Synthesis, study and antimicrobial activity of new homo, copolymer maleic anhydridse with moiety heterocyclic creatinine

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Abstract: The presented work involved the preparation of polymers homo and copoly containing heterocyclic (creatinine) derived from Maleic anhydride. The first step involved the preparation of maliamic acidcreatinine derivative, by reaction of maleic anhydride withcreatinine. The second step includes the polymerization of the prepared homo and copolypolymers by free radical polymerization using (BPO) as initiator with heating. The third step involves study of degredation analysis of polymer in different at 37°C using UV-Visible spectroscopy, thermal analysis and Swelling studies. polymers were characterized by softening points, with other physical properties, FTIR, H1-NMR spectra, and thermal analysis (DSC), screening of the antimicrobial activity of the prepared polymers was evaluated against 3 types of bacteria.

Keyword: polymer, creatinine, maleic anhydride, antimicrobial

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I. Introduction

Polymers are macromolecules built up by the linking together of large numbers of much smaller molecules(!). The small molecules that combine with each other to form polymer molecules are termed monomers, and the reactions by which they combine are termed polymerizations, Some reactive olefin monomers, with heteroatom groups being more reactive than alkyl or aryl groups, Heterocyclic monomers that are polymerized are lactones, lactams, and cyclic amines. Upon addition of an initiator, cyclic monomers go on to form linear polymers. The reactivity of heterocyclic monomers depends on their ring strain. Heterocyclic chemistry is one of the largest branches of organic chemistry (2-5) It is particularly important in nature because of the wide variety of industrial significance and ysiological activities There has been recent interest in developing, synthetic routes for preparation of significant. Biopolymers, havedrawn considerable attention ide signing ,hydrogel wit ideally applied biopolymer is alginicafford the past few decades, biodegradable polymers have been, applied as carriers for controlled delivery of low molecular weight drugs as well as bioactive proteins فريال 6,7]. Biodegradable polymers, either synthetic or natural, are capable of being cleaved into biocompatible by,products through chemical or enzyme-catalyzed hydrolysis. This biodegradable property makes it possible to, implant them into the body without the need of subsequent removal by the surgical operation Drugs formulated, with these polymers can be released in a controlled manner, by which the drug concentration in the target site is,enhanced. The release rates of the drugs from biodegradable polymers can be controlled by a number of factors, such as biodegradation kinetics of the polymers [8-10], ysicochemical properties of the polymers and drug,[11,12], thermodynamic compatibility between the polymers and drugs [13], and the shape of the devices [14-16]. Maleic anhydride is an excellent monomer which can provide reactive anhydride or carboxylic, groups with nucleoilic molecules. [17,18]. MAN has an extremely low tendency to homopolymerize in a radical polymerization condition; on the contrary, it copolymerizes with a variety of donor monomers[19]. The chemical, modification of synthetic polymers allows the control of their mechanical and thermal properties [20] and, expands their explicabilities, as introduction of MA on the non-polar backbone of polyolefins and rubbers has overcome the disadvantage of low surface energy of these polymers, improving their surface hydroilicityfor, the benefit of printing and coating applications, and adhesion with polar polymers(polyamides), [21-]. One of the, most common monomers in the polymer modification is MA and its isostructural analogues such as Nsubstituted, maleimides, fumaric, citraconic and itaconic acids and their esters amides, imides, and anhydrides of, these dicarboxylic acids [22]. synthetic polymers also constitute a wide area of materialscithe properties of pure, synthetic, polymers and pure biological polymers are often inadequate for producing materials with good chemical mechanical thermal, and biological characteristics, [23]. recently modification study was carried out with maleic anhydride as a grafted Copolymer[24,25]

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Experimental

A- chemicals: All chemicals used in this work were from BDH, Merk, Fluka and were used without further purification. Compounds prepared were characterized by (FT- IR ,1H-NMR & Mass spectral data). Melting points are uncorrected determined by using /SMP31 Thi-Qar University college of sciences

B-instruments: FTIR spectra were recorded using KBr discs on Shimadzu-8400 spectrootometer, Al-Shatra technical institute, Southern Technical University. H-NMR spectra were recorded on near magnetic resonance Bruker DRX System, 500 MHz Using TMS as internal standard and DMSO as solvent, Sharif University of Technology, Tehran, IranMass spectra by (Mass Selective Detector 5973 Network) 70 e/v, Sharif University of Technology, Tehran, Iran.

1-Synthesis of AmicAcid .[26-29]

In a 150 mL two-necked flask equipped with magnetic stirrer ,droop funnel and reflux condenser were placed (0.005mole 0.5 g) of maleic anhydride and (25mL) of a dry acetone. When all anhydride had been dissolved by stirring a solution of (0.005 mole , 0.5 g) of Creatinine in (25mL) of acetone was allowed to run through the dropping funnel dropwise for 30 min under cooling range (0-5 C 0). continued stirring under room temperature for 2 hrs. precipitatesettled out and filtered by suction filtration. Washed with solvent dried and recrystallized from ethanol pale browncolor,mp(171-173),85% yiled.

POLYMERIZATION:[30,31]

Polymerization of creatininemaleamicacid(homo polymerization)

0.3g. of prepared monomer was dissolved in 5ml of DMF, and 0.05% weight of dibenzoyl peroxide was added, under nitrogen atmosere, the polymerization tube was covered, heated by water bath at 900C about 1hr. The pal yellow polymer was obtained when the reaction mix. poured on ice water with 85% yield with s.p about 143-151 ^{0}C

Polymerization of creatininemaleamic acid with acrylonitrile(co-polymerization)

0.3g. of prepared monomer and acrylonitrile was dissolved in 10 ml of DMF and 0.05% weight of dibenzoyl peroxide was added, under nitrogen atmosere, the polymerization tube was covered, heated by water bath at 90° C about 1hr. The oily yellow polymer was obtained with 60% yield.

STUDIES:-

Swelling studies

swelling studies of polymer made as follows:-

Homo and co-Polymer was swollen in solution withPH 7 at 37 °C to determine the parameters of swelling and diffusion. Swollen metter removed from the water bath at regular intervals were dried superficially with filter paper..

Decomposition study:

A study was conducted to measure the speed decomposition of polymer, A mixture of polymer and solution prepared using 1.5 and 7.4 , where first solution was prepared of 0.1N(HCL), the second solution (7.4) of Phosphate buffer ,the mixture of polymer and solution was kept in a contener, 50 mg of polymer was added, kept at $37^{\circ}C$ without stirring, release sample was periodically drown with an analysis by UV spectra at 279,294 nm to determine the amount of release creatinine. A calibration curve was constructed with software built in the computerized UV otometer and the controlled release polymer were carried out in different value (4, 10) at 37 OC .result show in table 3,4, fig 1 ,2

The Biological activity [32-35]

The synthesized compounds were exposed to antimicrobial activity. Antimicrobial activities were observed for some of the compounds using strain of gram positive such as (Staphylococcus aurous), gram negative (E. coli). The antimicrobial activities of the synthesized compounds were studied by disc diffusion method. Bacterial inoculums were spread on Nutrient agar. After the inoculums dried, 6 mm diameter wells were made in the agar plate with a sterile cork borer. The synthesized compounds were dissolved in DMSO at concentration of 10⁻²M,Cefotaxime 10⁻² M was used as standard for the antibacterial activity. The Petri plates were incubated at 37°C for 24 hours. The Zone of inhibition was measured in mm and the results are listed in Table (5)

II. Results and Discussion

In this study the polymer was prepared containing creatininemoiety as pendant groups the homo and co-poly (maleamic acid) which acted as drug or antimicrobial agents. The reactions was explained as shown below:-

the structural elucidation of poly(creatininemaleamic acid) which functionalization chemistry, with carboxyl groups and lactam cycle along the backbone was analyzed using FTIR spectroscopy, and assignments for the characteristic groups were developed FTIR spectra of prepared polymer is given in Fig. (4) a new band appeared at 1710 cm-1 which confirmed the presence of carboxyl groups and the absorption of OH carboxylic was appeared at 3450-2962 cm-1 with comparing with Fig. (3) ofcreatininyl monomer alone. Supporting evidence for the structural elucidation was revealed by 1H NMR analysis, Fig. (7) indicated the content of protons at 2.6 ppm are assigned to –CH-CH- of succinamicd(7-8)ppm and the CH₂ ofcreatinien cycle was appeared at d(4.3) ppm in addition, the undetectable resonance of protons in the COOH may be due to the formation of hydrogen bonding between inter,or intra-molecules and the proton of carboxylic acid was appear at 9.3 ppm,wher" FTIR spectra of copolymer for LSe polyester is shown in the Figure 5. The pronounced peaks around 1734 cm-1 suggest the presence of carbonyl (C=O) groups from the ester. The bands centered at around 2931 cm-1 was assigned to methylene(-CH2-) groups for diacids/diols and observed in all the spectra of the polyester. The C-H symmetric stretching of aliatic-CH2- group observed at 2931cm-1. Strong vibrational mode observed at 1217cm-1 is associated with C-O-C asymmetric stretching mode aliatic ester."[11]

Differential Scanning Calorimetry (DSC)

The DSC scans were recorded at a heating rate of 10° C/min using a Perkin-Elmer Pyris I analyser. calibration standard Differential Indium was used as the ScanningCalorimetry (DSC)Analysis. DSC is a technique for determining the quantity of heat either absorbed or released when a substance undergoes a physical or chemical change. Several parameters.[10], The thermal transition of polymer show stability even 800° C, and the results shown in(fig.6).[18]

Table I Physical data of Polymers (homo,co)creatininylmaliamic

| Compound | S. po | oint ⁰ C | yield% | Rec.solvent |
|----------|---------|---------------------|--------|---------------|
| homo | 143-152 | - | 80 | DMF- WATER |
| Co-poly | oily | - | 60 | DMF- WATER |

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| Table II IR value data of polymers (homo,co-)creatininemaliamic | | | | | | | | |
|---|-----------|------------|--------------|----------|--------|--------------|------|--|
| Compound | C=O of β- | C=O of | C=O of amide | N-H bend | N-H of | $C \equiv N$ | λmax | |
| | lactam | carboxylic | | of amide | amine | | | |
| homo | 1770 | 1750 | 1650 | 1515 | 3290 | | 294 | |
| Co- | 1780 | 1755 | 1645 | 1520 | 3286 | 2255 | 279 | |

Table III absorbanceData(A) of homopolymer at =1-2 and =7.4 at λ_{max} 294 nm

| ` / | 1 J |
|----------|----------|
| A at 1-2 | A at 7.4 |
| 0.433 | 0.870 |
| 0.385 | 0.451 |
| 0.384 | 0.441 |
| 0.384 | 0.337 |
| 0.346 | 0.303 |

Note: the last reading take after 24 hr (over night)fig 4

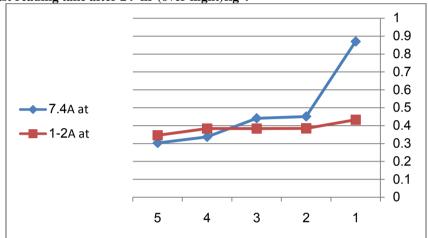


Fig1-speed decomposition of homo polymerin different buffer,pH value (pH1-2, pH 7.4) at 37°C

Table IV absorbanceData(A) of co-polymer at =1-2 and =7.4 at λ_{max} 279nm

| A at 1-2 | A at 7.4 |
|----------|----------|
| 0.983 | 0.167 |
| 0.362 | 0.166 |
| 0.361 | 0.165 |
| 0.353 | 0.114 |
| 0.557 | 0.257 |

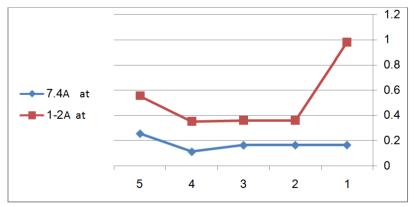


Fig-2-speed decomposition of co-polymer in different buffer,pH value (pH1-2, pH 7.4) at 37°C

Table V Antimicrobial activity of polymers (Zone of inhibition in mm)

| Samples/50 μg/ml | | | E.Coli S.aureus Pseudomonas | | | |
|------------------|------|------|-----------------------------|------|------|--|
| Homo polymer | | | 10mm | 12mm | 12mm | |
| С | | | - | = | - | |
| Co-polymer | 341 | 34mm | | 31mm | | |
| C | 28mm | 25mm | 26mm | | | |

note: C = CONTROL SOLVENT, and local bacteria using for test

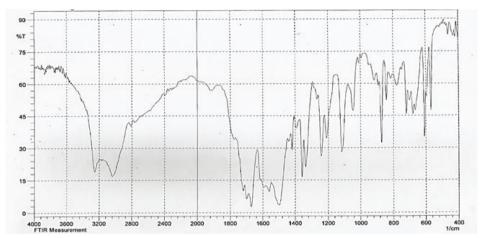


Fig 3-FTIR spectra of monomer (creatininylmaliamicacid

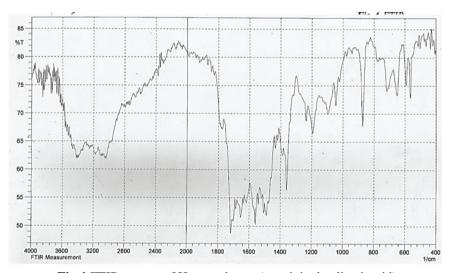


Fig 4-FTIR spectra of Homopolymer (creatininylmaliamicacid)

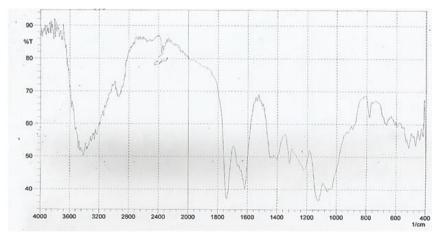


Fig 5-FTIR spectra of Copolyacrelonitryle with (creatininylmaliamicacid)

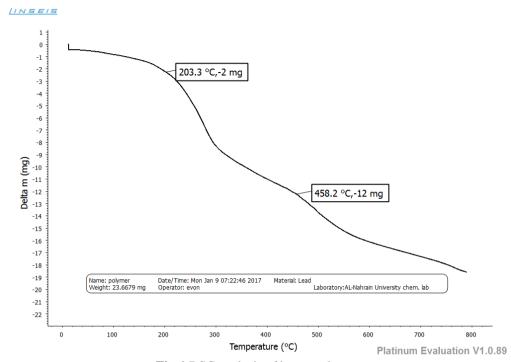


Fig 6-DSC-analysis of homo polymer

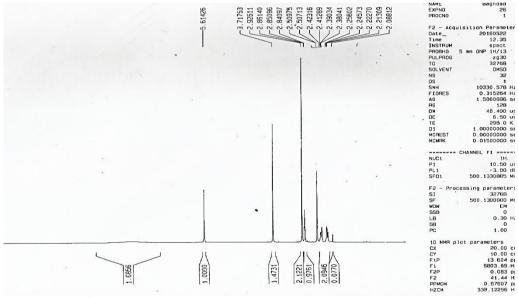


Fig-7–H-NMR spectra of Homopolymer (creatininylmaliamicacid)

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