

Chitosan and Graphene Oxide-based Nanocomposites for Water Purification and Medical Applications: A Review

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Chitosan (CS) is a natural polysaccharide isolated from insects, molluscs, and fungi. The specific properties of chitosan can be enhanced using physicochemical processes. The composites prepared using CS and graphene oxide (GO) contain active functional groups such as epoxide, carboxyl, and hydroxyl, which possess excellent biocompatibility, high adsorption capacity, and biodegradability. Their low cost and ease of scale-up make them employable for multiple applications in water-treatment plants, electronics, solar cells, and pharmaceuticals. This review provides an overview of sources, types, and properties of chitin, chitosan, and graphene oxide. The use of these composites for the preparation of antimicrobial drugs has been discussed here. The article also explores the applicability of such composites for removal of heavy metals (lead, copper, chromium, cobalt, mercury, *etc.*), dyes (methylene blue and other reactive dyes), and organic and inorganic contaminants (ofloxacin, naphthanol, phenol, and oil, *etc.*). The article highlights various knowledge gaps in the field and the scope of future work.

Keywords: Chitosan; Graphene oxide; Nanocomposite; Water purification; Medical applications

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INTRODUCTION

Polymer-based nanocomposites have applications in various fields such as wastewater treatment, electronics, energy, pharmaceuticals, and medicine. Low cost, high recyclability, biocompatibility, ease of degradability, and scalability make bio-based, commonly available, natural polymers such as chitosan (CS) materials of choice for multiple fields ranging from wastewater treatment to medicinal applications (Dutta *et al.* 2004; Cheung *et al.* 2015; Palza 2015). Their high affinity for metals, organic and inorganic particles coupled with the higher surface area makes them promising materials for the removal of contaminants from wastewater (Chen *et al.* 2013; Schwarz *et al.* 2016; Singh *et al.* 2019; Suri 2020; Suri *et al.* 2021; Upadhyay *et al.* 2021). In combination with graphene oxide (GO) and inorganic nanoparticles such as ZnO, Fe₃O₄, GO, and TiO₂, it is possible to introduce additional functional groups that can increase their chelation capacity, improve the surface area and active sites, and assist in further modification of their properties (Kyzas and Bikiaris 2015; Morsy 2015; Salam 2017; ZabihiSahebi *et al.* 2019).

The presence of heavy metals in soil and water is a significant threat to the ecosystem. Heavy metals, dyes, and inorganic pollutants are widespread through various sources and taken up by plants in the environment. As a result, they enter into the food chain and are ingested by humans and animals. Consistent efforts are being made to

effectively remove contaminated water and soil samples (Chauhan *et al.* 2020). Various heavy metals like copper, chromium, and mercury are removed from contaminated samples using composting, landfarming, coagulation, flocculation, membrane separation, and adsorption processes (Kaur *et al.* 2020). Attempts have been made in the literature to develop green adsorbents for wastewater treatment.

Cs and GO are promising materials with unique properties such as biocompatibility, safety, and antimicrobial capacity. They are also proven to be effective as drug delivery agents. Attempts on forming nanocomposites made from natural materials such as CS and GO in various ways have yielded properties even better than individual CS or GO used alone (Ates *et al.* 2020). Thus, the present literature study covers the essential aspects of CS and GO-based nanocomposites and their applicability in the specified areas. It explores the sources, types, and properties of CS and GO. The applications of modified CS-GO-based composites for removing heavy metals, dyes, and pollutants from wastewater are reviewed. An overview of its antimicrobial properties and potential as a medium for drug delivery is also presented. This information can serve as a reference document for technologists and scientists to research and develop new systems based on Cs-GO nanocomposites (Ahmad *et al.* 2017).

ORIGIN AND SOURCES OF CHITIN AND CHITOSAN

The word chitin is derived from the Greek word “Chiton”, meaning covering or envelope. It was discovered as a new polysaccharide and reported for the first time in 1811 by French professor Henri Braconnot as *fungine* while researching edible mushrooms. Later, in 1823, Antoine Odier named it chitine because of its unique presence in insects' cuticles (Crini 2019). Later, its structure was determined by Albert Hofman (Hofmann 1979). In 1859, Charles Rouget described deacetylation of chitin to obtain a modified form of chitin, which was later termed chitosan by Hoppe-Seyler in 1894 (Crini 2019). The polysaccharides chitin and chitosan are made up of amino sugars D-glucosamine and N-acetyl-D-glucosamine. The ratio of amino sugars D-glucosamine and N-acetyl-D-glucosamine indicates whether the polysaccharide is considered chitin or chitosan (Brigham 2017). Chitosan is a linear cationic copolymer consisting of alternating units of N-acetyl-glucosamine (20%) and D-glucosamine (~80%) obtained by deacetylation of chitin, whereas chitin is a linear polymer composed of repeated units of β -1,4-linked N-acetylglucosamine (50%) (Philibert *et al.* 2017; Hahn *et al.* 2020b). Both polymers have exclusive properties and find applications in various fields (Cheung *et al.* 2015; Younes and Rinaudo 2015). Chitin consists of huge crystalline nitrogen-containing polysaccharides with an estimated production of about 10^{11} to 10^{14} tons per annum (Bastiaens *et al.* 2019). Due to versatility and accessibility, chitin is the second most abundant polysaccharide after cellulose and hemicellulose.

The primary sources of chitin and chitosan are shown in Fig. 1. The most common source of chitin is shellfish and other aquatic invertebrates. These species are found in considerable quantities in the fish processing industry (Yadav *et al.* 2019). Chitin is located in the exoskeleton of sea animals such as annelids, molluscs, coelenterate, crustaceans (crabs, shrimps, and lobsters), and insects such as honey bee, silkworms, scorpions, spiders, ants, beetles, and cockroaches as supporting tissues of organisms (Kaur and Dhillon 2014). The content of chitin is highest in shrimp cuticle and squid pen waste, ranging from 15 to 40% as compared to other crustacean waste since shrimps have a thinner shell wall than

lobsters and crabs (Tharanathan and Kittur 2003; Wang *et al.* 2020; Tan *et al.* 2020; Khayrova *et al.* 2021; Kou *et al.* 2021). While insects are a viable substitute source for chitin, being an abundant species globally, a lower quantity of chitin is extracted from fungi such as *F. fomentarius* and *L. vellereus* (Bastiaens *et al.* 2019, Sagheer *et al.* 2009). A significant part of the head and shell waste is obtained only from shrimp (about 60%). At the same time, the cuticle can contain up to 80% chitin (Pal *et al.* 2014). The amount of chitin depends on the type of species, body part, season, and growth rate.

In contrast, chitosan is a partially deacetylated copied form of chitin (Letourneau *et al.* 1976). Chitosan content of 1 to 10% on dry biomass base has been found, with a reported degree of deacetylation of 83 to 94%. Chitosan is produced synthetically by converting chitin in the presence of a deacetylase enzyme (Dhillon *et al.* 2013).

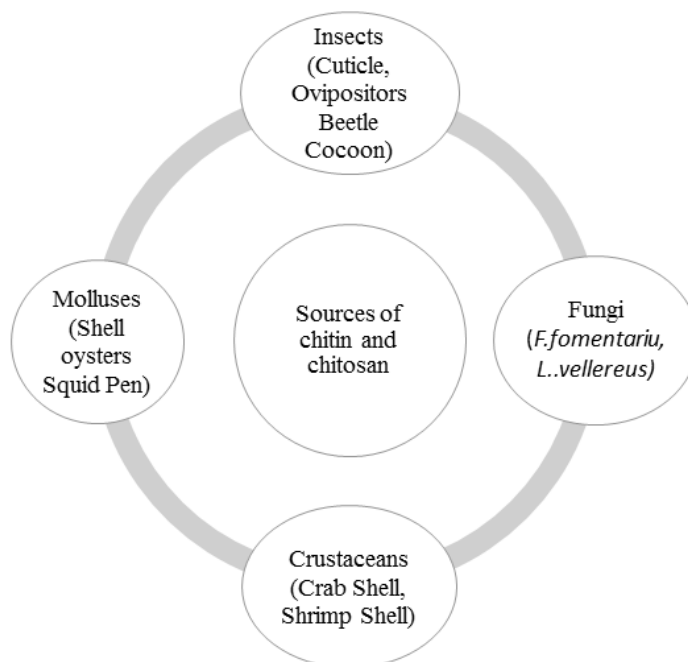


Fig. 1. Sources of chitin and chitosan

In addition to fungi, bacteria can be used for the production of chitosan using enzymatic deacetylation. Kaur and Dhillon (2014) isolated chitosan from soil bacteria (*Bacillus sp.* and *Serratia sp.*) through chitin's deacetylation. However, the efficiency of this process is affected by the insolubility and degradation of chitin (Aranaz *et al.* 2009). In addition, Kaur *et al.* (2012) suggested that the fast-growing bacteria or the isolated enzymes can be used for natural chitosan production. Over 10 billion tons of chitin are produced annually from the seafood processing industry (Casadidio *et al.* 2019). Chitin and chitosan are now extracted commercially in India, Japan, Poland, Australia, and the US. In the future, chitin's production rate will increase significantly due to increased waste generation of insects, exuviae, and exoskeletons. More research is required for surplus sources and utility of chitin/chitosan in the near term (Dutta *et al.* 2004).

Three types of crystal-like forms of chitin α , β , and γ (mixer of α and β chitin) are found in nature (Aam *et al.* 2010). Different types of chitin allomorphs are shown in Fig. 2. Of these, α chitin is typically isolated from the shells and the exoskeleton of crustaceans, insect cuticles, shrimps, crab, and lobsters (Kaur and Dhillon 2014; Yadav *et al.* 2019). In

α chitin, polysaccharide chains are organized in an antiparallel direction, allowing maximum bonding. Hence, α chitin is highly stable, having a high degree of crystallinity (80%). β chitin is usually found in squid pens, aphrodite chaetae, seaweeds, protozoa (Sagheer *et al.* 2009; Hajji *et al.* 2014). Here, polysaccharide chains are organized in parallel, with a crystallinity index of 70%. The target distance between the adjacent polymer chains makes it more reactive and soluble in solvents (Yadav *et al.* 2019). The γ -chitin exists mainly in fungi and yeast. It is recognized by two parallel and one antiparallel sheet arrangement.

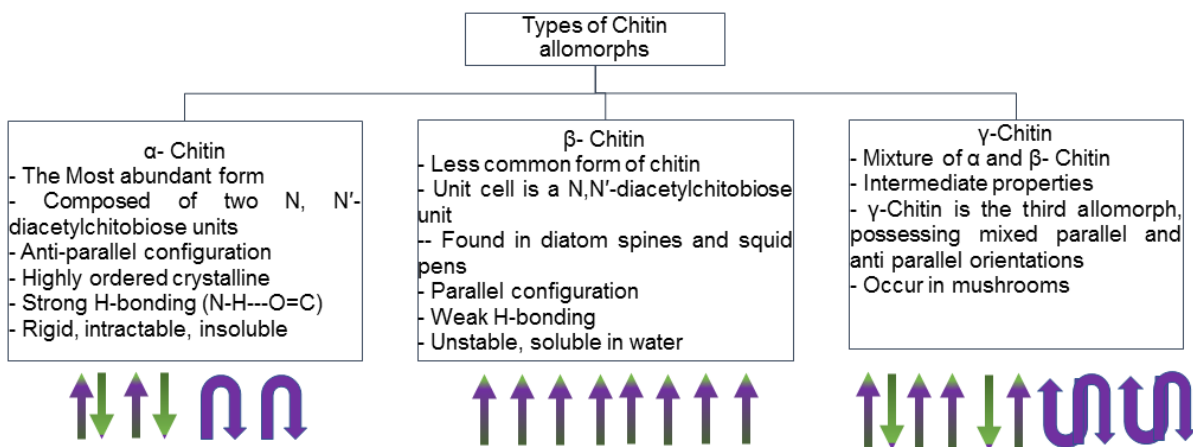


Fig. 2. Different types of Chitin Allomorphs

PROPERTIES OF CHITIN AND CHITOSAN

CS is more versatile than chitin due to amino groups at the C-2 positions (Dutta *et al.* 2004). Modification of chitin and CS help in the increasing of its solubility and versatility (Ishihara *et al.* 2012). Both chitin and CS can form films, chelate with metals, and moderate water permeability (Dutta *et al.* 2004). Due to low toxicity and digestibility, they can be used as food additives. Also, they are shown to be helpful to lower cholesterol in human blood (Knoor 1983). Both prevent the invasion of pathogens. However, chitosan is known to exhibit better properties and thus applicability than chitin. Chitin is a white, rigid, inelastic, nitrogenous polysaccharide (5 to 7% nitrogen). Being hydrophobic, it is insoluble in water and organic solvents. Due to its one reactive amine and two hydroxyl groups, chitosan has higher solubility, hydrophilic character, and adsorption capacity than chitin. CS is also soluble in aqueous solutions of acids. Solubility in aqueous solution gets further enhanced by N-alkylidinations and N-acylation (Muzzarelli 1997). It has an extensive semicrystalline structure. It has low immunogenicity and binds to mammalian and microbial cell aggregates (Dutta *et al.* 2004; Cheung *et al.* 2015). CS also displays mucoadhesive film-forming and chelating activity (Patra *et al.* 2018). The pH sensitivity and high biological compatibility of chitosan motivate researchers to explore its application in drug delivery systems, especially for cancer treatment (Dutta *et al.* 2004; Philibert *et al.* 2017; Yassue-Cordeiro *et al.* 2018; Ramachandran *et al.* 2019). Due to its long shelf-life and ability to form semipermeable, rigid, permanent and impermeable films, it can be used to wrap food products (Muzzarelli *et al.* 1986). CS is effective to remove endotoxins found in proteins as contaminants. Endotoxins are pH and heat stable and are not easily destroyable using normal sterilising conditions. CS can interact with endotoxins and help

in downstream protein purification processes (Wang *et al.* 2013a).

CS is biocompatible, non-toxic, biodegradable, and has broad-spectrum antimicrobial activities (Jiang *et al.* 2014). The antifungal activity of CS is higher than chitin (Limam *et al.* 2011). CS can promote bone formation because it can be shaped into structures and geometries suitable for cell in-growth and post-growth conditions (Figueiredo *et al.* 2015). With unique biological activities, including analgesic, antitumor, haemostatic, and antioxidant properties, CS finds application in the biomedical field (Brigham 2017; Casadidio *et al.* 2019).

METHODS OF EXTRACTION OF CS

The chitosan family consists of different D-N-deacetylated forms of chitin, and its properties are highly dependent on the degree of deacetylation and its molecular weight (Younes and Rinaudo 2015; Philibert *et al.* 2017). CS is extracted from chitin through demineralization, deproteinization, deacetylation, and decolourization using chemical or biological processes such as enzymatic treatment or fermentation. It can be synthesized through nitrogen and oxygen substitution, copolymerization, and numerous other methods (Wang *et al.* 2016, 2020).

The difference in alkali solution concentration, reaction time, temperature, chitin/alkali solution ratio, CS with different degrees of deacetylation can be obtained. The degree of acetylation represents the percentage of N-acetyl-d-glucosamine units to the total number of units. Hence for CS, the percentage degree of acetylation is below 50. Acetyl groups are removed arbitrarily, and a de-polymerization reaction occurs, which is indicated by changes in the molecular weight of CS. Various methods for the extraction of chitin and CS have been reported, although no standard practice has been implemented. However, demineralization and deproteinization are mostly chosen either using chemicals or biological agents. Few studies for extraction of chitin and chitosan using chemical and biological extraction are discussed below.

Different researchers have investigated various chemicals and solutions to extract CS and chitin. Deproteinization is a fundamental step used to extract protein, while demineralization is an acidic step used to remove the inorganic calcium carbonate (Khanafari *et al.* 2008; Tan *et al.* 2020). In the demineralization process, hydrochloric acid, nitric acid, sulfuric acid, acetic acid, and formic acid are used. In contrast, the alkaline method for deproteinization is commonly used for protein extraction (Brine and Austin 1981). Figure 3 explains the general procedure used for the extraction of chitin from crab and shrimp waste.

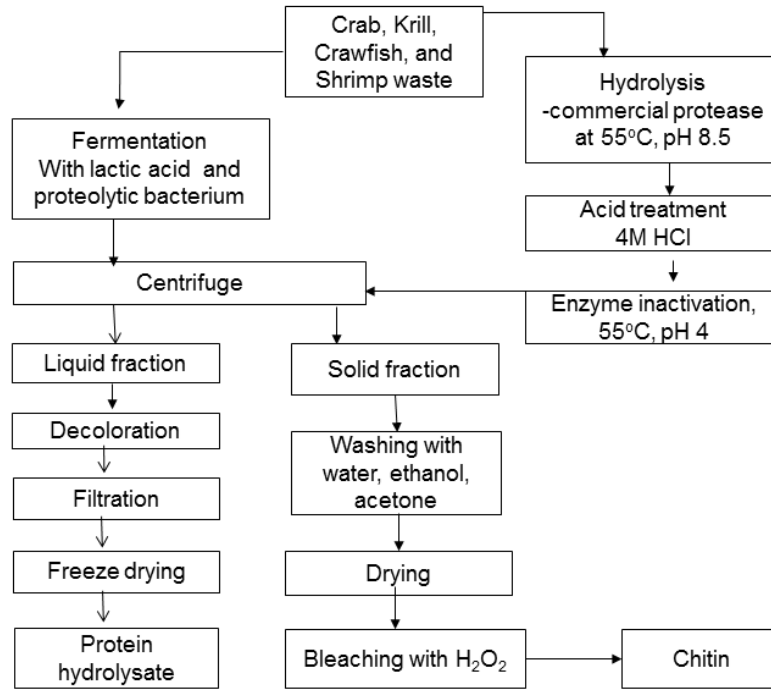


Fig. 3. Flowchart of procedure used for the extraction of chitin

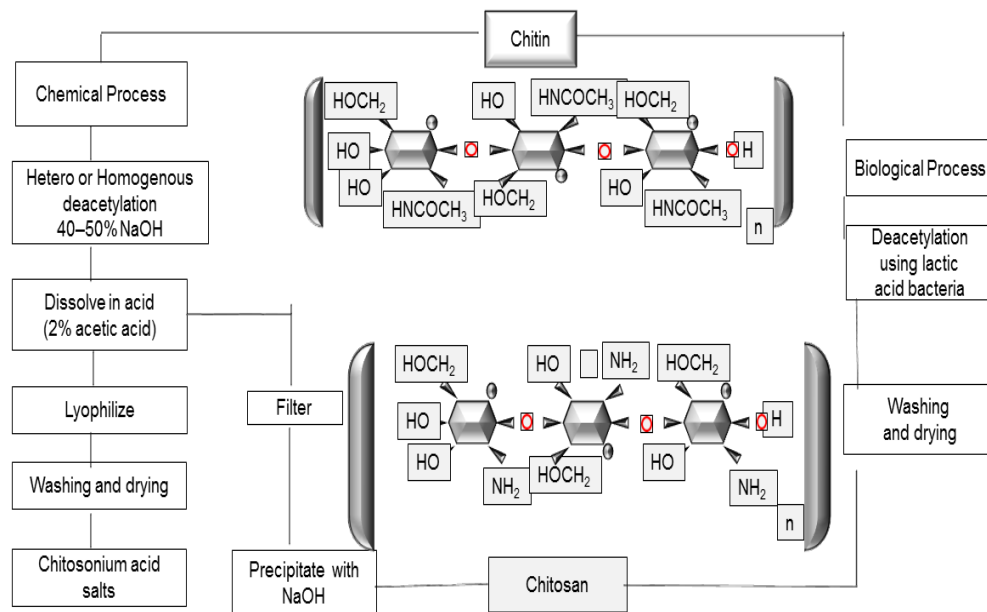


Fig. 4. General flowchart of procedure used for the biological extraction of chitin

Chitin can be processed chemically as well as biologically to produce chitin, as shown in Fig. 4. While deacetylation is done with sodium hydroxide in chemical processes, biological processes utilise lactic acid bacteria for the same. Researchers have tried to modified these processes using different agents under different reaction conditions. The extraction of chitin and CS is affected by the body parts and the type of organism used. Biological methods of CS extraction use fermentation processes for shrimp or other

organism waste. The solid and liquid fraction can be separated to allow for chitin extraction. Demineralization and protein removal steps can be done, as was in the chemical procedure. Nessa *et al.* (2010) extracted chitosan from sun-dried prawn shells by producing coarse particles in a centrifugal grinding machine. Some coarse particles were demineralized with 10% HCl acid at 27 °C for 22 h. Deproteinising was done using NaOH solution for 24 h at 70 °C, then decolourization and drying under vacuum. A crispy powder of chitin was produced. Deacetylation of chitin was attained using NaOH for 45-72 h. The resulting chitosan was rinsed with distilled water and oven-dried. The yield of chitin was 20%, and CS was 19.6%. Tarafdar and Biswas (2013) reported extraction using two different prawn shells and shrimp waste. In the first method, 10 g of prawn shell waste was washed and demineralized by adding 1.5 N HCl at 25 °C for 1 h followed by deproteinisation with 0.5 to 3% NaOH at 100 °C for 30 min. Deacetylation of chitin produced CS, which was prepared by treating with 42% aqueous NaOH at 95 °C for 1.5 h and washed then dried. In the second method, 5 g of shrimp waste was soaked then deproteinized in an aqueous NaOH solution at 25 °C for 21 h. The deproteinized shell was demineralized by 4% HCl at 25 °C for 12 h. The chitin was dried at ambient temperature. CS was obtained by treating chitin with 50% aqueous NaOH at 40 °C for three days. CS was dried at ambient temperature. The overall procedures used in chemical, biological methods investigated in literature using insects, shellfish and fungi are shown in Fig. 5.

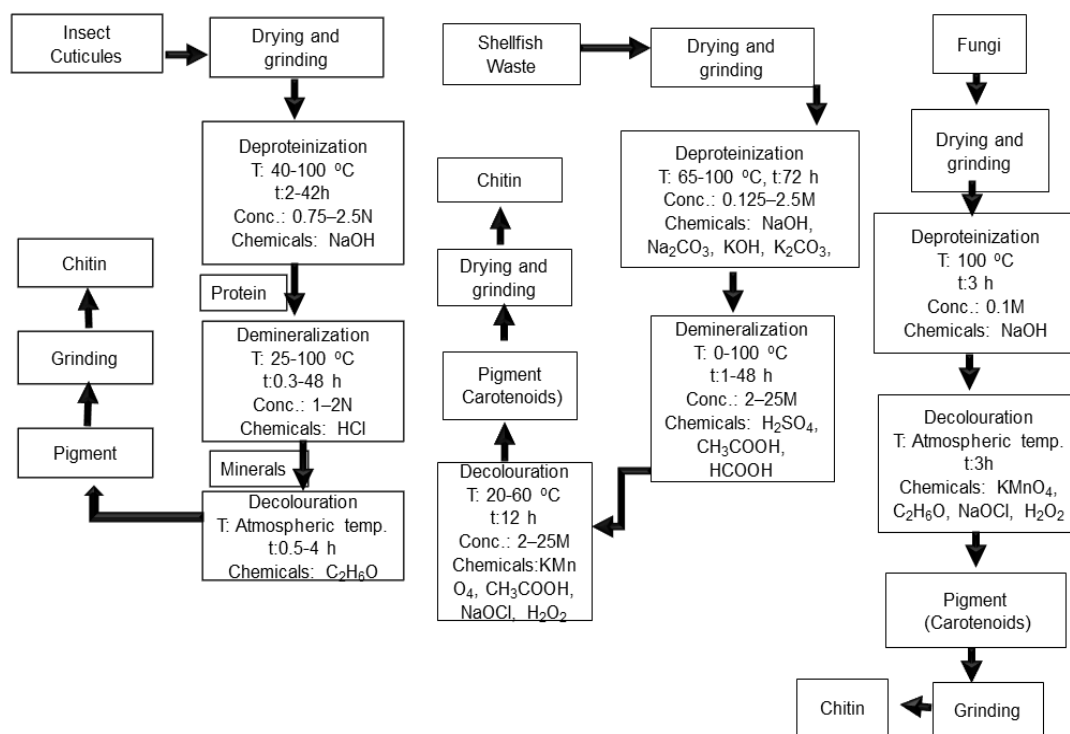


Fig. 5. Chemical and biological extraction of chitosan from chitin

Mohammed *et al.* (2013) extracted chitin and CS from prawn shells. Initially, frozen prawn shells were washed with boiled water then dried at 60 °C. The dried prawn shells were crushed and treated with 5% NaOH and refluxed at 60 °C for 2 h, followed by acetone treatment to remove colours at 25 °C for 24 h. To dissolve the calcium carbonate, these were further treated with a 0.5 or 1% HCl solution for 24 h at 25 °C. Chitosan was

obtained by deacetylation process using NaOH solution at elevated temperature and concentration. After the procedure, chitosan was washed several times with distilled water and dried at 60 °C in a vacuum oven. Figure 6 depicts the relative comparison of chemical and biological processes to produce chitosan. While chemical extraction methods are quick, consume less time, and are applied on a commercial scale, they consume more energy and make protein components unsuitable for animal feed. As an alternative, biological processes offer the advantage of environmental safety and the production of valuable by-products for animal feed. Investigations to reduce the processing time and cost is still underway.

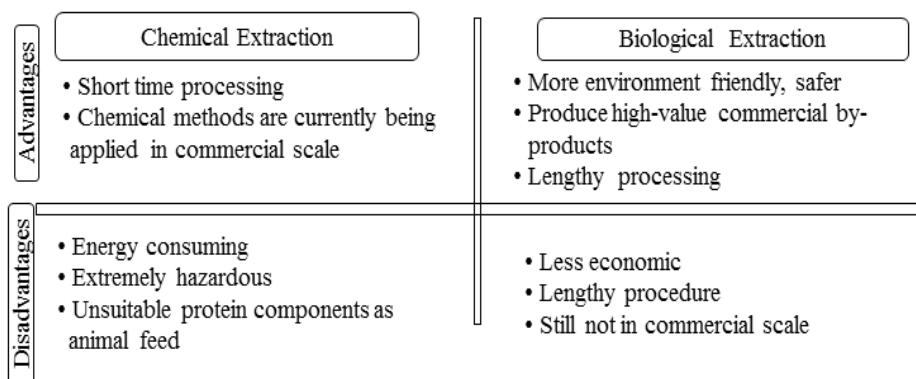


Fig. 6. Comparison of chemical and biological extraction methods

Table 1. Chitin and Chitosan Extraction from Different Species

Source	Chitin (%)	Chitosan (%)	Reference
<i>Bombyx mori</i>	15 to 20	70 to 80	(Zhang <i>et al.</i> 2000)
<i>Bombyx mori</i>	2.59 to 4.23	73% to 96.75	(Paulino <i>et al.</i> 2006)
<i>Mud crab Scylla tranquebarica</i>	Carapace: 10.74 Claw: 7.91; Legs; 14.62	6.59 4.12; 8.42	(Thirunavukkarasu and Shanmugam 2009)
<i>Zophobas morio</i>	35	25	(Mohammed <i>et al.</i> 2013)
<i>Leptinotarsa decemlineata</i>	7 to 20	67 to 72	(Kaya <i>et al.</i> 2014)
Shrimp	20	14.9	(Hajji <i>et al.</i> 2014)
Crab	10	5.3	(Hajji <i>et al.</i> 2014)
Cuttlefish	5	1.2	(Hajji <i>et al.</i> 2014)
<i>Fomitopsis pinicola</i>	30.11	71.75	(Kaya <i>et al.</i> 2015)
<i>Gryllus bimaculatus</i>	2.35	1.79	(Kim <i>et al.</i> 2017)
<i>Zophobas morio</i>	4.8 to 5.4	66 to 76	(Soon <i>et al.</i> 2018)
<i>Brachytrupes portentosus</i>	4.3 to 7.1	2.4 to 5.8	(Ibitoye <i>et al.</i> 2018)
<i>Macropipus holsatus</i>	12.23	9.52	(Pădurețu <i>et al.</i> 2019)
<i>Potamon algeriense</i>	8.27	5.89	(Fadlaoui <i>et al.</i> 2019)
<i>Hermetia illucens</i>	46	80	(Khayrova <i>et al.</i> 2019)
<i>Hermetia illucens</i>	9 to 30	18 to 29	(Khayrova <i>et al.</i> 2020)
<i>Hermetia illucens</i>	83 to 87	13 to 43	(Hahn <i>et al.</i> 2020a)

Table 1 lists previous work on chitin and chitosan extraction from different species (Bolat *et al.* 2010; Limam *et al.* 2011; Isa *et al.* 2012). Sagheer *et al.* (2009) extracted protein content in chitin from the crustaceans of Arab Gulf states. Similarly, Abdou *et al.* (2008) reported the production of chitin and its various derivative from crustaceans. Bolat *et al.* (2010) reported the extraction and characterization of chitin and chitosan from the crab. Though different experimental conditions are found in the literature, mild conditions are preferable to obtain chitin and chitosan with a high percentage of the degree of acetylation (Percot *et al.* 2003).

SPECIFIC PROPERTIES OF CS-GO COMPOSITES

Graphene oxide (GO), a novel 2D nanomaterial, is mainly produced by the modified Hummers method from natural graphite powder. For many years it attracted significant attention due to having various properties such as being readily exfoliated into monolayer sheets, hydrophilic functional groups on its basal planes and edges, interfacial interactivity with a target matrix, and electronic properties. Covalently grafting CS onto GO sheets improves the solubility of graphene. Many researchers grafted or crosslinked the CS onto GO. Various researchers synthesised or prepared the different types of CS-GO materials such as nanoparticles, green adsorbents, casting on membranes, hydrogel and aerogels. Some of the studies are discussed here to provide insight into the chemistry of CS-GO. Studies have shown the solid electrostatic interactions occurred between the cationic CS and the negatively charged GO composites (Suri and Khandegar 2021). Horse-radish-peroxidase (HRP) was adsorbed onto CS-GO through covalent bonding, which assisted in enhancing the mechanical strength of HRP at a reasonably low cost. The reusability studies specified that the HRP- CS-GO could be reused for a minimum of 5 cycles (Suri *et al.* 2020). Researchers have designed CS nanoparticles embedded into GO to be used for a drug delivery system. The XRD analysis showed that the composite has an amorphous structure bonded with intramolecular hydrogen bonding and excellent porous, rough surface morphology and flaky structure, as shown in Fig. 7 (Hosseini *et al.* 2021).

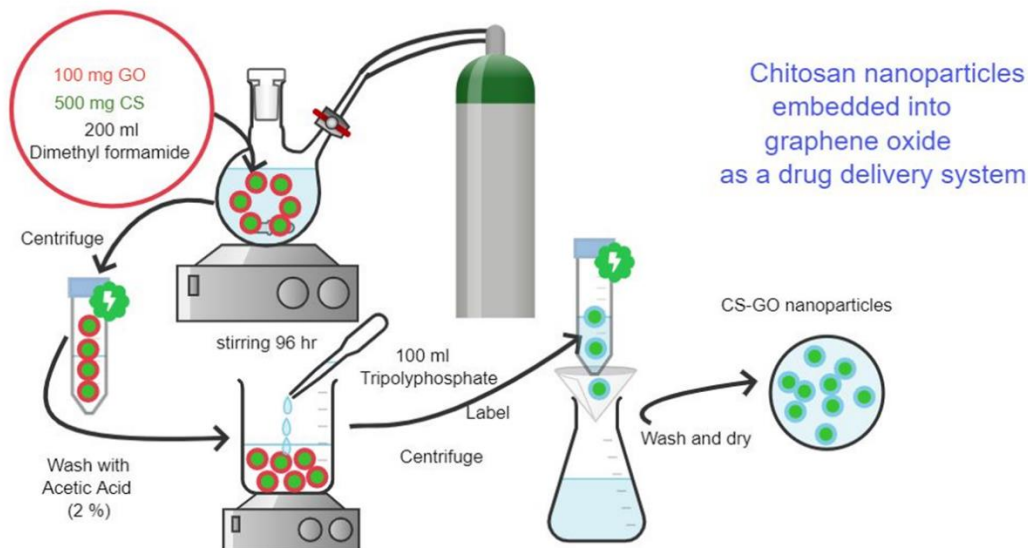


Fig. 7. Methods of synthesis of CS-GO for drug delivery system (Hosseini *et al.* 2021)

APPLICATIONS OF CHITOSAN COMPOSITES

Polyfunctional CS is an organic molecule that is used in a wide range of applications. The chemical modification of CS increases the selectivity and adsorption potential of the target; amendment adjusts the Lewis basicity by introducing functional groups. These groups are different in donor ability than the amine and hydroxyl groups. Improvement can be carried out with or without crosslinking agents. Various forms of chitosan and adsorbent synthesis techniques are given in Fig. 4. GO has distinctive properties including high conductivity, flexibility, and controllable permittivity with hydrophilicity (Kostarelos and Novoselov 2014). These properties facilitate the development of GO-based multifunctional biomedical devices. Therefore, GO has been developed in different sizes and forms by the chemical modifications process. GO is made by chemical exfoliation of graphite powders through strong oxidants (Wang *et al.* 2013b). It shows high specific surface area (*e.g.* 2,600 m²/g) (Sponza and Alicanoglu 2017). This property can help as a support medium to make the nanoparticles without accumulation in the adsorption process. Graphene consists of a single layer of sp² bonded carbon atoms. Graphene is the thinnest material and the most robust material due to its particular thermal, mechanical, optical, and electrical properties (Geim 2009). Oxygen functionalities in GO permit the interactions with the positive ion and provide active sites for the nucleation and growth of nanoparticles (Guria *et al.* 2016).

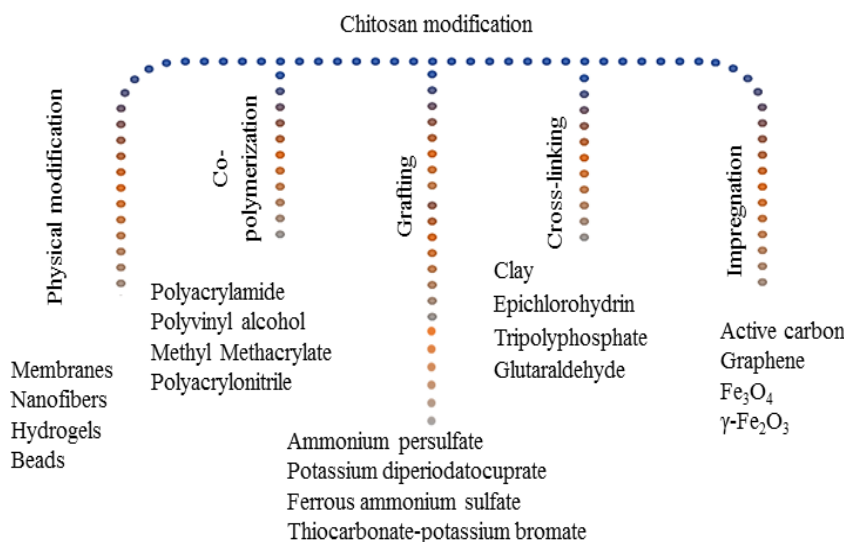


Fig. 8. Forms and modification of chitosan to improve its properties

GO has plenty of functional groups and high mechanical strength as the prospective material for removing heavy metals (Menazea *et al.* 2020). However, inter-functional solid bonds between graphene sheets result in static surface chemical properties, lower surface area, and poor aqueous solutions dispersion. Reduced adsorption efficiency limits its use in wastewater treatment (Li *et al.* 2009b). CS and GO-based nanocomposites can be used in several cosmetics, as a fixative in photography, and as an adsorbent and flocculating agent in wastewater treatment, in the paper and textile units agriculture sector (Dutta *et al.* 2004; Philibert *et al.* 2017). Chemical, physical, or combinations of both modes of modification are used for CS. The physical transformation comprises blending and conversion of chitosan's forms. The physical change leads to expansion of the polymer

chains of chitosan. As a result, internal sorption sites increase, and their crystalline state decreases. Blending chitosan with active carbon, graphene, Fe_3O_4 , $\gamma\text{-Fe}_2\text{O}_3$, and clay is another suitable method for synthesizing better strength, adsorption, sensitivity, and magnetism property nanocomposite (Badry *et al.* 2017; Wang and Zhuang 2017). Chitosan modifications by chemical treatments are either crosslinking or graft co-polymerisation. Chemical changes improve complex formation properties with metal ions and surfactants (Rinaudo 2006; Alves and Mano 2008). Different cross-linkers such as epichlorohydrin, tripolyphosphate, and glutaraldehyde are used to modify (Mojiri *et al.* 2019). Homogeneous crosslinking of chitosan improves metal binding capacity relative to heterogeneous cross-linking (Varma *et al.* 2004). Figure 8 shows the different forms and modifications of CS, which can help to strengthen its properties.

Biomedical Applications

A drug delivery system is defined as a formulation or a device that enables the administration of a therapeutic substance or active pharmaceutical ingredient in the body, improves its efficacy as well as safety by controlling the rate, time, and place of release of drugs (Jain 2020). Researchers have investigated the potential of nanoparticles such as dendrimers, liposomes, self-assembling peptides, water-soluble polymers, and polymeric micelles as effective drug delivery agents. They find unique applications for cancer treatment where nanoparticulate drug carriers improve cure efficacy by reducing off-target systemic toxicity and passive drug targeting to tumour cells or tissues (Yu *et al.* 2016; Edgar and Wang 2017). Graphene oxide (GO), reduced graphene oxide, and graphene quantum dots have shown high surface area, which provides sufficient drug loading capacity along with biocompatibility. Functionalization of GO or reduced GO with polyethylene glycol, mannose, and sulfonic acid conjugated to folic acid has improved biodegradability, biocompatibility, and thermosensitivity, facilitates pH-dependent and targets specific controlled release of the drug to tumour cells (Karki *et al.* 2020). Also, GO displays low uptake in the reticuloendothelial system and extended blood circulation time (Zhao *et al.* 2018). Still, these GO-based nanocomposite systems have limitations related to drug solubility (colloidal instability due to aggregation), controlled and targeted drug release to cancer cells, hemolytic properties, and *in vivo* toxicity (McCallion *et al.* 2016; Karki *et al.* 2020). Extensive research is being carried out to identify new and novel functional groups for GO modifications, where CS has emerged as an excellent helpful natural modifier of GO. CS-based nanocomposites are biocompatible, non-toxic, and display increased drug efficacy (Thakur and Thakur, 2014). Due to their small size, these nanoparticles or nanocomposites can move through blood-brain barriers, thus using efficient drug delivery systems (Li *et al.* 2018). Despite the above advantages, owing to its low solubility and poor mechanical properties, the application of CS is still minimal. Different formulations of CS and its composites with several other compounds are being worked upon to increase targeted drug action and efficacy for drug delivery systems (Ali and Ahmed 2018; Li *et al.* 2018; Ashrafizadeh *et al.* 2019).

Drug delivery system

Derivatized chitosan has been used to modify GO to prepare nanocomposites for drug delivery to cancerous cells. Rana *et al.* (2011) developed chitosan-graphene oxide nanocomposites. The chitosan-functionalized graphene oxides served as a drug delivery system, which was loaded with Ibuprofen and 5-fluorouracil. It displayed controlled release and long term biocompatibility in human lymphoblastic leukemia and MCF7-human breast

cancer cells. Another functionalized nanocarrier based on CS-GO nanocomposite was prepared by Bao *et al.* (2011) through the amidation process. CS-GO nanocomposite was applied to load the anti-cancer drug Camptothecin. Cell toxicity was analyzed against HepG2 and HeLa cancer cells using MTT assay, where a 50% growth inhibition concentration was observed at 29 μm . Thus, the novel nanocarrier was potent against cancer cells. An *in vitro* drug release experiment showed that 17.5% camptothecin could be released. This CS-GO nanocomposite camptothecin complex exhibited enhanced stability and a high loading capacity.

Yang *et al.* (2016) modified GO with carboxymethyl chitosan and further conjugated it with hyaluronic acid and fluorescein isothiocyanate. This conjugate combination was used as a drug delivery vehicle for doxorubicin, a model anti-cancer drug, to study *in vitro* release behaviour. It specifically targeted cancer cells and inhibited the growth of HeLa, CD44 overexpressing cells. It exhibited pH-dependent release and a drug loading efficiency of 95%. Pan *et al.* (2016) developed a GO-CS nanocomposite by conjugating GO with carboxymethyl chitosan, fluorescein isothiocyanate, and lactobionic acid to be used as a targeted anti-cancer drug delivery system; nanocomposite without LA conjugate was used as the control. Functional group lactobionic acid is recognized explicitly by asialoglycoprotein receptors overexpressed on cancerous hepatic cells. High drug loading content and efficiency of >96% was observed in SMMC-7721 cancer cells. The pH-sensitive release of the nanocomposite was followed, which is typical due to reduced pH microenvironment of cancerous cells. Higher uptake was observed using this nanocomposite for SMMC-7721 cancer cells through confocal microscopy and cell toxicity assays, strengthening its use as an anti-cancer drug delivery system. Further, Wang *et al.* (2018) synthesized a galactosylated CS-GO-doxorubicin drug delivery system against hepatocellular carcinoma. The action of the synthesized composite was investigated against HepG2 and SMMC-7721 hepatocarcinoma cell lines. After coating glycosylated-CS over the surface of GO, drug loading capacity was found to be 98%. Drug release to hepatic carcinoma cells was studied using cellular assays such as cell proliferation and cellular uptake assay. An *in vivo* anti-tumour efficacy study showed efficient inhibition of tumour cells as compared to control. The above drug conjugated system might be used as biomedicine and may effectively target liver cancer cells.

In another study, developed chitosan/sodium alginate products were functionalized using magnetic GO-based nanocomposites and were loaded with doxorubicin. The functionalized magnetic graphene oxide nano sheets have high drug loading efficiency (Xie *et al.* 2019). A nanocarrier system based on reduced-GO was loaded with doxorubicin and coated with CS for stabilization was developed, which displayed high biocompatibility and efficiency for entrapping doxorubicin (~65%) and depicted controlled release (~50% release in 48 h). Furthermore, it was demonstrated to precisely deliver doxorubicin intracellularly in PC-3 prostate cancer cells with cytotoxicity >65% (SreeHarsha *et al.* 2019). Researchers have also synthesized folic acid coupled CS and GO nanocomposites. These composites were used for loading polyphenol and fullerene. They depicted good drug loading and encapsulation efficiency, drug release property, and storage stability. Also, cytotoxicity analysis of this nanocomposite in human hepatic cell line MHCC97H, exhibited greater inhibition capacity than regular human hepatic non-cancerous cell line (Tao *et al.* 2019). Shi *et al.* (2016) also prepared carboxymethyl-CS-GO-based nanoparticles using the electrostatic droplet generation method, which effectively adsorbed gatifloxacin, ofloxacin bovine serum albumin, lysozyme, and doxorubicin hydrochloride (Tao *et al.* 2019). Very recently, Anirudhan *et al.* (2020) synthesized CS and folic acid

nanocomposite by conjugating through *N,N'*-dicyclohexylcarbodiimide coupling. Furthermore, itaconic acid and acrylic acid monomers were grafted to the hydroxyl groups of prepared nanocomposites to generate -COOH functional groups and combined with modified amine GO. Further, the anti-tumour drug was loaded onto the modified amine CS-GO conjugated folic acid nanocomposite through π - π stacking and hydrogen bonding interactions. The loading capacity of doxorubicin in the polymeric matrix was determined to be 95%. An advanced CS-GO hybrid-responsive system of enhanced biocompatibility, high doxorubicin-loading ability, and tumour-inhibition efficacy was developed.

Table 2. CS-GO Nanocomposites with Chemotherapeutic Activity Against Cancer Cells

Nanocomposite/Drug Composite	Drug Conjugated	Type of Cancer or Cancer Cell Line	References
CS-GO	Camptothecin	HepG2 and HeLa cell lines	Bao <i>et al.</i> (2011)
Chitosan-functionalized GO	Ibuprofen and 5-fluorouracil	CEM human lymphoblastic leukemia and MCF7-human breast cancer	Rana <i>et al.</i> (2011)
GO conjugated with Carboxymethyl chitosan, hyaluronic acid and fluorescein isothiocyanate	Doxorubicin	HeLa, CD44 over-expressed cells	Yang <i>et al.</i> (2016)
GO conjugated with carboxymethyl chitosan, fluorescein isothiocyanate, and lactobionic acid	Doxorubicin	SMMC-7721 cancer cells	Pan <i>et al.</i> (2016)
Functionalized GO with CS sodium alginate	Doxorubicin	MCF-7 Human Breast cancer cell line	Lei <i>et al.</i> (2016)
Galactosylated CS-GO-doxorubicin	Doxorubicin	HepG2 and SMMC-7721 hepatocarcinoma cell lines	Wang <i>et al.</i> (2018)
Dispersion of GO Ag nanohybrid particles in the chitosan hydrogel matrix	Doxorubicin	Human colon cancer cells (SW480)	Rasoulzadehza li and Namazi (2018)
Magnetic GO nanosheets functionalized with CS and sodium alginate through non-covalent layer-by-layer self-assembly	Doxorubicin hydrochloride	Human lung cancer cell line (A549)	Xie <i>et al.</i> (2019)
Folic acid coupled CS-GO	Ginkgo Biloba Leaves polyprenol and Fullerene (C60F)	MHCC97H	Tao <i>et al.</i> (2019)
Chitosan-based hybrid nanoparticle of DOX-loaded reduced form of GO	Doxorubicin	Prostate cancer cells PC-3	SreeHarsha <i>et al.</i> (2019)
Amine functionalized GO conjugated with chemically modified CS with folic acid	Doxorubicin	MCF7 and Hela cells	Anirudhan <i>et al.</i> (2020)

For hybrid preparation, deprotonated carboxyl of GO nanoparticles and the protonated amine of CS's backbone were allowed to self-assemble. The developed hybrid was loaded with doxorubicin and displayed charge reversal from negative charge during blood circulation (pH 7.4) to positive charge at tumour extracellular microenvironment (pH 6.5). Biocompatibility, cell toxicity, in-vitro triggered release, and intracellular uptake of the prepared nanocomposite were studied in HepG2 cells; cell viability of HepG2 cells upon treatment the nanohybrids were found to be around 96% (Zhao *et al.* 2018). Another drug delivery system was synthesized by Rasoulzadehzali and Namazi (2018) by dispersion of GO-Ag particles in the CS-based hydrogel matrix for controlled release of doxorubicin, and a sustained-controlled drug release profile was observed. Additionally, Lei *et al.* (2016) developed another nanocarrier with functionalized graphene oxide, chitosan, and sodium alginate. The composite displayed significant pH-dependent doxorubicin release behaviour and cytotoxicity in MCF7 anti-cancer cells.

Over the last decade, different research groups have used vivid strategies to generate CS-GO drug delivery systems with many drugs, as summarized in Table 2. These studies support the application of graphene-chitosan based material as a chemotherapeutic agent. These new drug delivery systems promise targeted delivery, controlled release, and enhanced therapeutic efficiency of many anticancer drugs for cancer treatment.

Antimicrobial properties

Microbial infection is responsible for a plethora of diseases. Anti-microbials are agents that target microbes such as bacteria, fungi, protozoans, and viruses. Antibiotics are used widely to treat bacterial infections; however, antibiotic resistance has become a significant public health challenge as multidrug-resistant microorganisms are steadily rising. Nanotechnology-based antibiotics or nanobiotics have promising results in targeting antibiotic resistance in several diseases. These nanoparticles act through multiple mechanisms, load multiple drugs onto a single nanoparticle, thereby reducing the chances of antibiotic resistance. Metal-containing nanoparticles, chitosan-containing nanoparticles and graphene-based nanoparticles, cationic liposomes, and dendrimers are a few examples (Pelgrift and Friedman 2013; Gómez-Núñez *et al.* 2020; Vassallo *et al.* 2020). Due to its structure, graphene has intrinsic anti-bacterial activity. Hence in nanoparticles based on graphene, bacterial cell walls and membranes are damaged by sharp edges of graphene, leading to the release of intracellular content followed by bacterial cell death. The generation of reactive oxygen species has been speculated as another cause (Gómez-Núñez *et al.* 2020).

The antimicrobial effectiveness of CS-containing nanoparticles is probably due to their cationic nature. These particles associate with negatively charged cell walls or membranes on bacterial cells, altering their permeability leading to efflux of cytoplasmic contents and eventually bacterial cell death. These nanoparticles can associate with negatively charged bacterial DNA and chelates metal ions necessary for the functioning of enzymes and membrane integrity, thereby exerting solid antimicrobial activity (Pelgrift and Friedman 2013; Yilmaz Atay 2019). CS-GO based nanocomposites display intense antimicrobial action against bacteria, fungi, and yeasts. Over the last decade, many research groups have developed several CS-GO-based nano-composites with different functional groups having potent anti-microbial activity. Keshvardoostchokami *et al.* (2020) synthesized GO-based silver nanocomposites, where CS was used as a substrate. A broad-spectrum antibacterial activity of these nanocomposites was observed against both Gram-positive (*C. glutamicum*) and gram-negative bacterium (*E. coli* strain DH5 α).

Approximately 5 μL (2 g/L) of the nanocomposite efficiently reduced 10^8 colony-forming units (CFU)/ml to zero. An antimicrobial biofilm from CS-iron oxide coated GO hydrogel (CH-GIO) displayed robustness and, during characterization, showed significantly improved mechanical and thermal properties. Hydrogel biofilm incredibly worked against both Gram-positive and negative bacterial species such as *S. aureus* (methicillin-resistant), *S. aureus*, *E. coli*, and against fungus *C. albicans*. The effect was assessed by agar diffusion and cell viability assay using MTT dye. 10^5 CFU/mL of bacteria were seeded onto agar plates. Upon treatment with CS-GO films, the cell population was reduced to zero, possibly due to the generation of reactive oxygen species when these particles enter through the cell membrane.

Further, post-treatment with CS-GO films, the cell viability of mouse L929 fibroblastic cells were 80 to 93% higher than controls, thus confirming the non-cytotoxic nature of the synthesized films. The least inhibitory concentration value of CS-GO against *E. coli* was 32 (g/ml) (Li *et al.* 2016). Another composite, AgO-CoO-CdO/Poly(alanine)-chitosan-reduced graphene oxide (PACSGO) nanocomposite was developed by Zhang *et al.* (2020). This composite worked as a nano-photocatalyst for the substantial degradation of organic dye compounds from water. Additionally, the group reported enhanced antibacterial activity against *S. aureus*, *E. coli*, *P. aeruginosa*, and *B. cereus* medium and increased inhibition zone value with the addition of the nanocomposites. This nanocomposite showed a higher percentage of scavenging activity of AgO-CoO-CdO/PACSGO than other composites.

de Faria *et al.* (2015), using GO and silver particles nanocomposites, fabricated electrospun mats, where the blend of CS and poly(lactide-co-glycolide) (PLGA) was used as biopolymeric fibre. These PLGA/chit-GOAg electrospun mats displayed a bacterial inactivation rate of $\sim 98\%$ for *E. coli* and *P. aeruginosa* compared to control (nonmodified control PLGA-chitosan). A lower inactivation rate of 79.41% was observed against *S. aureus* due to a thick peptidoglycan layer in the cell wall of Gram-positive bacteria. Additionally, Yang *et al.* (2019) synthesized an antibacterial nano agent by coating quaternized chitosan (QCS) on the surface of Fe_3O_4 nanoparticles-anchored GO. This nanocomposite killed pathogens upon generation of hyperthermia through photothermal near infra-red irradiation.

The anti-bacterial efficacy of CS-GO based film was also investigated in detail by Marin *et al.* (2019). Electrospun CS-GO-and polyvinyl alcohol-based composite nanofibrous membrane was synthesized and analyzed. With the percentage of GO at 1%, the scaffold membrane showed antibacterial properties against gram-positive bacteria, *B. cereus* and *S. aureus*, and gram-negative bacteria, *S. enterica* and *E. coli*. The anti-bacterial inhibitory effect of the electrospun nanocomposite scaffold might be due to GO, as its edges penetrate the cell membrane, leading to membrane rupture and lipid peroxidation. Thus, the nanofibrous composite membrane might be used as a scaffold in exposed wounds and infected areas where the risk of bacterial infection is grave. Khalil *et al.* (2020) also showed that GO-CS and GO-EDTA based nanocomposites display anti-microbial activities against *E. coli*, *S. aureus*, and *C. albicans*. There have been a few pre-clinical investigations on their wound healing applications (Moradi *et al.* 2021). For instance, a temporary skin graft was made of CS-polyvinyl pyrrolidone-GO nanosheets. Live and dead assays were performed in-vitro. Cell viability and bactericidal capacity of this membrane were increased by using GO in the sheet. Accelerated healing tests were done using the rat's skin. A period of 14 days was required for complete regeneration of skin at a large wound. Nanosheet made of CS-GO covering was able to heal the wound almost completely

(>99%) after 21 days following the injury (Mahmoudi *et al.* 2017).

CS-GO-PVA composite was prepared using synthesized guanidine modified GO with (0, 0.1, 0.2, 0.5, and 1.0 wt%). In-vivo experiments validated that CS-GO (0.5%)-PVC sheets had a high antibacterial activity (50% more than GO used alone). *In vivo* studies on mice had shown 41% faster wound healing capacity than the control (Chen *et al.* 2020). The synergistic effect was observed on combining CS-GO-poly(lactic acid) (PLLA) for wound healing. CS-PLLA-based scaffolds were coated with GO. Pig iliac endothelial cells and female Sprague-Dawley rats were used for cytocompatibility and wound healing investigations. GO-coated CS/PLLA scaffolds resulted in a 60.48% decrease in the size of the wound. The wound was recovered in 21 days that shows its effectiveness as a wound-healing material. The scaffold was an effective bactericidal agent against *E. coli* and *S. aureus* (Yang *et al.* 2021).

Environmental Applications

CS finds extensive application in removal of metals in water treatment, as the amine and hydroxyl groups present in it easily form chelates with metal ions. CS-based membranes are reported to be useful to treat wastewater (Thakur and Voicu 2016). CS's insolubility in water and alkaline solution can be modified by GO, an excellent adsorbent with a high surface area. The self-agglomeration of GO is a limiting factor for its more comprehensive application. Various researchers have attempted to synthesis composite material using CS-GO. The negative charges on the surface of aqueous dispersed G sheets bond chemically with the cationic group present in chitosan's amino polysaccharides. Carboxylic groups of GO react through the amine group of CS, in some cases creating an amide bond between them. Various modifying agents such as EDTA can be added to the composite to functionalize these composites further. There is a wide range of applications of these composites, varying from water purification to electrode development. Most studies have been performed using optimization of parameters including solution properties including pH, temperature, an initial amount of adsorbent, contact time, swelling behaviour, and adsorbent properties such as its amount, form, and recyclability. Many active sites and more adsorbate in the waste stream have led to a higher mass gradient between solution and adsorbate. The ease of separation of adsorbents from the waste stream is also an essential factor influencing the adsorbent's broad choice.

Removal of heavy metals

Various functional groups including epoxy, hydroxyl, and carboxylic in GO provide oxygen to bind with metal ions. Likewise, amino groups present in CS binds with metal ions and can adsorb them from contaminated effluent. However, various limitations, including low operation efficiency, stability, and aggregation, are related to graphene oxide and chitosan usage alone. Hence, scientists have performed studies to crosslink and functionalize CS with polar groups such as graphene oxide. The modification of CS-GO composite has been performed using various materials. Table 3 lists multiple metals, CS-GO, and their modified composites and maximum adsorption efficiency. The research work shows its potential for adsorption of metals like Cu, Cr, As, Pb, Hg, Au, Pd. Heavy metals such as Cr(VI) can lead to severe hazardous effects on the environment and human beings. While Cr(III) is an essential trace nutrient, it is limited in drinking water to 170 (mg/L) by WHO (Altundogan 2005).

On the other hand, Cr(VI), occurring as chromate and dichromate, is a mutagen and carcinogen. The World Health Organization limit of Cr (VI) in drinking water is 0.050

mg/L, an excess of which may lead to various skin diseases and even cancer in human beings (WHO 1997). Multiple researchers have investigated CS-GO nanocomposites as an adsorbent to remove chromium (Anush *et al.* 2020; Samuel *et al.* 2018). Samuel *et al.* (2018) showed that this composite on exposure to chromium contaminated water possesses an adsorption capacity of 104.2 mg/g at pH 2.0 for the contact time of 420 min recyclability of 10 times. Copper and chromium metals were removed using a composite of CS-GO modified using 3-(p-anisyl)-4-formylsydnone by Anush *et al.* (2020). They studied wastewater contaminated with copper and chromium and reported that the pseudo-second-order kinetic reaction that followed Langmuir isotherm-model was best for their study.

Furthermore, the metal removed was a vital function of initial metal concentration in the water to be treated. The maximum removal capacity for Cu (II) and Cr (VI) was found to be 111.1 and 142.8 (mg/g), respectively. The reaction was a thermodynamically favourable process, being spontaneous and endothermic. Arsenic is a heavy metal and a prominent constituent of industrial waste. The World Health Organization has set the limit of 10 ($\mu\text{g/L}$) in drinking water (Minatel *et al.* 2018). Arsenic is a toxic metal released into the ecosystem through industrial activities. Entering the food chain causes nephrotoxicity, diabetes, cardiovascular, pulmonary, and skin diseases (De Loma *et al.* 2019). Functionalized CS-GO composite can be used as a potential adsorbent for arsenic removal from the aqueous medium. Kumar and Jiang (2016) reported that GO and CS composite performance depends on pH of the solution. Their study's optimum pH was 4.3 to 6.5 for As (V) removal with endothermic, spontaneous metal adsorption process, which follows a pseudo-second-order kinetic model. The composite was able to adsorb As(III) and As(V) with the maximum adsorption capacity of 64.2 and 71.9 (mg/g), respectively, with material recyclability of 3 cycles.

CS-GO composite's functionalization using gadolinium resulted in improved efficiency to separate arsenic from aqueous medium. This composite's enhanced adsorption capacity of 252.1 (mg/g) showed its potential to treat arsenic-contaminated wastewater (Choi *et al.* 2020). The adsorption capacity of GO-Gadolinium without chitosan is much lower than the composite modified using chitosan 216.7 (mg/g) (Lingamdinne *et al.* 2021). Graphene oxide and reduced graphene oxide mixed with magnetic particles removed As(III) and As(V) from the aqueous solution. The studies showed that treatment time, initial arsenic concentration, pH, temperature, anions, and humic acid affected the composite's arsenic removal efficiency. Likewise, magnetically modified CS-GO nanoparticles were efficient for Ni (II) adsorption from waste-water with the maximum adsorption of 12.2 (mg/g) (Tran *et al.* 2019). Mercury is also a harmful metal and is one of the major pollutants in river water. Chitosan was investigated as filler material inside the GO matrix to form CS-GO nanocomposite. GO, and CS adsorption capacity improved from 381 mg/g to 397 mg/g upon nano filling magnetically modified CS into GO matrix (Kyzas *et al.* 2014). EDTA was used to alter the properties of the CS-GO composite. Magnetically modified CS-GO composites were functionalized using EDTA and were found to be potential adsorbents for metals like Pb(II), Cu(II), and As (III) (Shahzad *et al.* 2017a). The CS-GO composite was prepared by freeze-drying and investigated for selectivity of lead metal. The composite prepared by adsorption with 5 wt% GO in CSGO exhibited high mechanical strength and the maximum Pb (II) adsorption as 99 mg/g (He *et al.* 2011). Attempts were made to prepare the membrane by combining EDTA, GO, and CS to remove Pb (II). The adsorption efficiency using CS/EDTA/GO (0.3%) membrane was obtained at 889 (mg/g) (Croitoru *et al.* 2020). Studies on the adsorption capacity of this membrane for other heavy metals can also be undertaken.

Table 3. CS-GO Nanocomposites for Heavy Metal Removal

Adsorbent	Heavy Metal	Adsorption Capacity (mg/g)	Reference
CS with PVA	Cr (VI) Cu (II) Zn (II)	3.5 5.3 2.8	(Jaros <i>et al.</i> 2005)
Magnetic CS and GO	Au (III) Pb (II)	Au (III): 1076.6 Pb (II): 216.9	(Liu <i>et al.</i> 2012)
Cyclodextrin- CS and GO	Cr (VI)	61.31	(Li <i>et al.</i> 2013)
GOCS10	Cu (II), Pb (II)	Cu (II): 70 Pb (II): 90	(Chen <i>et al.</i> 2013)
Magnetic CS and GO	Pb (II)	76.94	(Fan <i>et al.</i> 2013)
Magnetic CS and GO	Hg (II)	381	(Kyzas <i>et al.</i> 2014)
β -cyclodextrin/magnetic GO/EDTA	Cr (VI)	68.41	(Wang <i>et al.</i> 2014)
Magnetic CS and GO	Cr (VI)	82.14	(Debnath <i>et al.</i> 2014)
CS and GO nanofibrous composite	Cu (II), Pb (II) Cr (VI)	Cu (II): 423.8 Pb (II) :461.3, Cr (VI) :310.4	(Najafabadi <i>et al.</i> 2015)
TGOCS	Cr (IV)	219.5	(Ge and Ma 2015)
CS and GO with disodium EDTA-2Na	Cr (VI)	86	(Zhang <i>et al.</i> 2016)
CS and GO	As (III) As (V)	As (III): 64 As (V): 72	(Kumar and Jiang 2016)
CS-GO and CS/rGO	Cu (II)	202 and 150	(Yan <i>et al.</i> 2016)
EDTA functionalized magnetic CS and GO	Pb (II), Cu (II) As (III)	Pb (II): 206.52 Cu (II): 207.26, As (III): 42.75	(Shahzad <i>et al.</i> 2017b)
Phosphorylated CS and GO	U(VI)	779.44	(Cai <i>et al.</i> 2017)
GO-CS aerogel	U(VI)	250	(Huang <i>et al.</i> 2017)
CS and GO	Cr (VI)	104.16	(Samuel <i>et al.</i> 2018)
Magnetic CS and GO	Cu (II)	217.4	(Hosseinzadeh and Ramin 2018)
Low molecular CS	Cu (II)	80	(Boamah <i>et al.</i> 2016)
Cross-linked CS and GO	Pb (II)	566	(Sharma <i>et al.</i> 2019)
CS and GO	Cr (VI)	178	(Moghaddam <i>et al.</i> 2019)
Grafted CS and GO	Cr (VI)	270.27	(Samuel <i>et al.</i> 2019)
GO-CS-PVA hydrogel	Cd (II) Ni (II)	Cd (II):172 Ni (II): 70	(Li <i>et al.</i> 2019)
Magnetic CS and GO	As (III)	45	(Sherlala <i>et al.</i> 2019)
Magnetic CS and GO	Ni (II)	12.24	(Tran <i>et al.</i> 2019)
GO-CS sponge	Co (II) Ni (II)	Co (II): 224.8 Ni (II) :423.7	(Di <i>et al.</i> 2019)
Membranes using CS-GO	Pb (II)	767	(Croitoru <i>et al.</i> 2020)
GO-CS Gadolinium composite	As(V)	252	(Choi <i>et al.</i> 2020)
CS and GO	Cd (VI) Cu (II) Pb (II)	Cd (VI): 48.7, Cu (II) : 60.7, Pb (II) : 32.3	(Li <i>et al.</i> 2020)
Modified CS with 3-(p-anisyl)-4-formylsydnone and GO	Cr (VI) Cu (II)	Cr (VI): 142.85 Cu (II): 111.11	(Anush <i>et al.</i> 2020)
GO-Gadolinium composite	As(V)	216	(Lingamdinne <i>et al.</i> 2021)
GO-CS-Fe ₃ O ₄	Cd (II)	84	(Parastar <i>et al.</i> 2021)

Bessa *et al.* (2020) reported Hg (II) removal efficiency (97%) from ultra-pure water with a small mercury dose using CS-GO composite. Simultaneously, using the tap water, river, and seawater, removal efficiency was reduced to about 81%, 13%, and 7%, respectively. Their study concluded that chlorine in an aqueous medium significantly reduced the removal efficiency of the composite due to the formation of steady chloro-mercury complexes. CS-GO nanocomposite could adsorb Pd(II) and Au(III). CS with 5 wt.% GO combination showed maximum adsorption 1077 (mg/g) for Au(III) and 216.9 (mg/g) for Pd(II) (Liu *et al.* 2012). Other studies highlighted hydrogel's potential obtained using CS-GO and polyvinyl alcohol (PVA) in various proportions, with the optimum being 1:2:4. Adsorption studies showed that the model followed Langmuir isotherm. Co-CS-PVA matrix showed the maximum adsorption capacity for Cd (II) and Ni (II) as about 172 and 70 mg/g, respectively, for the contact time of 16 h (Li *et al.* 2019). GO was mixed with 4-aminothiophenol and NaNO₂ to form GO-SH material, mixed with chitosan powder to obtain GO-CS-SH composite material. In multi-metal effluent, the material was found to be most suitable for adsorption of Cd(II), followed by Cu(II) and Pb(II) (Li *et al.* 2015). Similarly, studies on Go-Cs composite material without any functionalization resulted in the maximum adsorption capacity for Cu(II), Pb(II), and Cd(II) as about 60, 48, and 32 mg/g, respectively (Li *et al.* 2020). Menazea *et al.* (2020) studied the interactions of heavy metals with CS-GO and proposed a model using density functional theory and suggested that high heavy metal removal can be achieved by CS-GO combination.

Removal of dye

The textile and printing industry produces dye-containing effluent, which should be treated before discharge into water bodies. Approximately 1 to 15% of the dye is released into waste water during the dyeing process, eventually entering the ecosystem (Galindo 2001). Aromatic compounds in paints make their biodegradation an arduous task. Coloured substances in water inhibit the permeation of light and hinder the photosynthesis process. If ingested, these dyes can cause eye burns, nausea, vomiting, dyspnea, and cyanosis disease (Senthilkumaar *et al.* 2005).

Similar to the removal of metals, adsorption is the method of choice for dye removal. When used individually, CS and GO have shown a high affinity for the adsorption of dyes present in the wastewater (Chandel *et al.* 2020). However, composite formed using chitosan and graphene showed improved adsorption properties over several cycles of repeated usage. Table 4 depicts the various CS-GO composites like hydrogel, beads, spheres, and sponge investigated for dye removal from aqueous solution. However, the adsorption of methylene blue dye on GO adsorbent used alone was 287 mg/g. CS-GO composite material leads to adsorption capacity as high as 402.6 mg/g (Sabzevari *et al.* 2018a). Due to the unique properties of CS-GO, they can be used for the adsorption of methylene blue dye. A sponge prepared using chitosan (9%) with GO exhibited an adsorption capacity of 275.5 mg/ g for methylene blue during a filtration process (Qi *et al.* 2018b).

Due to the better removal ability of magnetic nanoparticles from an aqueous medium, synthesis and kinetic studies of magnetic nanoparticles for water treatment have been successfully reported. Tran *et al.* (2017b) demonstrated the enhanced adsorption efficiency of magnetically modified CS-GO composite. The studies reported the adsorption efficiency of 10% weight of GO in CS/Fe₃O₄/GO nanocomposite for MB dye after 95 h for five repeated operation cycles. Neves *et al.* (2020) prepared a magnetic CS-GO composite to remove basic brown four dye.

Table 4. CS-GO Nanocomposites for Dye Removal

Adsorbent	Dye	Adsorption Capacity (mg/g)	Reference
Magnetic CS grafted with GO	MB	95.31	(Fan <i>et al.</i> 2012a)
Chitosan/graphene oxide	MB	180.83	(Fan <i>et al.</i> 2012b)
CSGO10	MB	74	(Singh <i>et al.</i> 2019)
CS-GO	Reactive black 5	277	(Travlou <i>et al.</i> 2013a)
Magnetic CS-GO	Reactive black	391	(Travlou <i>et al.</i> 2013b)
Chitosan/graphene oxide 10	MB, Eosin Y	MB: 300 Eosin Y: 302	(Chen <i>et al.</i> 2013)
GO-CS/silica	CR	294.12	(Du <i>et al.</i> 2014)
Magnetic CS-GO	MO	398.08	(Jiang <i>et al.</i> 2016)
CS-GO foam	MO	173.3	(Ma <i>et al.</i> 2016)
GO was cross-linked with CS	Acid yellow 36 and Acid blue 74	Acid yellow 36: 68 Acid blue 74: 85	(Banerjee <i>et al.</i> 2017)
Magnetic GO-CS composite	MO	30	(Tran <i>et al.</i> 2017a)
CS-GO sphere	MO and Acid red 1	MO: 230.91 Acid red 1:132.94	(Zhang <i>et al.</i> 2018)
GO-CS sponge	MB	275.5	(Qi <i>et al.</i> 2018a)
GO and CS-GO composite	MB	GO composite: 287 CS-GO composite: 402	(Sabzevari <i>et al.</i> 2018b)
Cs-Go aerogel	Indigo Carmine and MB	Indigo Carmine: 377 MB:169	(Luna <i>et al.</i> 2019)
CS-GO sponge bionic filter	Crystal violet dye	98	(Zhou <i>et al.</i> 2018)
GO derivative containing quaternary ammonium salt and magnetic CS	Basic brown 4	650	Neves <i>et al.</i> (2020)
Polyacrylate CS-GO hydrogel	MB and Food yellow 3	MB~ 296 Food yellow 3: 280.3	Chang <i>et al.</i> (2020)
Cs reinforced Go-hydroxyapatite (CS@GO-Hap) matrix	CR, Acid Red 1 and Reactive Red 2	CR: 43.06 Acid Red 1: 41.32 Reactive Red 2 :40.03	Sirajudheen <i>et al.</i> (2020)

Composite graphene oxide functionalized using quaternary ammonium salt and magnetic chitosan provided the maximum adsorption capacity of 650 (mg/g) and 95(%). The study showed that in three successive adsorption and desorption cycles, the dye uptake efficiency was 64%. Graphene nanoplates, a graphene derivative crosslinked with chitosan to form composite spheres, were explored for the adsorption of methyl orange and acid red one dye. The high adsorption capacity for methyl orange and acid red 1 was 230.9 (mg/g) and 132.9 (mg/g), respectively, which showed the CS-GO sphere's potential as an efficient dye adsorbent. The adsorption capacity was maintained at 90% after five repeated usage cycles (Zhang *et al.* 2018). CS-GO bead adsorbent prepared by gel and GO and investigated by researchers for adsorption of various cationic and anionic dyes. For orange dye II, 84 (%) removal efficiency was observed with CS/Gel0.1GN.

Vo *et al.* (2020) prepared hydrogel materials in which chitosan chains crosslinks GO-nanosheets. GO and CS were mixed in solution in the above hybrid, followed by sonification to form hydrogel of GO-CS. While the higher GO level was beneficial for cationic dye's adsorption, higher chitosan hydrogel showed selectivity towards anionic dyes. The combination provides maximum loading of congo red dye about 176 (mg/g) (Kamal *et al.* 2016). Recently, biopolymer-based aerogels have been studied for application in adsorption and water purification processes. Chitosan-based aerogel has emerged as a potential material for the remediation of dye-contaminated water. Chitosan with active amine and hydroxyl groups has a strong tendency to bind with various types of dyes. Chitosan is a highly porous material that has limited application due to its high propensity to collapse. Besides acting as a good filler in the chitosan matrix, graphene oxide provides mechanical strength and enhances chitosan's adsorption efficiency. Aerogel formed using nanocomposite CS-GO aerogels exhibited high retention ability for Indigo Carmine and MB, as 377 and 169 (mg/g), respectively (Luna *et al.* 2019). Similarly, aerogel prepared using CS-GO was an effective adsorbent for Metanil Yellow dye with removal efficiencies between 91 and 96 (%). This dye's adsorption capacity was 431.0 (mg/g) at the adsorbent dose of 8 mg, concentration 400 (mg/L), for the contact time of about 35 min, maintained at a shaking speed of 175 rpm. The aerogel performed the adsorption and desorption up to 5 cycles, which resulted in an adsorption capacity of around 80 (%) (Lai *et al.* 2019). Qi *et al.* (2018b) synthesized GO-CS nanocomposite to remove MB in column study. Optimum loading for CS into GO solution was 9% in CS-GO hydrogel, which MB released efficiently.

Removal of organic and inorganic pollutants

Chitosan offers excellent potential for the removal of a variety of contaminants from effluents. Modified and unmodified chitosan-based nanocomposites have shown an effective adsorption potential to eliminate a variety of heavy metals, which include Cu (II) (Boamah *et al.* 2016; Modrzejewska *et al.* 2016), Pb(II) (Shahzad *et al.* 2017; Dinh *et al.* 2018), Cr(VI) (Moghaddam *et al.* 2019; Samuel *et al.* 2019; Naicker *et al.* 2020), As(V) (Shahzad *et al.* 2017a), Ni(II) (Di *et al.* 2019; Tran *et al.* 2019), and Hg(II) (Kyzas *et al.* 2014). Also CS used has been used for removal of inorganic nutrients (NO_3^- , NO_2^- , and PO_4^{3-}) (Jozwiak *et al.* 2017, 2019; Kumar *et al.* 2019) and organic pollutants (Jozwiak *et al.* 2017; Escudero-Oñate and Martínez-Francés 2018). Further, Table 5 summarizes CS composites that have been used to remove organic and inorganic contaminants from wastewater.

Table 5. Inorganic and Organic Contaminant Removal using CS

Pollutant	Adsorbent	Removal efficiency (%)	Reference
n-hexadecane	Spores of <i>Bacillus subtilis</i> LAMI008 were entrapped in 3 mm chitosan beads and cross-linked with 0.3% glutaraldehyde	100	Sar and Rosenberg (1983)
Nitrate	Protonated cross-linked chitosan gel beads by glutaraldehyde	77	Jaafari <i>et al.</i> (2001)
Hydrocarbons	<i>R. corynebacterioides</i> immobilized on CS	60	Gentili <i>et al.</i> (2006)
Fluoride	Chitosan loaded with titanium	89	Jagtap <i>et al.</i> (2009)
Diheptyl phthalate	Molybdate-impregnated chitosan beads	92.5	Barreto <i>et al.</i> (2010)
Insecticide (permethrin)	Modified CS with ZnO	99	Arayne <i>et al.</i> (2011)
Pesticide (Atrazine)	Cross-linked chitosan-silver nanoparticles	98	Saifuddin <i>et al.</i> (2011)
Oil	CS microspheres produced by ionic gelation of CS with sodium tripolyphosphate	90	Grem <i>et al.</i> (2013)
Hexadecane	Bacterial strain <i>B. pumilus</i> entrapped in chitosan	81.83	Costa <i>et al.</i> (2014)
Naphthalene	Carbon nanotubes mixed with CS/polyvinyl alcohol and cross-linked with silane	97	Bibi <i>et al.</i> (2015)
Oil	Zirconium-chitosan composites	79	Elanchezhiyan <i>et al.</i> (2016)
Hydrocarbons	Chitosan beads	99	Dellagnezze <i>et al.</i> (2016)
Oil	chitosan/magnesium-aluminum layered double hydroxide hybrid composite	78	Elanchezhiyan and Meenakshi (2017)
Oil	Superhydrophobic and superoleophilic chitosan sponge	99	Su <i>et al.</i> (2017)
Fluoride	Zr (IV) modified CS-GO	90	Zhang <i>et al.</i> (2017)
Oil	Amphiphilic sodium salt of oleoyl carboxymethyl chitosan	75-85	Doshi <i>et al.</i> (2018)
Naphthanol	Magnetic Cs-Go composite	99.8	Rebekah <i>et al.</i> (2020)
Ofloxacin	CS-GO	99	Suri <i>et al.</i> (2021)

Arayne and co-workers (2011) investigated the potential of raw and ZnO-modified CS beads to remove insecticide used for agriculture. They found that chitosan beads have outstanding adsorption and removed 49% and 99% of an insecticide (permethrin) when using raw and modified chitosan beads, respectively.

Danalioglu *et al.* (2017) prepared a novel magnetic activated carbon/chitosan composite to remove ciprofloxacin, erythromycin, and amoxicillin. They found the best fit with a Langmuir isotherm model, indicating that 90 (mg/g) ciprofloxacin, 178.6 (mg/g) erythromycin, and 526.3 (mg/g) amoxicillin had adsorbed. Danalioglu *et al.* (2017) compared the adsorption potential of magnetic CS with activated carbon for ciprofloxacin with alginate-Fe₃O₄ hydrogel fibre and GO-Ca alginate adsorbent. About 154 µg/g ciprofloxacin was adsorbed using magnetic alginate-Fe₃O₄ hydrogel fibre and 18.45 to 39.06 mg/g GO-Ca alginate adsorbent (Wu *et al.* 2013).

Saifuddin *et al.* (2011) removed pesticide (atrazine) using CS-Ag NPs. Maximum 98% atrazine (from 1 ppm solution) was removed from solution using a composite dosage of 2.0 g/L of Cs-Ag NPs. Other investigators have reported using chitosan beads modified with sodium alginate and calcium chloride to remove phenol and o-chlorophenol (Li *et al.* 2009a). They said that modification in CS enhanced its stability and its sorption capacity toward pollutants. The maximum sorption capacities of 108.7 (mg/g) and 97.1 (mg/g) were achieved for o-chlorophenol and phenol, respectively. Zr(IV) is oxyphilic and makes a bond with oxygen present in GO. Researchers have prepared zirconium-chitosan/graphene oxide membrane (80 µm thick) and evaluated its adsorption capacity for anions present in wastewater. Zr(IV) forms an electro-positive ion. It has a strong affinity for fluoride ions with more than 90% removal efficiency in the contact time of around 30 min. The membrane displayed high selectivity for bicarbonate and sulfate ions as well. The membrane with sufficient mechanical strength has been reported to have a high potential for water purification application (Zhang *et al.* 2017).

CS has been recognized as one of the most effective biopolymers for removing oil droplets from water. CS exhibits a unique assembly that is prone to natural functionalization and permits the modification of novel sportive materials with oil-enhanced selectivity and adsorption potential (Farzana and Meenakshi 2015; Elanchezhiyan *et al.* 2016). Nitrogen, phosphorus and potassium are the vital nutrients required for proper plant growth, but excessive fertilizers in the fields lead to leaching soil and water bodies. As per the World Health Organization, the limit NO³⁻ and PO₄³⁻ in drinking water is 40 and less than 0.5 mg/L, respectively. The excessive amounts of these components in drinking water are harmful and lead to various diseases such as methemoglobinemia and gastric cancer (Zheng and Wang 2010). Various researchers have explored chitosan composite for nutrient removal from water. Hybrid beads encapsulating chitosan on the graphene oxide functionalized using triaminotriazine have been used to remove nutrients from agricultural soil. These beads were tested on a non-aqueous medium enriched with nutrients such as NO³⁻ and PO₄³⁻. The retention capacity of beads for NO³⁻ and PO₄³⁻ was 58.5 and 61.4 mg/g, respectively, which was much higher than any of these individual adsorbents (Kumar *et al.* 2019). In agriculture, pesticides are commonly used to control insects and pests that cause severe damage to the crop. The associated toxic effects of chemical pesticides have prompted researchers to look for biopesticides as an alternative. However, biopesticides' hydrophobicity is the most significant limiting factor in their wide range of applications (Lao *et al.* 2010). The organic solvents used in high quantities to improve their solubility in an aqueous medium are known to be highly polluting in nature (Muda *et al.* 2020). Researchers are looking for eco-friendly water

solubilizing carrier agents to stabilize biopesticides in an aqueous medium chemically. The binding mechanisms for carrier agents include encapsulation, entrapment, adsorption, and ligands (Meredith *et al.* 2016).

Strontium-90 is found mainly in nuclear waste. Its longer half-life, as well as its biocompatibility, make it toxic for human health. Among numerous methods to treat nuclear waste, adsorption is considered an effective technique to remove strontium. The low adsorption capacity of existing adsorbents such as clay and synthetic hydroxyapatite prompted researchers to develop adsorbents with higher selectivity. It has been found that the maximum adsorption capacity of strontium on GO and chitosan was 146, 112 mg/g, respectively. However, its adsorption capacity for GO-chitosan composite was 180 mg/g (Rouby *et al.* 2018). Rotenone (1,2,12,12a-tetrahydro-8,9-dimethoxy-2-(1-methyl-ethenyl)-1-benzopyrano(3,5-b)furo(2,3-h)benzopyran-6(6h)-one) is a non-toxic organic pesticide for pests such as corn borers, apple and pea aphids, Mexican bean beetles, and household pests. Researchers have performed studies to improve the hydrophobicity of rotenone using a Cs-GO nano-composite using various Cs and GO proportions. The process parameters were optimized, resulting in 48.5 times higher dissolution of pesticide in Cs-GO nanocomposite than control (Muda *et al.* 2020). The effluent of the dye industry, as well as the pharmaceutical industry, contains 2-naphthol. The studies have shown the potential of magnetically modified Cs-GO composite for its adsorption. The maximum adsorption capacity was 169 mg/g with almost complete removal efficiency for the treatment time of 45 mins (Rebekah *et al.* 2020).

CHALLENGES AND FUTURE DIRECTIONS

CS and GO are natural materials that have shown excellent biomedical and adsorption capacity, which are further improved by combining the two materials. CS-GO, prepared as a nanocomposite, offers a large specific surface area and good electrical properties. These composites promise a bright future for many treatments, including cancer and tumour treatment. The enhanced therapeutic value of CS-GO composite materials has been evaluated using animal model-based studies. They are safe, efficient, and ensure drug delivery at specific locations in the body in a controlled manner. Although advancement has been achieved in *in-vivo* studies, much work lies in clinical level investigations. The ongoing research on CS-GO is not sufficient and must be enhanced. The analysis described in this review is still in its infancy. It is being continuously improvised, but stringent validation needs to be performed *in vivo* and *in-vitro*, followed by assessing long-term adverse effects through extensive clinical trials to obtain an efficient, viable, and safe product. Clinical trials can precisely identify and compare the environmentally most favorable options for many medicinal applications. More clinical data is required to understand the relative advantages and limitations of CS-GO composites compared to CS or GO used alone.

The mounting number of metals, dyes, and organic pollutants can lead to severe environmental degradation. The effect of the same on human health and the eco-system can be disastrous if not addressed soon. CS-GO-based hydrogel, aerogel, beads, and spheres have helped remove metals including Cu, Cr, Pb, Pd, As, Au, and Hg, which are otherwise severely toxic. Currently, the research work is focused on the treatment of individual metals or dyes. However, the waste material in the field is a mixture of various metals. Thus, there is an urgent need to study the efficiency of these composite materials

to treat multi-metal waste. Magnetically modified composite emerged as a potential adsorbent for metals, dyes, organic, and inorganic contaminants. Under laboratory conditions, complete removal of pollutants has been achieved in a few studies. The retention capacity of these composite was found to be closer to 95%. Scale-up studies of these composite can lead to commercial application. Furthermore, the reusability study must also be prioritized in future research to verify the effectiveness and successfully fabricated nanocomposites compared to conventional treatments.

CONCLUDING COMMENTS

This review has summarized the properties of chitosan (CS), graphene oxide (GO), and CS-GO materials. The application of chitosan and graphene oxide nanocomposite has been studied in the present review, especially its adsorption capability and antimicrobial activity. This unique category of chitosan-based products can be modified with various functional groups to control their hydrophobicity, as well as cationic and anionic properties. The literature that was surveyed in this work makes it clear that the chitosan is a low-cost biopolymer, environmental-friendly, and abundant in availability. However, the modified form of CS-GO has many applications and is cost-effective. Their unique attributes have allowed researchers to use them as a suitable medium to treat many diseases. Although the successful clinical translation of CS-GO composite has not been achieved yet, they have shown great promise, thereby providing hope for new treatment of various diseases soon. In the current scenario, the costs associated with synthesis of CS-GO may not be lowered than other commercially available nanocomposites; therefore, more well-organized and cost-effective ways are required to produce CS-GO. Future research could examine a low-cost CS-GO composite production technique without compromising environmental and health implications.

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