Polymer and heterocyclic compounds their utility and application as drug

Abstract

Polymers and heterocyclic compounds play an important role in pharmacy according to their functions as excipients and inert carriers of other pharmacological active compounds. This review article discuss the recent developments in using this compounds in pharmaceuticals applications as active compounds in both as co drug or drug carriers as well as in delivery systems concerning drug therapy. These polymers and heterocyclic compounds according to their chemical character play important rules in many biological system and that is why we are here discussing there importance in many biological events such as anti microbial, antihyperkalema and anti cancer agents.

Keywords: polymer, anticancer drug, polydiacetylenes

Introduction

Polymer as drug and drug carriers

It is important that the drug must reaches its site of action at a nearly the same concentration and that its therapeutic dose range have to be remains constant over a long period of time during to its pathway toward the desired target. However, the action of pharmaceutical agents was found to be effected by many factors, including degradation during its pathway, their interaction with other target cells which is not the desired one, and their inability of penetration to the body tissues because of their chemical structure morphology such as allyments or the functional groups are not well fit to join its target cell. So according to the above stated reasons , many chemical compounds are being studied to achieve a greater pharmacological importance; among the studied compounds are, drug carriers polymers and drug polymers in which them self acts as drug. These polymers are types of systems usually important for time- and distribution-controlled drug delivery. The mechanisms of these polymers used in controlled release process needs polymers having variety of physiological and chemical properties so that to perform its action as drug. Also for hetero cyclic compounds to reach its target have not to be effected and lose any group designed for this purposes. It was known that several types of polymers have been tested as drug delivery systems, including nano- and micro-particles, nano- and micro-spheres, capsomy part of its moleomes, and micelles.1,2

In all these systems, drugs can be encapsulated or joints to the polymer backbone. These drug polymer or drug carrier polymers have been used for a verity of treatments such as antineoplastic activity, bacterial infections and inflammatory desises, sequestrate in addition to vaccine. It was also known that many cancer drugs are nearly 90% heterocyclic compounds.3 These compounds were used as anticancer agents due to their being common and widely spread in nature, with numerous number cellular and mechanistic pathways for their interactions with differents sites at the same time .It is also worth to note that there are variety of metabolic pathways regarding using cellular cancer pathology which can be attributed to heterocyclic compounds. In this article we are going to introduce the most important heterocyclic and polymeric compounds incorporated to cancer and other therapy in both areas the market and also that which are in development in both of using polymer and heterocyclic chemistry and also discussing their properties that make them valuable as drugs.

Results and discussion

The slow release of drug using acrylic Itaconic co-polymer

This study was published elsewere4 and showed regular release of anti biotic (in vivo) within a period of 14 days on 4 groups of animals (rabbits) Figure 1. Polymer synthesis and conditions were illustrated below Scheme1. Drug-Polymer associated with Tuberculostatic Activity Based on Poly (N-Vinyl Pyrrrolidone-alt-Itaconic Anhydride and Novel Aminoacid Hydrazides was also investigated by Delia G, et al.,5 The polymer synthetic pathway can be shown as below Scheme 2 & Scheme 3. This polymer showed significant activities against Tuberculosis as it was stated above.

Figure 1 Polydacetylene (PDA) synthesis schema via self-assemble polymerisation of diacetylene monomers.

Polydiacetylenes as bio sensor

Usually polydiacetylenes in their streuctures act as a by-layer molecules similar to that of the cell wall which will used to diagnose a variety of common diseases, depending on polymer fictionnalization with suitable moiety such as sugar group or lipid or others so it will becomes bio sensor due to collar change phenomena created by the interaction of this group with malarial toxins or any other events. The color change is irreversible so that it can be used to diagnose several diseases the polymerization process can shown as below: For example, GM1 and GT1b gangliosides, which are present on the surface of intestinal cells and at neuromuscular junctions, were utilized for color-based detection of cholera toxin and botulinum neurotoxin. For this synthesis, 5% ganglioside lipid was used as a function that be used as a site of action against the targeted site, Using a higher concentration it was found that it will inhibits polymerization of the diacetylene and stops its formation and as are selt no diagnostic ability for this polymer.6 Brazilian researchers in October 2010 have synthesized...
10,12 pentacosadiynoic acid (PCDA) with N-[(2-tetradecan amide)-ethyl] ribonamide (TDER) vesicles to determine the colorimetric response induced by pathogenic bacteria (Staphylococcus aureus and Escherichia coli) (Figure 2).

The addition of bacterial supernatants caused a colorimetric transition in TDER/PCDA vesicles, even in diluted concentrations, which indicate that a certain chemical interactions occur between the vesicles and the released bacterial toxin compounds. This study was important for food packing and Food born bacteriatiopes. Japanese researchers in march 2016 studied imidazoyl Functionalized diacetylenes. In their study, They report the first example of polydiacetylenes (PDAs), where the PDA-based system acts as both a sensing probe and killer for bacteria. The contact of imidazolium and imidazole-derived PDA with various bacterial strains including MRSA (methicillin-resistant Staphylococcus aureus) and ESBL-EC (extended-spectrum β-lactamase-producing Escherichia coli) results in a color change from blue-to-red of the solution as well as a rapid disruption of the bacterial membrane, which is followed up by transmission electron microscopy and confocal microscope. The study of Zeta potential analysis was found to support the successful electrostatic interaction between PDA as a positively charge toward the negatively charged bacterial cell wall. This is an example of the dual activity of polymers which acts as biosensors and at the same time as (Bacterial killers).These dual activity of PDS was shown as low (Figure 3).

Scheme 1 Synthesis of Acrylic itaconic co polymer loded with antibiotics.

Scheme 2 The formation of amidizolonyl amino acid as aresidue of the polymer.

Biomolecule-functionalized smart polydiacetylene

Figure 2 Biomolecule-functionalized Polydiacetylene (PDA) based biomedical and environmental sensing.

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Scheme 3 Showing the steps of preparation of the copoly (N-vinyl Pyrrolidone-alt-tiaconic Anhydride)with amino acids.

Figure 3 showing the effects of polydiactelyne functionalized with imidazoylgroup on bacteria cell membrane.

Polymers as drugs

Sodium polystyrene sulfonate (KayexalateVR , SPS) is the first known type of Polymer which is the first synthetic polymers to be widely used as a clinical sequestrate, its strecture is shown in scheme.4 the salt of a polymeric acid, SPS is able to reversibly bind a verity of captions, including potassium. Since potassium is the most popular ion in the colon region as part of intestinal gastric tracts, and is reversibly absorbed in this lower part GI tract, So polymeric agent with capability to such potassium this ion will provide an effective means to reduce serum potassium(Elevated serum potassium)and as a result it will releave kidney diseases ,(hyperkalemia) is a disease with incorporated with patient suffering from CKD or cardiovascular conditions which will result in arrhythmia and finally to sudden death[8]. SPS was approved by the FDA for its use on the treatment of hyperkalemia in 1958 SPS which can be synthesized by the polymerization of styrene in in presence of cross linking agent such as divinylbenzene as shown below: Scheme 4 hows the synthetic steps of polystyrene sulfonate
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Heterocyclic compounds in anti-cancer drug design strategy

As it was stated in the introduction part of this review that heterocyclic compounds are widely spread in nature they have become so important for anti-cancer drug design and other therapeutic path...
ways. They represent a precursor and building molecules of a variety of drugs or co drug compounds depending on the active functional groups they have. Depending on the above facts these heterocyclic compounds becomes the bases of most drugs designing systems. The therapeutic time and ageing of drugs As well as enzymatic binding all are associated with heterocyclic moieties. So heterocycles are becomes good choice when designing molecules that will interact with targets and disrupt the biological pathways associated with cancer progression. Especially pathways related to cell growth and development are often targeted by such as anti-cancer therapies as stated above. Moreover there are versatile method by which can modified the stricture of heterocyclic rings with any additional group or element which finally willed allows them to cover a broad area of chemical interaction sites, and as a result will quiting them as excellent starting points for cancer therapeutic designing and developments. Finally and as a result of the stated above facts, Heterocyclic compounds play an important rules in drug designing and development and provides the market with a lot of new pharmaceutical compounds every year for both cancer and other therapeutic drugs.

It is worth to say that nearly 65% of the anti-cancer drugs granted market approved by FDA between 2010-2015 form the basis of many of the anti-cancer agent currently in development today are Nitrogen-based heterocycles. It was approved by the FDA between 2010 and 2015 that nitrogen based heterocyclic compounds forms two third of the commercial cancer drugs. Nitrogen heterocycles, in doles are among the most valuable, in research demonstrated their ability to induce cell death in a number of cancer cell lines. During the last few decades, indole and its derivatives have been shown to have biological pathways against cancer through the on growing research and developments. These include the prevention of cell signaling normal cell cycle progression, tumor vascularisation and DNA repair, as well as the ability to induce cellular oxidative stress and cell death. The most important was an early used commercially indole-based anticancer agents are vincristine and vinblastine which are recognized for their tubulin polymerization inhibition since the early-mid 1960s and both still of clinical importance today. Vincristine (Figure 4) is used for the treatment of acute lymphoblastic leukaemia and both Hodgkin’s and- non-Hodgkin’s lymphoma.

The development in heterocyclic compounds its utility and applications toward cancer screening studies showed a lot of works in this area of drug discovery as an examples coumarin and oxazine compounds. For coumarin containing heterocyles the following compounds showed antitumor activities against two cell lines breast carcinoma Michigan Cancer Foundation-7 (MCF-7) and hepatocellular carcinoma (HepG-2), at the National Cancer Institute, Cairo, Egypt using 5-fluorouracil as standard drug. The structure of the synthesized compounds were illustrated below. While for, oxygen
and nitrogen containing heterocycle (Chloro pyrano Oxazines) compounds and according to the following Scheme 8 together with the graphical curves (NCI report) showed remarkable results for these types of compounds. Scheme 8 shows the synthetic steps of Amino chloropyrano oxazine compounds and the graphical results showing the effects of these compounds on different types of cancer cells The above investigation revealed that the studied compounds kill about 89-100% cancer cell (6-8 types) at the same time.

Figure 8 shows the synthetic steps of Amino chloropyrano oxazine compounds and the graphical results showing the effects of these compounds on different types of cancer cells The above investigation revealed that the studied compounds kill about 89-100% cancer cell (6-8 types) at the same time.

Figure 5 Dericoumarin derivatives as anti tumor agents.

Acknowledgments

None.

Conflicts of interest

The author declares there is no conflict of interest.

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