

# Halloysite Nanotubes Modified by Chitosan as an Efficient and Eco-Friendly Heterogeneous Nanocatalyst for the Synthesis of Heterocyclic Compounds <sup>†</sup>

Diana Fallah Jelodar, Zoleikha Hajizadeh and Ali Maleki \*

Catalysts and Organic Synthesis Research Laboratory, Department of Chemistry, Iran University of Science and Technology, Tehran 16846-13114, Iran; diana92fallah@gmail.com (D.F.J.);

hajizadeh.zoleikha@gmail.com (Z.H.)

\* Correspondence: maleki@iust.ac.ir; Tel.: +98-21-7724-054-050; Fax: +98-2173-021-584

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**Abstract:** In this study, halloysite nanotubes (HNTs) are modified by chitosan as a natural cationic amino polysaccharide. Halloysite nanotubes/chitosan (HNTs/Chit) were characterized by Fourier transform infrared (FT-IR) spectroscopy and energy dispersive X-ray (EDX) analysis. Also, its performance as a heterogeneous catalyst was investigated in the synthesis of pyranopyrazole derivatives. Being a reusable and easily recoverable catalyst, eco-friendliness, high efficiency, and mild reaction conditions are some advantages of the present work.

**Keywords:** halloysite nanotubes; chitosan; nanocatalyst; heterocyclic compounds

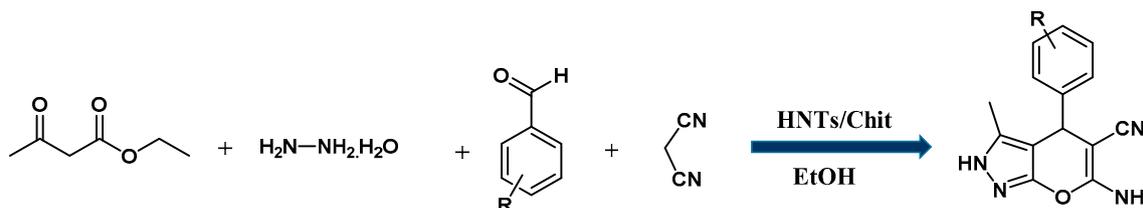
## 1. Introduction

Halloysite nanotubes (HNTs) are a natural aluminosilicate with a hollow tubular structure and the same ratio of tetrahedral and octahedral sheets [1]. HNTs are applied in various applications due to their special properties, such as green, nanotube morphology, accessibility, biocompatibility, porosity, and mechanical stability [2]. Moreover, different chemistry of the outer and inner halloysite nanotube leads to selective modification [3]. HNTs have been functionalized with organic and inorganic materials like chitosan, poly (ethylene imine), and alginate [4–6].

Chitosan (CS) is a natural cationic amino polysaccharide obtained by alkaline N-deacetylation of chitin [7]. Chitosan with the sheet structure like cellulose was achieved by a combination of  $\beta$ -1,4-linked 2-acetoamino-2-deoxy-D-glucopyranose and 2-amino-2-deoxy-glucopyranose units [8]. Recently, chitosan with excellent features such as non-toxicity, biocompatibility, hydrophilicity, antibacterial activity, and biodegradability has attracted more attention among scientists [9]. Chitosan has been applied in different applications including drug delivery systems, wound healing, and tissue engineering [10,11].

The synthesis of pyranopyrazole compounds with pharmacological and medicinal properties was recently suggested by new catalysts similar to magnetic Fe<sub>3</sub>O<sub>4</sub> nanoparticles and isonicotinic acid [12]. Multicomponent reactions (MCRs) with most noteworthy features such as atom economy, short reaction time, and simplicity in the synthesis of complex structures are applied in the synthesis of heterocyclic materials as an important class of organic compounds. It continues to our research on MCRs and nanomaterials and, due to the importance of heterocyclic compounds [13–31], herein,

HNTs/Chit as a green, reusable, and efficient nanocatalyst was used in the synthesis of pyranopyrazole derivatives in mild conditions (Scheme 1).



**Scheme 1.** The synthesis of pyranopyrazole derivatives by HNTs/Chit catalyst.

## 2. Experimental

### 2.1. General

HNTs, chitosan, and all other materials and solvents were obtained from Merck and Aldrich company. The FT-IR spectrum of the product was taken by a Shimadzu IR-470 spectrometer on a KBr pellet. EDX spectra were provided with a Numerix DXP-X10P. Melting points were measured with an Electrothermal 9100 apparatus and are uncorrected.

### 2.2. Synthesis of HNTs/Chit

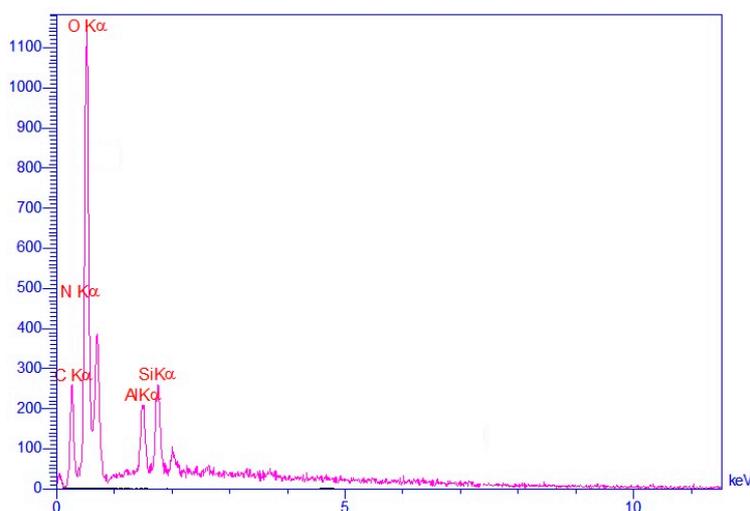
At first, 0.5 g of chitosan was added to 20 mL deionized water. Then, acetic acid solution (0.5 M) was added dropwise until the chitosan was completely dissolved. One gram of HNTs was dispersed in 20 mL deionized water and added to the chitosan solution. The mixture was stirred overnight. Subsequently, it was frozen into ice at  $-20\text{ }^{\circ}\text{C}$ .

### 2.3. General Procedure for the Synthesis of Pyranopyrazole Derivatives 5a-e

The mixture of ethyl acetoacetate (2 mmol), hydrazinehydrate (2 mmol), aromatic aldehyde (1 mmol), and malononitrile (1 mmol) was stirred in 5 mL of EtOH in the presence of HNTs/Chit (20 mg) under reflux condition for 30 min. The reaction progress was checked by thin-layer chromatography (TLC). After the completion of the reaction (as indicated by TLC), the catalyst was separated by filtration. The crude product was recrystallized from EtOH to yield a pure product.

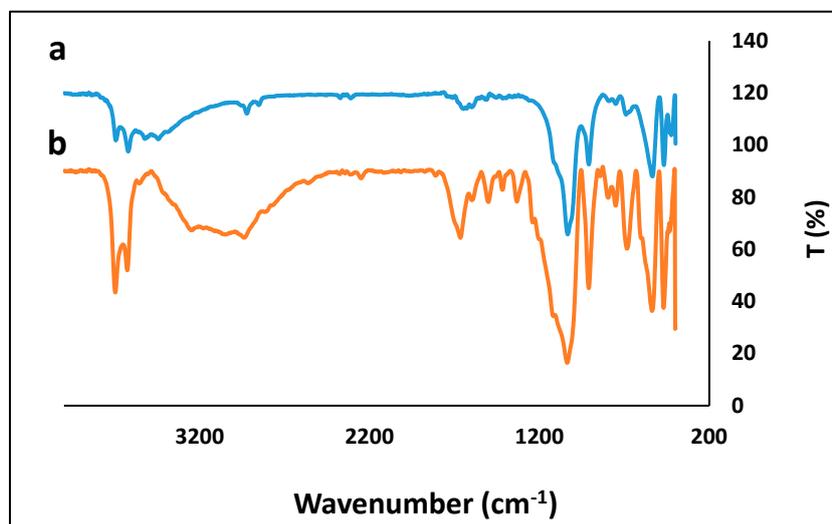
## 3. Results and Discussion

As can be seen in Figure 1, the result of the EDX analysis of HNTs/Chit nanocomposite confirms the presence of Al, Si, O, C, and N elements in the synthesis nanocatalyst.



**Figure 1.** EDX analysis of the HNTs/Chit nanocatalyst.

Furthermore, FT-IR spectroscopy was used as a common analysis. The FT-IR spectrum of the HNTs is shown in Figure 2a. The bands at 520, 460, 1030, and 910  $\text{cm}^{-1}$  are related to Al–O–Si, Si–O–Si, Si–O, and Al–OH, respectively. The stretching vibrations of inner-surface Al–OH are shown at 3690 and 3620  $\text{cm}^{-1}$ . As can be seen in the HNTs/Chit spectrum (Figure 2b), the vibrational stretching of C=N appeared at 1650  $\text{cm}^{-1}$ . Also, the absorption band at 1550  $\text{cm}^{-1}$  was related to the distortion vibration of N–H groups of chitosan.



**Figure 2.** FT-IR spectra of (a) HNTs and (b) HNTs/Chit.

### 3.1. Catalytic Application of HNTs/Chit in the Synthesis of Pyranopyrazole Derivatives

Repeatability of the efficiency of this strategy was confirmed by using different aromatic aldehydes with electron-withdrawing and electron-releasing substitutions and the synthesis of various pyranopyrazole derivatives under mild conditions with great yields. The results are summarized in Table 1.

**Table 1.** Synthesis of various pyranopyrazole derivatives.

Entry	R	Product	Yield <sup>a</sup> (%)	Mp (°C)
1	H	<b>5a</b>	95	244–246
2	3-NO <sub>2</sub>	<b>5b</b>	93	234–235
3	4-NO <sub>2</sub>	<b>5c</b>	91	247–249
4	4-Cl	<b>5d</b>	90	230–232
5	4-Me	<b>5e</b>	90	220–222

<sup>a</sup> Isolated yield.

## 4. Conclusions

In summary, the synthesis of nanocomposites based on natural and green materials is suggested in this research. Halloysite nanotubes were modified easily by chitosan and applied as an efficient nanocatalyst in organic reactions. Mild reaction conditions, reusability of the catalyst, and eco-friendliness are some of the advantages of this study.

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