

pH-SENSITIVE AND MAGNETICALLY RESPONSIVE SUPERABSORBENT NANOCOMPOSITE HYDROGEL

A thesis submitted in partial fulfillment of the requirement
for the degree of M.Sc. in chemistry

SUBMITTED BY

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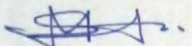
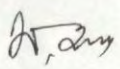
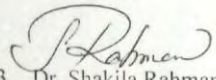
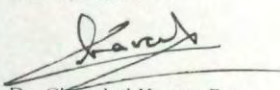
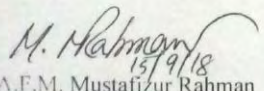


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The thesis entitled "pH-Sensitive and Magnetically Responsive Superabsorbent Nanocomposite Hydrogel" submitted by Labiba Bushra, Roll No.:1014032701, Session: October 2014, has been accepted as satisfactory in partial fulfillment of the requirement for the degree of Master of Science in chemistry on September 15, 2018.

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Dedicated to
Beloved Parents, Husband
&
My Honorable Supervisor

Table of Contents

List of Figures.....iv
.
List of Tables.....iv
List of Tables and Abbreviations of Technical Symbols and Terms.....iv

Acknowledgement.....v
Abstract.....v

List of Tables and Figures

Figures

Figure 1.1 One dimensional crosslinker	3
Figure 1.2 Two dimensional crosslinker	4
Figure 1.3 Three dimensional crosslinker	4
Figure 1.4 magnetic nanoparticles not covalent bonded	16
Figure 1.5 magnetic nanoparticles leave the polymer chain when magnetic field is applied	16
Figure: 3.1 Schematic representation of magnetic crosslinker.....	33
Figure 3.2 FTIR spectra of Fe ₃ O ₄ , silica coated MION, aminated MION and acrylated MION.....	35
Figure 3.4 kinetics study ofswelling.....	37
Figure 3.5 pH dependent swelling	38
Figure 3.6 Magnetic crosslinker dispersed in water and get attract to magnet.....	39
Figure 3.7 polymer nanocomposite hydrogel with magnetic crosslinker is attracted by magnet.....	39
Figure 3.8 Widescan spectra of aminated MION, acrylated MION.....	40
Figure 3.9 High resolution of C1s spectra of aminated MION	41
Figure 3.10 High resolution of C1s spectra of acrylated MION	41
Figure 3.11 high resolution of N1s spectra of aminated MION	42
Figure 3.12 High resolution of N1s spectra of acrylated MION	42
Figure 3.13 XRD spectra of Fe ₃ O ₄ , acrylated MION	43
Figure 3.14 XRD spectra of nanocomposite.....	44
Figure 3.15 (A) SEM image of MION at Resolution x 50,000	45
Figure 3.15 (B) SEM image of (MION) at Resolution x 100,000.....	46
Figure 3.17 (A) SEM image of Acrylated MION at Resolution x 50,000	47
Figure 3.17 (B) SEM image of Acrylated MION at Resolution x 100,000.....	47
Figure 3.18 (A) SEM image of nanocomposite (4 : 0.025 g) at Resolution x 50,000.....	48
Figure 3.18 (B) SEM image of nanocomposite at Resolution x 100,000.....	48
Figure 3.19 (A) SEM image of nanocomposite (4: 0.0125 g) at Resolution x 50,000.....	48
Figure 3.19 (B) SEM image of nanocomposite (4: 0.0125 g) at Resolution x 100,000.....	49
Figure 3.20 EDX spectra of Fe ₃ O ₄	50
Figure 3.21 EDX spectra of silica coated of MION	50

Figure 3.22 EDX spectra of acrylated MION.....	51
Figure 3.23 (A) EDX spectra of nanocomposite(4:0.0125g)	52
Figure: 3.24 Stress vs Strain curves of magnetic nano composite hydrogel, BIS PAA hydrogel and PAA(No crosslinker).....	51
Figure:3.25 Stress vs Strain curves of magnetic nano composite hydrogel with various amount of PAA, BIS PAA hydrogel and PAA(No crosslinker).....	52

Tables

Table 3.1 characteristic IR band of MION, Silica coated MION, Aminated MION, Acrylated MION.....	35
Table 3.2 Elemental composition of magnetic and nanocomposite compound.....	49

List of Tables and Abbreviations of Technical Symbols and Terms

1. Magnetic iron oxide nano particles (MION)
2. Poly (acrylic acid) (PAA)
3. N, N'-methylenebisacrylamide (MBA)
4. N. N'-methylenebis(acrylamide) (MBAAm)

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Abstract

Hydrogels are highly crosslinked polymer materials that are able to swell when immersed in aqueous solutions. Some hydrogels can be designed in such a way that they undergo some conformational changes in response to external stimuli such as temperature, pH, ionic strength, light, electrical signal and magnetic field etc. The hydrogels have been exploited in a wide variety of applications including drug delivery and tissue engineering, biosensors and microfluidic devices. Herein, we develop a novel strategy for combining pH - and magneto-responsiveness into one hydrogel system. Specifically, as pH -sensitive part of the hydrogel we used polyacrylamide, which was crosslinked with acrylic-functionalised silica coated magnetic nanoparticles to possess the magnetic responsiveness. Fe₃O₄ nanoparticles were synthesized and coated with silica, which provides the nanoparticles with a reactive surface for functionalization with -NH₂ group through the silanation with 3-aminopropyl triethoxysilane. The -NH₂ groups were then further functionalized by reacting with acrylic acid to generate magnetic nanoparticles containing acrylic group, which were finally crosslinked with PAA to give the desired nanocomposite hydrogel. The synthesized nanocomposite was characterized by x-ray photoelectron spectroscopy (XPS), x-ray diffraction (XRD), energy dispersive x-ray (EDX), scanning electron microscopy (SEM), and Fourier transform infrared (FTIR) analysis. The synthesized nanocomposite hydrogel absorbed water as high as 400 greater than the mass of itself and they showed super absorbance character. Moreover hydrogel showed pH -responsive swelling when dipped in water with variable pH. On the other hand, the as-synthesized hydrogel showed magnetic behavior owing to the presence magnetite nanoparticles in the form of 3D crosslinker. The prepared nanocomposite hydrogel also exhibited nice self-healing properties due to the presence of acrylic acid groups in the system.

Contents

Introduction.....	1
1.2.1 One Dimensional Crosslinker.....	3
1.2.3 Three Dimensional Crosslinker.....	4
1.3 Responsiveness of hydrogel.....	5
1.3.1 pH responsive hydrogel.....	5
1.3.2 Thermo-responsive hydrogel.....	5
1.3.3 Magneto- responsive hydrogel.....	6
1.3.4 light responsive hydrogel.....	8
1.3.5 Electric field responsive hydrogel.....	9
1.4 Multi responsive hydrogel.....	10
1.4.1 Magneto- and thermo-responsive hydrogel.....	10
1.4.2 pH- and Thermo-responsive hydrogel.....	10
1.5 Magnetic Nanoparticle synthesis method.....	11
1.5.1 Thermal decomposition method.....	11
1.5.2 Micro emulsion method.....	11
1.5.3 Co-precipitation method.....	11
1.6 Surface modification of Iron oxide nanocomponents.....	11
1.7 Magnetically responsive polymer nanocomponents.....	12
1.8 pH-responsive polymer nanocomponents.....	13
1.9. Swelling properties of hydrogel.....	13
1.11 Literature Review.....	15
1.12 Research Goal.....	16
Experimental.....	23
2.1 Materials and Instrument.....	24
2.1.1 Chemicals and Reagents.....	24
2.1.2 Materials.....	24
2.1.3 Instruments.....	25
2.2 Synthesis of magnetic crosslinker.....	25
2.2.1 Synthesis of Fe ₃ O ₄	25
2.2.2 Silica coated Fe ₃ O ₄	25
2.2.3 Silica coated Fe ₃ O ₄ functionalized by –NH ₂ group.....	26

2.2.4 Further functionalized by –CH=CH ₂ group.....	26
2.3 Synthesis of nanocomposite	27
2.4.1 Kinetics study of swelling	27
2.4.2 pH dependent swelling study.....	28
2.6 Field emission scanning electron microscopy (SEM)	29
2.7 Energy dispersive x-ray (EDX) spectroscopy	29
2.8 X-ray diffraction (XRD).....	29
2.9 X-ray photoelectron spectroscopy.....	29
2.10 Mechanical strength of hydrogel.....	30
Results and Discussion	32
3.1 Synthesis of magnetic nanocomposite.....	33
3.1.1 Synthesis of magnetic crosslinker	33
3.1.2 Polymer nanocomposite Synthesis	34
3.2 Functional group analysis using FTIR.....	34
3.4 X-ray photoelectron spectroscopy (XPS)	
3.4.1 High Resolution C1s Spectra.....	38
3.4.2 High resolution N1s spectra.....	40
3.5 X-ray diffraction analysis (XRD)	41
3.6 Surface morphology study using scanning electron microscope (SEM).....	42
3.7 Energy dispersive x-ray spectral analysis.....	45
3.8 Mechanical strength analysis.....	51
3.9 Swelling measurement.....	50
3.9.1 Kinetics study of swelling	50
3.9.2 pH dependent swelling study.....	51
Conclusion.....	52

Chapter 1

Introduction

1.1 Hydrogel

Hydrogels are highly crosslinked polymer materials that are able to swell but do not dissolve in water. The word “hydrogel”, first introduced in 1894 which the material described there was not a hydrogel as we describe it today. It was colloidal gel made with inorganic salts. First generation of hydrogel’s that form of a crosslinker involve the chemical of a monomer or polymer with an initiator is to develop material with high swelling, good mechanical properties [1]. Second generation of materials capable of response to specific stimuli, such as temperature, pH, ionic strength, concentration of specific molecules in solution [2]. Finally, a third generation of hydrogels focusing on the investigation and development of stereo complexed materials hydrogels cross linked by other physical interactions. The ability of hydrogels to absorb water arises from hydrophilic functional groups attached to the polymer backbone. The gel is a state that is neither completely liquid nor completely solid. These half liquid-like and half solid-like properties cause many interesting relaxation behaviors that are not found in either a pure solid or a pure liquid.

Hydrogels can be classified into two distinct categories, the natural and the synthetic hydrogels. Natural hydrogels include collagen, fibrin, and derivatives of natural materials such as chitosan, alginate and silk fibers [3]. Two main drawbacks of natural hydrogels, however, make their final microstructures and properties difficult to control re reducibility [4]. In contrast, synthetic hydrogels such as poly (ethylene glycol) di-acrylate, poly-acrylamide, and poly-vinylalcohol are more reproducible, although their final structure can also depend on polymerization conditions. During last two decades, natural hydrogels were gradually replaced by synthetic hydrogels which has long service life, high capacity of water absorption, and high gel strength[5]. Hydrogels can be synthesized from purely synthetic components. Also, it is stable in the conditions of sharp and strong fluctuations of temperatures. Hydrogels may absorb from 10-20% up to thousands of times their dry weight in water. Polymeric hydrogel networks may be formed by various techniques, however the most common synthetic route is the free-radical cross-linking co polymerization of a hydrophilic non-ionic monomer such as acrylic acid (AA) with a small amount of a cross-linker, e.g. In order to increase their swelling capacity, an ionic comonomer is also included into the reaction mixture. Since the monomers for hydrogel preparation are usually solid at the usual polymerization temperature, it is necessary to carry out the polymerization reactions in an aqueous solution. Hydrogel structure and,

thus, the hydrogel properties are closely related to the conditions under which the Hydrogel are formed, i.e., the cross-linker concentration, the initial degree of dilution of the monomers and the chemistry of the units building the network -structure[6]. They possess also a degree of flexibility very similar to natural tissue due to their large water content. Recently, hydrogels have been defined as two or multi-component systems consisting of a three-dimension.

1.2 Crosslinker of hydrogel

A cross-link is a bond that links on polymer chain to another. They can be covalent bonds or ionic bonds. Generally, which agents perform this joining work is called crosslinker. Crosslinkers contain at least two reactive groups. Reactive groups that can be targeted for crosslinking include primary amines, sulfhydryls, carbonyls, vinyl group, carbohydrates, carboxylic acids etc. When polymer chains are linked together by cross-linker, they lose some of their ability to move as individual polymer chains. Hence chemical crosslinking is mechanically and thermally stable. Crosslinking may be initiated by heat, pressure, change in pH, or radiation. Crosslinkers are chosen depending on chemical specificity, same or different reactive groups, spontaneously reactive or photo reactive groups etc. Crosslinkers are commonly used to modify nucleic acids, drugs and solid surfaces

1.2.1 One Dimensional Crosslinker

In this group of crosslinker the reactive groups are usually remain at the end of chain. It is also called 1D crosslinker. 1D crosslinker contains at least two active sites which are able to react with substrate or polymer matrix. Due to lower active sites comparatively less amount of polymer chains is linked up. Common use linear crosslinkers are N, N'-methylenebisacrylamide (MBA), bis (triethoxysilylpropyl) tetrasulfide, imido ester etc.

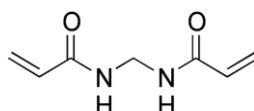


Figure 1.1 One dimensional crosslinker

1.2.2 Two Dimensional Crosslinker

Planar cross-linkers are normally called 2D cross-linker. Its reactive groups are on the upper and lower side of the surface. In case of linear crosslinker there are only two linking sites. But in case of 2D crosslinker many more active sites are available on the surface. These active sites interact with the coming group of polymer or other matrices and form an excellent composite material. 2D crosslinkers are able to link comparatively higher number of polymer chains. As a result, different physical and chemical properties are improved by using 2D crosslinker. Well known 2D crosslinkers included nanoclay, graphene oxide (GO) nanoparticles, functionalized GO etc. are widely used to form nanocomposite. These nanocomposites are used in wastewater treatment, electronic devices and many more sectors.

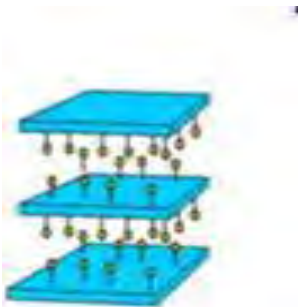


Figure 1.2 Two dimensional crosslinker

1.2.3 Three Dimensional Crosslinker

This class of crosslinkers added new dimension in the crosslinked composite materials. The shape of the 3D crosslinker may be spherical. Its surface area is comparatively higher than that of the linear or planar crosslinker. In case of 2D crosslinker only two sides contain active sites for the interaction with substrate but in case of 3D crosslinker, reactive sites are laying all sides of 3D crosslinker. Hence it contains a wide number of reactivites in a small surface area.



Figure 1.3 Three dimensional crosslinker

Due to this its reactivity increases dramatically. Since reactive sites are scattered, 3D crosslinkers are able to link with much more polymer chains. As a result of the produced nanocomposite become robust and improved chemical and mechanical properties of the nanocomposites.

1.3 Responsiveness of hydrogel

Typical stimuli are temperature, pH, electric field, light, magnetic field etc. And the responses can also be manifold: dissolution precipitation, degradation, drug release, change in hydration state, swelling/collapsing, hydrophilic/hydrophobic surface, change in shape, conformational change. This article will focus on temperature and magneto as external stimuli, since these systems are predominantly studied. Stimuli-responsive hydrogels are cross-linked, hydrophilic polymer networks that undergo a physicochemical transition in response to a change in external stimuli such as pH, temperature, light and analyte concentration.

1.3.1 pH responsive hydrogel

pH responsive polymers are materials which will respond to the changes in the pH of the surrounding medium by varying their dimensions. Such materials increase its size (swell) or collapse depending on the pH of their environment. This behavior is exhibited due to the presence of certain functional group in the polymer chain. There are two kinds of pH sensitive materials: one which have acidic group (-COOH, -SO₃H) and swell in basic pH, and others which have basic groups (-NH₂) and swell poly acrylic acid is an example of the former chitosan is an example of the latter. The mechanism of response is same for both, only the stimulus varies. In aqueous media of appropriate pH and ionic strength, the pendant groups ionize and develop fixed charges on the polymer network, generating electrostatic repulsive forces responsible for pH dependent swelling or de swelling of the hydrogel, thereby controlling the drug release. Small changes in pH can result in significant change in the mesh size of the polymeric networks.

1.3.2 Thermo-responsive hydrogel

Thermo-responsive hydrogels are that exhibit a drastic and discontinuous change of their physical properties with temperature. The term is commonly used when the property concerned is solubility in a given, solvent but it may also be used when other properties

are affected. Thermo-responsive polymers belong to the class of stimuli responsive hydrogel, in contrast to temperature-sensitive. Temperature-sensitive hydrogels are probably the most studied class of stimuli sensitive polymer in drug delivery research. These hydrogels are able to swell or de-swell as a result of change in the temperature of the surrounding fluid. Considering volume-phase transitions give us a volume-phase transition temperature (VPTT) instead of a critical solution temperature, there are still two main classifications of observed behavior, positive volume-phase transitions and negative volume-phase transitions. A positive volume-phase transition or “thermophilic” hydrogel will swell upon heating, increasing its volume. A negative volume-phase transition or “thermophobic” hydrogel will collapse upon heating, decreasing its volume. Thermophilic hydrogels typically maintain a collapsed conformation at temperatures below their VPTT while thermophobic hydrogels are able to swell at lower temperatures [7]. Thermo-responsive hydrogels that exhibit a drastic and discontinuous change of their physical properties with temperature. The term is commonly used when the property concerned is solubility in a given, solvent but it may also be used when other properties are affected. Thermo- responsive polymers belong to the class of stimuli responsive hydrogel, in contrast to temperature-sensitive.

1.3.3 Magneto- responsive hydrogel

Recently, there is a growing interest and effort to expand the range of available stimuli from temperature, pH which could be limited by mass transfer, to newly emerging remote electromagnetic triggers like light, electricity and magnetic field, to obtain responsive materials with more precise responsivity and higher efficiency. Magnetic nanocomponents are multi-component materials, containing nano sized magnetic materials. They response to an external stimulus i.e., magnetic field, different materials (e.g., gels, liquid crystals, renewable polymers, silica, carbon or metal organic frameworks) with various types of magnetic particles, offering exciting perspectives for fundamental investigation. Magnetic nanomaterials have attracted considerable attention because of their unique size-dependent properties, and many important technological applications. Magnetic nanocomponents usually consist of components exhibiting magnetic properties, like iron oxide such as Fe_3O_4 and Fe_2O_3 , pure metals such as ferrum Cobalt and nickel, their chemical compounds and alloys. Magnetic properties based on size, shape, structure, crystallinity, and chemistry of materials. In nature iron oxides exist in many forms like magnetite (Fe_3O_4), hematite ($\alpha\text{-Fe}_2\text{O}_3$) and maghemite ($\gamma\text{-(Fe}_2\text{O}_3)$). Among these oxides Fe_3O_4 particles are more

demandable due to surface efficiency. For the first time, the magnetite crystal structure had been studied in 1915 as one of the first mineral structures evaluated through X-ray diffraction method by Bragg and Nishikawa. Recently, Fe_3O_4 have been intensively investigated because of their super paramagnetic, high co reactivity and low Curie temperature. Magnetite (Fe_3O_4) is a kind of excellent magnetic material among iron oxides. It has a cubic inverse spinel structure. which oxygen forms a face centered cubic closed packing and Fe^{3+} ions occupy the tetrahedral sites and octahedral sites, and bears distinguished electric and magnetic properties on account of the hopping of electrons between Fe^{2+} and Fe^{3+} in the octahedral sites. Magnetite has been widely applied in numerous fields such as Ferro fluid, catalysis, pigment color, magnetic storage media, environment protection, cell separation, clinic diagnosis and therapy. Fe_3O_4 MNPs are also nontoxic and biocompatible. The name hematite is derived from the Greek word for blood *haima*, due to the red coloration found in some varieties of hematite. Hematite is the form of Iron iron(III) oxide (Fe_2O_3), hematite exhibits antiferromagnetic behavior below temperature of 260K. This temperature is known as Morin temperature. Above this temperature, hematite exhibits weak ferromagnetic behavior. Hematite is widely used in magnetic recording media, as a catalyst, gas sensors etc. These applications of iron oxide arise due to its low cost, thermal stability, its resistance to corrosion and its non-toxic nature. Among the various magnetic materials, the cubic spinel structured maghemite ($\gamma\text{-Fe}_2\text{O}_3$) represents an important class of magnetic transition metal oxide materials in which oxygen atoms form a face centered close-packed structure. $\gamma\text{-Fe}_2\text{O}_3$ is an ideal candidate for fabrication of luminescent and magnetic dual functional nanocomposites due to its excellent transparent properties. The existence of amorphous Fe_2O_3 and four polymorphs (alpha, beta, gamma and epsilon) is well established. At a temperature of 650°C , hematite turns into Fe_3O_4 with a high energy loss. Hematite has strongly antiferromagnetic properties. Gamma Fe_2O_3 (maghemite) is the ferrimagnetic cubic form of Fe(III) oxide and it differs from the inverse spinel structure of magnetite through vacancies on the cation sublattice. In time, at room-temperature, the maghemite turns into hematite crystalline structure. Maghemite has the same crystalline structure like Fe_3O_4 (magnetite). Maghemite Fe_2O_3 is biocompatible and therefore is one of the most extensively used biomaterials for different applications like cell separation, drug delivery in cancer therapy.

Hydrogels are three dimensional hydrophilic polymer networks that can swell in water and hold a large amount of water while maintaining the structure. A three dimensional

network is formed by crosslinking polymer chains. Crosslinking can be provided by covalent bonds, hydrogen bonding, Vander Waals interactions or physical entanglements. They experience reversible volume changes in response to external stimulus such as pH, temperature and ionic concentration. Stimuli responsive hydrogels often exhibit sharp volume phase transitions triggered by specific chemical or physical stimuli. Volume phase transitions has been vastly studied due to its potential in triggering the functions of movements such as deformation, volume change modulus changes, force generation.

Magneto responsive hydrogel belong to this recent trend and are of special interest, as they can realize temporal and remote control and have great potential, for instance for microfluidic systems in analytical or therapeutic devices. Fe^{3+} ions are the most wanted in formation of intracellular and macromolecular biologically active or magnetic resonance contrast materials because of their non-toxicity and their existence in nature in many tissues. For these in vivo applications, the magnetic nanoparticles need to exhibit high magnetic saturation. Biocompatible and non-toxic magnetic nanoparticles, well-dispersed in the solvent carrier, have a lot of potential for in vivo applications in which superparamagnetic particles are of interest because they do not retain any residual magnetism after removal of a magnetic field. Magnetic nanoparticle materials offer major advantages due to their unique size and physicochemical properties. Because of the widespread applications of magnetic nanoparticles in biotechnology, biomedical, material science, engineering, and environmental areas, much attention has been paid to the synthesis of different kinds of magnetic nanoparticles. Other biomedical applications of magnetic nanoparticles include drug delivery and magnetic resonance imaging. The large surface-to-volume ratio of MNPs provides abundant chemically active sites for biomolecule conjugation, dispersions of magnetic nanoparticles are attractive in separation applications as they offer high surface area and can be functionalized to selectively discriminate between different molecular or cellular species. A magnet is used to separate the magnetically targeted cells from the non-targeted cells, main advantages of these materials include the low interference, low background signal, no requirement of pre-treatment, and the fact that they can be small enough to be portable. [8-13]

1.3.4 light responsive hydrogel

Light-sensitive hydrogels have potential applications in developing optical switches, display units, and ophthalmic drug delivery devices. Light-sensitive hydrogels can be

separated into UV-sensitive and visible light-sensitive hydrogels. Unlike UV light, visible light is readily available, inexpensive, safe, clean and easily manipulated. Visible light-sensitive hydrogels were prepared introducing a light sensitive chromophore to poly(N-isopropylacrylamide) hydrogels. When light (e.g.488 nm) is applied to the hydrogel, the chromophore absorbs light which is then dissipated locally as heat by radiation less transitions, increasing the 'local' temperature of the hydrogel. The temperature increase alters the swelling behavior of poly (N-isopropylacrylamide) hydrogels, which are thermo-sensitive hydrogels. The temperature increase is proportional to the light intensity and the chromophore concentration. Light-sensitive hydrogels have potential applications in developing optical switches, display units, and ophthalmic drug delivery devices. Generally, upon the irradiation of light with certain wavelengths, these chromophores reveal changes in their different forms induces macroscopic changes in the structures and thus the properties of the hydrogels, including adhesion, color, permeability, hydrophilicity, ionizability, etc. Base on this, light-responsive hydrogels have been employed to manage size-selective mass transport (electron, gas, inorganic and organic ions) properties, such as color, molecular polarity, molecular size and shape, or transform between ionic state and nonionic state.[14]

1.3.5 Electric field responsive hydrogel

Electric current induced hydrogels are basically composed of polyelectrolyte and shrinks or swells in response to an applied electric field . Polymers contain a large number of the ionizable group on their backbone chain thus sensitive towards both pH and electricity. Electro-sensitive hydrogel, which are basically pH-sensitive hydrogel, are able to convert chemical energy to mechanical energy. Elctro-sensitive hydrogel, Electro-sensitive hydrogels undergo shrinking or swelling in the presence of an applied electric field. Sometimes, the hydrogels show swelling on one side and de swelling on the other side. The hydrogel shape change (including swelling, shrinking) depends on a number of conditions. If the surface of hydrogels is in contact with the electrode, the result of applying electric field to the hydrogel may be different from systems where the hydrogel is placed in water (or acetone–water mixture) without touching the electrode. The result will be different yet if the aqueous phase contains electrolytes. Electro-sensitive hydrogels have been applied controlled drug delivery.[15]

1.4 Multi responsive hydrogel

Responsive polymer hydrogels can largely and reversibly change their volume or shapes under slight external stimuli, including temperature, pH, ionic strength, special chemicals, redox, light, humidity, electric field, magnetic fields, etc. This property of responsive polymer hydrogels can be utilized in diverse applications, such as sensor, valve, on-off switches, biomimetic actuators, artificial muscles and so on. A variety of stimuli-responsive hydrogels has been emerging recently, putting this smart hydrogel's responsiveness into use by three dimensional (3D) structure is challenging. Stimuli-responsive polymers are smart polymers which can respond to external stimuli in the form of changing their own properties. [16]

1.4.1 Magneto- and thermo-responsive hydrogel

A specially designed stimuli responsive hydrogel was fabricated. into the network of chemical crosslinked polyacrylamide hydrogel is one of the important environmental factors in typical physiological, biological, chemical systems. AAm hydrogels were aggregated to form physical crosslinks due to the hydrophobic interactions when the pH value of the environment was above 5.0 and the temperature was low. Magnetic nanoparticles incorporate the polymer system but these magnetic nanoparticles are bonded with the polymer system by weak van der Waals force. When the magnetic field applied the magnetic nanoparticle aggregate and break down the weak forces.[17]

1.4.2 pH- and Thermo-responsive hydrogel

pH- and temperature-responsive hydrogels are the most widely investigated intelligent hydrogels because these two stimuli are the most common used external stimuli. This can be a change in conformation, change in solubility, alteration of the hydrophilic/hydrophobic balance or release of a bioactive molecule (e.g. drug molecule). This also includes a combination of several responses at the same time the combination of a pH-responsive system with a thermo-responsive polymer can further alter the hydrophilic/hydrophobic balance. This allows a polymer to become membrane active at a specific temperature and/or a specific pH. Stimuli-responsive hydrogels have recently captured a lot of attention due to their ability to respond to external triggers such as temperature and pH. This makes them attractive candidates for a wide range of biological and biomedical applications.[18,19]

1.5 Magnetic Nanoparticle synthesis method

1.5.1 Thermal decomposition method

Thermal decomposition Heat is used to cause chemical decomposition of organometallic compounds into monodisperse magnetic nanocrystals. This is achieved in the presence of organic solvents and stabilizing surfactants. Shape and morphology of resulting crystals are mainly controlled through the ratios of the used reagents, reaction temperature and time thermal decomposition complicated as a synthesis technique.[20]

1.5.2 Micro emulsion method

As micro emulsions are stable dispersions of two liquids, they can be used to synthesize nanoparticles through precipitation caused by the addition of solvents such as ethanol or acetone. The resulting nanoparticle-precipitate can be isolated through centrifugation or filtering. Examples of particulates obtained by microemulsion are gold-coated cobalt/platinum, metallic cobalt and cobalt/platinum alloys.

1.5.3 Co-precipitation method

The most important and established methods of magnetic nanoparticle synthesis are Co-precipitation. A simple and efficient technique to build iron oxides through adding a base to an aqueous $\text{Fe}^{2+}/\text{Fe}^{3+}$ salt solution at room temperature (or above) and under conditions of inert atmosphere. Co-precipitation very simple, ambient conditions 20 minutes, water needed. As a synthesis technique, co-precipitation is most used, as it is a very efficient and easy. Among various chemical methods for synthesis of different types of metal oxides, co-precipitation method has several advantages over other methods including, good homogeneity, low cost, high purity of product and not requiring organic solvents and heat treatment.

In this research, we aim to synthesize Iron Oxide(Fe_3O_4) nanomaterials via chemical co-precipitation method combined with nontoxic ferric chloride; ferrous chloride and the toxic sodium hydroxide as starting materials.

1.6 Surface modification of Iron oxide nanocomponents

Bare iron oxide nanoparticles tend to aggregate because of strong magnetic attractions among particles, particle-particle bipolar attraction, the van der Waal force and high

surface energy. On the otherhand when the magnetic nanoparticles are in contact of air, these particles are further oxidized to form alpha or gamma Fe_2O_3 . To avoid this problem surface of iron oxide nanoparticles are coated to improve their stability and biocompatibility and to achieve hydrophilicity and conjugating capability. Several coating materials including organic polymers (e.g. dextran, chitosan, polyethylene glycol), organic surfactants (e.g. sodium oleate and dodecylamine), inorganic metals (e.g. gold), inorganic oxides (e.g. silica oxide and carbon) etc. are used to modify the surface of the magnetic nanoparticles. Here this inorganic oxide such Si act as a stabilizer to prevent aggregation of magnetic iron oxide nanoparticle.

1.7 Magnetically responsive polymer nanocomponents

During the last decade, the development of magnetic polymer nanocomponents materials has been the source of discovery of spectacular new phenomena, with potential applications in the multidimensional fields. Among the broad spectrum of nanoscale materials being investigated for various environmental and biomedical applications, magnetic nanoparticles (MNPs) have gained significant attention due to their intrinsic magnetic properties, which makes them successful as magnetically recoverable catalysts, drug delivery agents, anticancer materials, magnetic resonance imaging devices, etc. Magnetic nanoparticles and nanocomposites have aroused significant scientific and technological interest because of their potential applications in the fields of biomedicine, information technology, magnetic resonance imaging, catalysis, telecommunication, and environmental remediation. Magnetic polymer nanocomponents generally comprise of magnetic nanoparticles embedded in polymer matrix. However, magnetic nanoparticles dispersed in composites usually have a strong tendency to form agglomerates for reduction of energy associated with high surface area-to-volume ratio of the nanosized particles. To avoid aggregation of magnetic nanoparticles, protection strategies have been developed to chemically stabilize the magnetic nanoparticle. The properties of magnetic particles remain same after stabilization. So its application in polymer nanocomposites is increasing day by day. [21]

1.8 pH-responsive polymer nanocomposites

Normally, pH -responsive hydrogels are prepared from polymers which exhibit acid group like carboxylic or sulphonic acid. To create these surfaces, the pH -responsive polymer, poly(acrylic acid) (PAA), is bonded with 3D crosslinker by free radical polymerization. As one of the most widely studied smart materials, pH -responsive hydrogels have attracted an increasing interest and been more and more studied in the past several years, because of the easy controllability of pH- and more importantly their applications in various fields.

The main feature of this kind of smart polymer is an ability to receive or release protons which responding to the pH changes. Usually, these polymers are polyelectrolyte where containing acid groups or basic groups. The acid groups accept protons at low pH values and release protons at high pH values. Thus, the polymer will swell with the increase of pH because of the electrostatic repulsion of the negatively charged groups. The examples of polyanions are poly(acrylic acid) (PAA) and poly(methacrylic acid) (PMAA). these smart polymers also change the solubility by changing the electrical charge of the polymer Molecule. So, these hydrogels need to decrease the electrical charge in the molecular level in order to transform from a soluble state to an insoluble state. There are several ways that can be used to decrease the polymer's electric charge which are decreasing pH, neutralizing the electric charge and reducing the hydrophilicity of the polymeric macromolecules. The stimuli-responsive polymers or smart-polymers can be developed by several ways including in response to temperature and pH which can make them applicable in several applications such as modified release of drugs, tissue engineering and biosensors.

1.9. Swelling properties of hydrogel

A crosslinked polymer hydrogels swell but not dissolve when water or a solvent enters it. The swelling properties, which usually use degree of swelling to define hydrogels, depend on many factors such as network density, solvent nature, polymer solvent interaction parameter. Because of the high water content and biocompatibility of hydrogels, many applications are related to biomedical usage. When a cross-linked polymer keep in a good solvent, it become swell rather than dissolve. Swelling degree define the change of dimension of polymer. A highly cross-linked polymer shows less degree of swelling. The swelling capacity of a polymer is determined by the amount of liquid material that can be

absorbed. The amount of water sorption is quantified by the degree of swelling the ratio of the swollen polymer volume or mass to that of the dry polymer. The swelling value can be calculated by using the following formula ,

$$\text{Swelling \%} = \frac{(W_s - W_d)}{W_d} \times 100\%$$

Where, W_d = Weight of polymer, W_s = weight of swollen polymer. The equilibrium swelling degree and the elastic modulus of hydrogels depend on the cross-link and charge densities of the polymer network as well as on the cross-linked polymer concentration after the gel preparation. Polymeric hydrogel networks may be formed by various technique; however, the most common synthetic route is the free-radical cross-linking copolymerization of a hydrophilic non-ionic monomer such as acrylic acid (AA) with a small amount of 3D cross-linker. In order to increase their gel formation capacity and swelling capacity bi functional crosslinker e.g, N, N'-methylenebis(acrylamide) (MBAAm) is also included into the reaction mixture. Since the monomers for hydrogel preparation are usually solid at the usual polymerization temperature, it is necessary to carry out the polymerization reactions in an aqueous solution. poly(acrylic acid) (PAA) hydrogels prepared from AA, 3D crosslinker, and MBAAm in aqueous solutions cross-linker concentration dependence of the cross-linking efficiency. MBAAm, that is the fraction of MBAAm forming effective cross-links. Increasing number of ionic groups in hydrogels is known to increase their swelling capacities. This is mainly due to the simultaneous increase of the number of counter-ions inside the gel, which produces an additional osmotic pressure that swells the gel.

1.10 Mechanical strength of hydrogel

Hydrogels are consist of a water-swollen network of crosslinked polymer chain. They can be made from synthetic polymers such as poly(vinyl alcohol)(PVA) or Poly(Acrylic acid)(PAA). Their biocompatibility, ease of fabrication and viscoelastic properties makes them highly suitable for use as constructs engineer tissues. One of the main limitations with using hydrogels to engineer tissues in vitro is their poor mechanical properties. Previous attempts at engineering tissues using hydrogel have produced tissues with significantly poorer mechanical strength than the real tissues. There are several reasons for this including the random alignment of fibres and high water content within the hydrogel.[22]

1.11 Literature Review

Stimuli-responsive hydrogels have found increasing attention, which is attributed to the manifold applications they can be used for. Several year intensive research was invested in stimuli-responsive hydrogel. Their stimuli-responsiveness led not only to novel responsive groups, which enabled the translation of an external physical impact into a change of a material property, but also to hydrogel that are equipped with more than one responsive group. The ability of hydrogels to absorb water arises from hydrophilic functional groups attached to the polymer backbone while their resistance to dissolution arises from cross-links between network chains. Water inside the hydrogel allows free diffusion of some solute molecules, while the polymer serves as a matrix to hold water together. Another aspect of hydrogels is that the gel is a single polymer molecule, that is, the network chains in the gel are connected to each other to form one big molecule on macroscopic scale. Stimuli-responsive hydrogels are cross-linked, hydrophilic polymer networks that undergo a physicochemical transition in response to a change in external stimuli such as pH, temperature, light, ionic strength, electric field, magnetic field and the presence of electrolyte. As a result, dual- or multi-stimuli responsive hydrogel are obtained, which is the starting point to more sophisticated applications, due to the variability that is introduced to the responsiveness. By using this property, it might be used in drug delivery, tissue engineering, biosensors, microfluidic devices, biomedical and application of agricultural field etc. Multi-responsive hydrogel can show more than two stimuli such as pH-, thermo- and light-responsive hydrogel, light-, redox- and temperature-responsive polymers, environmental-, pH- and temperature-responsive hydrogel, Thermo-responsive hydrogel undergoes conformational change in response to temperature change. As designed, the polymer exhibited a responsiveness towards pH, temperature and light and was considered to be triple stimuli- responsive. They show different swelling at different temperature. At a significant temperature range they absorb water and another temperature they squeeze water. Magneto-responsive-hydrogel contain magnetic nanoparticles. This magnetic nanoparticles, dispersed in the polymer system within weak van der Waals force to make the hydrogel magnetically responsive. Applied magnetic field causes the hydrogel to squeeze due to the attraction of the magnetic components to the magnet. At a significant temperature range they absorb water another temperature they squeeze water. By using magnetic property, it may separate from the aqueous solution. This behavior of the magnetic nanoparticles might make things complex when the hydrogel shows

regenerability. By using this both property it may use in drug delivery, tissue engineering, and in environmental pollutant remediation. Magnetic nanoparticles are dispersed in the polymer system within Weak Van Dar Waals force to make the hydrogel magnetically responsive. When the magnetic field is applied the magnetic nanoparticles aggregate breaking the weak forces and when it is taken off they usually be reluctant to get back to their original position.

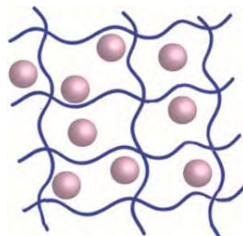


Figure 1.4 magnetic nanoparticles not covalent bonded

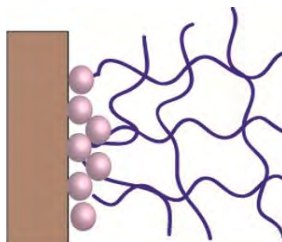


Figure 1.5 Magnetic nanoparticles leave the polymer chain when magnetic field is applied

This is our object. We also synthesized Fe_3O_4 and this magnetic nanoparticle also dispersed in the polymer system. But our magnetic nanoparticle dispersed in the polymer system within strong covalent bond. For this when we applied magnetic field the magnetic nanoparticle is not aggregate breaking the strong covalent bond and when the magnetic field is taken off they do not get back their original position.[23-47]

1.12 Research Goal

The interesting behavior of hydrogels has been exploited in a wide variety of applications including drug delivery and tissue engineering, biosensors and microfluidic devices. However, due to the random nature of the crosslinking reactions during the polymerization many polymer gels have poor mechanical properties. This is a significant drawback of the hydrogels restricting their applications specifically where high stress is required . Incorporating different materials such as ceramics, metals, silicates and magnetic particles

into the hydrogel matrix usually lead to improved properties, enhanced mechanical strength and structural stability. Hydrogel composite systems exploit the individual properties of the components into combined effects increasing the versatility of their applications. Herein, we aim to combine the pH- and magneto- responsiveness in one hydrogel system and study their properties.

The major goal of this research is to design a noble strategy for the preparation of a magneto- and pH- responsive nanocomposite hydrogel. Among different external stimuli for multi-responsive hydrogels, we choose pH- and magnetic field, which are very important in numerous applications including drug delivery and tissue engineering. As pH- sensitive part of the hydrogel we will use poly(Acrylic acid), which will be crosslinked with acrylamide-functionalized magnetic nanoparticles to introduce the magnetic responsiveness. The magnetically responsive hydrogels available in the literature are composite of polymer and Fe_3O_4 particles where the magnetic Fe_3O_4 particles dispersed in the polymer system are held by weak van der Waals force. In our proposed research, the acrylamide-functionalized silica coated Fe_3O_4 nanoparticles will be covalently bonded through cross-linking with the polymeric chains of the hydrogel. To achieve our goal first step is to synthesize acrylamide-functionalized silica coated Fe_3O_4 that will be used as a general precursor for crosslinking as well as introducing the magnetic properties into the hydrogel. We use this magnetic crosslinker to prepare superabsorbent polyacrylic acid hydrogel via in situ polymerization using potassium persulfate as initiator. The as-synthesized nanocomposite will be characterized by x-ray photoelectron spectroscopy (XPS), x-ray diffraction (XRD), scanning electron microscopy (SEM), energy dispersive x-ray (EDX), Fourier transform infrared (FTIR) analyses. Finally, we will investigate the swelling and magnetic behavior of the synthesized superabsorbent nanocomposite hydrogel in water. We expect that this method would be a general platform for the incorporation of magnetic responsiveness through crosslinking of the acrylamide-functionalized magnetic nanoparticles with other polymers like polyacrylamide, polyacrylic acid etc. [48-54]

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Chapter 2

Experimental

2.1 Materials and Instrument

2.1.1 Chemicals and Reagents

In this research, we used most of the chemicals of Merck, Germany and Sigma Aldrich. Some chemicals are of BDH. The chemicals and reagents used in this work were analytical grade. They were used without further purification. Distilled water was used as solvent to prepare most of the solutions of this work.

2.1.2 Materials

The chemicals and reagents which are used in this research are given below:

- i. Ethanol (Merck, Germany)
- ii. Toluene (Merck, Germany)
- iii. Sodium hydroxide (Merck, Germany)
- iv. Potassium per sulfate (BDH)
- v. Sulfuric acid (Merck, Germany)
- vi. Hydrochloric acid (analytical grade)
- vii. Acetone (analytical grade)
- viii. Ferrous chloride. $4\text{H}_2\text{O}$ (Sigma Aldrich)
- ix. Ferric chloride. $6\text{H}_2\text{O}$ (Sigma Aldrich)
- x. Ammonia solution (25%)
- xi. Tetraethyl orthosilicate (TEOS) (Sigma Aldrich)
- xii. γ -aminopropyl triethoxy silane (APTES) (Sigma Aldrich)
- xiii. N, N'-diethylformamide (DMF) (Merck, Germany)
- xiv. Methacrylic anhydride (Sigma Aldrich)
- xv. Acrylic acid (BDH)
- xvi. N, N'-methylenebisacrylamide (Sigma Aldrich)

xvii. Hydrogen peroxide (Org Germany)

2.1.3 Instruments

Analysis of the samples was performed using the following instruments:

- i. Fourier Transform Infrared Spectrophotometer (SHIMADZU FTIR-8400)
- ii. Field Emission Scanning Electron Microscopy (JSM-7600F, Tokyo, Japan)
- iii. X-ray Diffractometer (Philips, Expert Pro, Holland)
- iv. X-ray photoelectron Spectroscopy
- v. Centrifuge machine (Hettich, Universal 16A)
- vi. pH meter (Hanna, HI 8424, Romania)
- vii. Digital Balance (AB 265/S/SACT METTLER, Toieto, Switzerland)
- viii. Freeze dryer (Heto FD3)

2.2 Synthesis of magnetic crosslinker

2.2.1 Synthesis of Fe₃O₄

The magnetite (Fe₃O₄) nanocomponents prepared by the most popular, easy and widely used, chemical co-precipitation method. An iron salt solution was obtained by mixing 0.005mol 2.70g FeCl₃ and 0.0025mol 0.633g FeCl₂ such that the ratio of Fe³⁺ to Fe²⁺ in 50ml deoxygenated distilled water by passing N₂ gas about 10 min. Twenty milliliter of 1.5M NaOH was rapidly poured into the ferric salt solution under magnetic stirring at room temperature (RT). Orange color of iron salt solution was instantly changed and black precipitate found. After continuously stirring for 20min, the precipitate was separated by magnetically and washed with the deoxygenated distilled water four times by centrifugation. Some sample was taken for FTIR and oven dried at 50°C overnight. The naked Fe₃O₄ nanocrystals were obtained as the control sample (called S1).

2.2.2 Silica coated Fe₃O₄

Magnetic Fe₃O₄ nanocrystals were coated by hydrolysis of TEOS on the surfaces of the magnetic Fe₃O₄ nanocrystals in basic solution via stobers method. After washing with the anhydrous ethanol, the precipitate Fe₃O₄ was ultrasonically dispersed in a solution

containing 240ml ethanol and 60ml water, then loaded into a three-necked bottle. The pH value was adjusted to 9 with 25% ammonia solution. Here 300 mg magnetic nanoparticle were dispersed in the solution. 4ml TEOS was added under vigorous stirring. The resulting dispersion was mechanically stirred for 10 hours. After 10h, the ferrofluid was heated at 50°C to complete hydrolysis for another 12h, and a little was removed to prepare a SEM sample. The particles were again separated by centrifugation and washed with the deoxygenated distilled water and anhydrous ethanol, then vacuum-dried at 50°C overnight. The silica-coated Fe₃O₄ nanocrystals were prepared and the sample was called S2.

2.2.3 Silica coated Fe₃O₄ functionalized by –NH₂ group

After washing with the anhydrous ethanol in step 2, then washed with dimethyl formamide (DMF). Then the silica-coated Fe₃O₄ nanocrystals were ultrasonically dispersed in a solution containing 120ml DMF and 80ml toluene, then loaded into a three-necked bottle as above. Ten milliliter 10mL (3-aminopropyltriethoxysilane) APTES was added drop by drop into the three necked bottle under magnetic stirring. After hydrolyzing for 24hour at room temperature, the particles were collected by centrifugation and washed within toluene 4 times. The silica-coated Fe₃O₄ nanocrystals functionalized by –NH₂ groups were thus obtained and were called S3 .[1]

2.2.4 Further functionalized by –CH=CH₂ group

After washing with toluene for the last time in step 3, the particles were dispersed in 150 ml toluene and loaded into a three-necked bottle and heated at 120°C until half of the solution became 90 ml, so that the water was thoroughly removed. After reaching that volume heating stopped and kept for cool down at room temperature. Always covered the flask tightly because water absorbed by the sample. After that 6.7mL methacrylic anhydride was added to the sample and magnetically stirred about 4 hours. Finally, the sample was washed with toluene several times and kept in toluene and got acrylic modified magnetic iron oxide nanoparticles. It is very essential to keep the sample into water for future reaction. For this solvent exchange needed from toluene to water. Toluene and water are practically insoluble in each other. Toluene and acetone are miscible in all proportions, as are also acetone and water. For this reasons acetone was added to sample in toluene and after that solvent was removed. This was repeated several times and as a result toluene was almost completely removed. After that, water was added to the sample

in acetone. This was repeated several times and the sample was heated at 58 °C. As a result, acetone was removed and the sample retained in water. We were thus obtained, and this sample was called S4.

2.3 Synthesis of nanocomposite

PAAm / Fe₃O₄ nanocomposite was synthesized via *in situ* process. Firstly, sonicate S4 sample for 10 min. Take 1.25 mL acrylated MION that contain 6.25 mg magnetic crosslinker into a vial and another vial 10 mg of potassium per sulfate (KPS) dissolve with 2 ml distill water and mixed it. After that this sample sonicate 5 min under 30 °C temperature. 2g acrylamide was taken with 4 ml distill water into a test tube and add 1 mg N, N'-methylene bis acrylamide (MBA) with 2ml distill water on a hot plate and set temperature 65 °C. Then add drop wise the sample acrylated MION and KPS into the test tube. Here we maintain total volume of 10 mL. Finally, free radical polymerization was allowed to continue by adding KPS solution dropwise. This mixture was heated at 65 °C and continuously stirring at 1200 rpm for 4 hours [2,3]. After completion of the reaction, resulting nanocomposite was dried with vacuum oven. After dried this sample collect. we also synthesize that contain 3.125 mg Acrylated MION and without acrylated MION follow this procedure and also maintain total volume of 10 mL. To get such type of nanocomposite we synthesized nanocomposite varying the amount of Acrylamide, magnetic crosslinker. The varied ratio of Acrylamide, and magnetic crosslinker is 2:0, 2:0.00625, 2:0.00312. We also follow this ratio 4:0 g, 4:0.025 g, 4:0.05 g. For the synthesis of nanocomposite, we also synthesize without N, N'-methylene bis acrylamide, (MBA) to see the crosslinker effect.[4]

2.4 Swelling study of hydrogel

2.4.1 Kinetics study of swelling

The dried hydrogels were left to swell in distilled water at room temperature for different time interval. Firstly, cut the sample small pieces. Take the sample into 100mL beaker and add distill water. After 0.5 hour we keep the sample on a Petri disk and weight the sample by the digital balance and measure the swelling ratio. At fixed time interval that 1h, 2h, 4h, 8h, 16h, 22h swelling ratio was measured by following this process. Swollen gels were removed from swelling medium at regular interval of time and dried superficially with

filter paper, weight and placed in the same medium. The measurements were continued until a constant weight was reached for each sample. [5]

2.4.2 pH dependent swelling study

In this series of experiments, swelling ratio for the synthesized hydrogels was measured in different pH solutions ranged from 3 to 11. Since the swelling capacity of all “anionic” hydrogels is appreciably decreased by the addition of counter ions (cations) to the swelling medium, Therefore, stock NaOH (pH 10.0) and HCl (pH 1.0) solutions were diluted with distilled water to reach desired basic and acidic pHs, respectively. No buffer solutions were used. Measure the swelling ratio at different pH. Swollen gels were removed from swelling medium at regular interval of time and dried superficially with filter paper, weight and placed in the same medium. The measurements were continued until a constant weight was reached for each sample. The swelling ratio and also the swelling percentage was measured. Follow this procedure we measured the swelling ratio at pH range from 3 to 11.[6,7]

2.5 Fourier transform infrared analysis (FTIR)

The infrared spectra of the polymer nanocomposite, magnetic iron oxide nanoparticle, silica coated magnetic iron oxide, aminated magnetic iron oxide (silica coated iron oxide), acrylated magnetic iron oxide nanoparticle, polymer nanocomposite, sample recorded on an FTIR spectrometer in the region of $4000 - 400 \text{ cm}^{-1}$. For FTIR every step we keep some sample into vial and oven dried under 50°C . A small portion of dried sample grinded as much as possible. On the otherhand silica coated magnetic iron oxide nanoparticles, aminated magnetic iron oxide (silica coated magnetic iron oxide nanoparticles), acrylic modified magnetic iron oxide nanoparticles were dried and grinded into a mortar with a pestle. After synthesis polymer nanocomposite we cut the gel as small size and put into the vial for dried here only use freeze dryer. After dryer it also grinded as much as possible. The powder mixture was then compressed in a metal holder under a pressure of 8–10 tons to make a pellet. The pellet was then placed in the path of IR beam for measurements.

2.6 Field emission scanning electron microscopy (SEM)

The surface morphology of the synthesized magnetic iron oxide nanoparticles, silica coated MION, amino propyl(silica coated magnetic iron oxide nanoparticles), acrylic modified magnetic iron oxide nanoparticles, PAA/Fe₃O₄ nanocomposite was adopted using Field Emission Scanning Electron Microscopy (FE-SEM). For SEM sample preparation we use glass slide. For clean and removing organic substrate use piranha solution in 2 : 1 molar of sulfuric acid(H₂SO₄) and Hydrogen peroxide(H₂O₂) After drying the glass slide we put 1 or 2 drop from the more and more diluted sample. The completely freeze dried samples were glued on a conducting carbon strip. The sample loaded strip was then mounted to a chamber that evacuated to $\sim 10^{-3}$ to 10^{-4} tor and then a very thin platinum layer (few nanometers thick) were sputtered on the sample to ensure the conductivity of the sample surface. The sample was then placed in the main FESEM chamber to view its surface. The microscope was operated at an accelerating voltage of 5.0 kV. The system was computer interfaced and thus provides recording of the surface images in the computer file for its use as hard copy.

2.7 Energy dispersive x-ray (EDX) spectroscopy

Elemental analysis of the synthesized magnetic iron oxide nanoparticles(MION), silica coated MION, acrylic modified magnetic iron oxide nanoparticles, PAAm/Fe₃O₄ nanocomposite were performed by EDX spectra. The glass slide that contain these sample were placed on a 1 cm × 1cm conducting steel plate. The steel plate was then placed on a conducting carbon glued strip. The sample was then placed in the main FESEM chamber integrated.

2.8 X-ray diffraction (XRD)

Magnetic iron oxide nanoparticles, modified Fe₃O₄ were analyzed for their X-ray diffraction pattern in the powder state. The powder samples were pressed in a square aluminum sample holder (40mm × 40mm) with a 1 mm deep rectangular hole (20mm x 15mm) and pressed against an optical smooth glass plate. The upper surface of the sample was labeled in the plane with its sample holder. The sample holder was then placed in the diffractometer.

2.9 X-ray photoelectron spectroscopy (XPS)

XPS measurements were performed on samples that had been prepared within 4 days using the AXIS ULTRA spectrometer (Kratos Analytical). The base pressure in the analytical chamber was $<3 \times 10^{-8}$ Pa. The monochromatic AlK α source ($h\nu=1486.6\text{eV}$) was used at a power of 210 W. The photoelectron exit angle was 90° , and the incident angle was 35.3° from the plane of the surface. The analysis spot was $400 \times 700\mu\text{m}$. Survey scans were collected for binding energies from 1100 to 0eV with analyzer pass energy of 160eV and a step of 0.35eV. The high-resolution spectra were run with a pass-energy of 20eV and a step of 0.1eV. Relative sensitivity factors (RSFs) for different elements were as follows: 1 for C (1s), 1.8 for N (1s), 2.93 for O (1s), 0.26 for Si (2s), 0.27 for Si (2p). Only one set of XPS scans was performed on a given sample; therefore, XPS analysis before and after surface reactions were performed on different samples.

2.10 Mechanical strength of hydrogel

The tensile stress-strain curves were examined with a 500N load cell at room temperature. The measurement conditions were as follows: cross/head speed=30 mm/min, distance between two jaws=20mm, hydrogel size=5mm height*40mm length*8 mm weight. The initial cross section ($5 \times 8 \text{ mm}^2$) was applied to calculate the tensile strength and Young's modulus. The tensile strength and elongation at break were recorded and the modulus was determined from elongation of 50-150 %.[8]

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Chapter 3

Results and Discussion

3.1 Synthesis of magnetic nanocomposite

3.1.1 Synthesis of magnetic crosslinker

Crosslinker plays a vital role for linking one polymer chain to another polymer chain. There are three types of crosslinker. These are 1D crosslinker, 2D crosslinker, 3D crosslinker. Among these crosslinkers are commercially available and some are lab prepared. Among different crosslinkers 3D crosslinker has comparatively larger surface area and reactive sites than conventional linear or planar crosslinker. In our study we developed a magnetic 3D crosslinker. For magnetic property magnetic iron oxide nanoparticles (MION) were synthesized via co-precipitation method. Where we used 2:1 molar ratio of Fe^{2+} and Fe^{3+} salt and 1.5 M NaOH solution. During this reaction we deoxygenated water by passing N_2 gas so that produced magnetic iron oxide nanoparticles could not come in contact with O_2 . If these nanoparticles come in contact with air, it oxidized to Fe_2O_3 . For these reason coating of nanoparticles is very essential and this urgent work was done by coating with silica using tetraethylorthosilicate (TEOS) in basic medium (25% NH_3 solution). This was done in two steps. First step, after addition of TEOS stirred 10h without heat. Next step was carried out applying heat at 50°C for 12h so that completely coated with silica. Here act as a stabilizer, Prevent magnetic particle-particle bipolar interaction. Now our target is to add such group which would link up with polymer chain. For this purposes silica coated MION was functionalized with amino group followed by treated with APTES (3-aminopropyltriethoxysilane). In addition to the $-\text{OH}$ groups on the surface can easily covalent with $-\text{NH}_2$ group. Finally treated with methacrylic anhydride and found acrylic group at the end of the chain. This acrylic group is able to link polymer chain and acrylated MION act as magnetic crosslinker. [1-9]

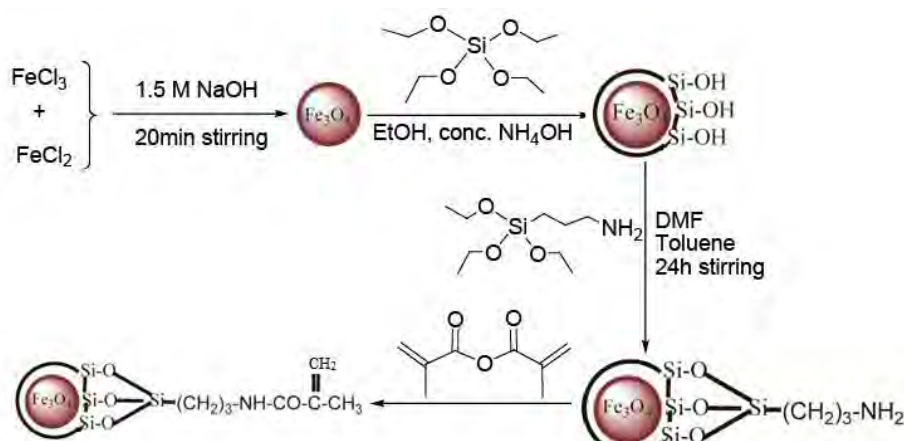


Figure: 3.1 Schematic representation of magnetic crosslinker

3.1.2 Polymer nanocomposite Synthesis

For the synthesis of nanocomposite, magnetic crosslinker were sonicated 10 min so that the aggregated particles apart from each other. After that 2g acrylic acid, 1 mg N, N'-methylenebisacrylamide, (MBA) dissolve and mixed with each other into the test tube. Again magnetic crosslinker with KPS sonicate 5 minutes and this case temperature kept below 30°C. After that this solution add dropwise into the test tube and this case temperature was 65°C. During sonication dissolved potassium per sulfate (KPS) which is as initiator added rapidly. After the addition of initiator KPS and magnetic crosslinker the polymerization start. As we use magnetic crosslinker which has manifold reactive sites hence many more polymer chains linked by the crosslinker. We use this magnetic crosslinker to prepare superabsorbent polyacrylamide hydrogel via in situ polymerization using potassium per sulfate as initiator.

Our target is to make such a nanocomposite which act as multi responsive that magneto- and thermo- responsive nanocomposite. For these highly reactive sites of magnetic crosslinker are able to link with many polymer chains that contain magnetic nanoparticle. For pH- responsiveness we use acrylic acid.[10-18]

3.2 Functional group analysis using FTIR

the FTIR spectra of magnetic iron oxide nanoparticles (MION) and modified MION. The 1st one is for MION. The strong absorption peak at 578 cm⁻¹ attribute to the vibration of Fe-O bond and this is the characteristic peak of naked Fe₃O₄. This also indicates the resonance with oxygen. The peak at 1628 cm⁻¹ indicates the presence of O-H vibration in H₂O and the peak at 3425 cm⁻¹ is attributed to the stretching vibrations of OH adsorbed on the surface of the Fe₃O₄ nanoparticle. The 2nd one is for silica coated iron oxide nanoparticle. The strong peaks at 1085 cm⁻¹ and 802 cm⁻¹ have been assigned to the asymmetric and symmetric linear stretching vibrations of Si-O-Si bonding. The bending vibration absorption peaks of Si-O-Si and Si-OH were observed at 461 cm⁻¹ and 962 cm⁻¹ respectively. These indicates that the successfully coating of silica layer another surface of magnetic iron oxide nanoparticles.

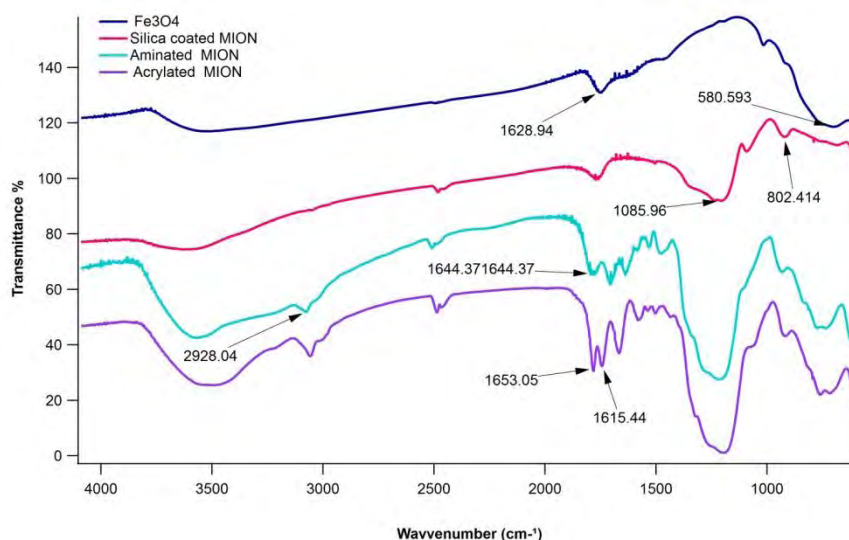


Figure 3.2 FTIR spectra of Fe_3O_4 , silica coated MION, aminated MION and acrylated MION

In the spectrum of aminated MION, the two bands at 3422 cm^{-1} and 1644 cm^{-1} can be referred to the N-H stretching vibration and NH_2 bending mode of free NH_2 group, respectively [5]. The antisymmetric C-H stretching vibrations appeared at 2935 cm^{-1} , and the bending vibration absorption peaks of $-\text{CH}_2$ and $-\text{CH}_3$ appeared in 1492 cm^{-1} and 1385 cm^{-1} , respectively; The C-N stretching vibrations appeared at 1335 cm^{-1} . The deformation vibration absorption peak of N-H appeared at 1553 cm^{-1} . These indicate the successful introduction of APTES to the surface of silica coated magnetic NPs. The 4th one is for acrylated MION. The sharp peak at 1615 cm^{-1} is the characteristic peak of C=C bond. From this we confirm that acrylic group is successfully added. All characteristic peaks of MION, Silica coated MION, Aminated MION, Acrylated MION that are found in our study are shown in table 3.1.

Table 3.1 Characteristic IR band of MION, Silica coated MION, Aminated MION, Acrylated MION

Wavenumber (cm^{-1})	Interpretation
580	Fe-O vibration
1627	O-H vibration in H_2O
3425	O-H stretching vibration in H_2O
804	Si-O-Si Symmetric stretching vibration

1080	Si-O-Si Asymmetric stretching vibration
962	Si-OH bending vibration
3422	N-H stretching vibration
1650	NH ₂ bending vibration
2935	C-H Anti symmetric stretching vibration
1492	-CH ₂ bending vibration
1385	-CH ₃ bending vibration
1335	C-N stretching vibration
1557	N-H vibration
1655	C=C vibration

3.3 Magnetic test of magnetic crosslinker and hydrogel

This test was performed two states like when the particles were in water and when the particles were dried. The magnetic test of the synthesized nanocomposite was carried out using magnetic bar. If the particles are magnetic then they are attracted by magnet and if the particles are not magnetic they are not attracted by the magnet. In the first case, the particles of our magnetic crosslinker which is dispersed in water strongly attracted by magnet. In the second case our magnetic crosslinker based nanocomposite hydrogel were in water but when a magnetic bar took to the vial then the particles were loosely attracted by the magnetic bar compare to our magnetic crosslinker. Because of our magnetic crosslinker based nanocomposite hydrogel also absorb a large amount of water and also create a big size. Our magnetic bar is not so strong as much as. So it is loosely attracted by the magnetic bar. Without magnetic crosslinker based hydrogel not attracted by the magnet.

That is shown in below.

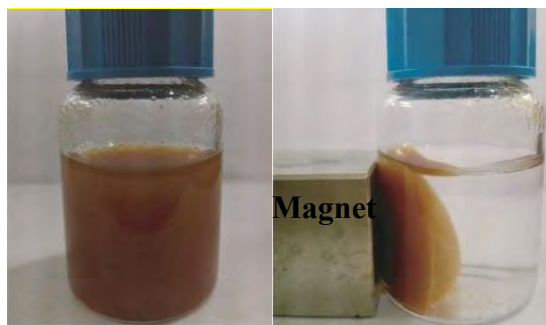


Figure 3.6 Magnetic crosslinker dispersed in water and get attract to magnet

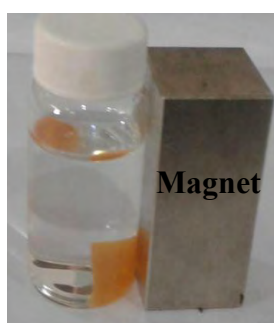


Figure 3.7 polymer nanocomposite hydrogel with magnetic crosslinker is attracted by magnet.

3.4 X-ray photoelectron spectroscopy (XPS)

We use the surface sensitive technique X-ray photoelectron spectroscopy (XPS) to conform the bonds that were formed in aminated MION, acrylated MION. The XPS spectra of aminated MION, acrylated MION is shown in Figure 3.8. In order to assess and confirm the bonds that were formed in aminated MION, acrylated MION, we use the most complicated, expensive and surface sensitive technique that is X-ray photoelectron spectroscopy.

From the Figure the XPS spectra of the two samples are similar because there is no elemental change occurred during aminated MION and the acrylation.

All XPS spectra of the presence of C1s, O1s, N1s, Si2s, Si2p. The spectrum of aminated MION the dominated species are O and Si. The species C and O are found at the binding energy 283.6eV and ~532.8eV, respectively. The intensity of N1s very small in aminated

and acrylated MION. The presence of nitrogen on the surface was detected from its characteristic emission peak at $\sim 400\text{eV}$ and is due to the amination of silica coated MION and amide formation in acrylated MION.

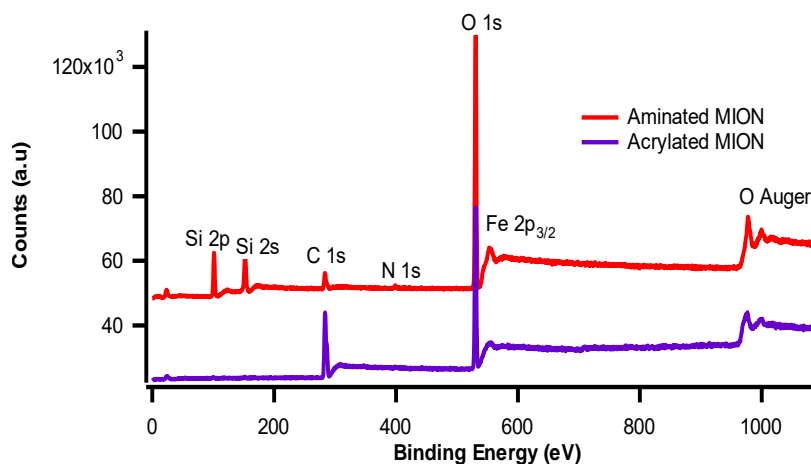


Figure 3.8 Widescan spectra of aminated MION, acrylated MION

The binding energy at 100.4eV and 151.2eV the existence of silicon on the surface as the magnetic iron oxide nanoparticles were coated with silica. All the two spectra have a common oxygen auger electron spectrum at binding energy 978eV . Widescan spectra give idea about the elemental composition on the surface only. But their bonding nature could not be known by this spectrum.

3.4.1 High Resolution C1s Spectra

Widescan spectra showed the presence of C but the valence state of C in the molecule could not be explained. For this high resolution of C1s spectra are taken. By this spectrum it is possible to know how the C bonded with other elements.

High resolution C1s spectra of aminated MION, are shown in figure 3.9. The C1s peak of aminated MION can be scattered into three components peak. According to the binding energy these peaks are assigned to the carbon of C-Si (284.3eV), C-C (285.3eV) and C-N (285.8eV) [6-8, 9]. The binding energy 284.5eV corresponds to a carbon atom bound only to silicon atom.

C-Si bond found due to the reaction between silanol groups of silica coated MION and 3-aminopropyltetraethylorthosilicate. Binding energy 285.3 ascribes to a carbon atom

bound only to other carbon atoms and hydrogen atoms. 285.8eV corresponds to a carbon bound to single nitrogen atom. This is due to the terminal amino group linked with methylene group.

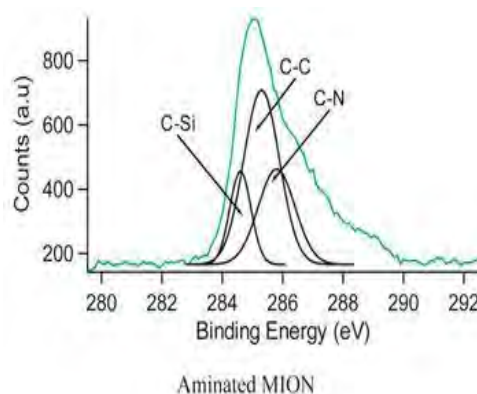


Figure 3.9 High resolution of C1s spectra of aminated MION

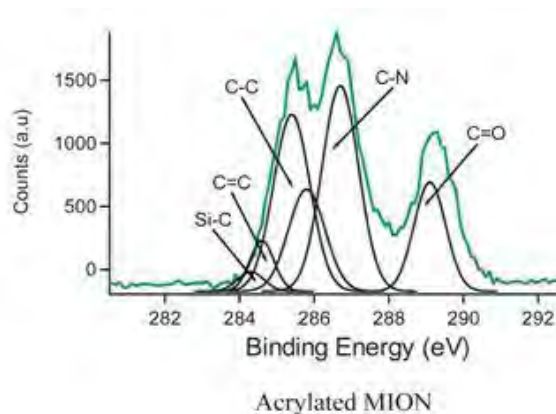


Figure 3.10 High resolution of C1s spectra of acrylated MION

After acrylation, there is significant changes appeared at the peak position. Now the peaks can be assigned into six components which are C-Si (284.3eV), C=C (284.6eV), C-C (285.4eV), N-C=O (286.6), C=O (288eV) and O-C=O (289.3eV) [10, 11]. An additional peak compare to aminated MION found at 284.6eV which ascribes the presence of C=C and it is the proof of the acrylation of the aminated MION. Another evidence for this is to the binding energy at 286.6eV for N-C=O which is mainly for the amide formation. The binding energy 289.3eV is due to the presence of O-C=O. This is most probably due to the presence of unreacted methacrylic anhydride.

3.4.2 High resolution N1s spectra

From the widescan spectrum it was seen that N was present but the bonding nature of N with other elements was unknown. The bonding nature of N with other elements high resolution of N1s spectra were carried out.

The high resolution N1s spectra of aminated MION are shown in Figure 3.11. In the case of aminated MION, there appears one peak at binding energy 400eV which describes the presence of primary amine at the end of chain.

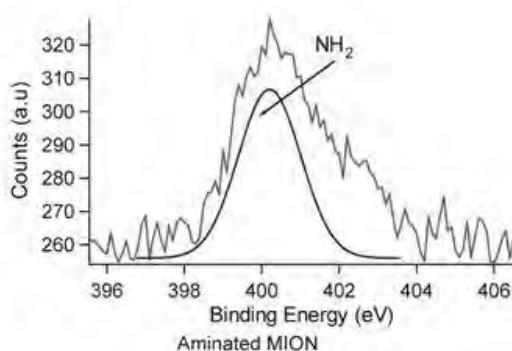


Figure 3.11 high resolution of N1s spectra of aminated MION

In the spectra of acrylated MION there are two scattered peaks appeared. First one is at binding energy 400eV and second one is for 402eV. Binding energy 400eV is attributed to a nitrogen atom linked to a carbon atom by a single bond (amide bond) [6], and the second peak at 402eV is assigned to the protonated amino group -NH_3^+ [7]. This indicates that all the amino groups in the aminated MION are not converted to amide.[19-22]

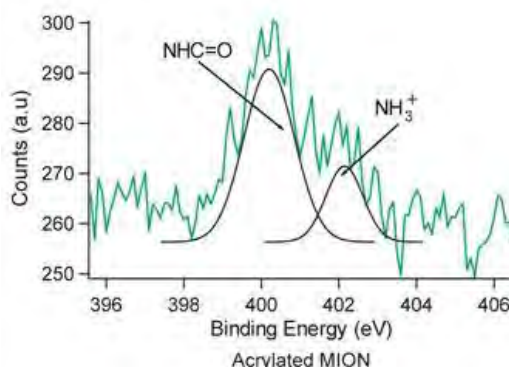


Figure 3.12 High resolution of N1s spectra of acrylated MION

3.5 X-ray diffraction analysis (XRD)

The crystalline properties and phase identification were characterized by X-ray diffraction using a Philips X'pert PRO X-ray diffractometer. The diffractogram was obtained by using Cu-K α radiation ($\lambda = 0.15406\text{nm}$) in the range 10 to 70 $^{\circ}$ C with steps of 0.021 and acquisition time of 1.0s/step.

The X-ray diffraction patterns of Fe₃O₄ and nanocomposite were shown in figure 3.13.

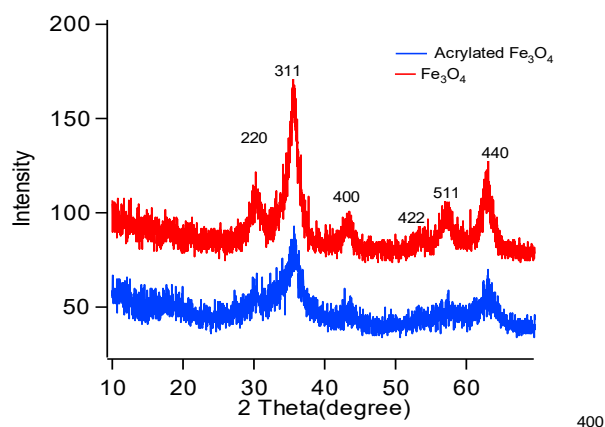


Figure 3.13 XRD spectra of Fe₃O₄, acrylated MION

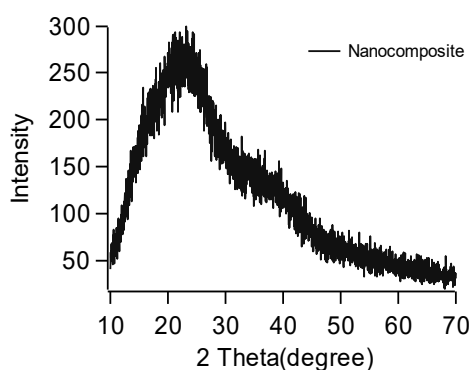


Figure 3.14 XRD spectra of Fe₃O₄ nanocomposite

The Fe₃O₄ particles have six diffraction peaks at $2\theta = 30.24^{\circ}$, 35.56° , 43.38° , 53.88° , 57.34° , and 62.82° . The peaks are associated with the crystallographic planes or Miller indices of (220), (311), (400), (422), (511), (440), respectively. It can be seen that, the peak position and relative intensities of the diffraction peaks are consistent with the characteristic peaks

of standard Fe_3O_4 crystal [Joint Committee on Powder Diffraction Standards (JCPDS card No.19-0629)] . The diffraction peaks of Fe_3O_4 were indexed cubic spinel structure.

Nanocomposite samples did not exhibit any sharp peaks in XRD patterns, with only a broad peak at $\sim 20^\circ$ attributed to the polymer networks.

From XRD data the average crystallite size was determined by Debye-Scherrer equation .

$$D_{hkl} = \frac{0.9\lambda}{\beta \cos\theta}$$

Where D_{hkl} is the average crystallite size, $\lambda = 0.15406$ nm corresponds to the wavelength of Cu $K\alpha$ radiation and β_{hkl} is full width half maximum (FWHM) in the 2θ scale of the diffraction peak from (hkl) crystal planes, and θ is the Bragg angle. The crystallite size was determined by taking the average of the sizes at the peaks D_{220} , D_{311} , D_{400} , D_{422} , D_{511} and D_{440} and it was found to be about 5.32nm. The lattice parameter, “a” was determined by the following equation

$$d_{hkl} = \frac{a}{\sqrt{h^2 + l^2 + k^2}}$$

Where d = interplanar spacing

The lattice parameter was found to be 8.36 Å, which is lower than the values reported JCPDS card No. 19-629(a=8.39), but the value is very close to the theoretical value [17, 18, 19]. The lattice strain was also determined as 0.0237. The second XRD pattern for the acrylic modified Fe_3O_4 and from the x-ray diffractogram found that peak position is remained same and it is the evident that there is no phase change occurred. The 3rd XRD pattern is for nanocomposite. From the pattern it is seen that, the nanocomposite has no sharp peak but at 20° attributed to the polymer networks.[23]

3.6 Surface morphology study using scanning electron microscope (SEM)

Surface morphology of synthesized magnetic iron oxide nanoparticles (MION), acrylated MION and magnetic crosslinker based nanocomposite was studied by FE-SEM.

FE-SEM of magnetic iron oxide nanoparticle is shown below:

From figure 3.15 it is seen that particles of MION are spherical in shape and highly uniform in size. The average particle size is about 10-15nm. Some particles are very small and about 3-5nm and some particles aggregate due to large surface area and also magnetic attraction among them. From high resolution image it is clearly seen that particles are finely arranged.

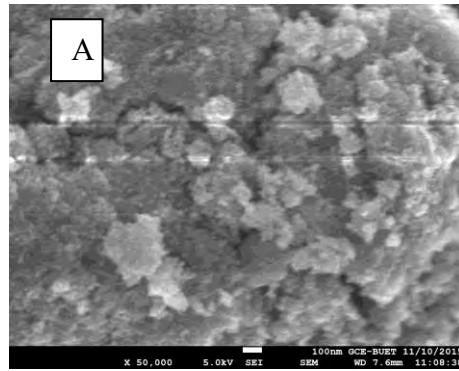


Figure 3.15 (A) SEM image of MION at Resolution x 50,000

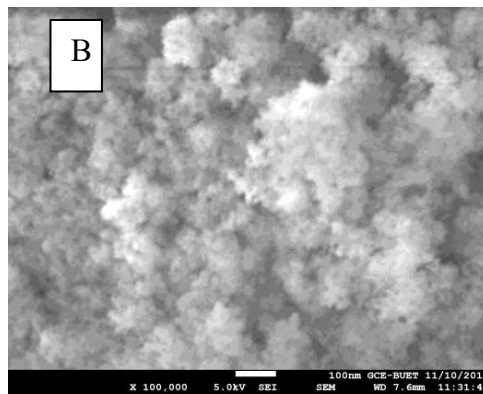


Figure3.15 (B) SEM image of (MION) at Resolution x 100,000.

Surface morphology of silica coated MION characterized by FE-SEM and the image shown in the figure3.16

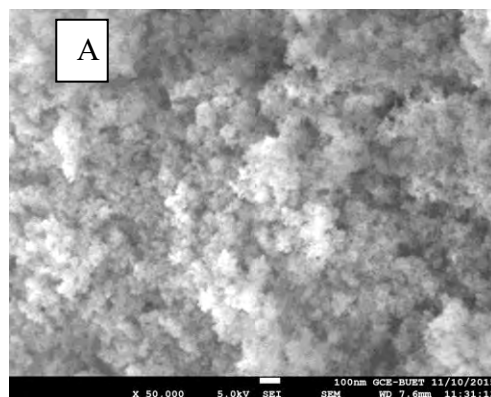


Figure 3.16 (A) SEM image of Silica coated MION at Resolution x 50,000

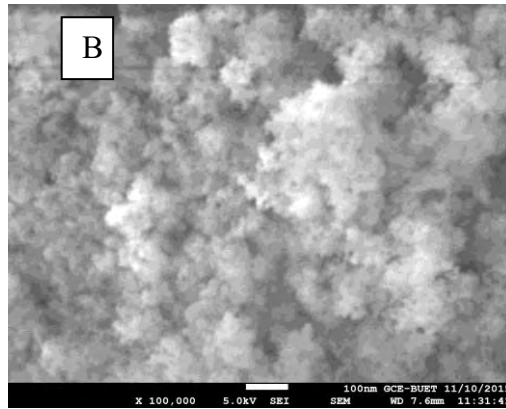


Figure 3.16 (B) SEM image of Silica coated MION at Resolution x 100,000.

From the figure 3.16 it is seen that the average size of silica coated MION is about 15-20 nm.

Surface morphology of acrylated MION characterized by FE-SEM and the image shown in the figure3.16

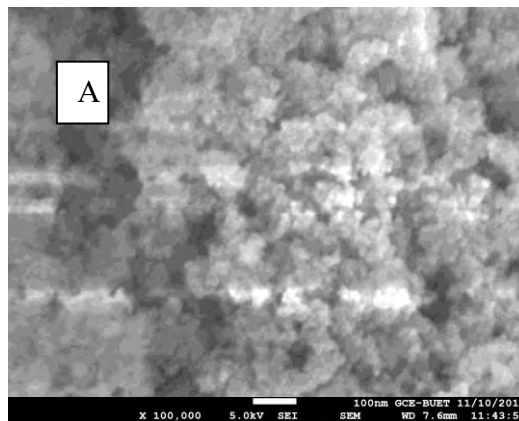


Figure3.17 (A) SEM image of acrylated MION (A) Resolution x 100,000

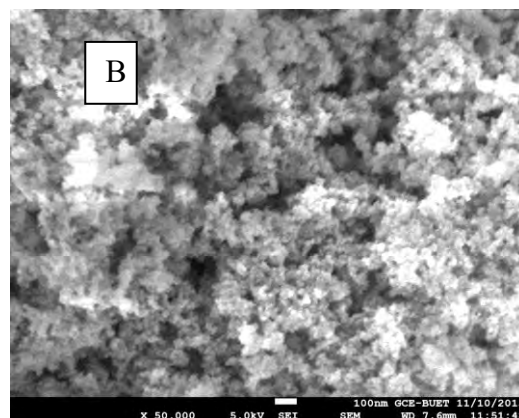


Figure 3.17 (B) SEM image of acrylated MION (A) Resolution x 50,000

From the figure 3.17 it is seen that particles of acrylated MION are cubic in shape and size is about 20-30nm. Particles are randomly arranged and also seen that they are one on another. The shape of the particles is all most similar which are seen from the high resolution image. [24,25]

3.7 Energy dispersive x-ray spectral analysis

Elemental analyses of the Fe_3O_4 , silica coated MION, acrylated MION and nanocomposite have been performed by Energy Dispersive X-ray (EDX) method. The EDX patterns are presented in Figure 3.20-23. The peaks observed at 0.277, 0.392, 0.525, 1.739 and 6.398 keV, for K lines of C, N, O, Si and Fe respectively. The percentages of Fe, O, Si, N, and C were determined from the intensity of the lines and are summarized in Table 3.2.

From EDX analysis we found mass% of the composition and atom% of the composition.

Table 3.2 Elemental composition of magnetic and nanocomposite compound

Sample	Iron (Fe) %	Oxygen (O) %	Silicon (Si)%	Nitrogen (N)%	Carbon (C)%
Fe_3O_4 (MION)	31	69			
Silica coated MION	14	42	43		
Acrylated MION	25	40	12	4	19

From the table it is seen that Fe_3O_4 has iron content 31% and oxygen content 69%. It is approximately equal to the theoretical value of Fe_3O_4 .

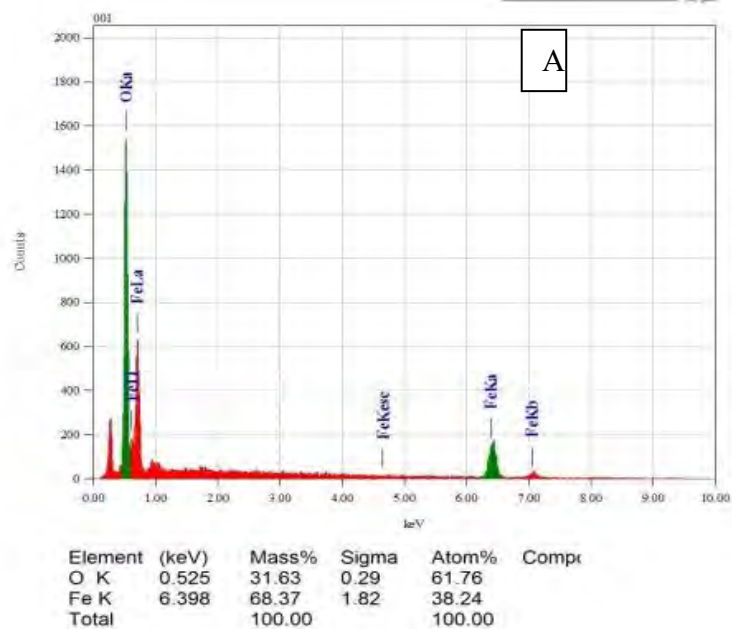


Figure 3.20 EDX spectra of Fe₃O₄

After silica coated of MION we expect we found a new element of Si, here we also found Si that is shown figure 3.21

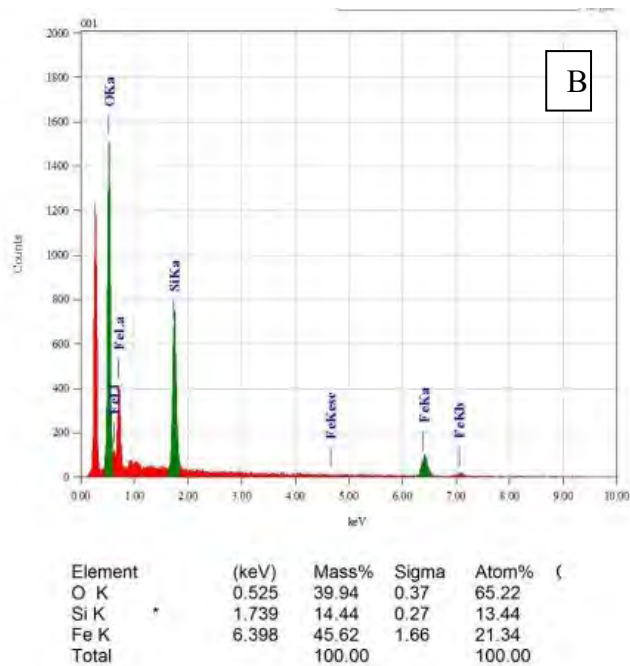


Figure 3.21 EDX spectra of silica coated of MION

Chapter 3: Results and Discussion

After acrylation we found acrylated MION and here we expect that two new element are found because of aminated MION and acrylate MION. For this we found as a new element C and N and we also found this. So our targeted all element are found by EDX analysis

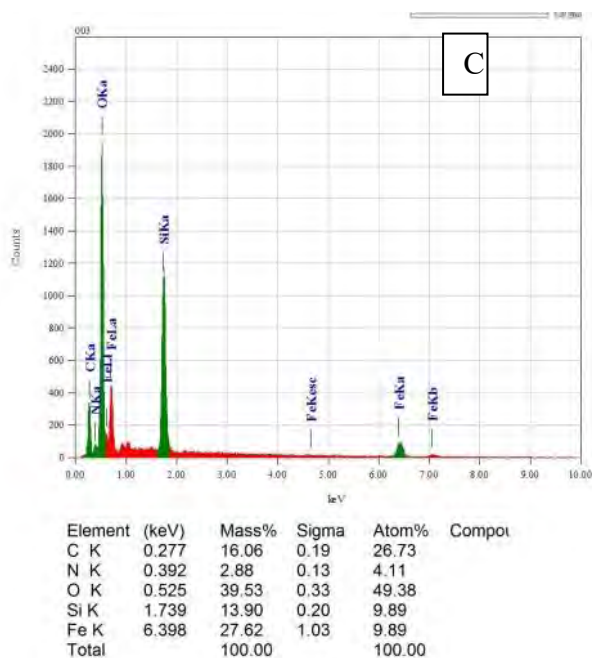


Figure 3.22 EDX spectra of acrylated MION

We synthesized different ratios of our magnetic crosslinker based hydrogel. This are 4:0.0125 g, 4:0.025. For this we found Fe and O amount are different. When we use 12 mg magnetic crosslinker then we found mass% of Fe 0.78 and O 34.07. Besides this when we use 25 mg of magnetic crosslinker we found mass% of Fe 1.05, O 42.44. This indicates when we increase magnetic crosslinker increase the mass% of Fe and O. EDX spectra of nanocomposite is given below[24-25]

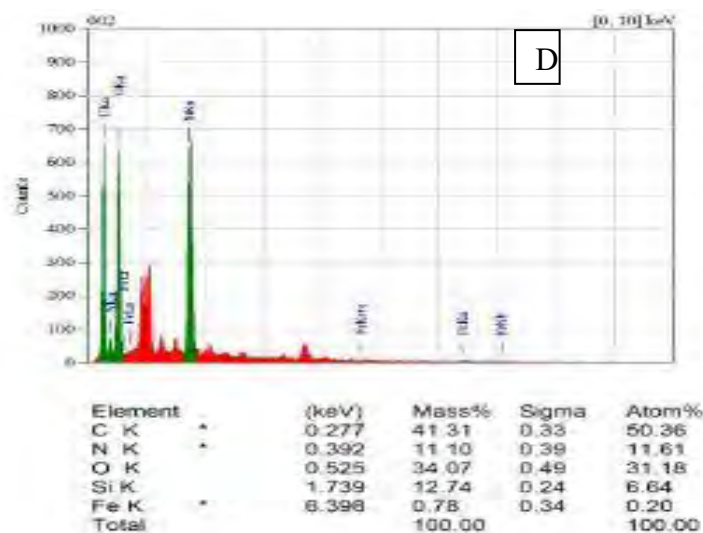


Figure 3.23 (A) EDX spectra of nanocomposite (4.0.0125g)

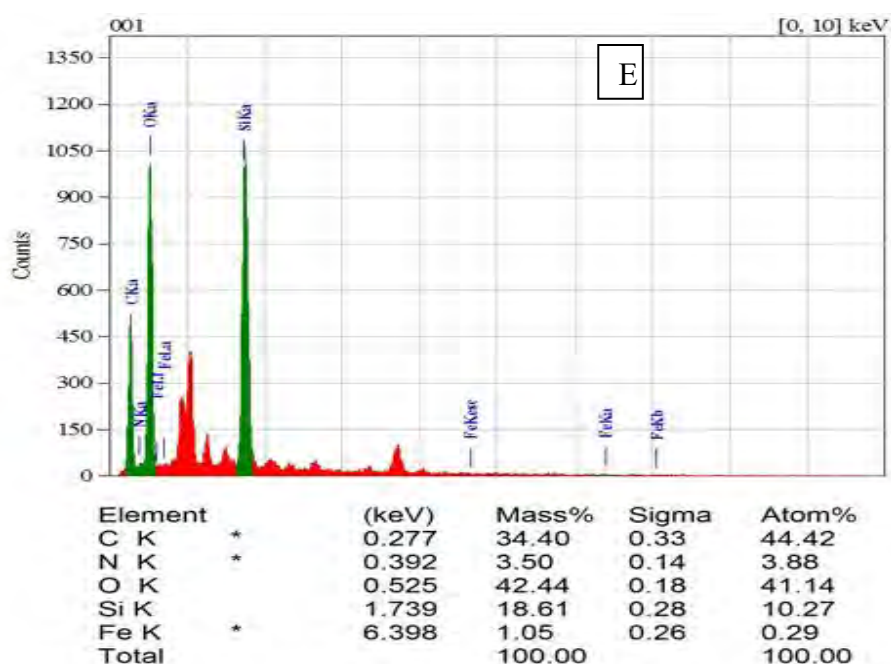


Figure 3.23 (B) EDX spectra of nanocomposite (4:0.025g)

3.8 Mechanical strength analysis

The result shows that a large change in hydrogel mechanical behavior depending on the increasing of crosslinkers dimension. Tensile strength and Young's modulus both are significantly dependent on the crosslinker. When we used synthesized 3D crosslinker to form hydrogel Tensile strength and Young's modulus both are significantly higher than the 1D or linear cross linker (BIS PAA).As in synthesized 3D cross linker covalent bond is

formed it can take high stress. For BIS PAA stress vs strain curve shows that its highest stress upto 0.1 MPa and break of elongation is 1000%. In the same time synthesized hydrogel containing 0.3% crosslinker stress vs strain curve shows that its highest stress more than 0.2 MPa and break of elongation is about 1700%. So here we see that our synthesized hydrogel shows significant higher stresses and also break of elongation then other linear or 1D dimensional crosslinkers hydrogel, which is a unique property.

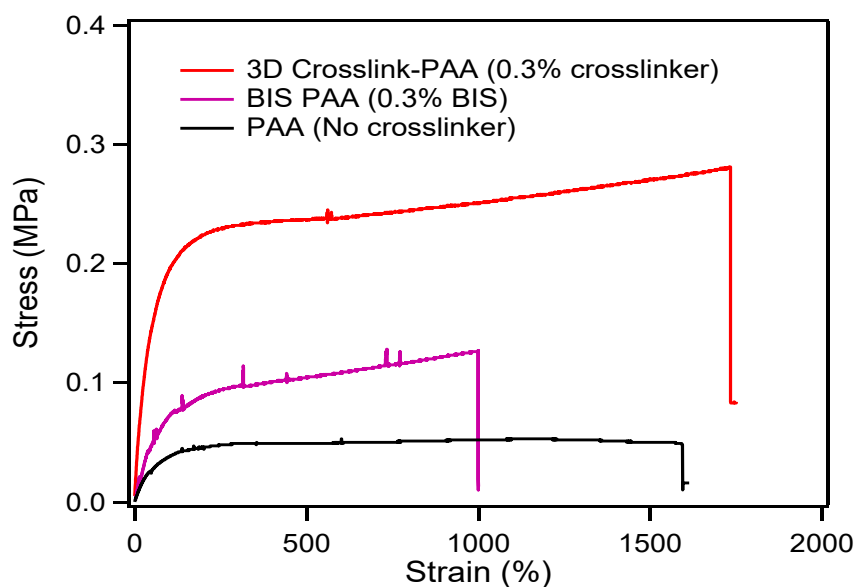


Figure: 3.24 Stress vs Strain curves of magnetic nano composite hydrogel, BIS PAA hydrogel and PAA(No crosslinker)

And also if we use higher amount of PAA then Tensile strength and Young's modulus both are significantly higher than low amount of PAA. Here more covalent bond may form with 3D crosslinker and PAA, so they show significantly higher stress and break of elongation also. If we compare this stress vs strain curve with other linear or 1D cross linker hydrogel we see same as before significant improvement of stress and elongation at break. [26]

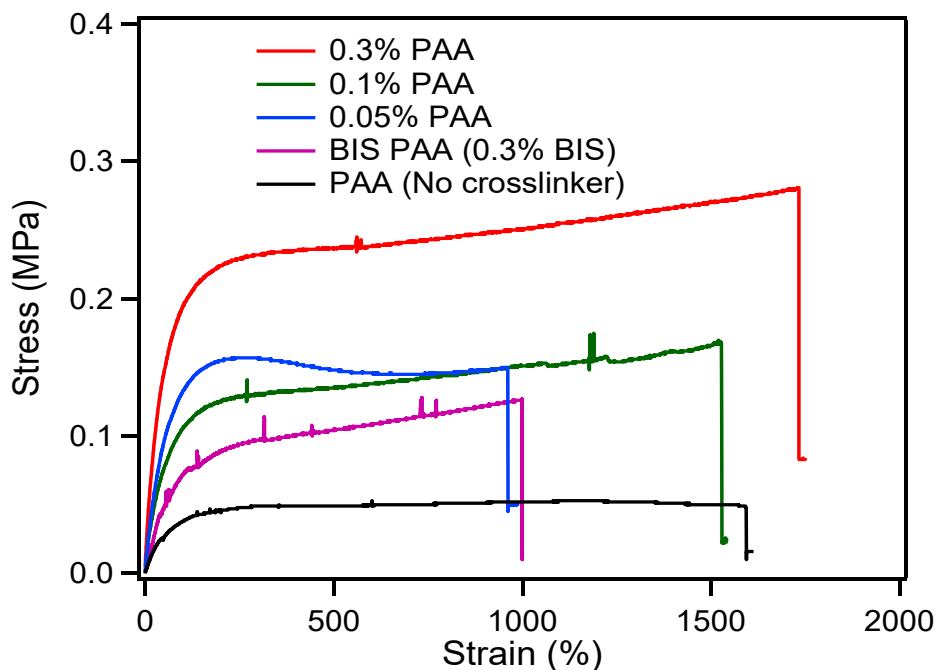


Figure:3.25 Stress vs Strain curves of magnetic nano composite hydrogel with various amount of PAA, BIS PAA hydrogel and PAA(No crosslinker)

3.9 Swelling measurement

3.9.1 Kinetics study of swelling

The favorable property of hydrogels is their ability to swell, when put in contact with a thermodynamically compatible solvent. When a hydrogel in its initial state is in contact with solvent molecules, the latter attacks the hydrogel surface and penetrates into the polymeric network of the gels. The swelling ratio is increasing with the time it keeping in the solvent increases. The highest swelling ratio shows the sample which is remain a long time like greater than 20 hours. It is rare that, when we kept the sample in solution it show slow swelling until 2 h ,but after this time swelling very rapidly. And then with the retention time it is swelling upto 400 times from the initial. We can say that our synthesized hydrogel is superabsorbent and thesis work in also unique.[27-30]

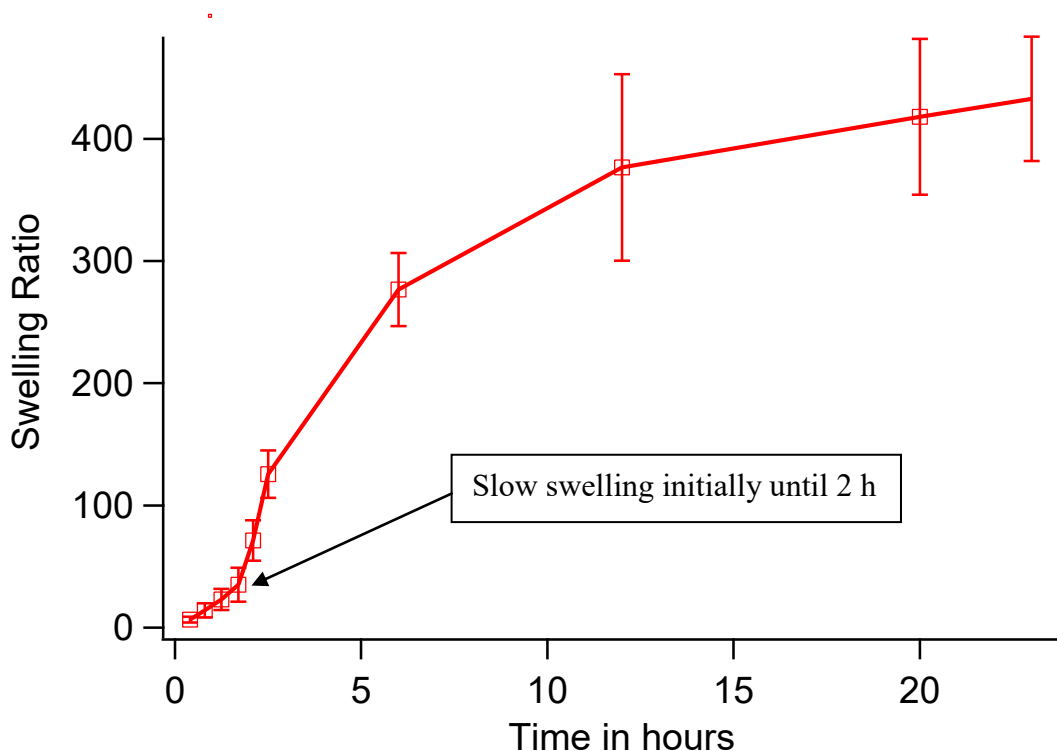


Figure 3.3 kinetics study of swelling

3.9.2 pH dependent swelling study

The pH dependent swelling behavior of the hydrogels was measured at various pH, pH range is from 3 to 11. Since the swelling capacity of all “anionic” hydrogels is appreciably decreased by the addition of counter ions (cations) to the swelling medium, Therefore, stock NaOH (pH 10.0) and HCl (pH 1.0) solutions were diluted with distilled water to reach desired basic and acidic pHs, respectively. No buffer solutions were used. Maximum swelling was obtained near pH 8. In acidic media, most acid groups are protonated, so decreased repulsion of anionic groups leads to a decreased swelling ratio. At higher pHs (5 - 8), some acid groups are ionized and the electrostatic repulsion between acid groups causes an enhancement of the swelling capacity. The reason of the swelling loss for the highly basic solutions is the charge screening effect of excess Na^+ in the swelling media, which shield the acidic anions and prevent effective anion-anion repulsion.[30]

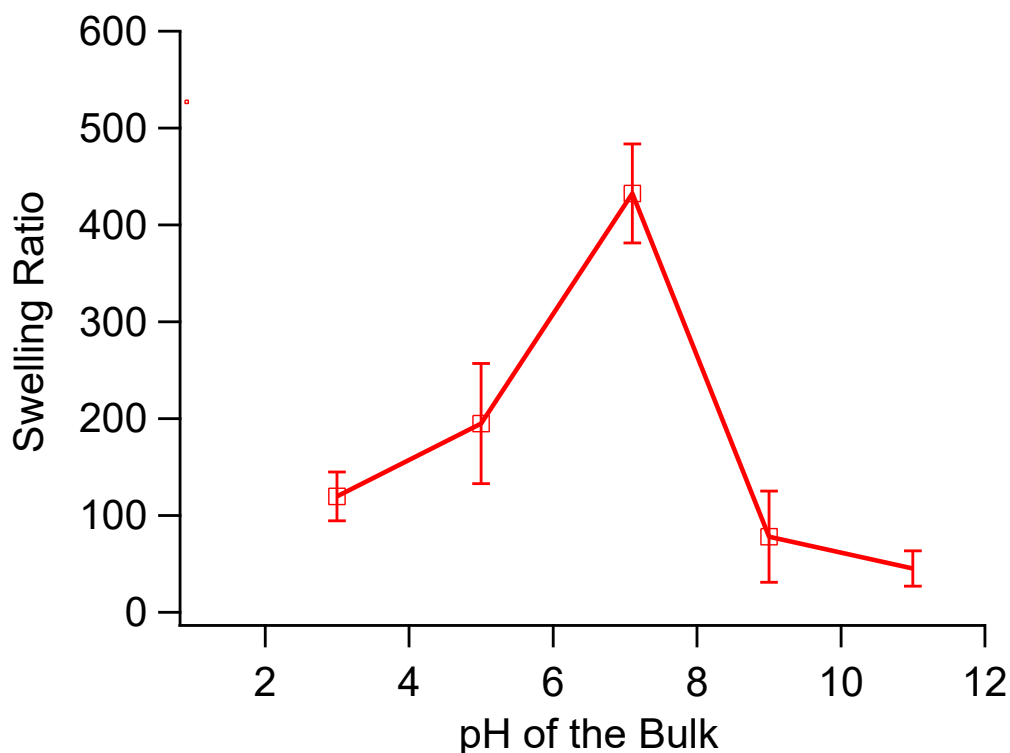


Figure 3.5 pH dependent swelling

Conclusion:

In this study, we synthesized magneto- and pH-responsive superabsorbent nanocomposite hydrogel by using a magnetic 3D crosslinker. We have introduced magnetic Fe_3O_4 nanoparticles into the hydrogel system and characterized by SEM and XRD analysis where the nanoparticle size was found uniform in size. Our targeted magnetic crosslinker as characterized by FTIR and XPS analysis revealed that acrylic groups ($-\text{NHCOCH}_3\text{CH}=\text{CH}_2$) were successfully covalently bonded to the silica-coated Fe_3O_4 . The elemental composition found by EDX and XPS supports that the synthesized material was our desired crosslinker. The high resolution of C1s results showed the presence of C-Si, C-C, C-N, C=O, C=C functional groups in the synthesized crosslinker. The high resolution of N1s spectra results also ascribed the formation of amide bond between the amino group of aminated MION and anhydride group of the methacrylic anhydride and confirmed that the formation of magnetic crosslinker. Next, we performed an in situ polymerization of acrylic acid presence of the acrylated MION crosslinker using KPS initiator. The

nanocomposite hydrogel exhibited excellent magnetic behavior confirming that our magnetic nanocomposite hydrogels are magneto- responsive. On the other hand, the swelling of the nanocomposite hydrogel was found to be pH- dependent, which proved the pH -responsive nature. From the kinetics study of swelling we assured that if increasing the retention time of the hydrogel in water increases the swelling ratio. We see that our synthesized hydrogl shows Tensile strength and Young's modulus both are significantly higher than other linear or 1D dimensional crosslinkers hydrogel, which is an unique property.

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