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Fabrication, Mechanical Testing, and Biocompatibility Assessment of Biopolymer Composites for Biomedical Implant Applications

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Abstract: The increasing demand for biocompatible and mechanically reliable materials in biomedical implant applications has driven extensive research into biopolymer-based composites as viable alternatives to conventional metallic implants. This study presents the fabrication, mechanical characterization, and biocompatibility assessment of novel biopolymer composites reinforced with bioactive fillers for load-bearing and non-load-bearing implant applications. Biopolymers such as polylactic acid and chitosan were selected due to their biodegradability, tunable mechanical properties, and favorable biological interactions. Composite samples were fabricated using solvent casting and compression molding techniques to ensure uniform filler dispersion and structural integrity. Mechanical testing including tensile strength, compressive strength, flexural behavior, and hardness was conducted following ASTM standards to evaluate suitability for implant environments. In vitro biocompatibility was assessed using cytotoxicity assays, cell adhesion studies, and surface morphology analysis. The results demonstrate that the incorporation of bioactive fillers significantly enhances mechanical performance while maintaining excellent cytocompatibility. The study confirms that properly engineered biopolymer composites can achieve mechanical properties comparable to cancellous bone while promoting favorable cellular responses. These findings highlight the potential of biopolymer composites as next-generation materials for orthopedic and dental implant applications, offering improved biological integration and reduced long-term complications compared to traditional implant materials.

Keywords: Biopolymer Composites, Biomedical Implants, Mechanical Properties, Biocompatibility, Bioactive Fillers

1. Introduction

The field of biomedical implants has undergone substantial transformation over the past few decades due to advancements in materials science and biomedical engineering. Conventional implant materials such as stainless steel, cobalt–chromium alloys, and titanium-based alloys have long been used because of their superior mechanical strength and corrosion resistance. However, despite their widespread clinical success, these materials present several limitations including stress shielding, poor biological integration, long-term ion release, and the necessity for secondary surgical removal in some cases [1]. These challenges have prompted researchers to explore alternative materials that can better mimic the mechanical and biological behavior of natural tissues. Biopolymers have emerged as promising candidates for biomedical implant applications due to their inherent biocompatibility, biodegradability, and versatility in structural design. Materials such as polylactic acid, polycaprolactone, chitosan, and alginate have demonstrated excellent biological performance in contact with living tissues [2]. However, the primary limitation of biopolymers lies in their relatively low mechanical strength and stiffness when compared to metallic counterparts, restricting their application in load-bearing implants. To overcome this limitation, the development of biopolymer composites reinforced with bioactive or ceramic fillers has gained significant attention. Biopolymer composites aim to combine the favorable biological characteristics of polymers with the mechanical reinforcement provided by fillers such as hydroxyapatite, bioactive glass, and calcium phosphates. These fillers not only improve mechanical performance but also enhance osteoconductivity and promote tissue regeneration at the implant–tissue interface [3]. The design of such composites requires careful optimization of filler content, dispersion, interfacial bonding, and processing techniques to achieve desired performance outcomes. Mechanical compatibility between implant materials and surrounding biological

tissues is a critical factor influencing implant longevity and success. Mismatch in elastic modulus can lead to stress shielding, resulting in bone resorption and implant loosening [4]. Therefore, biopolymer composites with tunable mechanical properties offer a strategic advantage by enabling customization according to specific anatomical and functional requirements. Additionally, surface properties such as roughness, wettability, and chemical composition play a crucial role in cell attachment, proliferation, and differentiation. This study focuses on the fabrication of biopolymer composites using biodegradable polymers reinforced with bioactive fillers, followed by systematic mechanical testing and *in vitro* biocompatibility evaluation. The objective is to assess their potential as implant materials capable of meeting both mechanical demands and biological performance criteria. By integrating experimental mechanical analysis with biological assessment, this research aims to contribute to the development of safer, more effective, and patient-specific implant materials for future biomedical applications.

2. Materials and Methods

The selection of materials for biopolymer composite fabrication was guided by considerations of biocompatibility, biodegradability, mechanical performance, and processing feasibility. Polylactic acid was chosen as the primary polymer matrix due to its established clinical use, predictable degradation behavior, and favorable mechanical properties. Chitosan was incorporated as a secondary biopolymer to enhance biological activity and improve cell–material interactions. Hydroxyapatite particles derived from synthetic sources were used as bioactive fillers to improve mechanical reinforcement and promote osteointegration [5]. Composite fabrication was carried out using a combination of solvent casting and compression molding techniques to ensure homogenous dispersion of fillers within the polymer matrix. Initially, polylactic acid was dissolved in an appropriate organic solvent under controlled stirring conditions. Chitosan was prepared separately in an acidic aqueous solution and blended with the polymer solution to form a uniform mixture. Hydroxyapatite particles were gradually introduced at varying weight percentages and dispersed using mechanical stirring followed by ultrasonic treatment to minimize agglomeration. The resulting composite slurry was cast into molds and allowed to evaporate under controlled environmental conditions to remove residual solvents. Dried samples were then subjected to compression molding at optimized temperature and pressure settings to enhance density and mechanical integrity. The fabricated specimens were machined into standardized dimensions for mechanical testing in accordance with ASTM protocols. Mechanical characterization included tensile, compressive, and flexural testing using a universal testing machine. Tensile tests were performed to determine ultimate tensile strength, elastic modulus, and elongation at break. Compression tests evaluated load-bearing capability, while flexural tests provided insights into bending resistance relevant to orthopedic applications. Hardness measurements were conducted using a Shore D hardness tester to assess surface resistance to deformation. Surface morphology and filler dispersion were examined using scanning electron microscopy to identify microstructural features influencing mechanical and biological behavior. For biocompatibility assessment, *in vitro* cytotoxicity tests were performed using fibroblast cell lines following ISO 10993 guidelines. Cell viability was quantified using MTT assays after predefined incubation periods. Cell adhesion and morphology were observed using fluorescence microscopy to evaluate cellular response to composite surfaces. Statistical analysis of experimental data was conducted to ensure reliability and reproducibility of results. Mechanical and biological performance metrics were compared across different composite formulations to identify optimal material compositions for biomedical implant applications.

3. Results and Discussion

The mechanical testing results revealed a significant improvement in the strength and stiffness of biopolymer composites with increasing hydroxyapatite content. Tensile strength values showed a marked increase compared to pure polymer matrices, indicating effective load transfer between the polymer and filler phases. The elastic modulus of optimized composites closely matched that of cancellous bone, suggesting reduced risk of stress shielding when used as implant materials [6]. Compression testing demonstrated enhanced load-bearing capacity, confirming suitability for applications subjected to compressive stresses. Flexural testing results further supported the mechanical robustness of the composites, with improved bending resistance attributed to uniform filler dispersion and strong interfacial bonding. Hardness measurements indicated increased surface durability, which is critical for minimizing wear and surface damage in implant environments. Scanning electron microscopy images confirmed homogenous distribution of hydroxyapatite particles and revealed limited microvoid formation, contributing to mechanical stability. Biocompatibility assessment showed excellent cellular response across all composite formulations. Cytotoxicity assays indicated high cell viability, exceeding acceptable thresholds specified by international standards. The presence of chitosan and hydroxyapatite contributed to enhanced cell adhesion and spreading, as observed in fluorescence microscopy images. Cells exhibited well-defined cytoskeletal structures and strong attachment to composite surfaces, reflecting favorable surface chemistry and topography [7]. The results demonstrate that the integration of bioactive fillers not only enhances mechanical performance but also promotes biological interactions essential for successful implant integration. The synergistic effect of polymer blending and filler reinforcement resulted in composites capable of

meeting mechanical demands while supporting cellular activity. These findings align with previous studies emphasizing the importance of composite design in achieving balanced mechanical and biological performance [8]. The study highlights the potential of biopolymer composites to address limitations associated with traditional implant materials. By tailoring composition and processing parameters, it is possible to engineer implant materials with properties optimized for specific clinical requirements. However, long-term in vivo studies and degradation behavior analysis are necessary to fully validate clinical applicability.

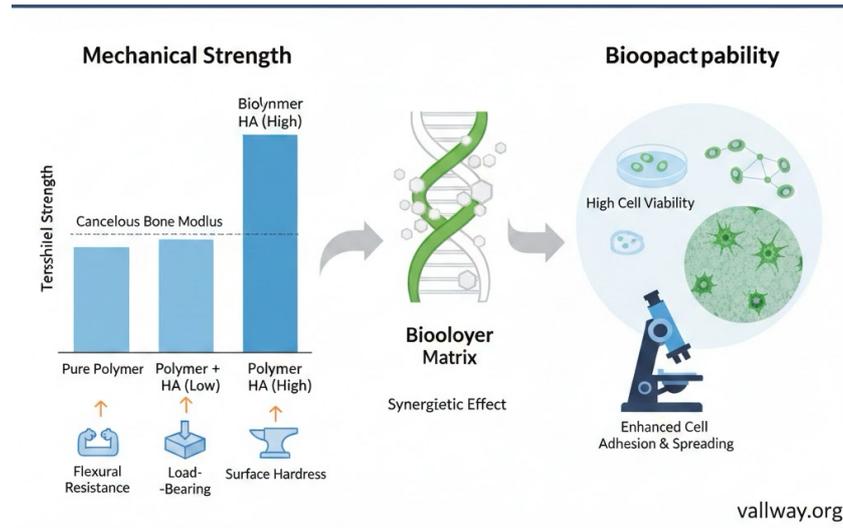


Fig. 1

4. Conclusion

This research demonstrates that biopolymer composites reinforced with bioactive fillers represent a promising class of materials for biomedical implant applications. The successful fabrication of polylactic acid–chitosan composites reinforced with hydroxyapatite highlights the feasibility of combining biodegradable polymers with bioactive reinforcements to achieve enhanced mechanical and biological performance. Mechanical testing confirmed significant improvements in tensile, compressive, and flexural properties, bringing them within the range of natural bone tissues and reducing the risk of stress shielding. The biocompatibility assessment further validated the suitability of these composites for biomedical use, with high cell viability, strong adhesion, and favorable cellular morphology observed during in vitro testing. The presence of hydroxyapatite played a critical role in promoting osteoconductive behavior, while chitosan contributed to improved surface bioactivity. These characteristics are essential for ensuring stable implant integration and long-term functionality. The findings underscore the importance of material design and processing optimization in developing next-generation implant materials. Biopolymer composites offer the flexibility to tailor degradation rates, mechanical properties, and biological interactions according to patient-specific needs. While the current study provides comprehensive experimental validation, future work should focus on in vivo performance evaluation, long-term degradation behavior, and clinical translation pathways. Overall, the study contributes valuable insights into the development of sustainable, biocompatible, and mechanically reliable implant materials. The demonstrated performance of the fabricated biopolymer composites positions them as strong candidates for orthopedic and dental implant applications, paving the way for safer and more effective biomedical solutions.

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