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A Schiff base complex of lanthanum on modified MCM-41 as a reusable nanocatalyst in the homoselective synthesis of 5-substituted 1*H*-tetrazoles†

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In this work, mesoporous MCM-41 was modified by a new Schiff-base formed from the condensation of triethylenetetramine and 5-bromosalicylaldehyde. Then, it was used for the stabilization of lanthanum metal (La-Schiff base@MCM-41) as a homoselective, reusable, efficient and biocompatible catalyst in the synthesis of 5-substituted 1*H*-tetrazole derivatives. The synthesized tetrazoles were characterized using ¹H NMR and FT-IR spectroscopy and methods to measure their physical properties. La-Schiff base@MCM-41 was characterized by using various techniques such as ICP, CHN, XRD, TGA, BET, FT-IR spectroscopy, SEM, EDS and WDX. This catalyst has good stability and a heterogeneous nature, enabling it to be easily recovered and reused several times without significant loss in catalytic activity. This present strategy has important advantages such as utilizing PEG as a green solvent, short reaction times, excellent yields, easy recycling of the catalyst and pure separation of the products. The recovered La-Schiff base@MCM-41 catalyst was characterized by using FT-IR spectroscopy, SEM and AAS.

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1 Introduction

Catalyst reusability is one of the principles of green chemistry and can increase the rate of a chemical process.^{1–6} Therefore, separation of used catalysts is one of the main challenges for chemists. On the other hand, a suitable, biocompatible and economical catalyst should have several properties such as simplicity of preparation, high catalytic activity, good selectivity, good stability, easy separation, and excellent reusability. The problems of recovering homogeneous catalysts and the low catalytic activity or low selectivity of heterogeneous catalysts have limited their application in industry and other sciences.^{3,7–9} To overcome these problems, nanomaterials or catalysts immobilized on nanomaterials seem to be ideal catalysts because nanomaterials have stability and a heterogeneous nature, enabling them to be easily recovered and reused like heterogeneous catalysts, and they also have high surface areas that can improve catalytic activity and selectivity in chemical procedures.^{5,10} Therefore, nanocatalysts exist at the border between heterogeneous and homogeneous catalysts because they have advantages from both, such as efficiency, selectivity and reusability.^{5,10,11} For example, various nanomaterials such

as nanopolymers,¹² carbon nanotubes,^{13–15} mesoporous silica,^{16–18} boehmite nanoparticles,^{19–22} graphene oxide nanosheets,²³ biochar nanoparticles,^{24,25} magnetic nanoparticles,^{7–9,26,27} metal–organic frameworks²⁸ *etc.* have been employed in chemistry and especially in catalysis applications. Amongst the nanomaterials, macroporous or mesoporous nanomaterials especially MCM-41 have been widely used as an ideal support for immobilization of various catalysts and other applications.^{29,30} MCM-41 has also been used in other fields such as drug delivery,^{31,32} extraction,³³ adsorption,^{34,35} sensors,^{36,37} supports for catalysts,^{38–40} and energy.⁴¹ This is because MCM-41 has unique properties, *e.g.* excellent thermal and chemical stability, high surface area (>1000 m² g^{−1}), easy surface functionalization, homogeneous hexagonal channel structure (1.5–10 nm pore diameters), large pore volumes (up to 1.3 ml g^{−1}), good biocompatibility and easy separation from the reaction mixture.^{38–43} The large specific surface area of MCM-41 leads to a high capacity of catalyst loading. Also, the high thermal and chemical stability of MCM-41 allows the application of MCM-41 under harsh conditions and various chemical conditions. Moreover, the large pore volume of MCM-41 allows the application of MCM-41 in the immobilization of organic ligands and metal complexes into its channels.³⁸ Therefore, we investigated a new Schiff-base complex of lanthanum-catalyst on MCM-41 (La-Schiff base@MCM-41) as an efficient, stable and recyclable nanocatalyst in the homoselective synthesis of 5-substituted 1*H*-tetrazole derivatives, because tetrazole

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derivatives have biological activity and they are also used as herbicides, analgesics, anti-HIV drug candidates, and antimicrobial, anti-inflammatory, anti-proliferative, and anticancer agents.^{44–52} For example, Candesartan, Irbesartan, Valsartan, Cilostazol, Losartan, Pranlukast and Pemioplast are several pharmacologically important tetrazole derivatives.^{53–55}

2 Experimental

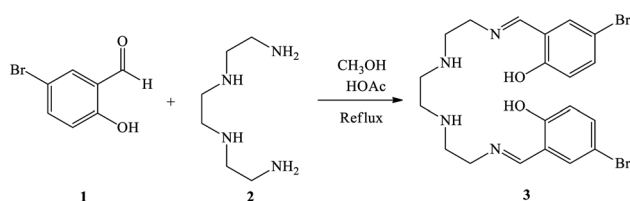
2.1 Materials and instruments

The chemical compounds and solvents for synthesis of the catalyst and tetrazoles used in this study were purchased from Merck, Aldrich, Fluka or Iranian companies and used without any purification.

The size and morphology of the MCM-41, nanocatalyst and recovered catalyst were studied using FESEM imaging using a MIRA3TESCAN-XMU or FESEM-TESCAN MIRA3 Scanning Electron Microscope. Moreover, this Scanning Electron Microscope was employed to determine the elemental composition (EDS and WDX) of the nanocatalyst. Besides, the exact concentration of lanthanum in the catalysts was measured by an inductively coupled plasma instrument (ICP analysis) model ELAN 6100 DRC-e from PerkinElmer Company and AAS using a 400p-novAA instrument from Analytik Jena Company. The concentration of carbon, hydrogen and nitrogen in the nanocatalyst was measured by an elemental analyzer (CHN analysis) from the British company CQOSTECH. The TGA diagram of the nanocatalyst was recorded by a NETZSCH STA 449F3 Thermal Analysis device under air in the temperature range of 30–800 °C. Powder XRD patterns of the MCM-41 and nanocatalyst were recorded with CuK α radiation at 40 kV and 30 mA by a PW1730 instrument from Philips Company. FT-IR spectra of the nanomaterials and tetrazoles were recorded in KBr pellets using a VRTEX70 model Bruker FT-IR spectrometer. Nitrogen adsorption isotherms of MCM-41 and the nanocatalyst were obtained using a BELSORP MINI II device by a standard gas manifold at 77 K. In addition, the nanomaterial samples were degassed using a BEL PREP VAC II device before analysis at 120 °C for 2 h. NMR spectra of the tetrazoles were recorded using a Bruker DRX-400 spectrometer at 100–400 MHz. Melting points of the tetrazoles were obtained with an Electrothermal 9100 instrument.

2.2 Synthesis of the Schiff base ligand

First, 2,5,8,11-tetraazadodeca-(1,11-diene-1,12-diyl)bis(4-bromophenol) as a new Schiff base ligand (3) was prepared by



Scheme 1 Synthesis of 2,5,8,11-tetraazadodeca-(1,11-diene-1,12-diyl)bis(4-bromophenol) as the Schiff base ligand (3).

the condensation of 5-bromosalicylaldehyde (1) with triethylenetetramine (2) according to the illustrated procedure in Scheme 1. In this regard, a solution of triethylenetetramine (1 ml, 6.7 mmol) in methanol (30 ml) was added drop-wise to a solution of 5-bromosalicylaldehyde (2.693 g, 13.4 mmol) in methanol (20 ml) under stirring. Glacial acetic acid (4 drops) was also added to the reaction mixture and refluxed for 4 h. The resulting yellow precipitate was filtered off, washed with methanol, and dried at room temperature.

2.3 Preparation of the catalyst

The modified MCM-41 by (3-chloropropyl)triethoxysilane (3-CPTMS@MCM-41) was synthesized according to the reported procedure in the literature.⁴⁰ To achieve immobilization of the Schiff base ligand (3) on MCM-41 (Schiff base@MCM-41), 1 g of 3-CPTMS@MCM-41 was refluxed with 3 (1 mmol) in toluene for 40 h. The obtained Schiff base@MCM-41 was isolated by simple filtration, washed with DMSO and hot ethanol and dried at room temperature. Finally, 1 g of Schiff base@MCM-41 was dispersed in ethanol, and then lanthanum nitrate (1 mmol) was added to the mixture and stirred for 24 h under reflux conditions. The obtained catalyst (La-Schiff base@MCM-41) was filtered, washed and dried at room temperature (Scheme 2).

2.4 General method for the synthesis of 5-substituted 1H-tetrazoles catalyzed by La-Schiff base@MCM-41

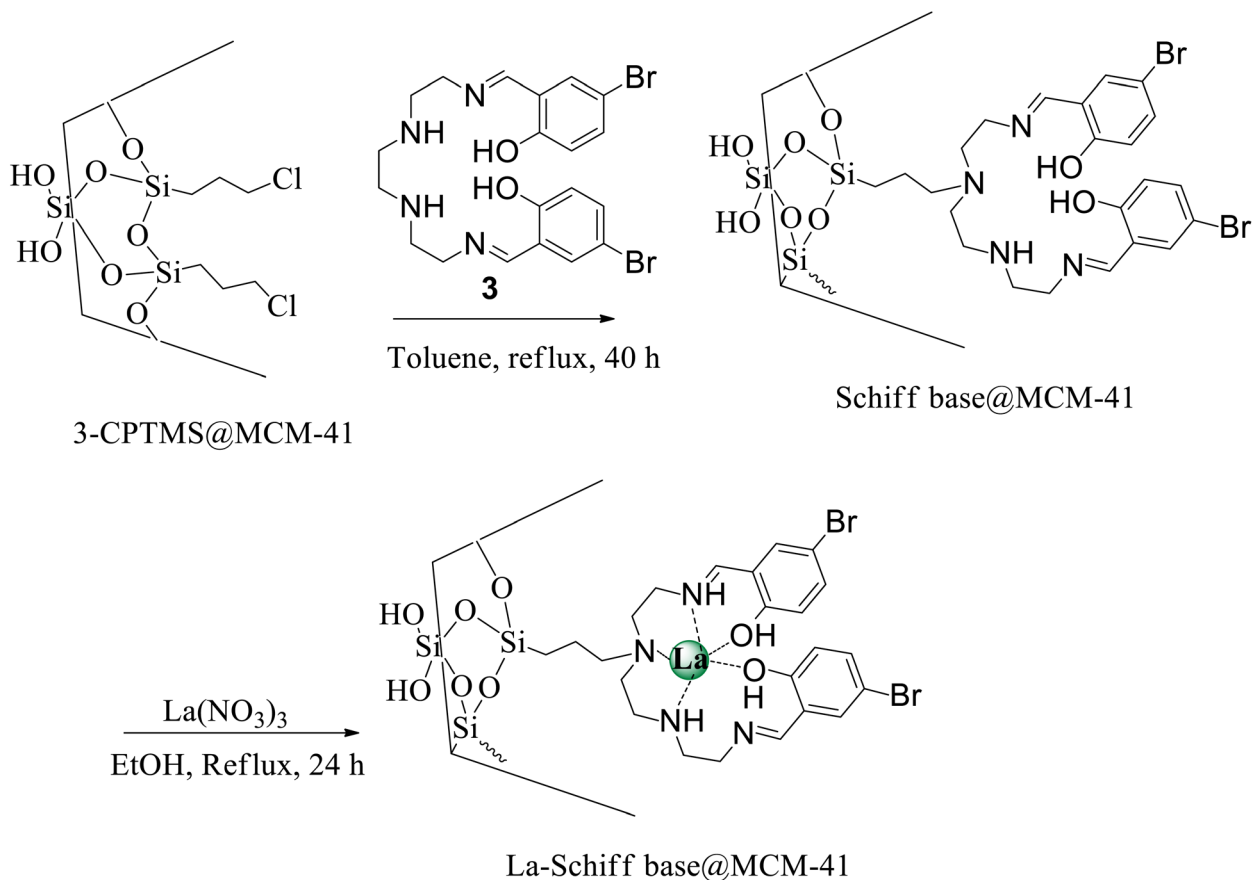
The [3 + 2] cycloaddition of sodium azide salt (NaN₃) with organic nitrile compounds was selected for the synthesis of heterocyclic tetrazoles in the presence of La-Schiff base@MCM-41 as catalyst. In this process, NaN₃ (1.4 mmol) and nitrile compounds (1 mmol) were stirred in the presence of 50 mg of La-Schiff base@MCM-41 in PEG-400 at 120 °C. The reaction was monitored by using TLC. At the end of the reaction, the mixture was cooled down to room temperature. Then, the reaction mixture was diluted by water and ethyl acetate and the La-Schiff base@MCM-41 catalyst was separated using simple filtration. Finally, 10 ml of HCl (4 N) was added to the solution and the tetrazole products were extracted from ethyl acetate. The organic solvent was evaporated and the products were dried using anhydrous sodium sulfate (Scheme 3). The tetrazole products were obtained with yields of 89–98% and all obtained tetrazoles were confirmed by ¹H NMR and FT-IR spectroscopy and physical property measurement such as melting point.

In order to show the reproducibility and reusability of the La-Schiff base@MCM-41 catalyst, the [3 + 2] cycloaddition of NaN₃ with benzonitrile was repeated under optimized conditions. After completion of the reaction, the catalyst was recovered by centrifugation, washed with water and ethyl acetate and further employed in the next cycle. The La-Schiff base@MCM-41 catalyst showed good reproducibility and can be reused up to 6 times.

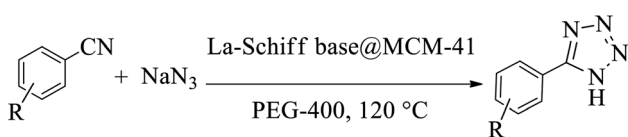
2.5 Spectral data

2.5.1 5-(4-Bromophenyl)-1H-tetrazole. ¹H NMR (400 MHz, DMSO): δ_{H} = 7.98–7.95 (d, J = 12 Hz, 2H), 7.83–7.81 (d, J = 12 Hz, 2H) ppm.





Scheme 2 Synthesis of La-Schiff base@MCM-41.



Scheme 3 General method for the synthesis of 5-substituted 1H-tetrazoles in the presence of La-Schiff base@MCM-41.

IR (KBr) cm^{-1} : 3423, 3090, 3064, 2966, 2852, 2768, 1638, 1611, 1559, 1482, 1431, 1385, 1276, 1252, 1157, 1119, 1054, 1018, 988, 872, 828, 742, 614, 451.

2.5.2 5-(3-Nitrophenyl)-1H-tetrazole. ^1H NMR (400 MHz, DMSO): δ_{H} = 18.44 (br, 1H), 8.82 (s, 1H), 8.48–8.45 (d, J = 12 Hz, 1H), 8.43–8.39 (d of q, J (d) = 12 Hz, J (q) = 2 Hz, 1H), 7.93–7.87 (t, J = 12 Hz, 1H) ppm.

2.5.3 2-(1H-tetrazol-5-yl)benzonitrile. ^1H NMR (400 MHz, DMSO): δ_{H} = 8.10–8.06 (t, J = 8 Hz, 2H), 7.95–7.89 (t, J = 12 Hz, 1H), 7.80–7.75 (t, J = 12 Hz, 1H) ppm.

IR (KBr) cm^{-1} : 3426, 3082, 3029, 2922, 2582, 2721, 2615, 2229, 2005, 1735, 1638, 1609, 1581, 1493, 1454, 1408, 1384, 1280, 1242, 1199, 1167, 1122, 1100, 1066, 1047, 1113, 999, 970, 856, 783, 756, 724, 705, 665, 585, 555, 518, 500.

2.5.4 5-(4-Nitrophenyl)-1H-tetrazole. ^1H NMR (400 MHz, DMSO): δ_{H} = 16.55 (br, 1H), 8.46–8.43 (d, J = 12 Hz, 2H), 8.31–8.28 (d, J = 12 Hz, 2H) ppm.

IR (KBr) cm^{-1} : 3212, 3069, 2903, 1957, 1822, 1717, 1604, 1512, 1337, 1108, 1059, 991, 857, 773, 691, 491, 443.

2.5.5 5-(4-Chlorophenyl)-1H-tetrazole. ^1H NMR (400 MHz, DMSO): δ_{H} = 16.99 (br, 1H), 8.06–8.03 (d, J = 12 Hz, 2H), 7.71–7.68 (d, J = 12 Hz, 2H) ppm.

IR (KBr) cm^{-1} : 3420, 3096, 3070, 2922, 2852, 2733, 1638, 1611, 1487, 1458, 1434, 1385, 1277, 1254, 1160, 1121, 1096, 1052, 1019, 988, 878, 829, 743, 709, 693, 623, 506, 464.

2.5.6 2-(1H-Tetrazol-5-yl)phenol. ^1H NMR (400 MHz, DMSO): δ_{H} = 8.00–7.98 (d, J = 12 Hz, 1H), 7.43–7.38 (t, J = 8 Hz, 1H), 7.08–7.06 (d, J = 8 Hz, 1H), 7.03–6.97 (t, J = 12 Hz, 1H) ppm.

IR (KBr) cm^{-1} : 3431, 3252, 3062, 2930, 2686, 2567, 1737, 1610, 1546, 1477, 1391, 1356, 1295, 1232, 1156, 1112, 1086, 1002, 941, 809, 741, 542.

2.5.7 5-(2-Chlorophenyl)-1H-tetrazole. ^1H NMR (400 MHz, DMSO): δ_{H} = 16.95 (br, 1H), 7.82–7.80 (d, J = 8 Hz, 1H), 7.73–7.70 (d, J = 12 Hz, 1H), 7.66–7.61 (t, J = 8 Hz, 1H), 7.58–7.54 (t, J = 8 Hz, 1H) ppm.

IR (KBr) cm^{-1} : 3418, 2823, 2708, 1660, 1651, 1634, 1602, 1565, 1556, 1539, 1471, 1441, 1409, 1385, 1369, 1243, 1163, 1129, 1076, 1060, 1039, 1020, 1007, 988, 943, 874, 777, 747, 732, 651, 486, 453, 434.

2.5.8 4-(1H-Tetrazol-5-yl)phenol. ^1H NMR (400 MHz, DMSO): δ_{H} = 16.51 (br, 1H), 10.15 (br, 1H), 7.87–7.84 (d, J = 12 Hz, 2H), 6.96–6.93 (d, J = 12 Hz, 1H) ppm.



2.5.9 5-Phenyl-1H-tetrazole. ^1H NMR (400 MHz, DMSO): $\delta_{\text{H}} = 16.88$ (br, 1H), 8.06–8.03 (m, 2H), 7.63–7.59 (m, 3H) ppm.

IR (KBr) cm^{-1} : 3423, 3055, 2985, 2901, 2832, 1813, 1638, 1611, 1562, 1486, 1466, 1411, 1384, 1288, 1256, 1163, 18084, 1057, 1034, 1015, 991, 959, 925, 784, 726, 703, 687, 619, 493, 462.

3 Results and discussion

3.1 Characterization of the catalyst

In the first step, MCM-41 with its surface modified by 3-chloropropyltriethoxysilane was obtained based on a new reported procedure.⁴⁰ Subsequently, 2,5,8,11-(tetraazadodeca-1,11-diene-1,12-diyl)bis(4-bromophenol), a new Schiff-base (ligand 3), complexed with lanthanum was stabilized on the chloro-

modified MCM-41 nanoparticles (La-Schiff base@MCM-41) as an efficient and reusable nanocatalyst in the homoselective synthesis of 5-substituted 1H-tetrazole derivatives. This is the first report of immobilization of 2,5,8,11-tetraazadodeca-1,11-diene-1,12-diyl)bis(4-bromophenol) on MCM-41. Also, this is the first report where a lanthanum complex of 2,5,8,11-tetraazadodeca-1,11-diene-1,12-diyl)bis(4-bromophenol) was used as a catalyst in the synthesis of organic compounds. Therefore, this catalyst could be effective for other organic condensation or cycloaddition reactions. A FESEM-TESCAN MIRA III Scanning Electron Microscope instrument was employed to study the size and particle morphology of the MCM-41 and La-Schiff base@MCM-41 catalyst (Fig. 1). The obtained images show that MCM-41 and La-Schiff base@MCM-

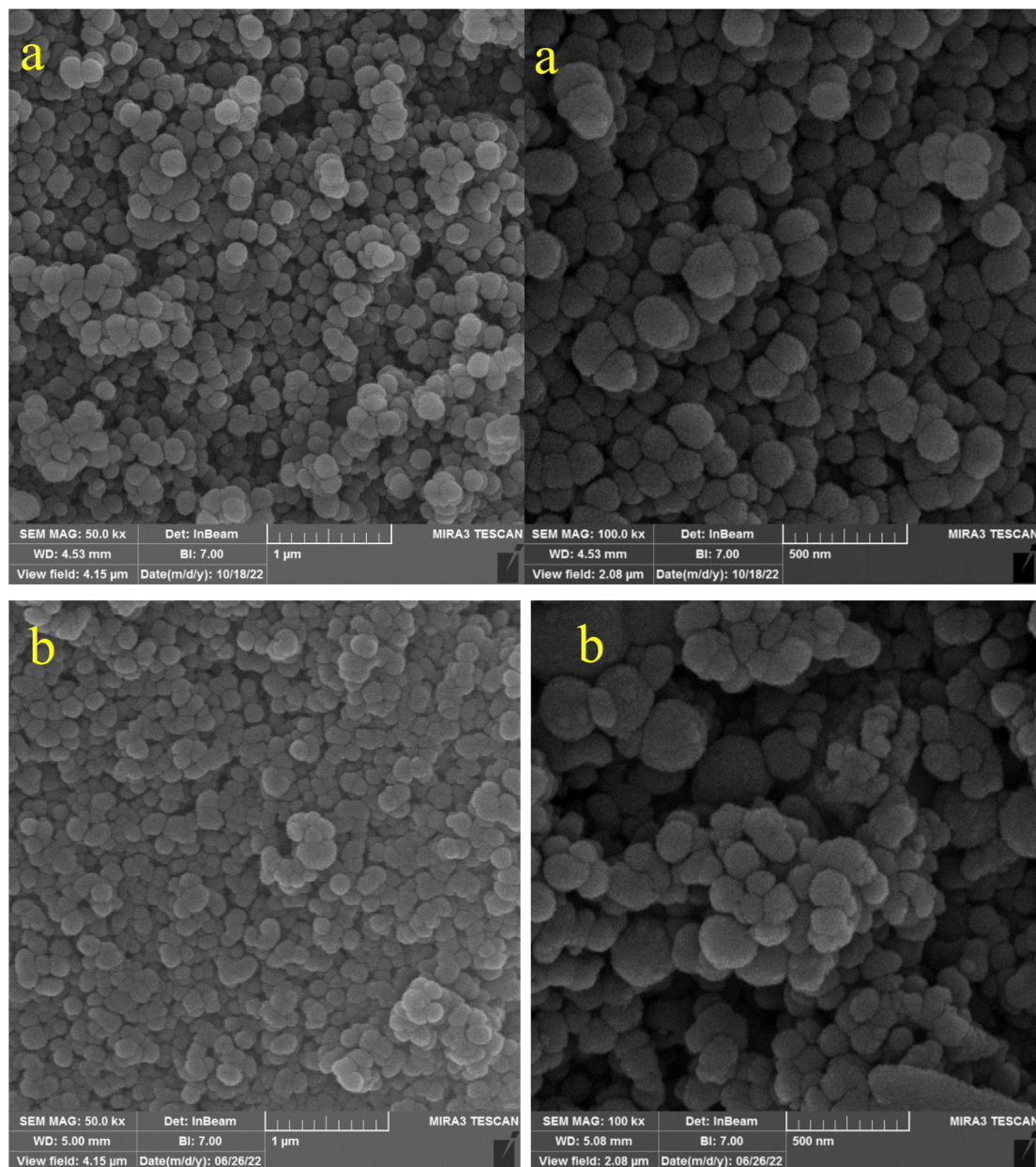


Fig. 1 SEM images of (a) MCM-41 and (b) La-Schiff base@MCM-41 catalyst.



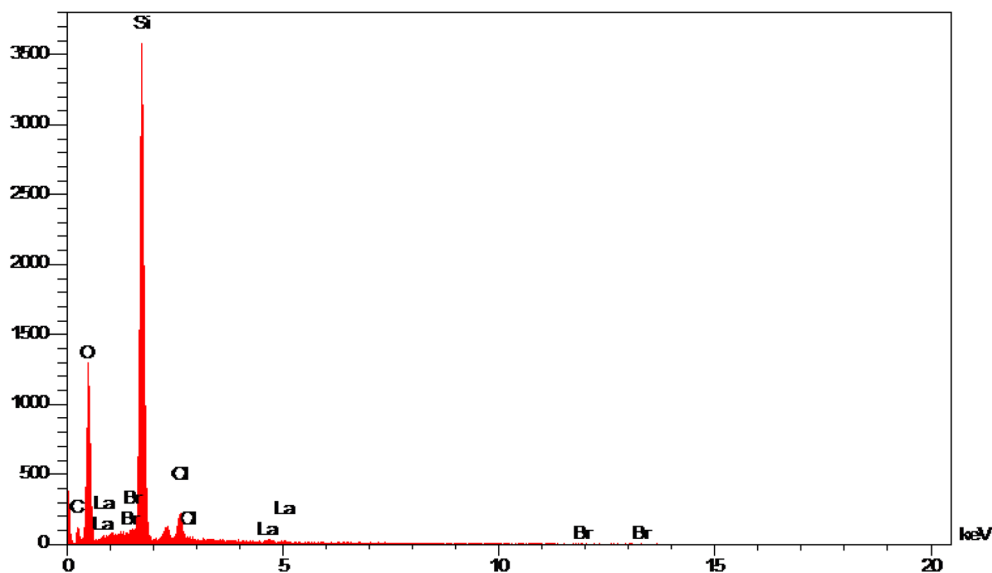


Fig. 2 EDS spectrum of the La-Schiff base@MCM-41 catalyst.

41 formed as uniform spherical shaped particles with quite homogeneous diameters of less than 100 nm. As can be seen in the SEM images, there is no significant difference in the shape and size of the MCM-41 particles and the catalyst, which indicates that the nanoparticles are stable during the modification.

The qualitative elemental composition analysis of the La-Schiff base@MCM-41 catalyst was performed using energy-dispersive X-ray spectroscopy (EDS) using a FESEM-TESCAN MIRA III Scanning Electron Microscope instrument (Fig. 2). The EDS diagram contains the special energy (keV) of each element in terms of intensity. A quantitative increase in the amount of each element leads to an increase in the intensity of

its peak in the EDS diagram. As shown in the EDS diagram of the La-Schiff base@MCM-41 catalyst, silicon and oxygen elements have the highest intensity, which are related to the skeleton of MCM-41. Also, the EDS diagram shows the presence of C, N and Br elements in the structure of the La-Schiff base@MCM-41 catalyst, which shows that the Schiff-base ligand is well stabilized on the MCM-41. Also, the EDS diagram shows La element in the structure of the La-Schiff base@MCM-41 catalyst, which indicates the successful stabilization of the lanthanum Schiff-base complex on the modified MCM-41. Furthermore, the EDS diagram of the La-Schiff base@MCM-41 catalyst shows the presence of chlorine

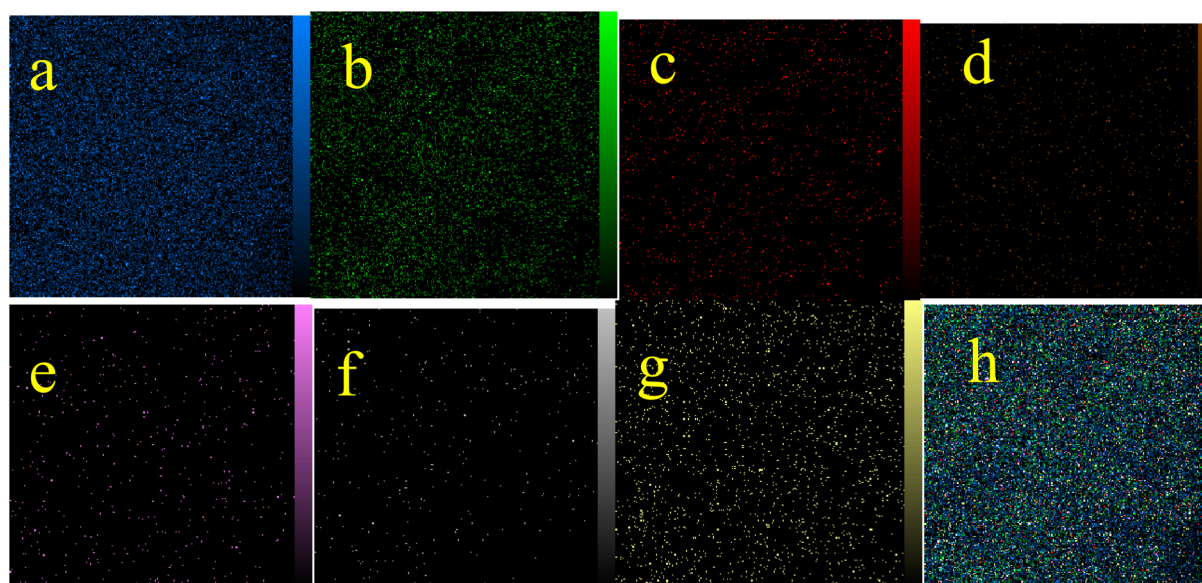


Fig. 3 Elemental mapping of (a) Si, (b) O, (c) C, (d) Br, (e) La, (f) N, and (g) Cl and (h) the combination of all elements for the La-Schiff base@MCM-41 catalyst.



element in its structure, which is related to some chlorine in 3-CPTMS@MCM-41 that was not replaced by the Schiff-base ligand **3**. Also, the results of elemental composition of the La-Schiff base@MCM-41 catalyst from EDS analysis were confirmed by wavelength dispersive X-ray spectroscopy (WDX) analysis, which is shown in Fig. 3. The obtained results from WDX clearly show a homogeneous distribution of Si, O, C, N, Br, Cl and La in the structure of the La-Schiff base@MCM-41 catalyst. The results of WDX, like EDS, show that silicon and oxygen are the most common elements of the La-Schiff base@MCM-41 catalyst.

Considering that lanthanum metal is the active site of the La-Schiff base@MCM-41 catalyst, the exact amount of lanthanum element in La-Schiff base@MCM-41 was obtained by ICP

analysis, and the value was $0.23 \times 10^{-3} \text{ mmol g}^{-1}$. Also, the content of carbon, hydrogen and nitrogen in La-Schiff base@MCM-41 was obtained using CHN analysis. The obtained results from CHN showed that this catalyst contains 9.17% carbon, 2.52% hydrogen and 0.22% nitrogen.

Volatile components can be measured by mass loss in TGA. Therefore, TGA can be employed to determine the organic and inorganic component ratio in a sample. For this, TGA was performed by gradually raising the temperature of the La-Schiff base@MCM-41 catalyst (Fig. 4). The TGA diagram of the La-Schiff base@MCM-41 catalyst indicated a small mass loss (3% weight) below 150 °C, which is attributed to the evaporation of adsorbed solvents.⁵⁶ Also, the TGA diagram of the La-Schiff base@MCM-41 catalyst indicated a notable mass loss (24%

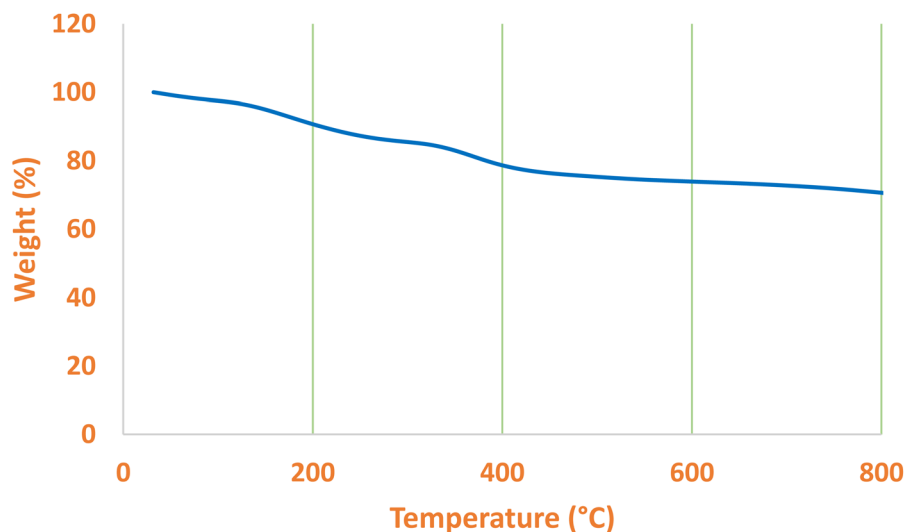


Fig. 4 TGA diagram of La-Schiff base@MCM-41 catalyst.

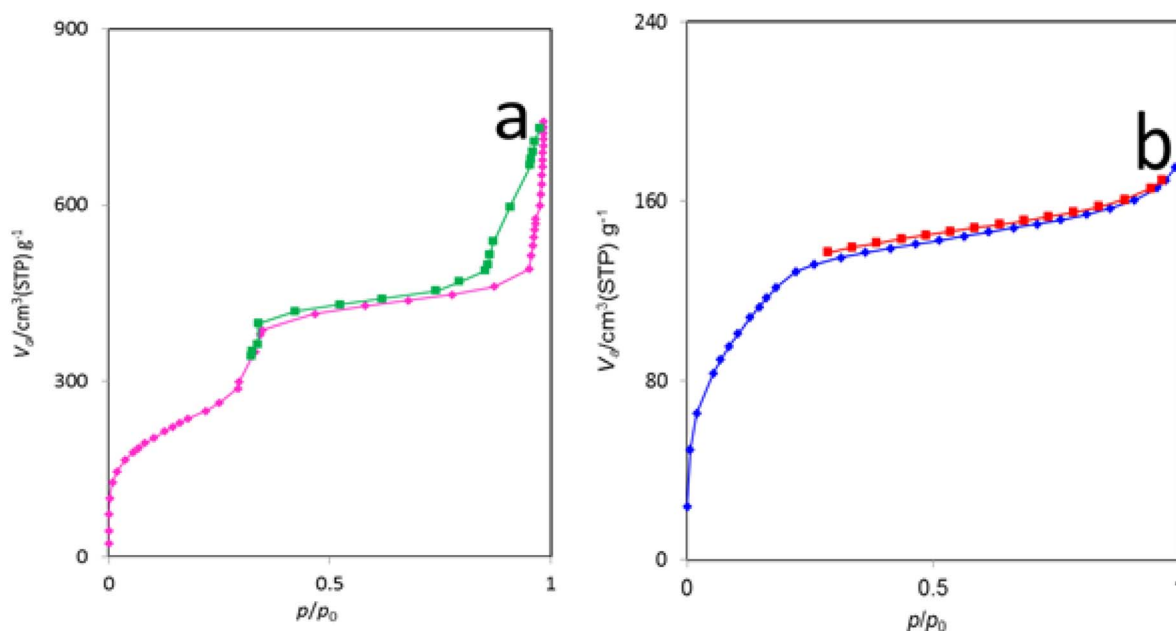


Fig. 5 N₂ adsorption–desorption isotherms of (a) MCM-41 and (b) La-Schiff base@MCM-41 catalyst.



Table 1 Textural properties of MCM-41 and La-Schiff base@MCM-41 catalyst

Entry	Sample	S_{BET} ($\text{m}^2 \text{g}^{-1}$)	Pore diameter (nm)	Pore volume ($\text{cm}^3 \text{g}^{-1}$)
1	MCM-41	854.13	5.37	1.15
2	La-Schiff base@MCM-41	452.58	2.381	0.269

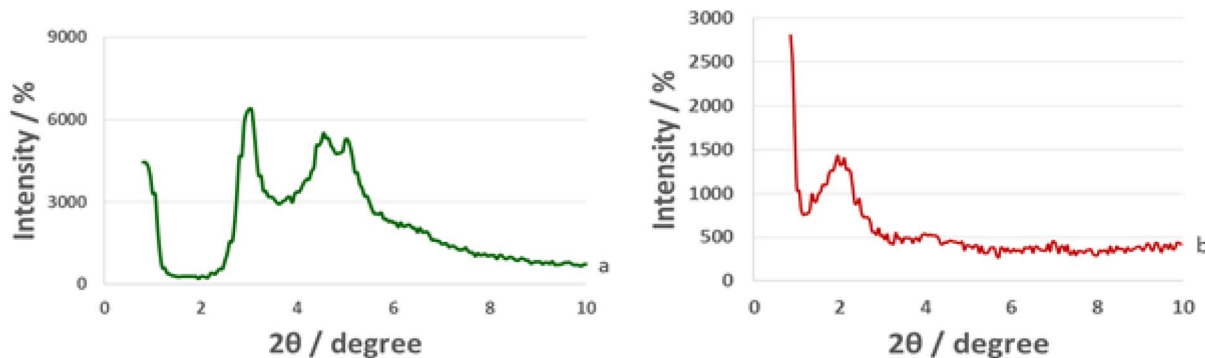


Fig. 6 Low angle XRD patterns of (a) MCM-41 and (b) La-Schiff base@MCM-41 catalyst.

weight) in the region of 150 °C–600 °C, which represented the decomposition of immobilized Schiff-base organic components on the surface of MCM-41.⁵⁷ Except for the evaporation of surface solvents, no weight loss was observed up to 160 °C, so this catalyst has thermal stability at least up to 160 °C.

The N_2 adsorption–desorption isotherms of MCM-41 and the La-Schiff base@MCM-41 catalyst are shown in Fig. 5. Based on the IUPAC classification, this catalyst displays a type IV isotherm, which represents the class of mesoporous materials.^{18,58} The BET surface area of La-Schiff base@MCM-41 is $452.58 \text{ m}^2 \text{g}^{-1}$ (Table 1, entry 2). Due to its high surface area, it can be used as a catalyst. Also, the average pore diameter and pore volume of La-Schiff base@MCM-41 were calculated by the BET method using N_2 adsorption–desorption and are 2.38 nm and $0.269 \text{ cm}^3 \text{g}^{-1}$, respectively (Table 1, entry 2). The BET surface area, average pore diameter and pore volume of MCM-41 are $854.13 \text{ m}^2 \text{g}^{-1}$, 5.37 nm and $1.15 \text{ cm}^3 \text{g}^{-1}$, respectively (Table 1, entry 1). The BET surface area, average pore diameter

and pore volume of La-Schiff base@MCM-41 are lower than those of the unmodified mesoporous MCM-41 due to the grafting of the Schiff-base complex on MCM-41 (Table 1).^{38,59} The reason for the decrease in BET surface area, average pore diameter and pore volume is that when organic groups and metal complexes are stabilized inside the pores of MCM-41, the pores of MCM-41 are filled, therefore the volume of the pores, diameter of the pores and the surface area are also reduced.

The small angle XRD patterns of MCM-41 and La-Schiff base@MCM-41 are shown in Fig. 6. The low angle XRD pattern of unfunctionalized MCM-41 has three peaks at 2θ values of about 2.95° (related to 100 reflections), 4.55° (related to 110 reflections) and 4.05° (related to 200 reflections), which represent the ordered hexagonal channels of MCM-41.⁴⁰ As is well known, functionalization and modifying the MCM-41 surface can reduce or fade the intensity of these peaks. As predicted, these peaks are present in the low angle XRD pattern

Table 2 Study to find the best reaction conditions for the synthesis of 5-substituted 1H-tetrazoles in the presence of the La-Schiff base@MCM-41 nanocatalyst

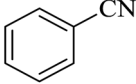
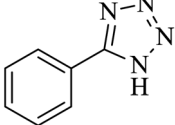
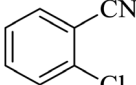
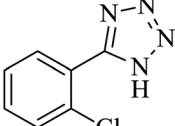
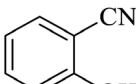
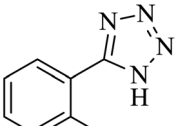
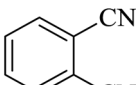
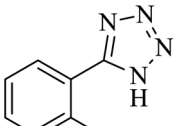
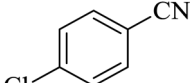
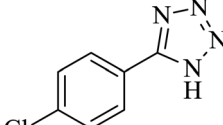
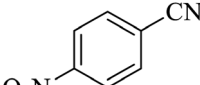
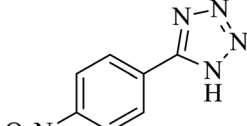
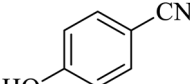
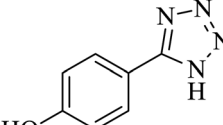
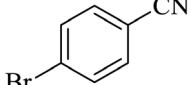
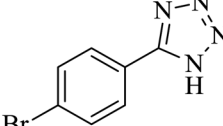
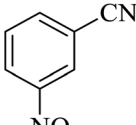
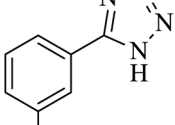
Entry	Amount of catalyst (mg)	Solvent	NaN_3 (mmol)	Time (min)	Temperature (°C)	Yield ^a (%)
1	—	PEG	1.4	140	120	NR
2	40	PEG	1.4	270	120	90
3	50	PEG	1.4	120	120	98
4	50	PEG	1.3	150	120	75
5	50	DMSO	1.4	120	120	68
6	50	H_2O	1.4	120	Reflux	55
7	50	<i>n</i> -Hexane	1.4	120	Reflux	Trace
8	50	Dichloromethane	1.4	120	Reflux	Trace
9	50	PEG	1.4	120	100	42

^a Isolated yield.



of La-Schiff base@MCM-41 with a lower intensity than those of the un-functionalized MCM-41, which is due to the grafting of the Schiff-base complex on MCM-41.⁶⁰ The reason for the decrease in intensity of the peaks is that when organic groups and metal complexes are stabilized inside the pores of MCM-41, the ordered hexagonal nature of the channels is reduced.

Table 3 Synthesis of 5-substituted 1*H*-tetrazole derivatives catalyzed by the La-Schiff base@MCM-41 nanocatalyst

Entry	Nitrile	Product	Time (min)	Yield (%)	Melting point
1			120	98	214–216
2			40	97	169–182
3			60	95	222–225
4			60	95	209–212
5			150	97	260–263
6			480	95	216–219
7			45	94	229–233
8			19h	89	259–262
9			410	91	149–152

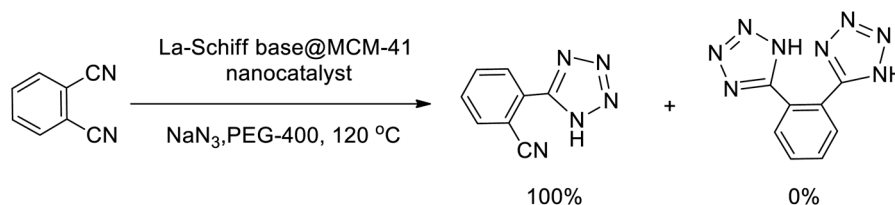


3.2 Catalytic study of the catalyst

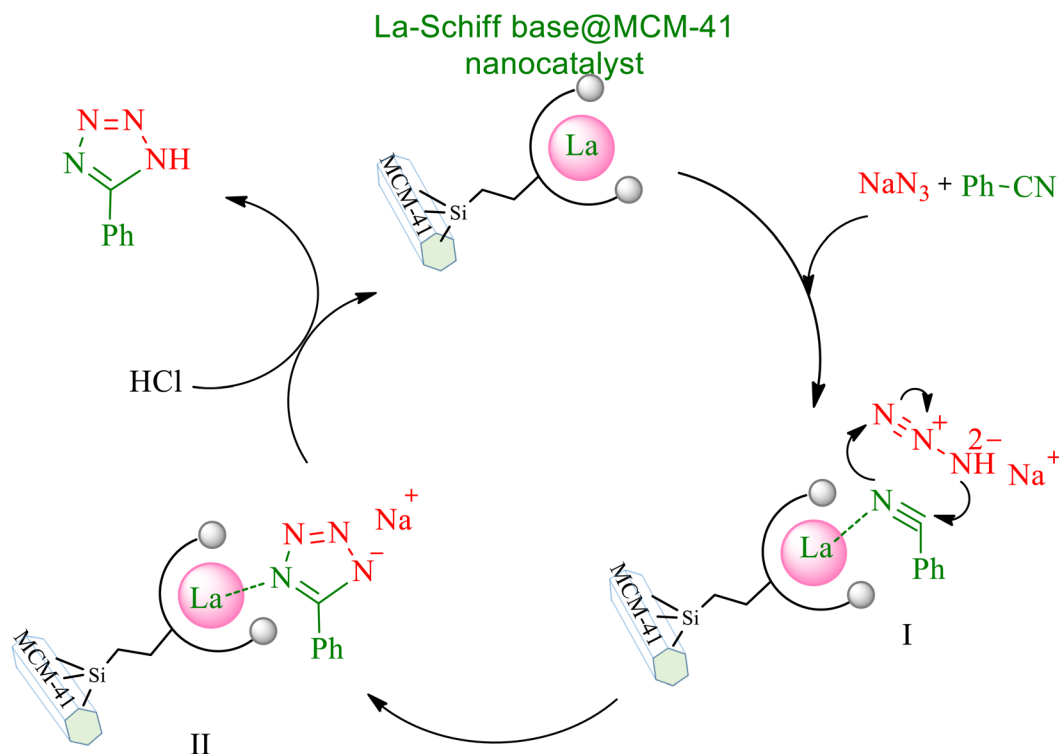
The catalytic ability of La-Schiff base@MCM-41 in organic chemistry was investigated in the synthesis of heterocyclic tetrazole compounds through the [3 + 2] cycloaddition of sodium azide (NaN_3) with benzonitrile derivatives. For this study, the reaction of NaN_3 with benzonitrile under various conditions was selected to find the best reaction conditions. In this regard, the effect of all parameters such as amount of NaN_3 , amount of La-Schiff base@MCM-41 catalyst, temperature and nature of solvent was examined in this reaction (Table 2). As shown in Table 2 (entry 2), 40 mg of La-Schiff base@MCM-41 catalyst cannot achieve completion of the reaction, while this reaction is completed when 50 mg of La-Schiff base@MCM-41 catalyst is used (Table 2, entry 3). Also, polar solvents provide better conditions for the synthesis of tetrazoles, therefore several polar solvents were examined to find the best reaction conditions for the [3 + 2] cycloaddition of NaN_3 and benzonitrile (Table 2, entries 4–6). Also, the reaction was investigated in other

solvents (such as dichloromethane and *n*-hexane), however the obtained results in these solvents were not favorable compared to the PEG-400 solvent (Table 2, entries 7 and 8). Because the boiling point of these solvents is low, the necessary temperature for the synthesis of tetrazole cannot be reached. Therefore, PEG-400 solvent was selected as the best solvent. Finally, the effect of temperature was examined (Table 2, entry 9), and the best results were obtained at 120 °C (Table 2, entry 3). The concentration of NaN_3 was also checked, and 1.4 mmol of NaN_3 was used for the synthesis of the tetrazole compounds.

In continuation, several tetrazole compounds were synthesized under the optimized conditions in the presence of the La-Schiff base@MCM-41 catalyst (Table 3). Amongst these, various benzonitrile derivatives having electron-donating or electron-withdrawing functional groups were investigated. In these studies, all 5-substituted 1*H*-tetrazoles were synthesized in tolerable yields in the presence of the La-Schiff base@MCM-41 catalyst. As shown in Scheme 4, this catalyst shows excellent homoselectivity in the synthesis of 5-substituted 1*H*-tetrazoles.



Scheme 4 Homoselectivity of the La-Schiff base@MCM-41 nanocatalyst in the synthesis of 5-substituted 1*H*-tetrazoles from the [3 + 2] cycloaddition of NaN_3 with dicyano substituted derivatives.



Scheme 5 Expected mechanism for the synthesis of 5-substituted 1*H*-tetrazoles in the presence of the La-Schiff base@MCM-41 nanocatalyst.



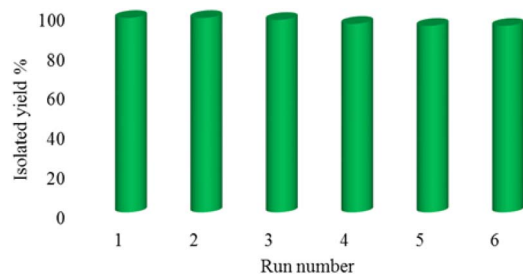


Fig. 7 The recoverability and reusability of the La-Schiff base@MCM-41 nanocatalyst in the synthesis of 5-phenyl-1*H*-tetrazole.

Further, in the reaction of NaN_3 with phthalonitrile having the same two cyano-functional groups, only mono-addition was observed (Scheme 4).

Based on previous literature,^{61,62} a suitable cyclic mechanism for the synthesis of 5-substituted 1*H*-tetrazoles in the presence of the La-Schiff base@MCM-41 catalyst is proposed in

Scheme 5. In this mechanism, intermediate **I** is formed from the interaction of the $\text{C}\equiv\text{N}$ bond with the catalyst, which then introduced intermediate **II**. Then, in the work-up step, intermediate **II** gives the final tetrazole product by addition of HCl and the catalyst is regenerated and can undergo another catalytic cycle of the reaction.

3.3 Reusability of the catalyst

The most important advantage of a heterogeneous catalyst over homogeneous catalysts is its recyclability. Accordingly, green chemistry recommends the use of heterogeneous catalysts because they are both environmentally friendly and economical. Therefore, the reusability of the La-Schiff base@MCM-41 catalyst was studied in the [3 + 2] cycloaddition of NaN_3 with benzonitrile under the optimized conditions. In this regard, the catalyst was separated by centrifugation after each cycle and further was employed in the next cycle without any activation. As shown in Fig. 7, the La-Schiff base@MCM-41 catalyst can be reused up to 6 times without a notable reduction of its catalytic efficiency.

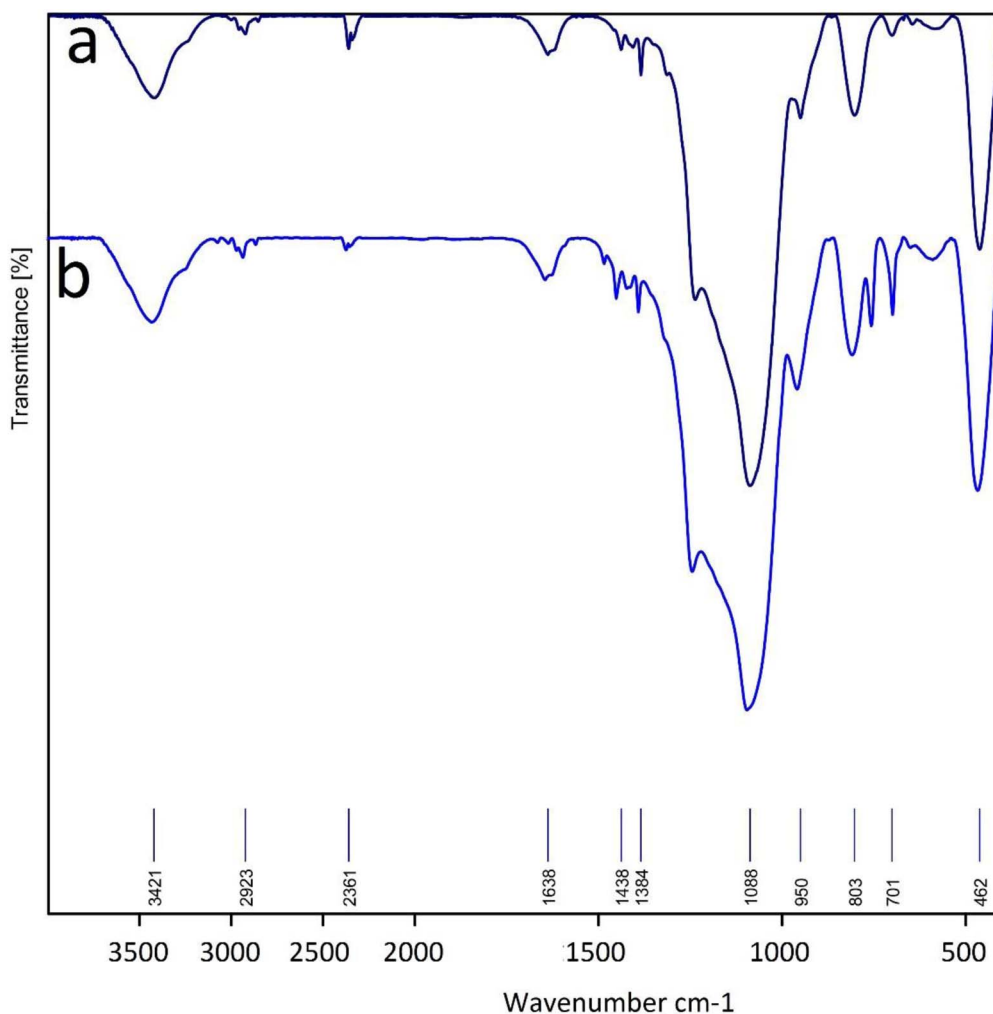


Fig. 8 FT-IR spectra of (a) La-Schiff base@MCM-41 before reuse and (b) recovered La-Schiff base@MCM-41.



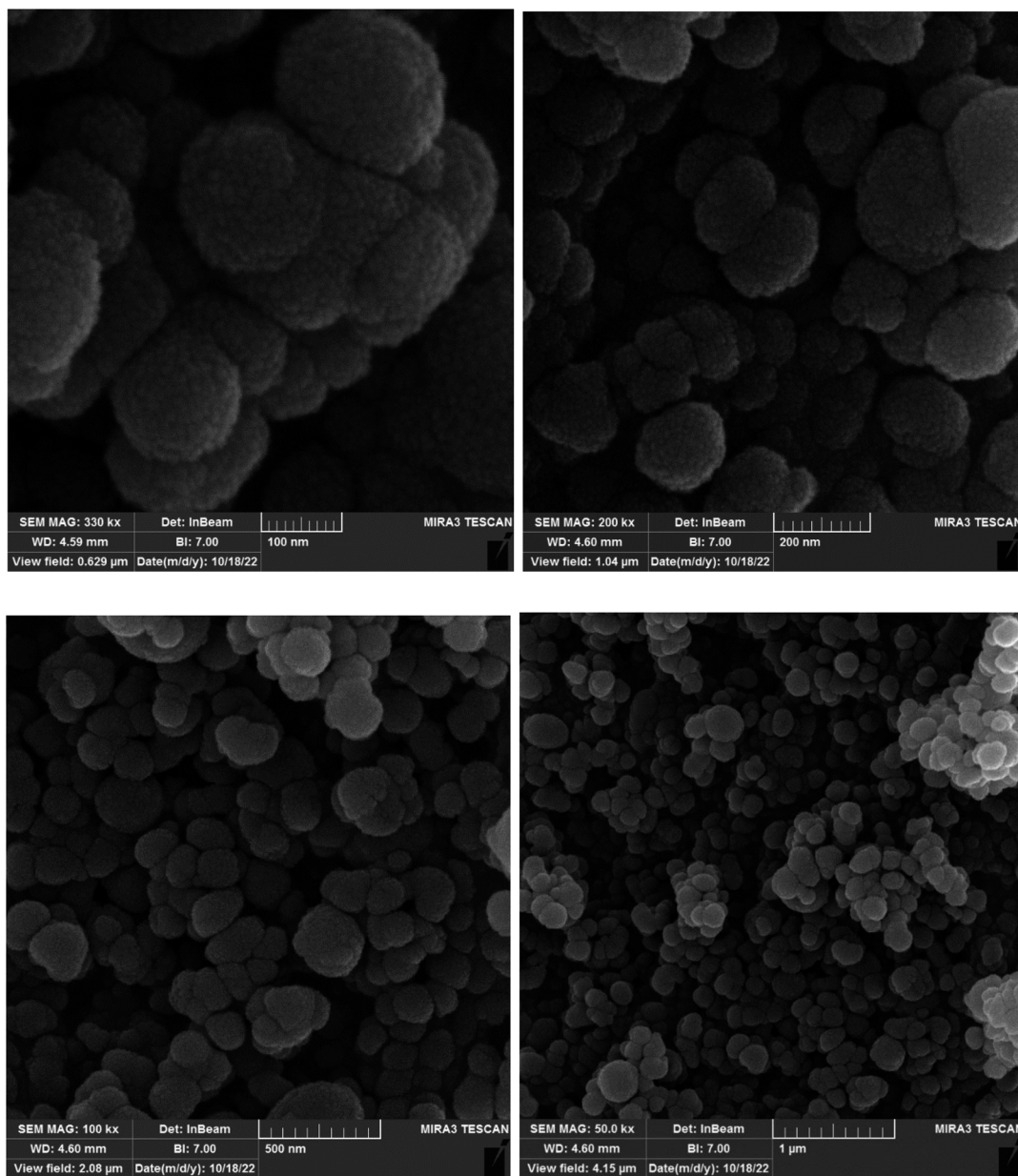


Fig. 9 FESEM images of recovered La-Schiff base@MCM-41 catalyst.

3.4 Characterization of the recovered catalyst

The recovered La-Schiff base@MCM-41 catalyst was characterized using FT-IR spectroscopy, SEM and AAS.

To investigate the heterogeneous nature of the La-Schiff base@MCM-41 catalyst, the reaction between NaN_3 and benzonitrile was performed under the optimized conditions. After completion of the reaction, the catalyst was removed by filtration. Then, the exact amount of leached lanthanum in the reaction solution was measured by atomic absorption spectroscopy (AAS). No significant amount of leached lanthanum was detected in the reaction solution. These results prove the heterogeneous nature and stability of the La-Schiff base@MCM-41 catalyst.

Also, the IR spectrum of the reused La-Schiff base@MCM-41 was compared to that of the fresh La-Schiff base@MCM-41 catalyst and a good similarity could be seen between them. The stability of La-Schiff base@MCM-41 was confirmed by the similar position and shape of the stretching vibrations in the FT-IR spectra of the recovered and fresh catalyst. These results provide strong evidence for the good stability of La-Schiff base@MCM-41 after reuse (Fig. 8).

In the IR spectra, three peaks at 462, 803, and 1088 cm^{-1} relate to the Si-O-Si vibrations.⁶³ The stretching vibration of N-H and O-H bonds can be observed at 3417 cm^{-1} .¹⁰ Also, stretching vibration of C-H bonds locates at 2923 cm^{-1} in the IR spectra.²¹ The bands at 701 cm^{-1} , 1438 cm^{-1} and 1638 cm^{-1}



Table 4 Comparison results of the La-Schiff base@MCM-41 nanocatalyst with other catalysts in the synthesis of 5-phenyl-1H-tetrazole

Entry	Catalyst	Time (h)	Yield (%)	TON	TOF (h ⁻¹)	Ref.
1	MCM-41	2.5	Trace	—	—	18
2	CoY zeolite	14	90	—	—	64
3	Cu-Zn alloy nanopowder	10	95	—	—	65
4	B(C ₆ F ₅) ₃	8	94	18.8	2.35	66
5	Fe ₃ O ₄ @SiO ₂ /salen Cu(II)	7	90	225	32.14	67
6	Fe ₃ O ₄ /ZnS HNSs	24	81.1	4.70	0.195	68
7	Pd-isatin-boehmite	8	94	21.3	2.6	69
8	Mesoporous ZnS	36	86	2.5	0.07	70
9	AgNO ₃	5	83	8300	1660	71
10	CuFe ₂ O ₄	12	82	2.05	0.17	72
11	Nano ZnO/Co ₃ O ₄	12	90	—	—	73
12	Pd-SMTU@boehmite	2.5	95	31.65	12.66	74
13	Cu-TBA@biochar	7	98	125.6	18.4	10
14	L-Cysteine-Pd@MCM-41	3	98	33.79	11.26	18
15	Ni-MP(AMP) ₂ @Fe-biochar	3.8	92	1066	280.7	24
16	Cu(II)-adenine-MCM-41	5	92	35.5	7.1	40
17	Pd-Arg@boehmite	7	97	28.7	4.1	57
18	Cu-DABP@Fe ₃ O ₄ /MCM-41	2	99	60	30	75
19	La-Schiff base@MCM-41	2	98	85 217	42 608	This work

correspond to the C-Br, aromatic C=C and C=N bonds, respectively.¹⁰

The FESEM images of the recovered La-Schiff base@MCM-41 were obtained using an electron microscope (model MIRA3TESCAN-XMU). The FESEM images of the recovered La-Schiff base@MCM-41 are shown in Fig. 9. As shown, no significant change could be observed in the FESEM images of this catalyst after reuse in terms of shape or particle size. The SEM images of the recovered catalyst showed a good similarity with the SEM images of the fresh catalyst. Therefore, this catalyst is stable under the reaction conditions of the synthesis of tetrazoles.

3.5 Comparison of the catalyst

Table 4 is provided to show the practicality of the La-Schiff base@MCM-41 catalyst in comparison with some reported catalysts for the [3 + 2] cycloaddition of NaN₃ with benzonitrile. As reported, the [3 + 2] cycloaddition of NaN₃ with benzonitrile in the presence of unfunctionalized MCM-41 without any metal loading provided no 1H-tetrazole yield (Table 4, entry 1). However, the La-Schiff base@MCM-41 catalyst exhibited a 98% yield of 1H-tetrazole within 120 min, which is superior to some previously reported catalysts in terms of reaction times and yields. As shown in Table 4, the TON and TOF values of the La-Schiff base@MCM-41 catalyst are much higher than those of previous catalysts. TON and TOF are two important factors to evaluate the efficiency of catalysts. Therefore, one of the most important innovations of this work is the very high TON and TOF values of this La-Schiff base@MCM-41 catalyst compared to other catalysts. Previous methods using hazardous solvents and extremely long reaction times have been limited in the synthesis of 5-phenyl-1H-tetrazoles with biological properties. Now, using PEG as a green solvent, a short reaction time, excellent yield, easy recycling of the catalyst and pure separation of the products are the important advantages of the present strategy over previous strategies.

4 Conclusions

In conclusion, we synthesized a new Schiff-base complex of lanthanum on MCM-41 (La-Schiff base@MCM-41). La-Schiff base@MCM-41 was characterized using various techniques such as ICP, CHN, XRD, TGA, BET, FT-IR spectroscopy, SEM, EDS and WDX. Then, the catalytic application of La-Schiff base@MCM-41 was studied in the homoselective synthesis of 5-substituted 1H-tetrazole derivatives. All products were obtained in good yields. Also, the recyclability of La-Schiff base@MCM-41 was described, which showed good recyclability in the synthesis of 5-substituted 1H-tetrazoles.

Conflicts of interest

There are no conflicts to declare.

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