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Data Article

Dataset for synthesis of conducting polymers nanocomposites based on aniline and 4-amino-benzylamine catalyzed by chromium (III) exchanged maghnite (Algerian MMT) via in situ polymerization



Fatima Zohra Zeggai ^{a, b}, Mohammed Belbachir ^b,
Benjamin Carbonnier ^c, Abdelkader Rahmouni ^{b, *},
Bachari Kaldoun ^a

^a Centre de Recherche Scientifique et Technique en Analyses Physico-chimiques (CRAPC), BP 38Bou-Ismaïl-RP, 42004, Tipaza, Algeria

^b Polymer Chemistry Laboratory, Oran1 University Ahmed Benbella, BP 1524, El'Menouer, 31000, Oran, Algeria

^c Paris-Est University, UMR 7182. 2-8, F-94320, Thiais, France

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ABSTRACT

In this data we report on conductors polymers nanocomposites synthesized by in situ polymerization of aniline (ANI) and/or 4-aminobenzylamine (4-ABA) in presence of chromium montmorillonite (MMT-Cr⁺³) and ammonium persulfate as oxidizing agent. Homopolymers and copolymers (PANI-co-4-ABA) were prepared at various initial monomer composition and were characterized by Fourier transform Infrared (FT-IR) and UV-vis spectroscopy, X-ray diffraction (XRD) and cyclic voltammeter. The data describes the behavior of the corresponding homopolymers Poly (4-ABA) and (PANI) and showed that the in-situ polymerization produced real nanocomposites containing aniline and 4-aminobenzylamine units and films of products exhibit good electrochemical properties.

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* Corresponding author.

E-mail address: ramaek23@yahoo.fr (A. Rahmouni).

Specifications Table

Subject	Polymer chemistry
Specific subject area	Synthesis and characterization of nanocomposites catalyzed by maghnite-H+(Algerian MMT) via in situ polymerization
Type of data	Table, Image and Figure
How data were acquired	SEM, NMR, FTIR, XRD,TGA,UV, Cyclic Voltammogram
Data format	Raw and analyzed
Experimental factors	Synthesis and characterization of new nanocomposites under effect of heterogeneous catalyst called maghnite-H+ (Algerian MMT) exchanged with chromium (III) via in situ polymerization. The obtained nanocomposite was characterized and discussed by several methods such as (XRD, FTIR, Electrical and electrochemical conductivity, SEM, HNMR).
Experimental features	Maghnite (Algerian MMT) was used as heterogeneous catalyst for synthesis of organic and polymeric materials.
Data source location	Republic algerian democratic and popular
Data accessibility	Data are supplied with this article
Related research article	A. Rahmouni and M. Belbachir. Molecular structure of PANI and its homologue PANI-PEO ₂₀₀₀ catalyzed by Maghnite-H+ (Algerian MMT): Synthesis, characterization and physical and chemical properties. Polymer Bulletin (2019) 76:4677–4701. https://doi.org/10.1007/s00289-018-2620-7 .

Value of the Data

- The data in this article will be informative to synthesis of polymeric materials under effect of heterogeneous catalyst called maghnite (Algerian MMT).
- By using these data, researchers can make comparisons with other polymerization like (cationic polymerization, anionic polymerization, radical polymerization, polymerization by emulsion) and comparisons between (homogenous catalyst and heterogeneous catalyst).
- Strategy for this method of synthesis employed in this Data article can be used as a reference for future studies in the electronic domain.
- The Data obtained in this work can be effectively applied for the synthesis of conductor polymers nanocomposites under effect of heterogeneous catalyst (Algerian-MMT) via in situ polymerization.
- The data can be highlighted for further studies in development of better strategy for synthesis of conducting polymer especially for electronic and electrical domain.

1. Data

The data described in this paper provides formation of nanocomposite structures used in electronic domain catalyzed by maghnite-Cr³⁺. PANI/MMT-Cr³⁺, Poly (4-ABA/MMT-Cr³⁺) and Poly (ANI-co-4-ABA)/MMT-Cr³⁺ nanocomposites were successfully synthesized under effect of modified

Table 1Elementary compositions wt. % of chromium (Cr³⁺) and sodium (Na⁺) exchanged sample raw-maghnite (Algerian MMT).

Compositions wt.%	SiO ₂	Al ₂ O ₃	Fe ₂ O ₃	CaO	Na ₂ O	MgO	K ₂ O	TiO ₂	Cr ₂ O ₃	Pert in fire
Raw-MMT	69.39	14.67	1.16	0.30	0.50	1.07	0.79	0.16	0.00	11.96
MMT-Na	70.75	14.46	1.05	0.19	2.61	1.01	0.78	0.14	0.00	09.01
MMT-Cr	71.01	14.06	1.00	0.14	0.15	0.98	0.71	0.15	2.61	09.35

Table 2

Peak maximum and d-spacing of protonated and the nanocomposites intercalated into sodium montmorillonite.

Samples	Peak maximum	Basal spacing	Interlayer spacing
	2θ max (deg)	d ₍₀₀₁₎ (Å°)	Δd (Å°)
MMT-Na	6.96	12.94	–
MMT-Cr	6.06	14.67	1.73
P(4aba_co-ani)/MMT-Cr (20/80)	5.65	15.63	2.69
P(4aba_co-ani)/MMT-Cr (80/20)	5.52	16.04	3.1
P(4aba_co-ani)/MMT-Cr (50/50)	5.64	15.71	2.77
P4ABA/MMT-Cr	5.56	15.88	2.94
PANI/MMT-Cr	5.64	15.71	2.77

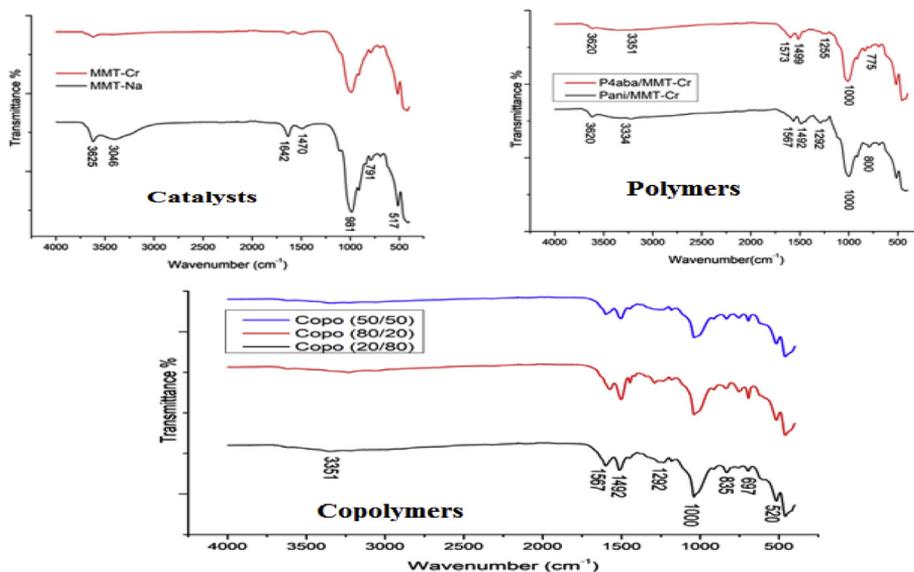


Fig. 1. FT IR-spectra of the MMT-Na, MMT-Cr and the nanocomposites poly (4-ABA/MMT-Cr), PANI/MMT-Cr and their copolymers Poly (4-ABA-co-ANI/MMT-Cr).

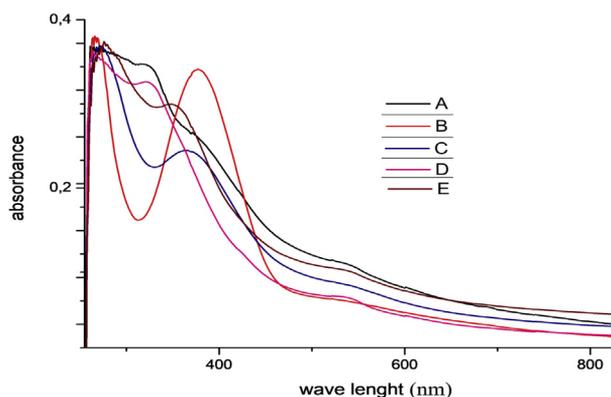


Fig. 2. UV-Vis spectra of the homo and copolymer nanocomposites doped with MMT-Cr, A: Poly (4-ABA/MMT-Cr), B: Poly (ANI/MMT-Cr), C: Poly (4-ABA-co-ANI)/MMT-Cr; 80/20, D: Poly (4-ABA-co-ANI)/MMT-Cr; 20/80, E: Poly (4ABA-co-ANI)/MMT-Cr; 50/50).

clay layered called maghnite- Cr^{3+} (Algerian MMT- Cr^{3+}) by in situ polymerization route in the presence of oxidizing agent. The formation of polymers and copolymers was confirmed by FTIR, XRD, ^1H NMR, ATG, electrical conductivity and Uv-Visible measurements [1,2]. Table 1 describes elementary compositions wt. % of chromium (Cr^{3+}) and sodium (Na^+) exchanged sample raw-maghnite (Algerian MMT). Table 2 describes Peak maximum and d-spacing of protonated and the nanocomposites intercalated into sodium montmorillonite. Fig. 1 FT IR-spectra of the MMT-Na, MMT-Cr and the nanocomposites poly (4-ABA/MMT-Cr), PANI/MMT-Cr and their copolymers Poly

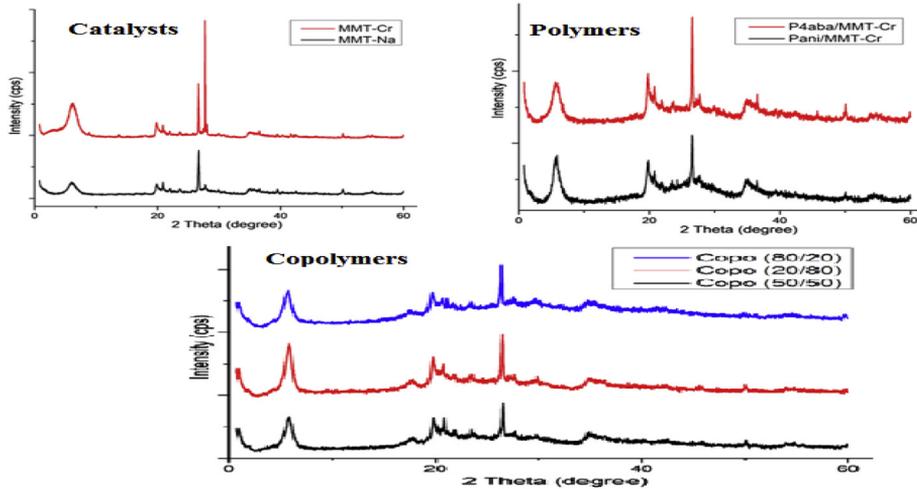


Fig. 3. X-ray diffraction patterns of two montmorillonite (MMT-Na and MMT-Cr), and the nanocomposites (PANI/MMT-Cr, Poly (ani-co-4aba)/MMT-Cr, P4ABA/MMT-Cr).

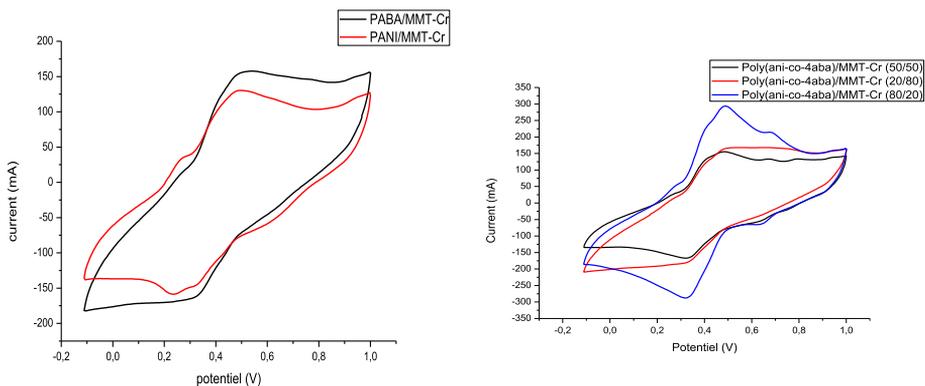


Fig. 4. Cyclic voltammogram recorded of polymer and copolymer films formed in 1.0 M HClO₄ on graphite carbon electrode.

(4-ABA-co-ANI/MMT-Cr³⁺). Fig. 2 describes UV–Vis spectra of the homo and copolymers nanocomposites doped with MMT-Cr³⁺, A: Poly (4-ABA/MMT-Cr), B: Poly (ANI/MMT-Cr), C: Poly (4-ABA-co-ANI)/MMT-Cr³⁺: 80/20, D: Poly (4-ABA-co-ANI)/MMT-Cr³⁺: 20/80, E: Poly (4ABA-co-ANI)/MMT-Cr³⁺: 50/50). Fig. 3 describes X-ray diffraction patterns of two montmorillonite (MMT-Na and MMT-Cr), and the nanocomposites (PANI/MMT-Cr, Poly (ani-co-4aba)/MMT-Cr, P4ABA/MMT-Cr). Fig. 4 describes cyclic voltammogram recorded of polymer and copolymer films formed in 1.0 M HClO₄ on graphite carbon electrode. Fig. 5 describes TGA curves of PANI prepared in the presence of Maghnite-H⁺ (0.25 M). Fig. 6 describes TGA curves of Poly (4-ABA-co-ANI)/MMT-Cr synthesized in the presence of Maghnite-H⁺ (0.25 M). Fig. 7 describes ¹H-NMR spectra of (PANI) obtained by the intercalated method between Aniline and Maghnite-Cr³⁺ (black powder).

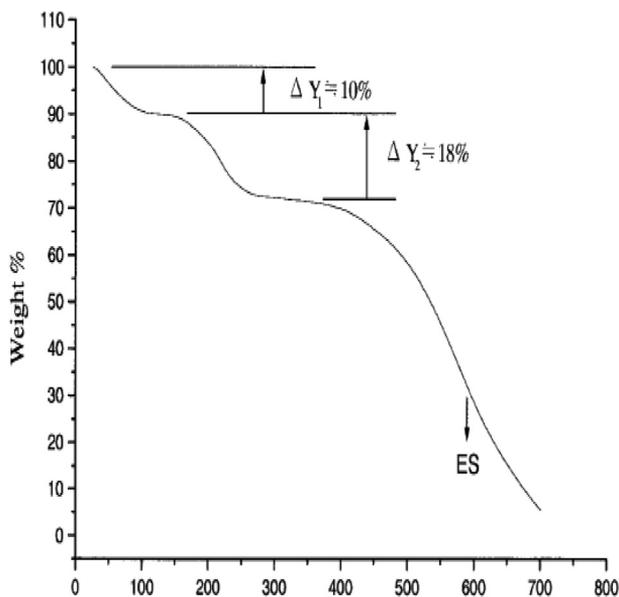


Fig. 5. TGA curves of PANI prepared in the presence of Maghnite-H⁺ (0.25 M).

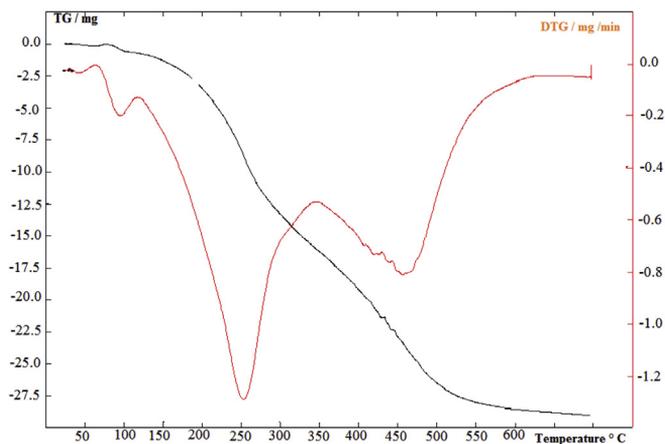


Fig. 6. TGA curves of Poly (4-ABA-co-ANI/MMT-Cr) prepared in the presence of Maghnite-H⁺ (0.25 M).

Fig. 8 describes ¹H-NMR spectra of the block copolymer poly (aniline)-b-poly (4-aminobenzylamine) catalyzed by Maghnite-Cr³⁺ by in situ polymerization. Fig. 9 describe proposed mechanism of homopolymer (PANI) catalyzed by Maghnite-H⁺ by in situ polymerization. Fig. 10 describe proposed mechanism of homopolymer poly (4-aminobenzylamine) catalyzed by Maghnite-H⁺ by in situ polymerization. Fig. 11 describe proposed mechanism block copolymer poly (aniline)-b-poly (4-aminobenzylamine).

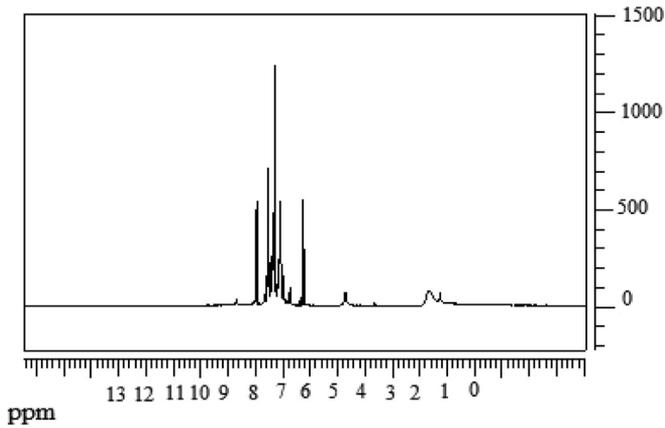


Fig. 7. ^1H -NMR spectra of (PANI) obtained by the intercalated method between Aniline and Maghnite- Cr^{3+} (black powder).

2. Experimental design, materials, and methods

2.1. Preparation of Maghnite- Cr^{+3} (MMT- Cr^{+3})

The raw-clay sample (Raw-MMT) was washed with distilled water to remove impurity [3,4]. The obtained montmorillonite (10 g) was crushed for 20 min using a Prolabo ceramic balls grinder. The greatest proton saturation of the <2 mm fractions of clay were obtained by first saturating with Na^+ ions using 1 M NaCl solution and to confirm the absence of chloride we use the silver nitrate

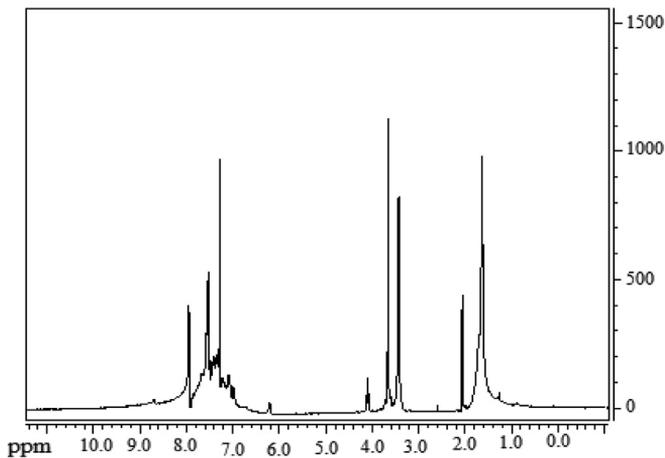


Fig. 8. ^1H -NMR spectra of the block copolymer poly (aniline)-b-poly (4-aminobenzylamine) catalyzed by Maghnite- Cr^{3+} by in situ polymerization.

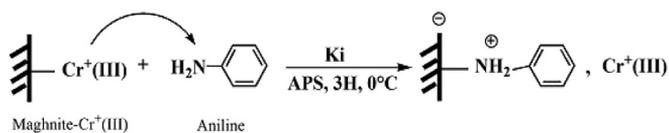
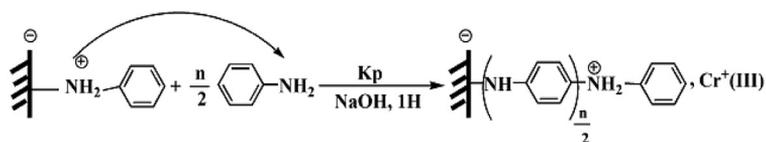
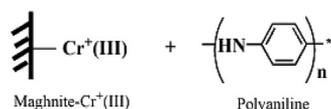
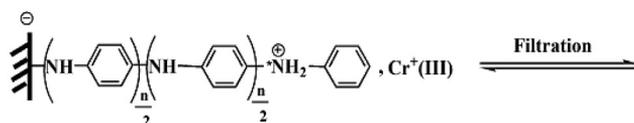
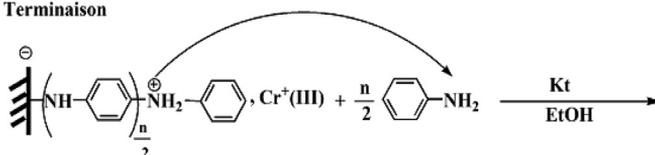
Initiation**Propagation****Termination**

Fig. 9. Proposed mechanism of homopolymer (PANI) catalyzed by Maghnite-H⁺ by in situ polymerization.

[5,6]. To obtain MMT-Na⁺ with chromium intercalated (MMT-Cr⁺³), the MMT-Na⁺ was dispersed into a 1 M CrNO₃ solution and stirred for 24 h and then the solid was recovered by centrifugation and washed with abundant water [7,8]. The catalyst composition was determined by X-ray fluorescence, the obtaining data are listed in Table 1 [9,10].

2.2. Synthesis of polymers/Maghnite-Cr⁺³

Polymer/MMT-Cr⁺³ nanocomposites have been prepared by In-Situ process and the synthesis procedure is briefly described as follows:

The monomers were added by various feed mole fractions. In all cases, the mole ratio of oxidant to the total monomer was defined.

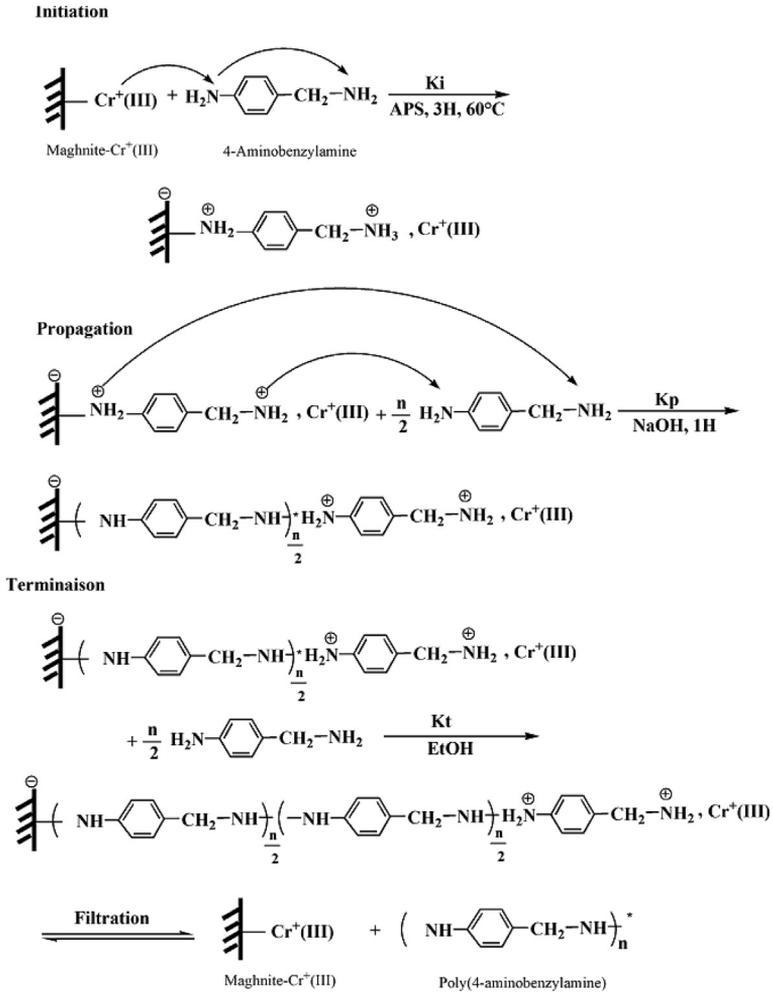
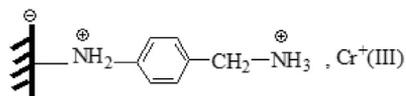
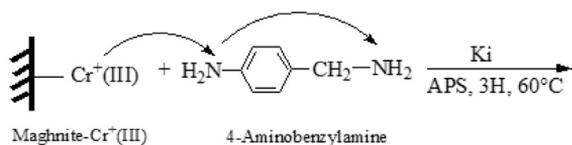
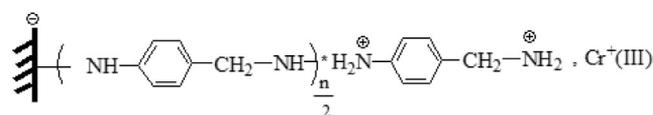
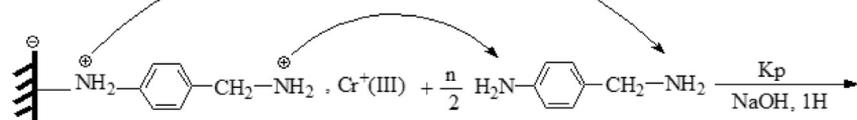


Fig. 10. Proposed mechanism of homopolymer poly(4-aminobenzylamine) catalyzed by Maghnite-H⁺ by in situ polymerization.

Initiation



Propagation



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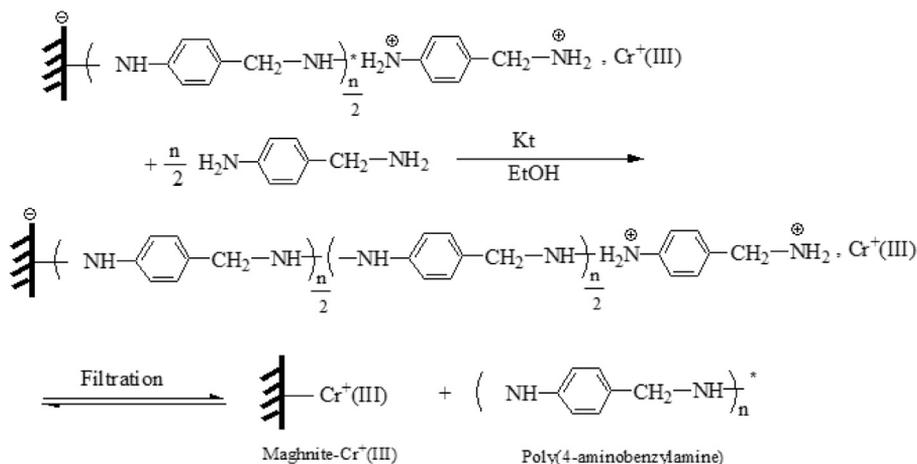


Fig. 11. Proposed mechanism block copolymer poly (aniline)-b-poly (4-aminobenzylamine) catalyzed by Maghnite-H+ by in situ polymerization.

Acknowledgments

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Conflict of Interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

Appendix A. Supplementary data

Supplementary data to this article can be found online at <https://doi.org/10.1016/j.dib.2020.105161>.

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