis of 3,4-dihydropyrimidin-2(1H)-one by Fe$_3$O$_4$@SiO$_2$-APTS-EDTA-asparagine nanocatalyst (1) were calculated and the results are summarized in Table 3$^{79,80}$. Hence, several parameters of the green approach such as environmental factor (E factor), process mass intensity, reaction mass efficiency, carbon efficiency, and atom economy were evaluated and compared to the ideal values$^{81}$. As presented in Table 3, all calculated values are close to the ideal values and were reported in supporting information.
Reusability of the Fe$_3$O$_4$@SiO$_2$-APTS-EDTA-asparagine nanocatalyst (1)

One of the critical scales in catalytic processes is reusability and recyclability of the catalyst. For evaluation of this parameter, the model reaction was examined using Fe$_3$O$_4$@SiO$_2$-APTS-EDTA-asparagine (1) for four runs. At the end of the reaction, the catalyst 1 was removed using an external magnet and the recycled catalyst was washed with dry toluene, dried and used in a subsequent model reaction. The obtained results are summarized in Fig. 11. Considering the results of isolated yields of products, the catalytic activity of Fe$_3$O$_4$@SiO$_2$-APTS-EDTA-asparagine nanocatalyst (1) after four runs is slightly reduced, which demonstrates proper conservancy of the catalytic activity after recycling.

Comparative study of Fe$_3$O$_4$@SiO$_2$-APTS-EDTA-asparagine nanocatalyst (1) and other catalysts for the Biginelli reaction

In order to compare the optimal catalytic activity and reaction conditions of the Fe$_3$O$_4$@SiO$_2$-APTS-EDTA-asparagine nanocatalyst (1) with previously reported catalysts for the three-component Biginelli reaction, we compared reaction conditions and yield of desired product (5a) in Table 4. As it can be observed from data in Table 4, all catalytic systems are capable of producing the desired product with satisfactory yields but Fe$_3$O$_4$@SiO$_2$-APTS-EDTA-asparagine nanocatalyst (1) in terms of yield and time factors, the reaction temperature, solvent and amount of catalyst loading demonstrates better performance than the other catalysts. Furthermore, additional advantage of this protocol is its easy separation from the crude products by using an external magnet compared to the most of reported heterogeneous catalytic systems.

Experimental

Chemicals and Instrumentation

Ferric chloride (FeCl$_3$.6H$_2$O), Ferrous chloride tetrahydrate (FeCl$_2$.4H$_2$O), (3-aminopropyl) triethoxysilane (APTS, 99%), tetraethyl orthosilicate (TEOS, 99%), ammonia (25 wt%), EDTA (MW = 292.24 g/mol), and asparagine (MW = 132.12 g/mol) were purchased from Merck and used without further purification. Urea, ethyl acetoacetate and aromatic aldehydes were purchased from international chemical companies including Merck and Sigma-Aldrich. The analytical TLC experiments were accomplished using Merck Kieselgel 60 F-254 Al-plates and then visualized by UV light and iodine vapour. Melting points of the products were measured on an Electrothermal 9100 apparatus and uncorrected. The functional groups of the samples were identified by FT-IR spectroscopy on a Perkin Elmer, Frontier FT-MIR spectrometer in the range of 600-4000 cm$^{-1}$ using KBr discs. The morphology of the nanocatalyst was observed by FESEM
TESCAN-MIRA3 and TEM Philips EM 208S. TGA curves of the \( \text{Fe}_3\text{O}_4@\text{SiO}_2\)-APTS-EDTA-asparagine (1) were recorded by Bahr company STA 504. X-ray diffraction (XRD) pattern of the catalyst 1 was taken by the Bruker D8 Advance device. The composition of the catalyst was determined by energy-dispersive X-ray (EDX) spectroscopy using a Numerix DXP-X10P instrument. Magnetization measurements were carried out on a BHV-55 vibrating sample magnetometer (VSM). \(^1\)H NMR spectra of the isolated products were recorded at 500 MHz using a Varian-INOVA spectrometer.

**General procedure for preparation of the magnetic \( \text{Fe}_3\text{O}_4 \) nanoparticles**

Preparation of \( \text{Fe}_3\text{O}_4 \) nanoparticles were according to a reported general method. In this procedure, in a 100 mL round-bottomed flask \( \text{FeCl}_3\cdot6\text{H}_2\text{O} \) (4.6 g, 0.017 mol) and \( \text{FeCl}_2\cdot4\text{H}_2\text{O} \) (2.3 g, 0.011 mol) were dissolved in deionized water (60 mL) and stirred for 30 min. Subsequently, aqueous \( \text{NH}_3 \) (10 mL) was added dropwise into the mixture and heated to 40°C under \( \text{N}_2 \) atmosphere for 2 h. The black solution was poured from the reaction vessel and \( \text{Fe}_3\text{O}_4 \) MNPs precipitates were separated from the solution using an external magnet, washed five times with deionized water and EtOH, and dried in the oven at 50°C for 24 h.

**General procedure for preparation of the silica-coated magnetic nanoparticles (\( \text{Fe}_3\text{O}_4@\text{SiO}_2 \))**

In accordance to the modified Stöber method, silica-coated \( \text{Fe}_3\text{O}_4 \) nanoparticles (\( \text{Fe}_3\text{O}_4@\text{SiO}_2 \)) were produced by a solvothermal reaction. For this purpose, the \( \text{Fe}_3\text{O}_4 \) MNPs (1.0 g) were dispersed in 30 mL of distilled water and ultrasonicated for 30 min. Then, a mixture of aqueous \( \text{NH}_3 \) (2 mL) and EtOH (40 mL) were added dropwise to the magnetite mixture and ultrasonicated for 30 min. Then, a mixture of TEOS (2 mL) and EtOH (40 mL) were added slowly to the suspension solution under continuous stirring for 24 h at 60°C. Eventually, the \( \text{Fe}_3\text{O}_4@\text{SiO}_2 \) core-shell MNPs were collected using an external magnet, washed with deionized water and EtOH and dried in the oven at 50°C for 5 h.

**Modification of the \( \text{Fe}_3\text{O}_4@\text{SiO}_2 \) NPs by (3-aminopropyl) triethoxysilane (\( \text{Fe}_3\text{O}_4@\text{SiO}_2\)-APTS)**

\( \text{Fe}_3\text{O}_4@\text{SiO}_2 \) core-shell MNPs were modified with (3-aminopropyl) triethoxysilane (APTS) using a typical modified method. Briefly, the \( \text{Fe}_3\text{O}_4@\text{SiO}_2 \) NPs (1.0 g) were ultrasonicated in 30 mL dried toluene. Subsequently, APTS (2.0 mL) was added to the magnetic mixture and stirred at 105°C for 24 h. After washing with dry toluene, the obtained MNPs separated and dried at 60°C for 12 h in a vacuum oven to prepare the \( \text{Fe}_3\text{O}_4@\text{SiO}_2\)-APTS MNPs.

**Preparation of the EDTA functionalized magnetic nanoparticles (\( \text{Fe}_3\text{O}_4@\text{SiO}_2\)-APTS-EDTA)**
In a round-bottom flask, magnetic Fe$_3$O$_4$@SiO$_2$-APTS NPs (1.0 g) were added to dry toluene (25 mL) and dispersed with ultrasonic for 15 min. Then, EDTA dianhydride (1.0 g) - synthesized according to Repo et al.$^{86}$ - and acetic anhydride added to the mixture and stirred at 80°C under N$_2$ atmosphere for 24 h. The magnetic Fe$_3$O$_4$@SiO$_2$-APTS-EDTA NPs were washed five times with EtOH followed by drying at 60°C for 6 h in a vacuum oven.

**Preparation of the asparagine grafted on the EDTA-modified Fe$_3$O$_4$@SiO$_2$ core-shell magnetic nanoparticles (Fe$_3$O$_4$@SiO$_2$-APTS-EDTA-asparagine, 1)**

In the last step, the magnetic Fe$_3$O$_4$@SiO$_2$-APTS-EDTA NPs were dispersed in 25 mL of dry toluene and asparagine (1.0 g) was added to the magnetic mixture and stirred under reflux conditions and N$_2$ atmosphere for 24 h. Magnetic precipitates were separated using an external magnet and after drying in the oven, brown powder of Fe$_3$O$_4$@SiO$_2$-APTS-EDTA-asparagine nanocatalyst (1) was obtained.

**General procedure for the synthesis of 3,4-dihydropyrimidin-2(1H)-one (5a-5t) catalyzed by Fe$_3$O$_4$@SiO$_2$-APTS-EDTA-asparagine nanomaterial (1)**

A mixture of urea (2, 1.0 mmol), aromatic aldehyde (3, 1.0 mmol), ethyl or methyl acetoacetate (4, 1.0 mmol), and Fe$_3$O$_4$@SiO$_2$-APTS-EDTA-asparagine (1, 10 mg) were added under solvent-free conditions for an appropriate time indicated in Table 2. After completion of the reaction, as monitored by TLC [eluent: n-hexane: EtOAc: 3:1], the catalyst was separated using an external magnet and the residue was concentrated to result in the crude product. Finally, the crude product was recrystallized from EtOH to obtain the pure product.

**Conclusion**

In summary, the novel and thermally stable asparagine grafted on EDTA-modified Fe$_3$O$_4$@SiO$_2$ core-shell magnetic nanoparticles (Fe$_3$O$_4$@SiO$_2$-APTS-EDTA-asparagine) was prepared for the first time. The Fe$_3$O$_4$@SiO$_2$-APTS-EDTA-asparagine heterogeneous nanocatalyst was used for highly efficient, facile, and green and sustainable synthesis of 3,4-dihydropyrimidin-2(1H)-one derivatives in a one-pot and three-component protocol through cyclocondensation of alkyl acetoacetate, urea and various aldehydes under solvent-free conditions. Consistency with the ideal values of green chemistry parameters, easy work up procedure, good to excellent yields in shorter reaction times, fast separation and recyclability of the catalyst are the additional advantages for its application in academic and industrial purposes.
Declarations

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**Tables**

**Table 1.** Optimization of conditions in the model reaction of urea (2), 4-chlorobenzaldehyde (3a), ethyl acetoacetate (4a), under different conditions in the presence of Fe3O4@SiO2-APTS-EDTA-asparagine nanocatalyst (1).

![Chemical structure](Eq. 1)
| Entry | Catalyst                                      | Solvent          | Temperature (°C) | Time (min) | Yield<sup>b</sup> (%) |
|-------|-----------------------------------------------|-------------------|------------------|------------|------------------------|
| 1     | -                                             | Solvent-free      | r.t              | 30         | Trace                  |
| 2     | Fe<sub>3</sub>O<sub>4</sub>@SiO<sub>2</sub>-APTS-EDTA-asparagine | Solvent-free      | r.t              | 30         | 85                     |
| 3     | Fe<sub>3</sub>O<sub>4</sub>@SiO<sub>2</sub>-APTS-EDTA-asparagine | Solvent-free      | 60               | 20         | 95                     |
| 4     | Fe<sub>3</sub>O<sub>4</sub>@SiO<sub>2</sub>-APTS-EDTA-asparagine | EtOH              | r.t              | 20         | 65                     |
| 5     | Fe<sub>3</sub>O<sub>4</sub>@SiO<sub>2</sub>-APTS-EDTA-asparagine | EtOH              | Reflux           | 20         | 85                     |
| 6     | Fe<sub>3</sub>O<sub>4</sub>@SiO<sub>2</sub>-APTS-EDTA-asparagine | MeOH              | r.t              | 40         | 45                     |
| 7     | Fe<sub>3</sub>O<sub>4</sub>@SiO<sub>2</sub>-APTS-EDTA-asparagine | MeOH              | Reflux           | 40         | 55                     |
| 8     | Fe<sub>3</sub>O<sub>4</sub>@SiO<sub>2</sub>-APTS-EDTA-asparagine | H<sub>2</sub>O    | r.t              | 30         | 60                     |
| 9     | Fe<sub>3</sub>O<sub>4</sub>@SiO<sub>2</sub>-APTS-EDTA-asparagine | H<sub>2</sub>O    | Reflux           | 30         | 65                     |
| 10    | Fe<sub>3</sub>O<sub>4</sub>@SiO<sub>2</sub>-APTS-EDTA-asparagine | EtOH/H<sub>2</sub>O (1:1) | r.t              | 20         | 63                     |
| 11    | Fe<sub>3</sub>O<sub>4</sub>@SiO<sub>2</sub>-APTS-EDTA-asparagine | EtOH/H<sub>2</sub>O (1:1) | Reflux           | 20         | 70                     |
| 12    | Fe<sub>3</sub>O<sub>4</sub>@SiO<sub>2</sub>-APTS-EDTA-asparagine | DMF               | r.t              | 30         | 55                     |
| 13    | Fe<sub>3</sub>O<sub>4</sub>@SiO<sub>2</sub>-APTS-EDTA-asparagine | DMF               | Reflux           | 30         | 65                     |
| 14    | EDTA                                          | Solvent-free      | 60               | 20         | 75                     |
| 15    | Asparagine                                    | Solvent-free      | 60               | 20         | 65                     |

<sup>a</sup>Reaction conditions: urea (2, 1 mmol), 4-chlorobenzaldehyde (3a, 1 mmol), ethyl acetoacetate (4a, 1 mmol), Fe<sub>3</sub>O<sub>4</sub>@SiO<sub>2</sub>-APTS-EDTA-asparagine (1) and solvent (3 mL, if not otherwise stated). <sup>b</sup>Isolated yield.

Due to technical limitations, table 2 is only available as a download in the Supplemental Files section.
Table 3. Measurement of green chemistry metrics for compound 5a

| Entry | Parameters of the green approach | Ideal value | Calculated values |
|-------|----------------------------------|-------------|------------------|
| 1     | E factor                         | 0           | 0.16             |
| 2     | Atom economy (AE %)              | 100%        | 89.1%            |
| 3     | Carbon efficiency (CE %)         | 100%        | 96%              |
| 4     | Process mass intensity (PMI)     | 1           | 1.16             |
| 5     | Reaction mass efficiency (RME %) | 100%        | 85.5%            |

Table 4. Comparative results of catalysts for the synthesis of 5a

| Entry | Catalyst                                      | Catalyst loading | Reaction conditions | Time (min) | Yield (%) |
|-------|-----------------------------------------------|------------------|---------------------|------------|-----------|
| 1     | Zn(II)-framework                              | 10 wt %          | Solvent-free/60 °C  | 120        | 91        |
| 2     | PANI-FeCl₃                                    | 200 mg           | CH₃CN/Reflux        | 1440       | 83        |
| 3     | MCM-41-APS-PMDANHSO₃H                         | 15 mg            | Solvent-free/80 °C  | 35         | 96        |
| 4     | Fe₃O₄@SiO₂-APTM-Fe(OH)₂                       | 10 mg            | Neat/80 °C          | 15         | 95        |
| 5     | zirconium (IV)-salophen perfluoroctanesulfonate | 0.05 mmol      | Solvent-free/90 °C  | 30         | 96        |
| 6     | Fe₃O₄@SiO₂-APTS-EDTA-asparagine               | 10 mg            | Solvent-free/60 °C  | 20         | 95        |

Figures
Figure 1

Schematic preparation of Fe3O4@SiO2-APTS-EDTA-asparagine (1), as a heterogeneous nanocatalyst, for the synthesis of 3,4-dihydropyrimidin-2(1H)-one 5 derivatives.
Figure 2

FT-IR spectra of the Fe3O4 (a), Fe3O4@SiO2 (b), Fe3O4@SiO2-APTS (c), Fe3O4@SiO2-APTS-EDTA (d) and Fe3O4@SiO2-APTS-EDTA-asparagine (1, e).
Figure 3

The EDX spectra of the magnetic Fe3O4@SiO2-APTS-EDTA-asparagine nanomaterial (1).
Figure 4

FESEM images of the magnetic Fe3O4@SiO2-APTS-EDTA-asparagine nanocatalyst (1).
Figure 5

TEM images of the magnetic Fe3O4@SiO2-APTS-EDTA-asparagine nanomaterial (1).
Figure 6

XRD patterns of the magnetic Fe₃O₄@SiO₂-APTS-EDTA-asparagine nanocatalyst (1).
Figure 7

VSM pattern of Fe3O4 (red curve) and magnetic Fe3O4@SiO2-APTS-EDTA-asparagine nanocatalyst (1, green curve).

Figure 8

TGA curve of the magnetic Fe3O4@SiO2-APTS-EDTA-asparagine nanomaterial (1).
Figure 9

Effect of solvent and the amount of Fe3O4@SiO2-APTS-EDTA-asparagine nanocatalyst (1) on the model reaction.
Figure 10

The proposed mechanism for the synthesis of 3,4-dihydropyrimidin-2(1H)-one derivatives using ethyl acetoacetate or methyl acetoacetate in the presence of Fe3O4@SiO2-APTS-EDTA-asparagine nanocatalyst (1).
Figure 11

Reusability of the Fe3O4@SiO2-APTS-EDTA-asparagine nanocatalyst (1) for the synthesis of 5a.

Supplementary Files

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