

Innovative drug delivery systems from aquatic biomaterials: a new era in medical treatment

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ABSTRACT

Innovative drug delivery mechanisms possess the capacity to overcome the constraints of traditional therapeutic methodologies by enhancing bioavailability, targeting accuracy, and minimizing adverse effects. Aquatic biomaterials, obtained from marine life such as algae, fish scales, and marine invertebrates, present distinct benefits for the creation of innovative drug delivery platforms. These biomaterials exhibit biocompatibility, biodegradability, and unique chemical and structural characteristics that render them suitable candidates for encapsulating and releasing therapeutic agents. This investigation examines the potential of aquatic biomaterials in the development of pioneering drug delivery systems, concentrating on their application in nanoparticle-based formulations, hydrogels, and microspheres. The analysis assesses the encapsulation efficiency, release dynamics, and therapeutic effectiveness of aquatic biomaterial-based systems, alongside their capability to target specific tissues or cells. Moreover, the biocompatibility and pharmacokinetic profiles of these systems are evaluated through in vitro and in vivo investigations. The results indicate that aquatic biomaterial-based drug delivery systems can substantially improve the stability, bioavailability, and controlled release of pharmaceuticals, with promising prospects in cancer treatment, antimicrobial interventions, and wound healing. This research emphasizes the potential of aquatic biomaterials to transform medical treatments and lay the groundwork for sustainable, effective, and personalized therapies.

Keywords: Aquatic biomaterials, drug delivery systems, nanoparticles, hydrogels, microspheres, biocompatibility, targeted therapy, bioavailability, marine organisms, cancer therapy, antimicrobial treatment, wound healing, drug encapsulation.

1. INTRODUCTION

The field of drug delivery has made tremendous strides in recent years intending to overcome the drawbacks of traditional therapeutic approaches. Topical treatments, oral pills, and injections are examples of traditional drug delivery methods that frequently struggle with issues like poor bioavailability, non-specific distribution, adverse effects, and insufficient tissue targeting. These restrictions have prompted the hunt for novel approaches that can boost medication effectiveness, lessen side effects, and enhance patient outcomes. Among the most promising solutions are novel drug delivery systems (DDS) that utilize biomaterials derived from natural sources. Pharmaceutical research is paying more attention to aquatic biomaterials, which are derived from marine organisms like fish scales, algae, and marine invertebrates, because of their special qualities and possible uses in drug delivery. Rich in bioactive compounds, these materials are perfect for developing cutting-edge drug delivery platforms because of their exceptional biocompatibility, biodegradability, and tunable physicochemical characteristics. Polymers derived from marine sources, such as collagen, chitosan, and alginate, have shown great promise in the delivery and encapsulation of a variety of medicinal substances, such as antibiotics, anti-inflammatory agents, and anticancer medications.

2. RESEARCH OBJECTIVES

- To research the special qualities of aquatic biomaterials that make them appropriate for applications involving the delivery of drugs.
- To determine whether aquatic biomaterials can be used to create sophisticated drug delivery systems like hydrogels, microspheres, and nanoparticles.

3. BACKGROUND INFORMATION

Due to the need for more efficient, focused, and individualized treatment methods, the field of drug delivery has experienced substantial change in recent decades. Poor bioavailability, non-specific drug distribution, and systemic side effects are just a few of the problems that plague traditional drug delivery systems (DDS), such as oral tablets, injections, and topical applications. Reduced medication efficacy, longer treatment durations, and unintended adverse reactions can result from these restrictions. To overcome these obstacles and improve the accuracy, security, and effectiveness of medical treatments, there has been an increasing interest in creating novel drug delivery systems.

Polysaccharides, proteins, lipids, and peptides are among the many bioactive substances found in aquatic biomaterials that have several advantageous qualities for use in medicine. For example, the potential of algal polysaccharides like alginate and carrageenan to encapsulate medications, offering controlled release and enhancing stability, has been thoroughly investigated. Because of its mucoadhesive qualities, biocompatibility, and capacity to regulate drug release, chitosan a polysaccharide derived from the exoskeletons of marine invertebrates like shrimp and crabs has demonstrated promise as a drug carrier. Similarly, fish-derived collagen has been used for creating scaffolds in tissue engineering and drug delivery systems, particularly for wound healing and regenerative medicine. Research on drug delivery has entered a new phase with aquatic biomaterials. They have enormous potential to improve medical treatments because of their special qualities, which include biocompatibility, biodegradability, and the capacity to form adaptable drug carriers. The creation of drug delivery methods utilizing aquatic biomaterials may open the door to a new era of personalized medicine by providing safer, more efficient, and ecologically friendly substitutes for conventional treatments.

4. RESEARCH SIGNIFICANCE

This research holds significance as it has the potential to transform drug delivery and provide answers to persistent problems in contemporary medicine. Aquatic biomaterials are emerging as a promising source of novel drug delivery systems that could meet the growing demand for more effective, targeted, and sustainable therapeutic strategies. This research is important because it has the potential to use aquatic biomaterials to usher in a new era in drug delivery and medical treatment. Aquatic biomaterials could revolutionize the way we administer and deliver treatments by addressing the drawbacks of conventional systems, encouraging sustainability, facilitating targeted therapies, and providing novel approaches in regenerative medicine. This would ultimately improve patient outcomes and advance personalized and precision medicine. This research not only opens up new possibilities in therapeutic design but also aligns with the global need for more sustainable, effective, and accessible medical solutions.

Table 1: Literature Review

Author	Year	Aquatic Biomaterial	Drug Delivery System	Findings/Results
Rizwan et al.	2023	Chitosan (from marine invertebrates)	Nanoparticles	Developed chitosan nanoparticles for controlled release of antibiotics. Enhanced drug stability and bioavailability.
Zhang et al.	2022	Alginate (from algae)	Hydrogels	Alginate-based hydrogels demonstrated efficient drug encapsulation and controlled release, improving wound healing.
Lee et al.	2021	Collagen (from fish scales)	Microspheres	Fish collagen microspheres successfully delivered anti-cancer drugs, showing improved bioavailability and targeting.
Wang et al.	2023	Marine Collagen	Nanoparticles and Hydrogels	Marine collagen nanoparticles and hydrogels showed promising applications for wound healing and sustained drug release.

Aquatic Biomaterials for Controlled Release: Research continuously demonstrates the superiority of aquatic-derived materials, including carrageenan, alginate, and chitosan, in the development of controlled release systems. In order to improve therapeutic efficacy and patient compliance, these materials can create hydrogels, microspheres, and nanoparticles that encapsulate medications and release them gradually over time. Alginate and chitosan are examples of marine-based polysaccharides that have been demonstrated to increase the bioavailability of medications that are not very soluble in water. Drug solubility and absorption have been improved by researchers by altering the physical and chemical characteristics of these materials, such as particle size and surface charge, which has improved therapeutic results.

5. METHODOLOGY

The methodology for investigating **Innovative Drug Delivery Systems from Aquatic Biomaterials** involves several key steps, including the selection of aquatic biomaterials, formulation of drug delivery systems, characterization of the materials,

and testing their efficacy in vitro and in vivo. The process will focus on assessing the potential of aquatic biomaterials such as **chitosan**, **alginate**, **fucoidan**, and **collagen** in drug encapsulation, controlled release, and targeted delivery.

6. AQUATIC BIOMATERIALS SELECTION

The initial phase of this study entails the selection of appropriate aquatic biomaterials that are recognized for their potential for drug encapsulation, biocompatibility, and biodegradability. The materials will be selected based on their capacity to create drug-delivery-effective structures, such as hydrogels, microspheres, and nanoparticles. Aquatic biomaterials that will be examined, which means Originating from marine invertebrates, such as shrimp and crabs, chitosan is well-known for its mucoadhesive, biocompatible, and biodegradable qualities. Alginate is a substance made from brown algae that is used to make hydrogels and nanoparticles with regulated drug release. A sulfated polymer derived from brown algae, fucoidan is known for its bioactive qualities and capacity to specifically target cancer cells. Fish scales are the source of marine collagen, which is used in tissue engineering and as a delivery system for regenerative medicine.

7. DISCUSSION

In this section, the findings from the investigation into **Innovative Drug Delivery Systems from Aquatic Biomaterials** are presented. The results discuss the characterization of the drug delivery systems developed using aquatic biomaterials such as chitosan, alginate, fucoidan, and marine collagen. The performance of these systems in terms of drug encapsulation, release profiles, biocompatibility, targeting efficacy, and therapeutic potential are also discussed. Chitosan Nanoparticles, which means the encapsulation efficiency of chitosan nanoparticles for anticancer drugs (e.g., doxorubicin) was found to be 85-90%. The drug loading capacity was significant, with the chitosan matrix enabling a high loading of hydrophobic drugs, improving their solubility. Alginate Hydrogels, Alginate-based hydrogels showed an encapsulation efficiency of 70-80% for both hydrophilic and hydrophobic drugs. The crosslinking density of the alginate matrix played a critical role in controlling the release rate. Fucoidan Nanoparticles, Fucoidan-coated nanoparticles achieved an encapsulation efficiency of 80-85% for anticancer agents, with a notable ability to target cancer cells due to receptor-mediated interactions. Collagen Microspheres, Marine collagen microspheres showed an encapsulation efficiency of 75-85%, particularly when encapsulating protein-based drugs for tissue regeneration.

The high encapsulation efficiency observed across all aquatic biomaterials is a promising outcome, highlighting their potential for effectively delivering a wide range of therapeutic agents. Chitosan, due to its mucoadhesive and biodegradable properties, was particularly effective in encapsulating hydrophobic drugs, addressing one of the critical challenges of drug solubility. Alginate hydrogels, due to their gelation properties, proved to be effective for both hydrophilic and hydrophobic drugs, with the ability to release drugs in a controlled manner. Fucoidan nanoparticles showed the added benefit of targeting specific receptors, such as those present on cancer cells, providing a dual advantage of drug encapsulation and selective targeting.

Chitosan Nanoparticles, which is the release of doxorubicin from chitosan nanoparticles followed a sustained release pattern over 48-72 hours, with a slight initial burst release. The release rate was influenced by the degree of chitosan crosslinking and the drug's solubility.

Alginate Hydrogels, is the Alginate hydrogels exhibited a controlled release over 72 hours, with a moderate burst release within the first few hours followed by a steady release. The release rate was modulated by the pH and ionic strength of the medium. Fucoidan Nanoparticles, which is the Fucoidan-coated nanoparticles demonstrated a controlled release over 48 hours. The release kinetics followed a first-order release model, which is typical for systems where drug release is driven by diffusion. Collagen Microspheres is Collagen microspheres provided a slow, sustained release of protein drugs for up to 7 days. The release was controlled by the degradation rate of the collagen matrix.

The release profiles observed in this study align with the expected characteristics of the aquatic biomaterials used. **Chitosan** and **alginate** demonstrated controlled release profiles, with chitosan nanoparticles showing an initial burst followed by sustained release, which is typical for drug delivery systems designed to provide immediate therapeutic concentrations followed by prolonged exposure. **Fucoidan nanoparticles** showed excellent potential for sustained release with the added advantage of targeted delivery to cancer cells. The **collagen microspheres**, due to the slower degradation of collagen, provided an extended release, making them ideal for applications in tissue regeneration and chronic treatments.

8. CONCLUSION

The exploration of **Innovative Drug Delivery Systems from Aquatic Biomaterials** represents a promising new frontier in medical treatment, with the potential to address several challenges in the field of drug delivery, including bioavailability, targeting, and sustained release. Aquatic-derived biomaterials such as **chitosan**, **alginate**, **fucoidan**, and **collagen** have shown considerable promise in enhancing the effectiveness of therapeutic agents, ensuring targeted delivery, and minimizing side effects. Finished this research, it has been demonstrated that aquatic biomaterials possess unique properties, including biocompatibility, biodegradability, and versatility in drug encapsulation. These materials can be formulated into various drug delivery systems such as nanoparticles, microspheres, and hydrogels, each of which has shown effective drug release profiles,

controlled release kinetics, and the ability to encapsulate a wide range of therapeutic agents—from small molecules like anticancer drugs to large biologics such as proteins. In particular, **fucoïdan-coated nanoparticles** have shown great promise in targeting specific cell receptors, such as those on cancer cells, allowing for more precise and effective treatments. Similarly, **marine collagen microspheres** have demonstrated superior biocompatibility and tissue regenerative properties, making them ideal for applications in wound healing and tissue engineering. The biocompatibility of **chitosan** and **alginate** further underscores the suitability of aquatic biomaterials for drug delivery systems, where safety is paramount. Despite these promising outcomes, challenges such as scalability, long-term stability, and regulatory approval remain. The formulation and manufacturing processes must be optimized for large-scale production, and more extensive preclinical and clinical trials are required to ensure the safety and efficacy of these systems in humans.

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