

1 *Review*

# 2 Environmental Applications of Chitosan Derivatives and 3 Chitosan Composites

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## 7 Abstract

8 Chitosan, a naturally abundant and biodegradable biopolymer derived from chitin  
9 found in crustacean shells, has emerged as a promising material for addressing  
10 environmental challenges. Its reactive amino and hydroxyl groups enable diverse  
11 interaction mechanisms, making it effective for removing heavy metals, dyes,  
12 pharmaceuticals, and other contaminants from water. However, the limitations of native  
13 chitosan, such as poor solubility and mechanical strength, necessitate strategic  
14 modifications. This review comprehensively examines recent advances in chitosan  
15 derivatives and composites, focusing on modern modification strategies (chemical,  
16 physical, and composite formation) that enhance stability, selectivity, and efficiency. It  
17 explores the design principles of high-performance composites and the multifaceted  
18 mechanisms of pollutant removal, including adsorption, catalysis, membrane filtration,  
19 and flocculation. Critical practical challenges related to scalability, regeneration, lifecycle  
20 sustainability, and real-world implementation are critically assessed. Furthermore,  
21 emerging trends that integrate circular economy principles, seafood waste valorization,  
22 and digital optimization through the use of artificial intelligence are highlighted. By  
23 consolidating current knowledge, this review aims to bridge the gap between laboratory  
24 innovations and large-scale environmental applications, guiding the development of  
25 intelligent, scalable, and ecologically responsible solutions based on this remarkable  
26 biopolymer.

27 **Keywords:** Chitosan; Chitosan derivatives; Chitosan composites; Biopolymer  
28 composites; Environmental remediation; Pollutant removal; Sustainable materials;  
29 Circular economy

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## 31 1. Introduction

32 The escalating global crisis of environmental pollution, fuelled by the relentless  
33 discharge of industrial effluents, agricultural runoff, pharmaceutical residues, and the  
34 pervasive issue of improper waste management, has created an urgent and intensifying  
35 demand for remediation technologies that are not only effective but also sustainable,  
36 efficient, and environmentally benign. Traditional synthetic polymers and chemical  
37 treatments, while historically prevalent, often fall short due to their non-renewable  
38 origins, potential for secondary pollution, and limited long-term ecological  
39 compatibility[1]. In this critical context, biopolymer-based materials, derived from  
40 renewable and often waste resources, have emerged as frontrunners in the quest for  
41 viable solutions[2]. Among these natural polymers, chitosan stands out with exceptional  
42 prominence, recognized for its remarkable versatility, abundance, and inherently eco-  
43 friendly characteristics[3,4]. As a deacetylated derivative of chitin[5], the second most

abundant biopolymer on Earth after cellulose, chitosan is not merely a byproduct of nature's vast structural inventory but a functional material with immense potential for addressing contemporary environmental challenges[6]. Its linear, cationic polysaccharide structure (Figure 1), composed of randomly distributed  $\beta$ -(1  $\rightarrow$  4)-linked D-glucosamine and N-acetyl-D-glucosamine units, grants it a unique chemical identity, particularly the presence of reactive primary amino ( $-\text{NH}_2$ ) and hydroxyl ( $-\text{OH}$ ) groups[7]. These functional moieties are the cornerstone of chitosan's exceptional reactivity, enabling a diverse array of interaction mechanisms as electrostatic attraction, complexation, hydrogen bonding, ion exchange, and physical adsorption, that render it highly effective in capturing and immobilizing a broad spectrum of pollutants from aqueous environments, including heavy metal ions, organic dyes, pharmaceuticals, endocrine disruptors, and microbial contaminants[8–10].

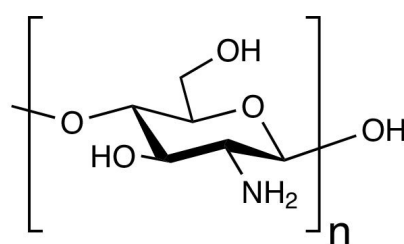


Figure 1. Chitosan structure

The significance of chitosan is further amplified by its primary source: the vast quantities of crustacean shell waste generated by the global seafood industry. Annually, an estimated 6 to 8 million tons of shrimp, crab, and lobster shells are discarded, presenting a significant disposal challenge and environmental burden[11]. However, this waste stream simultaneously represents an underutilized reservoir of high-value raw material. The transformation of this waste into chitosan exemplifies a pivotal shift toward circular economy models, where environmental liabilities are reimagined as resources for designing advanced functional materials[5,12]. While crustacean shells remain the dominant source, the exploration of alternative origins such as fungi (e.g., *Aspergillus niger*, *Mucor rouxii*), insects (like black soldier flies and crickets), and even bacterial cell walls offers promising avenues for more sustainable and seasonally independent production[13,14]. The industrial journey from waste shell to functional biopolymer involves a well-defined, albeit traditionally energy-intensive, three-step process: demineralization using dilute acid to remove calcium carbonate, deproteinization via alkaline treatment to eliminate proteins, and finally, deacetylation under strong alkaline conditions (40 - 50% NaOH) at elevated temperatures to cleave acetyl groups from chitin[15]. The degree of deacetylation (DD), typically ranging from 55% to over 90%, is a crucial parameter defining the final properties, with higher DD values correlating with increased amino group content, enhanced cationic character, and improved reactivity, particularly in acidic environments where protonation renders the polymer soluble[16]. Molecular weight (MW), which varies widely from tens of thousands to several million Daltons, significantly influences viscosity, mechanical strength, solubility, and diffusion kinetics, thereby impacting its applicability in various formats and conditions[17,18].

82 Despite its inherent advantages as biocompatibility, non-toxicity, biodegradability,  
83 antimicrobial activity, film-forming ability, and mucoadhesive properties[19], native  
84 chitosan faces intrinsic limitations that constrain its direct, large-scale deployment in  
85 environmental systems. Its insolubility in neutral and alkaline pH conditions[20],  
86 susceptibility to acid hydrolysis[21], low mechanical strength, tendency to swell and  
87 disintegrate in aqueous media, limited surface area, and non-selective binding  
88 behaviour necessitate strategic modifications to unlock its full potential. Consequently,  
89 extensive research has been devoted to overcoming these hurdles through a range of  
90 engineering strategies, including chemical derivatization, physical structuring, and  
91 composite formation. Chemical modifications have yielded derivatives like  
92 carboxymethyl chitosan (CMCS), which improves solubility across a wider pH range  
93 and exhibits ampholytic properties[22]; quaternized chitosan, which maintains its  
94 positive charge even at neutral pH, enhancing its utility in diverse applications[23]; and  
95 thiolated chitosan, which introduces redox activity and superior metal chelation  
96 capabilities, particularly for soft metals[24]. Physical processing techniques, such as  
97 cryostructuring, freeze-drying, and solution casting, enable the precise control of  
98 morphology and porosity, thereby tailoring the material for specific functions[25].  
99 However, perhaps the most impactful approach has been the development of composite  
100 materials, where chitosan is synergistically combined with various other components.  
101 Integration with inorganic fillers, such as diatomite, sepiolite, and layered silicates,  
102 enhances both the ion-exchange capacity and thermal stability of the material[26]. The  
103 incorporation of carbonaceous materials, such as biochar, graphene oxide, and carbon  
104 nanotubes, dramatically increases the surface area and introduces conductive pathways,  
105 thereby boosting both adsorption and catalytic capabilities[27]. Magnetic nanoparticles,  
106 such as  $\text{Fe}_3\text{O}_4$ , not only reinforce the structure but also facilitate easy separation after  
107 use, a crucial advantage for practical applications[28]. Combining chitosan with other  
108 biopolymers, such as alginate or lignin, further enhances mechanical properties and  
109 reduces costs[29,30]. These hybrid systems leverage the combined strengths of their  
110 constituents, often yielding performance that surpasses the sum of their parts. As  
111 demonstrated by composites capable of not only adsorbing but also catalytically  
112 reducing toxic species, such as Cr(VI), to less harmful forms, this approach yields  
113 enhanced performance[31].

114 The mechanisms by which chitosan-based materials remove pollutants are  
115 multifaceted and often work in concert. Adsorption, driven primarily by electrostatic  
116 interactions and complexation involving the amino and hydroxyl groups, is the  
117 dominant pathway[32], particularly effective for capturing both anionic and cationic  
118 species under varying pH conditions[33]. Catalytic functionalities can be integrated into  
119 composites, enabling active degradation of pollutants beyond simple capture[34].  
120 Chitosan's film-forming ability and charge properties also make it an excellent candidate  
121 for membrane filtration, offering antifouling characteristics beneficial for ultrafiltration  
122 and nanofiltration processes[35,36]. Furthermore, its natural cationic nature allows it to  
123 function as an effective flocculant, aggregating negatively charged colloids in turbid  
124 water, providing a green alternative to synthetic coagulants[37]. While laboratory  
125 investigations have consistently demonstrated the efficacy of chitosan and its  
126 derivatives, the transition from bench-scale success to widespread real-world  
127 application is fraught with challenges. Issues related to scalability, maintaining  
128 performance under variable and often harsh environmental conditions, efficient  
129 regeneration without causing secondary pollution, and ensuring overall lifecycle  
130 sustainability must be addressed[31,38]. Traditional fabrication methods, such as  
131 lyophilization, although effective, are often energy-intensive and challenging to scale up

132 economically[39]. The economic viability of chitosan technology is closely tied to the  
133 cost-effective sourcing of raw materials, making the use of unrefined seafood waste  
134 more attractive than purified commercial grades[40]. Recognizing these challenges, the  
135 field is increasingly embracing green synthesis methods, such as enzymatic processing  
136 and microbial fermentation, which operate under milder conditions, reduce chemical  
137 consumption, and minimize hazardous waste generation, aligning production more  
138 closely with sustainability goals[41]. Looking forward, the trajectory of chitosan  
139 technology is being further shaped by the integration of digital innovation. Artificial  
140 intelligence and machine learning are being harnessed to predict performance, optimize  
141 formulations, and enable real-time process monitoring. Concepts like digital twins and  
142 blockchain traceability are beginning to ensure transparency and efficiency across the  
143 entire value chain, from waste sourcing to final disposal[42]. This confluence of material  
144 science, green chemistry, composite engineering, and digital technology underscores the  
145 evolving narrative of chitosan, positioning it not just as a promising biopolymer but as a  
146 cornerstone material for the development of scalable, intelligent, and ecologically  
147 responsible solutions essential for global environmental protection in the 21st  
148 century[20].

149 Given the growing urgency of environmental degradation and the need for  
150 sustainable alternatives to synthetic polymers, this review provides a comprehensive  
151 synthesis of recent advances in chitosan-based materials[43]. By consolidating  
152 knowledge on its origin, structural characteristics, preparation methods, green synthesis  
153 routes, key derivatives, composite formulations, and functional properties, we aim to  
154 highlight the critical role of chitosan in addressing pressing ecological challenges[44].  
155 The necessity of this review lies in bridging the gap between laboratory-scale  
156 innovations and real-world implementation, guiding future research toward scalable,  
157 intelligent, and ecologically responsible solutions grounded in one of nature's most  
158 abundant and adaptable biopolymers: chitosan.

159 Based on the previous literature, review papers related to chitosan and its  
160 derivatives and composites primarily focus on their applications in the biomedical field,  
161 with some also exploring agricultural applications[45]. There is no comprehensive  
162 literature available that provides a systematic summary of recent knowledge related to  
163 the synthesis and environmental applications of chitosan and its derivatives, as well as  
164 composites. Therefore, this review aims to provide a comprehensive synthesis of recent  
165 advances in chitosan-based materials for environmental applications. We critically  
166 examine modern modification strategies, design principles of high-performance  
167 composites, the underlying mechanisms of pollutant removal, practical implementation  
168 barriers, and cutting-edge trends that integrate sustainability and digitalization. By  
169 consolidating current knowledge and identifying critical research gaps, this work aims  
170 to guide the development of scalable, intelligent, and ecologically responsible solutions  
171 for global environmental protection, based on one of nature's most abundant and  
172 adaptable biopolymers: chitosan. Moreover, several applications of chitosan derivatives  
173 and chitosan-based composites are mentioned.

## 174 **2. Modification Strategies and Design Principles for Enhanced** 175 **Environmental Performance of Chitosan**

### 176 *2.1. Modern Strategies for Modifying Chitosan to Enhance Stability, Selectivity, and Efficiency*

177 The inherent functional richness of chitosan, stemming from its abundant primary  
178 amino ( $-NH_2$ ) and hydroxyl ( $-OH$ ) groups, renders it a highly reactive biopolymer with  
179 significant potential for environmental applications[46]. However, native chitosan

180 exhibits several limitations that restrict its practical utility, including poor solubility at  
181 neutral and alkaline pH levels[47], low mechanical strength[35], acid lability[29],  
182 swelling-induced disintegration in aqueous environments, a limited specific surface  
183 area[22,48], and non-selective interaction with pollutants[49]. To overcome these  
184 drawbacks and tailor chitosan for targeted remediation tasks, extensive research has  
185 been devoted to developing modern modification strategies. These approaches aim to  
186 enhance three critical performance parameters: stability, selectivity, and efficiency,  
187 thereby transforming chitosan from a naturally occurring polysaccharide into a high-  
188 performance, application-specific material[50].

189 One of the most widely employed methods to improve structural integrity and  
190 chemical resistance is chemical crosslinking[51]. This technique involves the formation  
191 of covalent bonds between chitosan chains using bifunctional agents such as  
192 glutaraldehyde, epichlorohydrin (ECH), genipin, or bisphenol A diglycidyl ether  
193 (BADGE)[51,52]. Crosslinked chitosan matrices exhibit enhanced durability under acidic  
194 conditions where unmodified chitosan would otherwise dissolve, making them suitable  
195 for prolonged use in industrial wastewater treatment systems[53]. For instance, ECH-  
196 cross-linked chitosan grafted with 2,4-dichlorobenzaldehyde has demonstrated high  
197 efficiency in Pb(II) ion removal due to increased availability of active binding sites and  
198 improved mechanical resilience[54]. The use of natural cross-linkers, such as genipin, is  
199 increasingly favored over synthetic alternatives due to their lower toxicity and greater  
200 biocompatibility, aligning with the principles of green chemistry and expanding their  
201 applicability in eco-sensitive domains[55].

202 In parallel, graft copolymerization represents a powerful strategy for enhancing  
203 both functionality and adsorption capacity[56]. By introducing vinyl monomers, such as  
204 acrylic acid, acrylamide, or polyethyleneimine (PEI), onto the chitosan backbone,  
205 researchers can precisely engineer the surface chemistry to target specific classes of  
206 contaminants[56]. PEI-functionalized chitosan aerogels, for example, display exceptional  
207 affinity for Cr(VI) anions even in the presence of competing ions, owing to the high  
208 density of protonated amine groups that facilitate strong electrostatic attraction[57].  
209 Similarly, thio-urea-modified chitosan microspheres synthesized via microfluidic  
210 techniques exhibit rapid kinetics and high specificity for soft metal ions, such as Cd<sup>2+</sup>  
211 and Hg<sup>2+</sup>, due to the formation of stable coordination complexes by sulphur-containing  
212 ligands[58]. These chemically engineered derivatives enable the selective capture of  
213 pollutants in complex effluents, significantly improving removal efficiency and  
214 regeneration potential.

215 Surface functionalization further refines chitosan's performance by introducing  
216 molecular recognition sites that enhance selectivity toward target pollutants[59].  
217 Reactions such as Schiff base formation, where aldehydes like salicylaldehyde react with  
218 chitosan's amino groups to form imine linkages (-C=N-), can create selective chelating  
219 centres for transition metals[60]. Functionalization with amino acids, such as 4-  
220 aminobenzoic acid, adds carboxylate groups that increase coordination capacity with  
221 cationic species like Cu<sup>2+</sup> and Ni<sup>2+</sup>[61]. Additionally, integration of ionic liquids or  
222 surfactant moieties imparts dual functionality, combining hydrophobic interactions with  
223 electrostatic forces to improve dye adsorption, as observed in systems designed for  
224 tartrazine removal[62]. Such targeted modifications transform chitosan from a broadly  
225 reactive biopolymer into a precision material capable of discriminating between  
226 structurally similar contaminants.

227 An efficient approach involves the creation of composite systems, where chitosan is  
228 combined with inorganic, carbonaceous, or polymeric materials to achieve synergistic  
229 enhancements[63]. The synthesis of organic-inorganic composites has proven successful

in producing materials with superior properties compared to individual components. For example, incorporating silica-based materials such as diatomite, sepiolite, or nanosilica not only increases porosity and ion-exchange capacity but also stabilizes the chitosan matrix against swelling and degradation[64]. Studies have shown that chitosan–nanosilica composites (ChNS) and chitosan–silica gel composites (ChSG), prepared through mechanosorption and geometric modification of silica, exhibit significantly enhanced adsorption efficiency, especially under acidic conditions (pH = 2), highlighting the importance of stabilizing chitosan in aggressive environments[65].

Similarly, integration with carbon-based nanomaterials, including graphene oxide (GO), carbon nanotubes (CNTs), and biochar, not only improves mechanical strength and electrical conductivity but also dramatically increases specific surface area and electron transfer rates, which are beneficial for both adsorption and photocatalytic degradation processes[66]. Magnetic nanoparticles, such as Fe<sub>3</sub>O<sub>4</sub> or CoFe<sub>2</sub>O<sub>4</sub>, are embedded within chitosan matrices to combine excellent adsorption properties with the ease of magnetic separation, enabling efficient recovery and reuse without the need for filtration or centrifugation[30,67]. Furthermore, hybrid systems like MOF/chitosan composites leverage the enormous surface area and tunable pores of metal-organic frameworks with chitosan's film-forming ability and processability, yielding highly efficient adsorbents for heavy metals such as Pb(II) and U(VI)[68].

Physical modifications also play a crucial role in overcoming the intrinsic limitations of chitosan. Techniques such as solution casting, granulation, freeze-drying, and cryostructuring enable the transformation of chitosan into various morphologies, including beads, films, flakes, microspheres, aerogels, and membranes, each tailored for specific applications[69]. These physical forms help mitigate defects such as low porosity and low specific surface area, allowing better exposure of functional groups during the adsorption process[70]. For instance, cryostructured composites made from chitosan and silicate minerals exhibit highly interconnected porous networks that are ideal for pollutant uptake. During freezing, ice crystals formed act as templates for macropores, facilitating the rapid diffusion of contaminants into the bulk material[71].

Moreover, chemical alterations such as quaternization have been shown to broaden the solubility spectrum of chitosan from exclusively acidic media (pH < 6) to neutral and basic conditions (pH 4–9)[32]. Quaternary ammonium groups, either introduced directly onto the chitosan backbone or via spacers, impart a permanent positive charge regardless of solution pH, enabling operation across a broader range of environmental conditions[72]. A reported increase in the aqueous solubility of chitosan, up to 80 mg/mL, was achieved through quaternization, demonstrating its potential for formulation in diverse aqueous systems[73].

Another emerging trend is the amplification and modification of complex systems through the incorporation of multifunctional additives. Adding components such as montmorillonite, polycaprolactone, or essential oils not only enhances mechanical and barrier properties but also introduces new functionalities like antimicrobial activity and antioxidant release[74]. Bio-polymer/clay composites, for example, have drawn considerable attention due to the synergistic enhancement of physical and chemical properties compared to pure chitosan[75]. The inclusion of varying amounts of chitosan enables the fine-tuning of composite characteristics, which is crucial for future biomedical and environmental applications[76].

Finally, recent studies emphasize the need to balance the benefits of modification with cost-effectiveness and reusability. While crosslinking improves stability, it may reduce the number of accessible functional groups, thereby affecting adsorption capacity[77]. Therefore, finding an optimal equilibrium between structural

reinforcement and functional availability is critical for maximizing overall performance. Future development should focus on exploring more economical and convenient preparation methods, reducing material costs, improving reuse rates, and minimizing secondary pollution from regeneration processes[78].

In summary, modern strategies for modifying chitosan encompass a broad spectrum of physical and chemical techniques, including crosslinking, grafting, compositing, and quaternization, all aimed at enhancing stability, selectivity, and efficiency. By rationally designing these modifications based on pollutant characteristics and application requirements, researchers are creating next-generation chitosan-based materials that are not only scientifically advanced but also practically viable for real-world environmental remediation.

## 2.2. Fundamental Design Principles Guiding the Development of High-Performance Chitosan Composites

The development of high-performance chitosan composites is rooted in a systematic and multidisciplinary approach that integrates materials science, polymer chemistry, and environmental engineering[79]. These composites are not merely physical mixtures but rationally engineered systems designed to overcome the inherent limitations of native chitosan, such as poor mechanical strength, low thermal stability, acid solubility, swelling-induced disintegration, and limited surface area[80], while amplifying its advantageous properties, including biocompatibility, biodegradability, cationic character, and rich functional group chemistry[81]. The successful design of these advanced materials relies on several fundamental principles: interfacial compatibility and adhesion, morphological control and architectural engineering, functional synergy between components, stabilization through crosslinking, and application-driven tailoring[82].

A cornerstone of composite design is achieving strong interfacial compatibility and adhesion between the chitosan matrix and the reinforcing or functional filler. Without adequate bonding, phase separation, agglomeration, and poor stress transfer can occur, leading to structural weaknesses and reduced performance[83]. To enhance compatibility, fillers such as diatomite, sepiolite, nanosilica (NS), silica gel (SG), graphene oxide (GO), metal oxides (e.g., ZnO, Fe<sub>3</sub>O<sub>4</sub>, TiO<sub>2</sub>), or clay minerals (e.g., montmorillonite, MMT; natural zeolite, NZ) are often surface-modified or integrated via chemical linkers[84]. For instance, studies have demonstrated that chitosan–nanosilica composites (ChNS) and chitosan–silica gel composites (ChSG) synthesized through mechanosorption and geometric modification of silica exhibit significantly improved adsorption efficiency, particularly under acidic conditions (pH = 2), where unmodified chitosan would typically dissolve[49,85]. The use of cross-linkers, such as glutaraldehyde (GA), further strengthens the interface by forming covalent bonds between chitosan chains and the filler surface, as evidenced in composites like ChNS\_1\_GA and ChSG\_1\_GA, which exhibit enhanced structural integrity and resistance to degradation[86].

Closely tied to interfacial engineering is the principle of morphological control and architectural design[87]. The physical form and internal architecture of a composite directly influence its functional performance, especially in applications involving mass transfer, such as adsorption or catalysis[88]. Techniques such as solution casting, drop granulation, freeze-drying, cryostructuring, electrospinning, and hot-water stretching enable precise manipulation of porosity, pore size distribution, specific surface area, and mechanical resilience[89]. For example, cryostructured composites prepared from chitosan and silicate minerals exhibit highly interconnected porous networks that

329 facilitate the rapid diffusion of pollutants into the bulk material, making them ideal for  
330 the efficient uptake of persistent pharmaceuticals, such as carbamazepine (CBZ) and  
331 ciprofloxacin (CIP)[59,90]. Similarly, scanning electron microscopy (SEM) analyses of  
332 chitosan–SiO<sub>2</sub> composites reveal well-dispersed morphologies with smooth surfaces and  
333 minimal agglomeration, indicating good integration and homogeneity at the  
334 microscale[91]. In fibre-based systems, wet-spinning followed by immersion in  
335 dopamine lye has been shown to create a protective polydopamine (PDA) coating  
336 around chitosan fibres, significantly improving their mechanical properties and enabling  
337 potential use in biomedical and load-bearing applications[92].

338 Another critical design principle is the establishment of functional synergy between  
339 components, where each constituent contributes uniquely to the overall capability of the  
340 system[93]. Rather than simply combining materials, modern composites are engineered  
341 so that the whole performs better than the sum of its parts[94]. Magnetic nanoparticles  
342 such as Fe<sub>3</sub>O<sub>4</sub> or CoFe<sub>2</sub>O<sub>4</sub> are embedded within chitosan matrices not only to reinforce  
343 mechanical properties but also to enable easy magnetic separation after pollutant  
344 binding, a feature essential for practical wastewater treatment operations[95].  
345 Photocatalytic materials, such as TiO<sub>2</sub> or ZnO, introduce light-responsive degradation  
346 capabilities, allowing chitosan composites to act not only as passive adsorbents but also  
347 as active agents in breaking down organic dyes, including methyl orange and reactive  
348 red 120, under UV irradiation[96]. Antimicrobial functionalities are similarly enhanced  
349 by incorporating silver nanoparticles, curcumin, or rosemary extract, which work  
350 synergistically with chitosan's intrinsic biocidal properties[87,97]. Hybrid systems, such  
351 as chitosan-alginate@Fe/Mn mixed oxide nanocomposites, exemplify this synergy by  
352 first reducing toxic Cr(VI) to less harmful Cr(III) through redox reactions and then  
353 immobilizing the resulting cation through complexation with the amino groups of  
354 chitosan[51].

355 Stabilization through crosslinking plays a decisive role in determining the  
356 durability and operational lifespan of chitosan composites[98]. Crosslinking agents such  
357 as glutaraldehyde, epichlorohydrin (ECH), genipin, vanillin, or tripolyphosphate (TPP)  
358 create a three-dimensional network that resists dissolution in aqueous environments,  
359 particularly under acidic conditions[52]. This stabilization is crucial for maintaining  
360 structural integrity during prolonged exposure to industrial effluents[99]. Moreover,  
361 natural cross-linkers, such as genipin, offer lower cytotoxicity compared to synthetic  
362 alternatives, making them suitable for eco-friendly and biomedical applications[100].  
363 Studies on porous chitosan composites stabilized using various methods have shown  
364 that crosslinking not only enhances mechanical strength but also affects pycnometric  
365 density and specific surface area, which are vital parameters for both implant materials  
366 and adsorbents[101].

367 Ultimately, the design of chitosan composites should be guided by the intended  
368 application, ensuring that the material meets the functional, economic, and sustainability  
369 criteria[95]. For environmental remediation, composites are optimized for high porosity,  
370 selectivity toward target pollutants (e.g., Pb(II), Cd(II), As(V), and anionic dyes), and  
371 regenerability[2,102]. In biomedical contexts, emphasis is placed on cytocompatibility,  
372 controlled release, antibacterial action, and mechanical mimicry of natural tissues. For  
373 food packaging, barrier properties against moisture and gases, as well as antioxidant  
374 and antimicrobial functions, are prioritized[103]. The inclusion of varying amounts of  
375 chitosan enables the fine-tuning of composite characteristics, for example, adjusting the  
376 chitosan/carbon ratio in aerogel composites (Car/CSAs) to maximize methylene blue  
377 (MB) removal efficiency[104]. Similarly, in dental restorative materials, the addition of  
378 nano-chitosan to adhesive and flowable composite resins has been evaluated for its

379 impact on mechanical phenomena, solubility, and antimicrobial activity, demonstrating  
380 how even minor modifications can yield significant functional improvements[105].

381 Thus, the rational design of high-performance chitosan composites follows a  
382 holistic framework that balances interfacial adhesion, structural architecture, functional  
383 integration, chemical stabilization, and end-use requirements. By leveraging these  
384 fundamental principles, researchers are creating innovative, sustainable, and  
385 multifunctional materials that fully exploit the potential of one of nature's most versatile  
386 biopolymers, such as chitosan, paving the way for scalable and environmentally  
387 responsible solutions across diverse fields.

### 388 *2.3. Key Mechanisms Underlying Pollutant Removal via Adsorption, Catalysis, Membrane* 389 *Filtration, and Flocculation*

390 The effectiveness of chitosan-based materials in environmental remediation stems  
391 from their ability to engage multiple mechanisms for pollutant removal, each operating  
392 through distinct physicochemical pathways. These mechanisms, such as adsorption,  
393 catalysis, membrane filtration, and flocculation, which are not mutually exclusive but  
394 often function synergistically within engineered composites, enable efficient, selective,  
395 and scalable treatment of complex wastewater streams[106]. The integration of these  
396 processes allows for the simultaneous targeting of diverse contaminants, including  
397 heavy metals, organic dyes, pharmaceuticals, radionuclides, and microbial agents[107].

398 Adsorption is widely recognized as one of the most effective, economical, and  
399 environmentally friendly methods for removing pollutants from aqueous systems[108].  
400 It involves the accumulation of contaminant molecules (adsorbates) onto the surface of a  
401 solid material (adsorbent) through various intermolecular forces[109]. Chitosan and its  
402 derivatives are particularly effective adsorbents due to their abundant functional groups,  
403 primarily primary amino ( $-NH_2$ ) and hydroxyl ( $-OH$ ) groups, which participate in  
404 multiple interaction mechanisms[9,29,110]. Under acidic conditions ( $pH < 6$ ), the  
405 protonation of amino groups generates positively charged  $-NH_3^+$  sites that facilitate  
406 strong electrostatic attraction with anionic species such as chromate [Cr(VI)], arsenate  
407 [As(V)], sulfate dyes, uranyl ions [ $UO_2^{2+}$ ], and other oxyanions[51,61,63]. For cationic  
408 pollutants like Pb(II), Cd(II), Cu(II), Ni(II), and Zn(II), adsorption occurs via  
409 complexation or chelation, where metal ions coordinate with lone electron pairs on  
410 nitrogen and oxygen atoms, forming stable five- or six-membered rings[111,112]. This  
411 mechanism is further enhanced in modified chitosan systems, such as those  
412 incorporating carboxylate, phosphate, or thiol groups, which increase the number of  
413 available binding sites and improve selectivity[113].

414 In addition to electrostatic and coordination interactions, hydrogen bonding, van  
415 der Waals forces, and  $\pi$ - $\pi$  stacking contribute to the overall adsorption process,  
416 particularly when chitosan is combined with carbonaceous materials such as graphene  
417 oxide (GO) or biochar[114]. The presence of oxygen-containing functional groups on GO  
418 enhances its affinity for both organic and inorganic pollutants, making it highly effective  
419 for the adsorption of radionuclides such as Eu(III), Pu(IV), and Am(III)[115,116].  
420 Furthermore, ion exchange plays a significant role in composites containing inorganic  
421 fillers such as zeolites, layered silicates, or hydroxyapatite, where target metal ions  
422 replace mobile counterions (e.g.,  $Ca^{2+}$ ,  $Na^+$ ) during the uptake process[117].

423 The intraparticle diffusion model reveals that the adsorption process typically  
424 proceeds through three distinct stages: bulk diffusion, film diffusion, and pore  
425 diffusion[118]. Bulk diffusion refers to the transport of pollutants from the bulk solution  
426 to the external surface of the adsorbent. Film diffusion involves the movement of  
427 contaminants across the boundary layer surrounding the particle, while pore diffusion

governs the penetration of adsorbates into the internal porous network. In diatomite-chitosan composites, for example, physical adsorption, electrostatic interaction, and surface complexation collectively enhance vanadium [V(V)] removal efficiency[119]. Notably, some chitosan-based materials also exhibit reductive adsorption, where functional groups such as hydroxyl oxygen atoms reduce toxic V(V) to less harmful V(IV), thereby combining chemical transformation with immobilization[120].

Catalysis represents an advanced mode of pollutant removal that surpasses passive capture, enabling the active degradation of persistent organic pollutants[121]. While native chitosan is not inherently catalytic, it serves as an excellent support matrix for catalytically active species such as metal nanoparticles (e.g., Fe/Mn mixed oxides, TiO<sub>2</sub>, ZnO), enzymes, or conjugated polymers[122]. Chitosan-alginate@Fe/Mn mixed oxide nanocomposites, for instance, demonstrate high efficiency in reducing toxic Cr(VI) to less hazardous Cr(III) through redox reactions, followed by immobilization via complexation with chitosan's amino groups[40,123]. This dual functionality, simultaneous reduction and adsorption, significantly improves detoxification outcomes. Similarly, chitosan-supported photocatalysts, such as TiO<sub>2</sub> or SnO<sub>2</sub>, can degrade organic dyes under UV or visible light irradiation, leveraging the biopolymer's film-forming ability and dispersion capacity to prevent nanoparticle aggregation[124]. In more sophisticated systems, chitosan-based aerogels or cryogels act as hosts for noble metal catalysts, facilitating reactions such as the oxidation of phenolic compounds or the breakdown of antibiotics like tetracycline without generating harmful byproducts[125].

Membrane filtration utilizing chitosan-based membranes has gained increasing attention due to the polymer's excellent film-forming properties, tunable porosity, and antimicrobial characteristics[126]. These membranes operate through a combination of size exclusion, charge repulsion, and adsorptive retention, depending on the membrane architecture and surface chemistry[127]. Composite membranes made from chitosan and graphene oxide (CSGO), for example, exhibit improved flux, reduced fouling, and high rejection rates for dyes and heavy metals, making them suitable for greywater treatment and non-potable reuse applications[128]. The incorporation of GO increases mechanical strength and specific surface area, while chitosan contributes positive surface charge and biofouling resistance. Moreover, pH-responsive swelling behavior enables smart membrane operation, where pore size and permeability can be modulated by adjusting the solution pH, making them ideal for selective separation processes[129]. In some configurations, such as mixed matrix membranes (MMMs) or membrane adsorbers, embedded adsorbent particles (e.g., activated carbon, MOFs) provide additional binding sites, transforming the membrane into a multifunctional platform capable of both filtration and adsorption[130,131].

Flocculation is another vital mechanism in which chitosan acts as a natural cationic flocculant, neutralizing negatively charged colloidal particles and suspended solids in wastewater[132]. Upon addition to contaminated water, chitosan chains adsorb onto particle surfaces, compressing the electrical double layer and promoting aggregation through charge neutralization and polymer bridging[133]. This results in the formation of larger flocs that settle rapidly under gravity, facilitating easy removal of turbidity, organic matter, and even pathogens[134]. The effectiveness of chitosan as a flocculant depends on its degree of deacetylation, molecular weight, and solution pH[23,51,135]. High-DD chitosan offers more amino groups for charge interactions, while a higher molecular weight enhances chain extension and inter-particle bridging capability[136]. Unlike synthetic flocculants such as polyacrylamide, chitosan is biodegradable and non-toxic, making it an environmentally favourable alternative[135]. It has been successfully used in treating textile effluents, food processing wastewaters, and algal blooms, often

478 outperforming conventional coagulants in terms of reducing sludge volume and  
479 improving the clarity of treated water[137].

480 The synergy between these mechanisms is increasingly exploited in multifunctional  
481 chitosan composites. For example, a single system may combine adsorptive capture,  
482 catalytic degradation, and magnetic separation, enabling complete pollutant elimination  
483 with minimal operational complexity[138]. The design of such integrated platforms  
484 relies on a deep understanding of interfacial phenomena, mass transfer dynamics, and  
485 reaction kinetics, all of which are informed by advanced characterization techniques and  
486 modelling approaches[139].

487 In summary, chitosan-based materials employ a multifaceted approach to pollutant  
488 removal, integrating adsorption, catalysis, membrane filtration, and flocculation in a  
489 manner that is both complementary and synergistic[109]. By rationally engineering the  
490 chemical structure, morphology, and functional architecture of these composites,  
491 researchers can tailor their performance to specific environmental challenges. This  
492 mechanistic versatility underscores chitosan's role as a cornerstone material in the  
493 development of next-generation, sustainable water purification technologies[140].

#### 494 *2.4 Practical Challenges Related to Scalability, Regeneration, Lifecycle Sustainability, and Real-* 495 *World Implementation*

496 Despite the extensive research and promising laboratory-scale performance of  
497 chitosan-based materials for environmental applications, their transition from benchtop  
498 innovation to widespread industrial and municipal deployment faces a series of  
499 interconnected practical challenges[20]. These include issues related to the scalability of  
500 synthesis, regeneration, and reusability, as well as lifecycle sustainability and real-world  
501 operational constraints, all of which must be addressed to ensure that chitosan  
502 composites are not only scientifically practical but also economically viable,  
503 environmentally sound, and technically feasible in complex, dynamic  
504 environments[141].

505 One of the most significant barriers to large-scale adoption is the scalability of  
506 fabrication methods. Many high-performance chitosan composites are synthesized using  
507 techniques such as freeze-drying, electrospinning, layer-by-layer assembly, or  
508 cryostructuring, which offer excellent control over morphology, porosity, and functional  
509 integration[22,47,142]. However, these processes are inherently complex to scale up due  
510 to high energy demands, batch-to-batch variability, and low throughput. For example,  
511 aerogels and cryogels with ultra-high surface areas and 3D porous networks  
512 demonstrate exceptional adsorption capacity in controlled settings; however, their  
513 production requires specialized equipment, such as lyophilizers or supercritical CO<sub>2</sub>  
514 dryers, making them impractical for continuous manufacturing at industrial levels[143].  
515 Similarly, graft copolymerization, magnetic nanoparticle incorporation, or surface  
516 functionalization often involves multi-step procedures, toxic crosslinking agents (e.g.,  
517 glutaraldehyde), or stringent reaction conditions that complicate process  
518 standardization and raise concerns regarding safety and waste management[144]. To  
519 enable mass production without compromising performance, there is an urgent need for  
520 simple, reproducible, and environmentally friendly fabrication routes, such as solution  
521 casting, drop granulation, dip-coating, or air-spray deposition, that can be seamlessly  
522 integrated into existing water treatment infrastructure[145].

523 Closely linked to scalability is the issue of mechanical and chemical stability under  
524 real operating conditions. In practical wastewater treatment scenarios, chitosan-based  
525 materials must endure fluctuating pH levels, high ionic strength, microbial activity,  
526 abrasive flow dynamics, and prolonged exposure to organic solvents or

527 surfactants[136,146]. Native chitosan dissolves readily in acidic environments (pH < 6),  
528 severely limiting its applicability in effluents rich in mineral acids or organic acids  
529 commonly found in textile, mining, or electroplating industries[51]. Although  
530 crosslinking improves acid resistance, it may reduce the availability of active binding  
531 sites and slow down adsorption kinetics[77,147]. Furthermore, repeated swelling and  
532 deswelling cycles during use and regeneration can lead to structural fatigue, cracking, or  
533 disintegration over time[29]. Field trials have shown that while chitosan-diatomite,  
534 chitosan-sepiolite, or chitosan-alginate composites perform well initially, their efficiency  
535 declines after extended exposure to complex matrices containing oils, suspended solids,  
536 or competing ions[148]. Therefore, developing mechanically robust, chemically stable,  
537 and fouling-resistant composites remains a critical challenge for long-term  
538 implementation[149].

539 Another crucial factor affecting the economic viability of chitosan-based systems is  
540 regeneration and reusability. An ideal adsorbent should not only exhibit high removal  
541 capacity but also be efficiently regenerated and reused over multiple cycles to minimize  
542 material consumption and operational costs[150]. Several studies confirm that certain  
543 chitosan composites can retain significant functionality after regeneration. For instance,  
544 PEI-functionalized chitosan hydrogels maintained about 66% of their initial adsorption  
545 capacity after four cycles when eluted with 1 M HCl, indicating good recovery  
546 potential[151]. However, regeneration protocols often require large volumes of strong  
547 acids, bases, or organic solvents, raising concerns about secondary pollution, resource  
548 consumption, and handling hazards[152]. The necessity for extensive washing, where 1  
549 g of sorbent may require up to 0.3 L of water, further questions the ecological  
550 justification of regeneration, especially in water-scarce regions[153]. As a result,  
551 alternative end-of-life strategies are being explored, such as converting spent sorbents  
552 into construction materials, soil amendments, or catalyst supports, thereby shifting from  
553 circular reuse to integrated valorisation[154].

554 From a broader perspective, the lifecycle sustainability of chitosan composites must  
555 be evaluated holistically. While chitosan itself is derived from renewable sources,  
556 primarily seafood waste, and is biodegradable, the addition of synthetic polymers,  
557 nanoparticles, or non-biodegradable fillers (e.g., graphene oxide, carbon nanotubes,  
558 polyvinyl alcohol) can alter its environmental footprint[155]. Energy-intensive extraction  
559 and deacetylation processes, involving concentrated NaOH and HCl, contribute to  
560 greenhouse gas emissions and generate alkaline/acidic wastewater, thereby  
561 undermining the overall green credentials of the final product[31]. A comprehensive  
562 lifecycle analysis (LCA) should therefore assess all stages from raw material sourcing  
563 and processing to application, regeneration, and final disposal or recycling, to ensure net  
564 environmental benefit[156]. Emphasis should be placed on minimizing chemical usage,  
565 reducing energy input, and maximizing recyclability and biodegradability across the  
566 entire value chain[157].

567 Economic feasibility remains a key factor in determining real-world adoption.  
568 Although it is a byproduct of the fishing industry, purified chitosan can be more  
569 expensive than conventional adsorbents, such as activated carbon or ion-exchange  
570 resins[158]. Cost-effectiveness improves significantly when unrefined seafood waste  
571 (e.g., shrimp shells, crab exoskeletons) is used directly as feedstock, or when composites  
572 are designed for dual functions, such as simultaneous oil-water separation and heavy  
573 metal capture, or combined flocculation and disinfection[159]. Integration with existing  
574 technologies, such as fixed-bed columns, membrane modules, or transformer oil  
575 regeneration units, also enhances feasibility[31]. Pilot-scale testing in actual industrial  
576 effluent streams has demonstrated promising results, showing that chitosan composites

577 can extend equipment lifespan, reduce sludge volume, and lower hazardous waste  
578 generation[160]. Moreover, the use of composite adsorbents for regenerating insulating  
579 transformer oil contributes to both environmental protection and energy resilience by  
580 minimizing the need for frequent oil replacement and disposal[161].

581 In summary, while chitosan-based materials hold immense promise for sustainable  
582 environmental remediation, their widespread implementation requires overcoming  
583 multifaceted challenges beyond laboratory performance[162]. Addressing scalability,  
584 ensuring robustness under variable conditions, optimizing regeneration protocols,  
585 conducting rigorous lifecycle assessments, and aligning with circular economy  
586 principles are essential steps toward transforming these innovative materials from  
587 scientific curiosities into practical, economically viable, and environmentally responsible  
588 solutions for global pollution control[163]. Future research must focus on bridging the  
589 gap between scientific potential and real-world impact, ensuring that the full benefits of  
590 chitosan technology are realized not only in journals but also in rivers, soils, industries,  
591 and communities worldwide[164].

#### 592 *2.5. Emerging Trends: Circular Economy Integration, Seafood Waste Valorisation, and Digital* 593 *Optimization Using Artificial Intelligence*

594 The field of environmental remediation using chitosan-based materials is  
595 undergoing a profound transformation, driven by global imperatives for sustainability,  
596 resource efficiency, and technological innovation. Among the most significant emerging  
597 trends are the integration of chitosan technology within circular economy frameworks,  
598 the systematic valorisation of seafood processing waste as a primary feedstock, and the  
599 application of digital tools, particularly artificial intelligence (AI) and machine learning  
600 (ML) for material design, process optimization, and system monitoring. These  
601 interconnected advancements are shifting the paradigm from linear, extractive models  
602 of production to regenerative, intelligent systems that align scientific progress with  
603 ecological responsibility[165,166].

604 A cornerstone of this evolution is the integration of chitosan into circular economy  
605 principles, a model that emphasizes closed-loop systems, waste minimization, and the  
606 continuous reuse of resources[167]. Chitosan exemplifies this approach by transforming  
607 an abundant industrial byproduct, such as crustacean shells from fisheries and  
608 aquaculture, into high-value functional materials for water treatment, food packaging,  
609 biomedical applications, and agriculture[168]. Globally, over 6-8 million tons of shellfish  
610 waste are generated annually, often ending up in landfills or being incinerated, which  
611 contributes to environmental pollution and greenhouse gas emissions[169]. By  
612 redirecting this waste stream into biorefineries for chitin and chitosan extraction,  
613 industries can simultaneously reduce their ecological footprint and create new economic  
614 opportunities[170]. Furthermore, end-of-life strategies for spent chitosan composites are  
615 increasingly designed with circularity in mind: exhausted adsorbents loaded with heavy  
616 metals can be processed for metal recovery via acid elution and precipitation. At the  
617 same time, biodegradable matrices can be composted or repurposed as soil conditioners,  
618 thereby closing nutrient loops and supporting sustainable agriculture[154]. The  
619 European Union's new circular economy action plan explicitly promotes biopolymers,  
620 such as chitosan, as revolutionary alternatives to conventional synthetic polymers that  
621 persist in ecosystems, underscoring the policy-level recognition of their transformative  
622 potential[171].

623 Closely linked to this shift is the valorisation of seafood waste, which has emerged  
624 as a critical driver of sustainable chitosan production. Traditionally, shrimp, crab,  
625 lobster, and other crustacean shells were considered low-value residues requiring costly

626 disposal. However, recent innovations in green extraction technologies, such as  
627 enzymatic deproteinization, microbial fermentation, and mild demineralization using  
628 organic acids, have enabled the more efficient and eco-friendly recovery of high-purity  
629 chitin and chitosan[172]. Enzymatic methods, in particular, offer significant advantages  
630 over conventional acid-alkali treatments by reducing energy consumption, chemical  
631 usage, and wastewater generation[173]. For instance, proteolytic enzymes such as  
632 alcalase or papain selectively break down proteins without damaging the chitin  
633 structure, yielding higher-quality products with a lower environmental impact[174].  
634 Moreover, studies involving *Bacillus sp.* XT7 and other microorganisms demonstrate  
635 how fermentative processes can integrate deproteinization and demineralization steps  
636 under mild conditions, further enhancing process sustainability[175]. This trend extends  
637 beyond crustaceans; research on black tiger shrimp (*Penaeus monodon*) heads has  
638 demonstrated the use of integrated approaches for recovering both protein hydrolysate  
639 and chitosan, moving toward zero-waste biorefinery models[176]. Similarly, fungal  
640 biomass and insect exoskeletons are being explored as alternative sources, diversifying  
641 supply chains and reducing pressure on marine ecosystems[177].

642 Parallel to these advancements in bioresources, a new frontier is emerging through  
643 the application of artificial intelligence (AI), machine learning (ML), and data-driven  
644 modeling in the development and deployment of chitosan-based systems[178].  
645 Traditional approaches to designing chitosan composites rely heavily on empirical  
646 experimentation, a time-consuming and resource-intensive process that is prone to  
647 inefficiencies[22,179]. In contrast, AI-driven methodologies enable predictive analytics,  
648 real-time monitoring, and adaptive control across the entire lifecycle of chitosan  
649 materials[180]. Machine learning algorithms have been successfully applied to optimize  
650 biochar synthesis parameters, accurately forecasting yield and performance based on  
651 feedstock composition, pyrolysis temperature, and activation conditions, which is an  
652 approach directly transferable to chitosan composite fabrication[181]. Similarly, neural  
653 networks and regression models can simulate adsorption isotherms, kinetic behavior,  
654 and mechanical properties under varying pH, ionic strength, and contaminant  
655 concentrations, allowing researchers to identify optimal formulations without the need  
656 for exhaustive laboratory testing[182].

657 Advanced computational techniques, such as adaptive neuro-fuzzy inference  
658 systems (ANFIS) and artificial neural networks (ANNs), are proving particularly  
659 effective in modeling complex adsorption processes[183]. For example, ANNs have been  
660 used to predict the removal efficiency of methylene blue dye using modified clay,  
661 outperforming traditional response surface methodology (RSM) in accuracy and  
662 flexibility[184,185]. These models can account for nonlinear interactions between  
663 variables such as contact time, initial concentration, pH, and adsorbent dosage, which  
664 are difficult to capture with conventional statistical methods[186]. Furthermore, AI  
665 integration extends to operational decision-making in wastewater treatment plants  
666 where chitosan-based adsorbents are deployed. Intelligent sensors, combined with data  
667 analytics platforms, can monitor effluent quality in real-time, dynamically adjusting  
668 dosage rates, contact times, and regeneration cycles to maximize efficiency and minimize  
669 chemical consumption[160].

670 Digital twins, as virtual replicas of physical treatment systems, can simulate long-  
671 term performance, forecast maintenance needs, and evaluate the environmental impact  
672 of different operational scenarios[187]. When coupled with blockchain-enabled  
673 traceability systems, these technologies ensure transparency across the chitosan supply  
674 chain, from raw material sourcing to final disposal[188]. This level of digital integration

not only accelerates innovation but also enhances accountability and public trust in green technologies[189].

Moreover, AI is playing a crucial role in optimizing emerging hybrid systems such as magnetic nanocomposites, photocatalytic membranes, and stimuli-responsive hydrogels[190]. For instance, multi-criteria decision-making tools, such as fuzzy AHP-TOPSIS, are being used to select the best colour removal process using carbon-based adsorbents under real-world conditions, balancing technical performance with economic feasibility[191]. Similarly, AI-assisted analysis of georeferenced medical and viral dispersion data is paving the way for integrated epidemic control systems, demonstrating how digital infrastructure can support broader sustainability goals beyond environmental remediation[192].

In summary, the convergence of circular economy thinking, seafood waste valorisation, and AI-driven optimization represents a paradigm shift in the field of chitosan science[141]. These trends collectively move the discipline away from isolated, linear research efforts toward integrated, intelligent, and sustainable solutions[44]. As global pressures on natural resources intensify, the ability to transform waste into high-performance materials using data-informed, eco-conscious methods will be essential, not only for advancing scientific knowledge but also for building resilient and regenerative environmental systems[193]. The continued evolution of chitosan technology in this direction promises to position it at the forefront of the green materials revolution, offering scalable, innovative, and sustainable solutions to some of the world's most pressing environmental challenges[20].

### 3. Practical Environmental Applications of Chitosan Derivatives and Composites

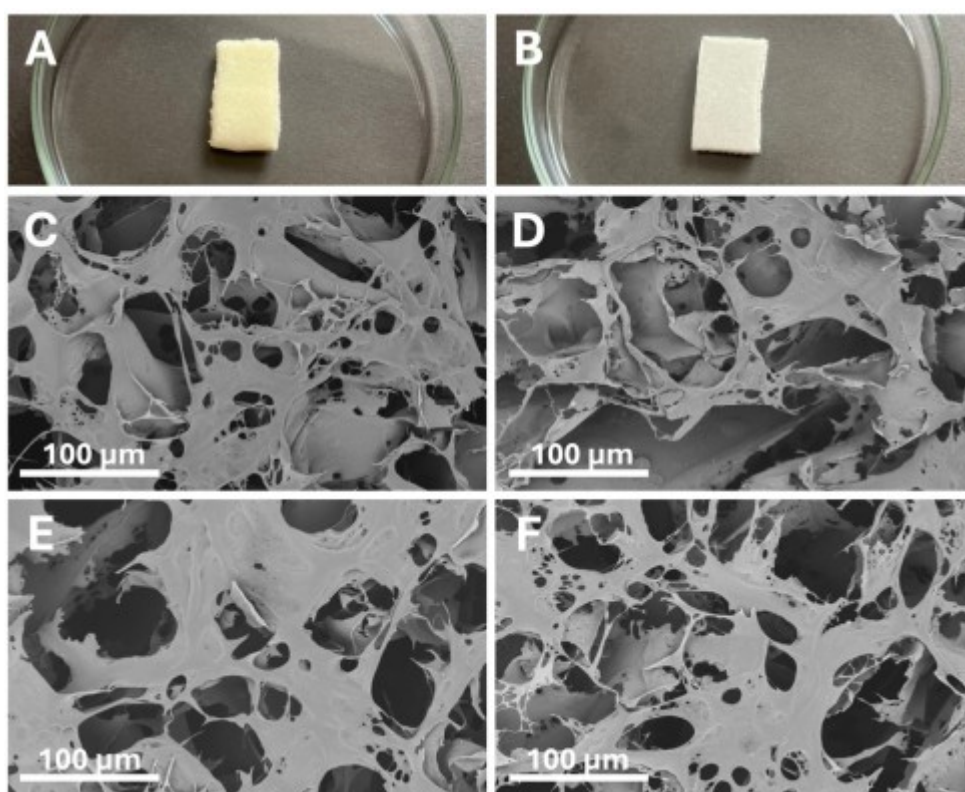
#### 3.1. Water Treatment

Chitosan and its derivatives have emerged as pivotal materials in modern water treatment technologies due to their exceptional physicochemical properties, biocompatibility, and sustainable origin. Derived from the deacetylation of chitin, abundant in crustacean shells, chitosan is a linear cationic polysaccharide with reactive amino and hydroxyl groups that enable multifunctional interactions with a wide range of aquatic pollutants[194]. Its application spans heavy metal removal, dye decolorization, organic contaminant adsorption, and microbial control, making it one of the most versatile biopolymers for environmental remediation[195].

##### 3.1.1. Heavy Metal Removal

The removal of toxic heavy metals such as Pb(II), Cd(II), Cu(II), Cr(VI), As(V), and Zn(II) from aqueous systems is a critical aspect of water purification, particularly in industrial effluents and contaminated groundwater[196]. Chitosan-based materials excel in this domain through several mechanisms, including coordination, ion exchange, and electrostatic interactions. Under acidic conditions ( $\text{pH} < 6$ ), the protonation of primary amine groups ( $-\text{NH}_3^+$ ) facilitates strong electrostatic attraction toward anionic species like chromate ( $\text{CrO}_4^{2-}$ ) and arsenate ( $\text{AsO}_4^{3-}$ )[51,61]. For cationic metals, the lone electron pairs on nitrogen and oxygen atoms coordinate with metal ions, forming stable five- or six-membered chelate rings[61,197]. Ion exchange also contributes significantly, especially when chitosan is combined with inorganic fillers such as zeolites or layered silicates, where target metal ions replace mobile counterions during the uptake process[63].

To enhance performance, chitosan is often integrated into composite structures. Hydroxyapatite-chitosan composites leverage the high affinity of calcium phosphate for divalent cations like  $Pb^{2+}$  and  $Cd^{2+}$ , while chitosan provides mechanical stability and additional binding sites[198]. Similarly,  $Fe_3O_4$ -chitosan magnetic composites offer an excellent adsorption capacity combined with ease of separation via external magnetic fields, enabling efficient recovery and reuse without the need for filtration[199]. Graphene oxide (GO)-chitosan composites offer ultra-high surface area and  $\pi$ - $\pi$  stacking capabilities, improving sorption kinetics and capacity for both metallic and organic contaminants. These hybrid systems demonstrate synergistic effects, outperforming individual components in real wastewater matrices[138]. Visualization of chitosan sponge and chitosan sponge with GO, characterized by numerous pores of various sizes, is shown in Figure 2.



**Figure 2.** The representative photographic images of the (A) sponges made of microcrystalline chitosan sponge and (B) sponge made of microcrystalline chitosan with the addition of GO. SEM image of the outer surface of the sponge made of microcrystalline chitosan sponge with and without the addition of GO, where (C) microcrystalline chitosan sponge surface before sterilization, (D) microcrystalline chitosan sponge + GO sponge surface before sterilization, (E) microcrystalline chitosan sponge surface after sterilization (25 kGy), (F) microcrystalline chitosan sponge + GO sponge surface after sterilization (25 kGy). Magnification 1500 $\times$ , presented scale bar indicates 100  $\mu m$ [200].

Adsorption efficiency is typically evaluated using kinetic and equilibrium models. Kinetic studies commonly follow pseudo-second-order behaviour, indicating chemisorption as the rate-limiting step, while diffusion models reveal contributions from film and intraparticle transport[114,201]. Equilibrium data are best described by the Langmuir isotherm, suggesting monolayer adsorption, or the Freundlich model, which

751 indicates heterogeneous surface binding. Maximum adsorption capacities reported in  
752 the literature vary widely, depending on the composite design, from hundreds to over  
753 1000 mg/g, highlighting the tunability of these materials[147,202].

754 Regeneration and reuse are essential for economic viability. Spent chitosan  
755 composites can be eluted with dilute acids (e.g., HCl) or chelating agents (e.g., EDTA),  
756 recovering up to 80% of the initial capacity after multiple cycles. However, repeated  
757 exposure to harsh regeneration protocols may degrade the polymer matrix, necessitating  
758 robust crosslinking strategies[49,90].

759 A notable case study involves an EDTA-functionalized magnetic chitosan  
760 nanocomposite, designed for selective and efficient capture of Pb(II) and Cd(II)[49]. The  
761 incorporation of ethylenediaminetetraacetic acid (EDTA) moieties enhances chelation  
762 strength, while Fe<sub>3</sub>O<sub>4</sub> nanoparticles allow magnetic retrieval. This system exhibits rapid  
763 kinetics, high selectivity even in multi-metal solutions, and effective regeneration using  
764 mild acid washes, demonstrating its potential for scalable deployment in industrial  
765 wastewater treatment[199,203].  
766

### 767 3.1.2 Dye Removal

768 Industrial effluents from textile, paper, and printing sectors contain persistent  
769 synthetic dyes, which are classified as anionic (e.g., Reactive Orange 16, Methyl Orange),  
770 cationic (e.g., Methylene Blue), and azo dyes (e.g., Reactive Red 120) that pose significant  
771 ecological risks due to their toxicity and resistance to degradation[49,114]. Chitosan  
772 effectively removes these dyes primarily through electrostatic adsorption, where  
773 protonated amino groups attract oppositely charged dye molecules. Anionic dyes are  
774 readily captured under acidic conditions, whereas cationic dyes require modification of  
775 chitosan's surface charge[204].

776 Crosslinking agents such as glutaraldehyde, epichlorohydrin, or bisphenol A  
777 diglycidyl ether improve chemical stability and prevent dissolution in aqueous  
778 media[51]. Functionalization strategies, including quaternization, introduce permanent  
779 positive charges, thereby extending the applicability of chitosan to neutral and alkaline  
780 environments. Quaternized chitosan derivatives exhibit enhanced affinity for anionic  
781 dyes regardless of pH, offering operational flexibility[205].

782 Chitosan is processed into various forms, including composite films, beads, and  
783 hydrogels, to optimize contact area and mass transfer. Beads prepared via dropwise  
784 gelation exhibit high reusability in fixed-bed columns, while hydrogels offer swelling-  
785 controlled release and enhanced accessibility of functional groups[206]. Films  
786 incorporating graphene oxide or clay minerals exhibit superior mechanical strength and  
787 barrier properties, making them suitable for membrane-based filtration[128].

788 Despite their efficacy, challenges remain. High ionic strength and competing anions  
789 (e.g., SO<sub>4</sub><sup>2-</sup>, Cl<sup>-</sup>) can interfere with dye adsorption, reducing efficiency[207].  
790 Additionally, some systems suffer from slow kinetics due to limited pore diffusion,  
791 particularly in dense matrices. Addressing these limitations requires tailored porosity  
792 engineering and the use of stimuli-responsive designs[208].  
793

### 794 3.1.3. Organic Pollutant Adsorption

795 Beyond metals and dyes, chitosan composites are increasingly used to remove  
796 emerging organic pollutants, including pharmaceuticals, pesticides, and endocrine  
797 disruptors (e.g., bisphenol A, ibuprofen, carbamazepine). The dominant mechanisms  
798 include  $\pi$ - $\pi$  stacking between aromatic rings in contaminants and graphitic domains in

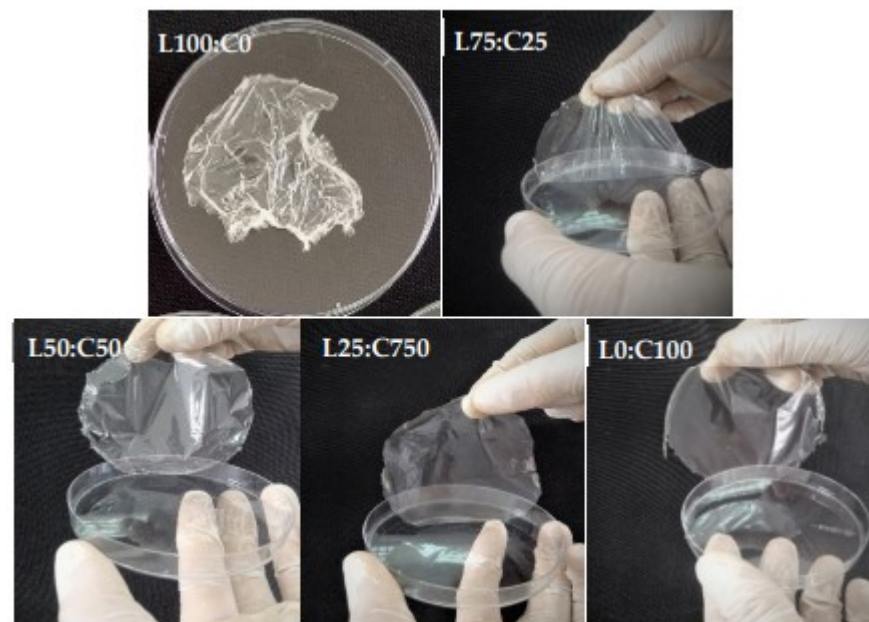
799 carbonaceous supports, as well as hydrogen bonding involving –OH and –NH<sub>2</sub>  
800 groups[209].

801 Graphene oxide-chitosan composites are particularly effective due to GO's large  
802 surface area and oxygen-containing functional groups, which enhance dispersion and  
803 interaction with polar organics[210]. In more advanced systems, molecularly imprinted  
804 polymers (MIPs) based on chitosan templates create selective recognition sites for  
805 specific analytes, mimicking the specificity of antibodies[211]. However, challenges  
806 persist in achieving high selectivity in complex mixtures and ensuring complete  
807 regeneration without template leaching.

### 808 3.2. Waste Management

#### 809 3.2.1. Biodegradable Packaging

811 Chitosan-based films and coatings represent a sustainable alternative to petroleum-  
812 derived plastics in food packaging. Their inherent antimicrobial activity, film-forming  
813 ability, and biodegradability make them ideal for preserving perishable goods[212].  
814 These films act as barriers against moisture, oxygen, and microbial contamination,  
815 thereby extending shelf life. However, pure chitosan films exhibit high hydrophilicity,  
816 limiting their moisture resistance[213,214]. This drawback is mitigated by blending with  
817 hydrophobic polymers or incorporating nanofillers such as silver nanoparticles, clay  
818 minerals, and lignin, which improve tensile strength, reduce water vapor permeability,  
819 and impart additional bioactivity[29]. As a progressive composite for these applications,  
820 chitosan-levan films (Figure 3) seem to be the most suitable choice, offering favourable  
821 mechanical performance, as well as antioxidant and antimicrobial properties.



823  
824 **Figure 3.** Visual appearance of chitosan-levan films[215].

825 Silver nanoparticles exhibit potent antibacterial effects, while layered silicates, such  
826 as montmorillonite, enhance thermal stability and gas barrier properties. Lignin, a  
827 natural polyphenolic compound, provides antioxidant functionality and UV  
828 protection[216]. Despite promising lab-scale results, commercial adoption faces hurdles  
829 related to cost, scalability, and regulatory approval. Large-scale production remains  
830 constrained by inconsistent raw material quality and energy-intensive processing

831 methods. Nevertheless, pilot-scale implementations in fruit and seafood packaging show  
832 tangible benefits, supporting gradual market integration[217].

### 834 3.2.2. Composting Additives

835 In solid waste valorization, chitosan serves as a composting additive that  
836 accelerates the decomposition of organic matter and reduces environmental  
837 emissions[11]. It stimulates microbial activity by acting as a nitrogen-rich substrate for  
838 beneficial bacteria and fungi[218]. Moreover, chitosan helps retain nitrogen in compost  
839 by forming complexes with ammonium ions, minimizing volatilization losses. Field  
840 trials indicate that chitosan-coated compost amendments reduce greenhouse gas  
841 emissions, particularly methane and nitrous oxide, by modulating microbial  
842 pathways[219].

843 Optimal dosages range between 0.5 - 2% w/w, beyond which inhibitory effects may  
844 occur. Long-term soil applications demonstrate improved aggregate stability and  
845 nutrient retention, although excessive use may alter the soil microbiota[95,220,221].  
846 Sustainable sourcing from seafood waste further enhances the circular economy appeal  
847 of chitosan in waste-to-resource strategies.

### 849 3.3. Soil Remediation

#### 850 3.3.1. Heavy Metal Immobilization

851 In contaminated soils, chitosan facilitates the immobilization of heavy metals  
852 through complexation, precipitation, and ion exchange. Reducing metal bioavailability  
853 prevents plant uptake and groundwater leaching. Biochar/chitosan hybrids are  
854 particularly effective, as they combine the high surface area and cation exchange  
855 capacity of biochar with the chelating functionality of chitosan[158]. These composites  
856 stabilize metals like Pb, Cd, and Cu in rhizosphere zones, promoting safer agricultural  
857 practices[95].

858 Field-scale applications demonstrate reduced metal mobility over extended periods,  
859 though long-term stability under varying pH and redox conditions requires further  
860 study. Integration into soil amendments offers a low-cost, eco-friendly strategy for  
861 rehabilitating mining-affected lands[222].

#### 863 3.3.2. Pesticide Remediation

864 For pesticide-contaminated soils, chitosan functions via sorption mechanisms,  
865 trapping organophosphates and chlorinated compounds through hydrogen bonding  
866 and hydrophobic interactions[223]. Chitosan-clay hybrids increase sorption capacity by  
867 expanding interlayer spacing and providing additional binding sites. Some formulations  
868 incorporate enzymes, such as laccase or peroxidase, which enable the enzymatic  
869 degradation of pesticides into less toxic metabolites[224]. Despite these advantages, real-  
870 world implementation is hindered by variability in soil composition, moisture content,  
871 and microbial interference, necessitating site-specific optimization.

### 873 3.4. Air Purification

874 Chitosan is playing a growing role in capturing pollutants from the air, targeting  
875 volatile organic compounds (VOCs), NO<sub>x</sub>, and SO<sub>2</sub>. Chitosan-MOF composites utilize  
876 the high porosity and tunable pores of metal-organic frameworks to adsorb gaseous  
877 pollutants selectively[40,138]. Additionally, activated carbon derived from chitosan  
878 pyrolysis offers a green route to porous adsorbents with high surface areas. These

materials can be integrated into HVAC filters or wearable masks for indoor and personal air purification[59]. While promising, scalability and cost-effectiveness remain key considerations for widespread deployment. Another example is chitosan-perlite composite, which was found to be much more effective for nitrogen adsorption over pure chitosan[225], as demonstrated in Figure 4.

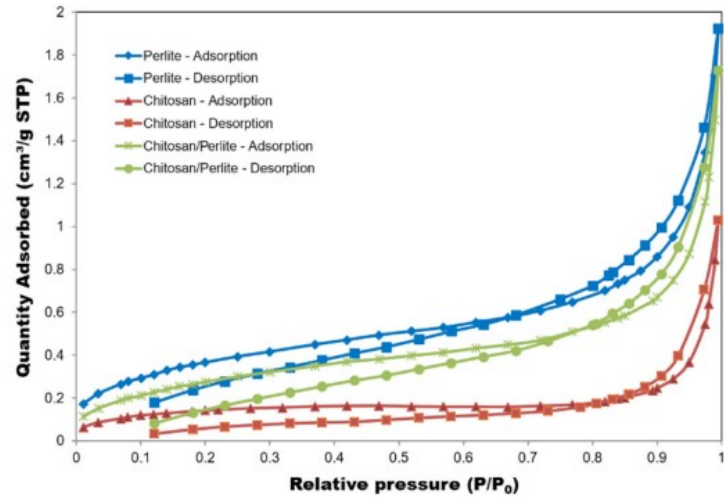


Figure 4. Nitrogen adsorption/desorption isotherms for perlite, chitosan, and chitosan/perlite composite[225].

### 3.5. Antimicrobial Activity

Chitosan is widely used in surface coatings for textiles, medical devices, and food packaging due to its intrinsic antimicrobial properties[226]. Mechanisms include the disruption of microbial cell membranes through electrostatic interactions with negatively charged phospholipids and the inhibition of DNA replication after cellular internalization[227,228]. Chemical modifications such as grafting N-halamines or quaternary ammonium salts enhance biocidal activity. However, concerns about nanoparticle leaching and the development of microbial resistance warrant careful formulation and lifecycle assessment[229,230]. A particularly interesting substrate is the composite based on chitosan, alginate, and silver nanoparticles, which is enriched with natural bee products (Figure 5). Such materials have interesting antioxidant, antifungal, and antibacterial properties[231].



903 **Figure 5.** Schematic interpretation of film preparation based on chitosan, alginate, silver  
904 nanoparticles, and natural bee products[231].

### 905 3.6. Oil Spill Cleanup

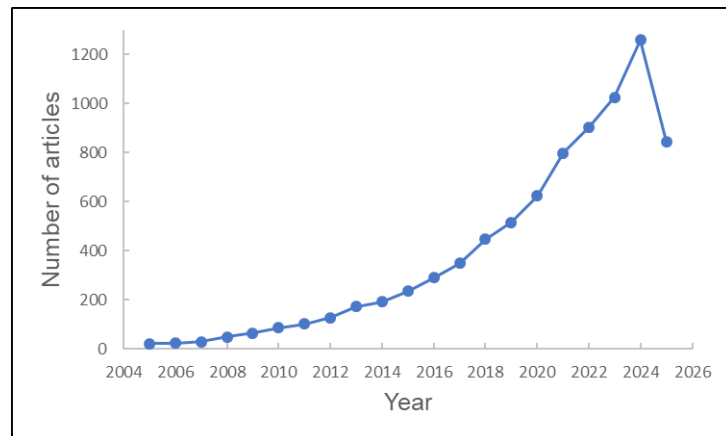
906  
907 For marine oil spills, modified chitosan aerogels and foams serve as highly  
908 absorbent materials. Hydrophobically modified chitosan exhibits selective uptake of  
909 non-polar hydrocarbons while repelling water[162,232]. These porous structures offer  
910 high absorption capacities (up to 30 times their weight), reusability through squeezing  
911 or solvent extraction, and good mechanical durability[146]. Challenges include  
912 maintaining performance under turbulent sea conditions and preventing secondary  
913 pollution upon degradation[233,234].

### 914 3.7. Other Environmental Applications

915  
916 Chitosan finds diverse uses in agricultural films, erosion control mats, biosensors,  
917 controlled-release fertilizers, desalination membranes, and gas separation  
918 membranes[20]. Its biodegradability supports temporary ground cover solutions, while  
919 its responsiveness enables innovative delivery systems. In membrane technology,  
920 chitosan-based thin-film composites show promise in forward osmosis and CO<sub>2</sub>/N<sub>2</sub>  
921 separation, contributing to clean energy and water security[3].

## 922 4. Discussion

923  
924 The environmental applications of chitosan and its derivatives have emerged as a  
925 cornerstone in the development of green, biodegradable, and cost-effective solutions for  
926 pollution mitigation[160]. Figure 6 represents the number of scientific papers published  
927 annually in the last 20 years.



929  
930 **Figure 6.** The number of scientific articles on environmental applications of chitosan retrieved  
931 from ISI WoS. The value for 2025 is by the end of July.

932 As evidenced by the body of research analyzed in this review, chitosan-based  
933 materials, particularly composites and chemically modified forms, demonstrate  
934 exceptional performance in the removal of heavy metals, organic dyes, pharmaceutical  
935 residues, and other contaminants from aqueous environments[9]. However, while  
936 laboratory-scale studies consistently report high adsorption capacities (often exceeding  
937 100–200 mg/g for metals like Pb(II), Cd(II), and Cr(VI)), the transition from benchtop

938 innovation to real-world implementation remains hindered by several interrelated  
939 scientific, engineering, and economic barriers[235]. A central theme across the reviewed  
940 literature is the strategic modification of native chitosan to overcome its inherent  
941 limitations: solubility at low pH, poor mechanical strength, and limited selectivity under  
942 competitive ion conditions[69]. The integration of chitosan with inorganic nanoparticles  
943 (e.g., TiO<sub>2</sub>, ZnO, Fe<sub>3</sub>O<sub>4</sub>), clays (e.g., montmorillonite), carbon nanomaterials (graphene  
944 oxide, CNTs), or biopolymers (cellulose, alginate) significantly enhances stability,  
945 surface area, and functional group density[110]. For instance, magnetic chitosan  
946 composites (e.g., Fe<sub>3</sub>O<sub>4</sub>/chitosan) not only improve separation efficiency but also  
947 facilitate regeneration and reuse, which is critical for circular economy models[158].  
948 Previous studies highlight that hybridization with metal oxides introduces additional  
949 adsorption mechanisms, including electrostatic attraction, ion exchange, and surface  
950 complexation, thereby broadening the range of treatable pollutants. Moreover, chemical  
951 derivatization such as quaternization, crosslinking with glutaraldehyde, or grafting with  
952 functional groups (-NH<sub>2</sub>, -COOH, -SH) alters the protonation behavior and binding  
953 affinity of chitosan[73,77,236]. Quaternized chitosan derivatives, for example, remain  
954 positively charged across a wide pH range, enabling effective anion removal (e.g.,  
955 fluoride, arsenate) even in acidic conditions where unmodified chitosan dissolves[46,73].  
956 This is particularly relevant for arsenic remediation, where recent studies report  
957 enhanced As(V) uptake via ligand exchange with surface hydroxyl groups on chitosan-  
958 TiO<sub>2</sub> composites[51,155]. Kinetic and isotherm modeling dominate the analytical  
959 framework in most studies, with pseudo-second-order kinetics and Langmuir isotherms  
960 frequently cited as best fits[237]. While these models suggest chemisorption and  
961 monolayer adsorption, their overuse without complementary spectroscopic validation  
962 (e.g., XPS, FTIR, EXAFS) risks oversimplification[49,238]. Several papers rely solely on  
963 model fitting without probing actual binding mechanisms, leading to potentially  
964 misleading conclusions about adsorption pathways. For example, attributing Cr(VI)  
965 removal entirely to electrostatic attraction overlooks the well-documented reduction of  
966 Cr(VI) to Cr(III) by amine groups, followed by coordination of the reduced species, a  
967 dual mechanism confirmed in multiple spectroscopic studies[51,203]. Furthermore, the  
968 majority of isotherm experiments are conducted using single-solute systems, which  
969 poorly represent real wastewater matrices containing multiple competing ions, natural  
970 organic matter, and suspended solids[239]. Only a minority of the reviewed studies  
971 evaluate performance in multi-pollutant or real industrial effluents, revealing significant  
972 capacity drops due to fouling or ion competition[240,241]. This discrepancy between  
973 idealized and realistic conditions underscores the need for more environmentally  
974 relevant testing protocols. Despite claims of "reusability," many studies report only 3–5  
975 regeneration cycles, often with >20% capacity loss by the final cycle[90,118,195]. The use  
976 of strong acids (HCl) or bases (NaOH) for desorption raises concerns about long-term  
977 structural integrity and secondary pollution. In some cases, the regeneration process  
978 itself generates hazardous waste streams, undermining the environmental benefits of the  
979 original adsorbent[110,242]. Additionally, few studies address the end-of-life fate of  
980 spent chitosan composites. While chitosan is inherently biodegradable, the presence of  
981 non-biodegradable components (e.g., synthetic polymers, metal nanoparticles) may  
982 inhibit decomposition or lead to nanoparticle leaching during disposal[243]. A truly  
983 sustainable chitosan composite must be evaluated not only on adsorption efficiency but  
984 also on lifecycle impacts, including raw material sourcing (e.g., shrimp vs. fungal  
985 chitosan), energy consumption during synthesis, reusability, and final disposal. The shift  
986 toward waste-derived chitosan (e.g., from crustacean shells, squid pens, or fungal  
987 biomass) aligns with circular economy principles and reduces reliance on finite

resources. However, the scalability of such green synthesis routes remains a challenge, particularly in regions lacking seafood processing infrastructure. The overwhelming majority of research remains confined to batch adsorption experiments at a small scale (<100 mL)[145,244]. Continuous flow systems, column studies, and pilot-scale trials are underrepresented, despite their relevance to industrial water treatment. Only a few studies explore chitosan-based membranes or fixed-bed reactors, which are more suitable for large-scale deployment[160,245]. Even then, issues such as membrane fouling, pressure drop, and mechanical degradation under prolonged operation are rarely addressed in depth. Moreover, the physical form of the material, bead, film, membrane, or powder, affects handling, flow dynamics, and applicability[246,247]. For instance, chitosan films show promise in air filtration and volatile organic compound (VOC) capture, while magnetic nanoparticles enable rapid separation in slurry systems[135]. Yet, the lack of standardized performance metrics across different formats makes comparative evaluation difficult. An analysis of funding sources reveals a strong concentration of research in Asia (China, India, Malaysia, Iran) and Europe, with significant support from national science foundations and regional development programs (e.g., NSFC, YUTP, EU-NextGenerationEU). This reflects strategic investment in green materials science but also highlights disparities in global research capacity. Nearly 10% of the reviewed studies report no external funding, suggesting that some innovations arise from under-resourced institutions, potentially limiting reproducibility or scale-up potential. Another critical gap is the inconsistent reporting of data availability and experimental details. While open-access publishing has increased transparency, many authors restrict raw data access to "upon request," hindering meta-analyses and independent validation. Similarly, detailed synthesis protocols, especially for nanocomposites, are often omitted, making replication challenging. The future of chitosan-based environmental materials lies not merely in incremental performance improvements but in the development of intelligent, multifunctional systems. Emerging trends include stimuli-responsive composites (pH-, temperature-, or light-sensitive), self-cleaning photocatalytic hybrids (e.g., chitosan-TiO<sub>2</sub>), and integration with digital monitoring (IoT-enabled sensors)[248,249]. Additionally, combining adsorption with catalytic degradation (e.g., Fenton-like systems using Fe-chitosan composites) offers a pathway beyond mere pollutant sequestration toward complete mineralization[250]. Finally, policy alignment and regulatory acceptance are essential for widespread adoption. Standardization of testing protocols, certification of biodegradability, and lifecycle assessment frameworks will be crucial in transitioning chitosan composites from academic curiosities to commercially viable environmental technologies.

## 5. Limitations and Future Perspectives

Despite the significant advancements in the development and application of chitosan-based materials for environmental remediation, several critical limitations hinder their widespread industrial adoption and long-term sustainability. These challenges span the entire lifecycle of chitosan products from raw material sourcing and synthesis to performance under real-world conditions, regeneration, and end-of-life management. Addressing these constraints is essential for transitioning chitosan technology from laboratory-scale innovation to large-scale, practical implementation.

One of the primary limitations lies in the scalability of synthesis methods. Many high-performance chitosan composites are fabricated using energy-intensive, batch-dependent techniques such as freeze-drying, cryostructuring, or layer-by-layer

assembly[22,251,252]. These processes are challenging to scale up due to high operational costs, low throughput, and the requirement for specialized equipment such as lyophilizers or supercritical CO<sub>2</sub> dryers. Furthermore, the use of toxic crosslinking agents, such as glutaraldehyde, enhances mechanical stability but raises environmental and health concerns, particularly regarding the disposal of residual chemicals and spent adsorbents[52,253]. Although greener alternatives, such as genipin or enzymatic crosslinking, are being explored, their high costs and limited availability restrict their broad application.

Another persistent challenge is the instability of native chitosan under neutral and alkaline pH conditions, where it loses its solubility and cationic character due to the deprotonation of amino groups[69]. This significantly limits its effectiveness in natural water systems and industrial effluents that operate outside the acidic range. While chemical modifications, such as quaternization or carboxymethylation, can extend the functional pH range, these derivatization processes often require multiple reaction steps, thereby increasing complexity and production costs[254]. Moreover, excessive crosslinking or functionalization may reduce the accessibility of active binding sites, thus compromising adsorption capacity and kinetics.

The mechanical durability of chitosan-based materials under dynamic environmental conditions is also a concern. In continuous flow systems, such as fixed-bed columns or membrane filtration units, repeated swelling and deswelling cycles can lead to structural fatigue, particle disintegration, and loss of porosity[22,255,256]. This issue is particularly evident in hydrogels and aerogels, which demonstrate excellent adsorption performance in batch experiments but struggle with mechanical resilience in real-world applications. Additionally, fouling caused by organic matter, colloidal particles, or microbial biofilms can block active sites and reduce efficiency over time, necessitating frequent cleaning or replacement[257]. However, some promising chitosan composite systems are already recognized. For example, chitosan/TiO<sub>2</sub> membrane exhibits a much lower degree of swelling than pure chitosan in water and buffers[258], as seen in Table 1.

Table 1. Degree of swelling of chitosan membrane and chitosan/TiO<sub>2</sub> composite membrane in water and different aqueous buffer solutions at pH 6, 7, and 8[258]. The same letters denote statistical similarity of samples with variation of the solution. Data expressed as mean and standard deviation, with 95% confidence according to Tukey's test.

Sample	Degree of Swelling (%)			
	H <sub>2</sub> O	pH 6	pH 7	pH 8
Chitosan	481.59 ± 74.26 <sup>a</sup>	138.20 ± 6.51 <sup>b</sup>	171.17 ± 40.92 <sup>b</sup>	180.50 ± 38.53 <sup>b</sup>
Composite	376.25 ± 8.31 <sup>a</sup>	115.42 ± 10.11 <sup>c</sup>	158.81 ± 7.77 <sup>b</sup>	131.20 ± 5.95 <sup>b</sup>

Regeneration and reusability, while theoretically feasible, present practical and ecological trade-offs. Although many chitosan composites can be regenerated using acidic or basic eluents, the repeated use of strong chemicals may degrade the polymer matrix, leach functional components, or generate secondary waste streams that require treatment[259,260]. For instance, regenerating metal-loaded composites with acids produces concentrated metal-laden effluents that must be managed responsibly. Furthermore, the economic feasibility of regeneration depends on balancing recovery

1081 efficiency with operational costs, including water and energy consumption during  
1082 washing and drying cycles[90,261].

1083 From a lifecycle sustainability perspective, conventional chitosan production via  
1084 chemical demineralization and deacetylation generates substantial amounts of acidic  
1085 and alkaline wastewater, contributing to environmental pollution[160]. Although  
1086 enzymatic and microbial extraction methods offer a more sustainable alternative by  
1087 reducing chemical usage and energy input, they are currently less efficient and more  
1088 time-consuming than traditional processes[262,263]. Scaling up these green synthesis  
1089 routes requires further optimization of microbial strains, enzyme immobilization  
1090 techniques, and bioreactor designs to achieve industrial throughput and cost-  
1091 effectiveness.

1092 Looking ahead, the future of chitosan-based environmental technologies lies in  
1093 integrating advanced material design, digital innovation, and principles of the circular  
1094 economy. The development of multifunctional composites such as magnetic,  
1095 photocatalytic, or stimuli-responsive systems will enable not only pollutant capture but  
1096 also degradation, sensing, and controlled release. For example, integrating chitosan with  
1097 metal-organic frameworks (MOFs) or graphene oxide can enhance selectivity and  
1098 catalytic activity, while the use of magnetic nanoparticles facilitates easy separation and  
1099 reuse[9,264].

1100 Artificial intelligence (AI) and machine learning (ML) are poised to revolutionize  
1101 the field by enabling predictive modeling of adsorption behavior, optimizing composite  
1102 formulations, and supporting real-time monitoring in treatment plants[186]. Digital  
1103 twins and blockchain-based traceability systems could ensure transparency across the  
1104 supply chain, further driving the sustainable adoption of chitosan-based materials for  
1105 environmental remediation[265].  
1106

## 1107 **6. Conclusions**

1108 The escalating environmental challenges of the 21st century demand innovative,  
1109 sustainable, and practical solutions. This review comprehensively examines the potential  
1110 of chitosan, a naturally abundant and biodegradable biopolymer, along with its  
1111 numerous derivatives and composites, as a cornerstone material for addressing these  
1112 pressing issues. Chitosan's unique chemical structure, featuring reactive amino and  
1113 hydroxyl groups, provides the foundation for its remarkable versatility in interacting  
1114 with a broad spectrum of pollutants, including heavy metals, dyes, organic  
1115 contaminants, and microorganisms. However, the inherent limitations of native  
1116 chitosan, such as poor solubility outside acidic conditions, limited mechanical strength,  
1117 and susceptibility to swelling, necessitate strategic modifications to realize its full  
1118 potential in practical applications.

1119 Modern modification strategies, encompassing chemical derivatization  
1120 (crosslinking, grafting, quaternization, functionalization), physical structuring (forming  
1121 beads, films, aerogels, membranes), and, most significantly, composite formation, have  
1122 proven highly effective. These approaches allow for the precise tailoring of chitosan-  
1123 based materials to enhance stability, selectivity, and efficiency. The creation of  
1124 composites by integrating chitosan with inorganic fillers (silica, clays, metal oxides),  
1125 carbonaceous materials (graphene oxide, biochar, CNTs), magnetic nanoparticles  
1126 ( $\text{Fe}_3\text{O}_4$ ), other biopolymers (alginate, lignin), and even metal-organic frameworks  
1127 (MOFs) has led to the development of hybrid materials with synergistic properties that  
1128 often surpass the sum of their components. These advanced materials can be engineered

1129 for specific functions, such as targeted adsorption, catalytic degradation, efficient  
1130 membrane filtration, or effective flocculation.

1131 The mechanisms underlying pollutant removal by chitosan-based systems are  
1132 multifaceted and often work in concert. Adsorption, driven by electrostatic interactions,  
1133 complexation, ion exchange, and hydrogen bonding, remains the dominant mechanism.  
1134 However, the integration of catalytic functionalities enables the active degradation of  
1135 pollutants, while the material's film-forming ability and surface charge properties make  
1136 it suitable for antifouling membrane applications. Furthermore, its cationic nature allows  
1137 it to function effectively as a green flocculant. This mechanistic diversity highlights the  
1138 adaptability of chitosan platforms in addressing complex environmental matrices.

1139 Despite the overwhelming success demonstrated in laboratory and pilot-scale  
1140 studies, the transition from bench-scale innovation to widespread real-world  
1141 implementation faces several practical challenges. Issues related to the scalability of  
1142 often energy-intensive fabrication methods (like lyophilization), ensuring long-term  
1143 stability and performance under variable and harsh environmental conditions,  
1144 developing efficient and environmentally benign regeneration protocols, and conducting  
1145 thorough lifecycle sustainability assessments must be addressed. Economic viability is  
1146 closely linked to the cost-effective sourcing of raw materials, favouring the use of  
1147 unrefined seafood waste over purified chitosan. Embracing green synthesis routes, such  
1148 as enzymatic processing and microbial fermentation, is crucial for aligning production  
1149 with sustainability goals.

1150 Looking ahead, the field is being shaped by exciting emerging trends. The  
1151 integration of chitosan technology within circular economy models is paramount,  
1152 transforming seafood processing waste into high-value materials and designing end-of-  
1153 life strategies for spent composites that involve metal recovery or composting. The  
1154 valorisation of seafood waste using green extraction techniques further enhances the  
1155 sustainability profile. Perhaps most significantly, the infusion of digital innovation,  
1156 particularly artificial intelligence (AI) and machine learning (ML), is revolutionizing the  
1157 design, optimization, and operational monitoring of chitosan-based systems. These tools  
1158 enable predictive modelling, accelerate material discovery, optimize process parameters,  
1159 and facilitate real-time control, paving the way for intelligent and efficient  
1160 environmental solutions.

1161 Thus, chitosan and its derivatives/composites represent an up-and-coming class of  
1162 materials for sustainable environmental protection. The extensive research and  
1163 development efforts highlighted in this review demonstrate the significant progress  
1164 made in overcoming the limitations of native chitosan and engineering high-  
1165 performance materials tailored for specific environmental applications. From water  
1166 treatment and soil remediation to air purification and waste management, the versatility  
1167 of chitosan-based systems is evident. While challenges remain in scaling up production,  
1168 ensuring robustness, and achieving full economic competitiveness, the convergence of  
1169 advanced material science, green chemistry, circular economy principles, and digital  
1170 optimization presents a clear pathway forward. Continued research focusing on  
1171 practical implementation, lifecycle assessment, and the integration of innovative  
1172 technologies will be essential. By addressing these aspects, chitosan-based technologies  
1173 can move beyond laboratory curiosities to become integral components of scalable,  
1174 intelligent, and ecologically responsible solutions, playing a vital role in safeguarding  
1175 the environment for future generations.

1176  
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## 1188 Abbreviations

1189 The following abbreviations are used in this manuscript:

DD	Degree of deacetylation
MW	Molecular weight
CMCS	Carboxymethyl chitosan
ECH	Epichlorhydrin
BADGE	Bisphenol A diglycidyl ether
PEI	Polyethyleneimide
ChNS	Chitosan-nanosilica composites
ChSG	Chitosan-silica gel composites
GO	Graphene oxide
CNTs	Carbon nanotubes
NS	Nanosilica
SG	Silica gel
GA	Glutaraldehyde
CBZ	Carbamazepine
CIP	Ciprofloxacin
SEM	Scanning electron microscopy
PDA	Polydopamine
TPP	Triphosphosphate
Car/CSAs	Chitosan/carbon ratio in aerogel composites
MB	Methylene blue
CSGO	Chitosan and graphene oxide membrane
MMMs	Mixed matrix membranes
MOFs	Metal organic frameworks
LCA	Lifecycle analysis
AI	Artificial intelligence
ML	Machine learning
ANFIS	Adaptive neuro-fuzzy inference systems
ANNs	Artificial neural networks
RSM	Response surface methodology
EDTA	Ethylenediaminetetraacetic acid
MIPs	Molecularly imprinted polymers
VOCs	Volatile organic compounds

1190

1191

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107 [Sem zadejte text.]

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