


 Cite this: *RSC Adv.*, 2020, 10, 20231

Received 2nd April 2020

Accepted 19th May 2020

DOI: 10.1039/d0ra02979c

rsc.li/rsc-advances

Application progress of enhanced coagulation in water treatment

 Hongmei Cui,^{ab} Xing Huang,^{id}^a Zhongchen Yu,^{*ab} Ping Chen^{ab} and Xiaoling Cao^a

Water industries worldwide consider coagulation/flocculation to be one of the major treatment methods for improving the overall efficiency and cost effectiveness of water and wastewater treatment. Enhancing the coagulation process is currently a popular research topic. In this review article, the latest developments in enhanced coagulation are summarized. In addition, the mechanisms of enhanced coagulation and the effect of process parameters on processing efficiency are discussed from the perspective of ballast-enhanced coagulation, preoxidation, ultrasound, and composite coagulants. Finally, improvements and new directions for enhanced coagulation are proposed.

1. Introduction

Coagulation is one of the most common phenomena in nature and artificially enhanced water treatment systems. Coagulation is a process in which small particles are combined into larger aggregates (flocs) and dissolved organic matter is adsorbed onto particulate aggregates so that these impurities can be removed in the subsequent solid/liquid separation processes.¹ The mechanism for the removal of organics *via* coagulation has three main aspects:² (1) positively charged metal ions and negatively charged organic colloids are electrically neutralized, destabilized and aggregated; (2) metal ions and soluble organic matter molecules form insoluble complexes and precipitates; and (3) physical and chemical adsorption of organics occurs on the surface of alum. As environmental pollution problems become more serious and water quality standards become more stringent, conventional coagulation technology has clearly failed to meet people's requirements for water quality safety. Enhanced coagulation and optimized coagulation are improved based on existing water treatment process facilities and by considering the operating conditions of previous and subsequent process flows to achieve the effect of advanced water and wastewater treatment.

The concept of enhanced coagulation has a long history and was discussed in a paper in the American AWWA Journal in 1965. The enhanced coagulation proposed by the American Hydraulic Association in the 1990s refers to the improvement of organic matter removal rates by increasing the amount of coagulant in the conventional coagulation treatment process of water while ensuring the turbidity removal effect process.³ This concept of enhanced coagulation is based on an increase in the

amount of coagulant added or the coagulation process controlled by the reaction pH conditions. Optimized coagulation is proposed based on enhanced coagulation, which is a coagulation process with multiple goals: maximizing particulate matter and turbidity removal, maximizing TOC and DBP precursor removal, reducing residual coagulant content, reducing sludge production, and minimizing production costs.^{4,5}

In recent years, the intensity of research on enhanced coagulation has been unprecedentedly increased, with many scholars conducting more in-depth research on enhanced coagulation. For example, studies have attempted to gain an in-depth understanding of the characteristics of organic matter in water bodies, identify organic matter removal rules to establish an organic matter removal mode, and summarize the mechanisms of enhanced coagulation removal of organic matter. Previous studies found that increases in efficiency are affected not only by the dose and pH of the coagulant but also by the nature and distribution of the organic matter and particulate matter in the water, including the temperature, hydraulic condition, and coagulant morphology.^{6–8} With the continuous expansion of modern coagulation treatment processes, the concept of pollutants in water bodies has become increasingly extensive, the treatment difficulty has become increasingly difficult, and ways to strengthen coagulation have become more diverse. For example, permanganate oxidation-assisted coagulation has been widely used to improve turbidity and natural organic matter (NOM) and algae removal in the coagulation–filtration process.^{9,10} Ballast microsand and magnetic powder are very mature technologies,^{11,12} including “Actiflo” technology and “CoMag” technology. Researchers have further enhanced the application of ballast technology in wastewater treatment by modifying or changing the dosing materials based on this principle. Jessica¹³ studied the potential impact of adding powdered activated carbon or organoclay (OC) adsorbent and

^aSchool of Civil Engineering and Architecture, Northeast Petroleum University, China

^bKey Laboratory of Disaster Prevention and Mitigation, Projective Engineering of Heilongjiang Province, Daqing 163318, China


ferric chloride (FeCl_3) coagulant together on the coagulation, and the results showed that the turbidity of the clarified water sample was lower when OC was added during the coagulation process and powdered activated carbon removed more than half or almost all of the dissolved aromatic compounds. Kadouche¹⁴ used synthetic hydroxyapatite in combination with chitosan and showed that it can effectively remove 88–95% of copper and zinc ions. Chitosan significantly improves the speed of coagulation and precipitation. Another popular topic of research is the development and utilization of new coagulants, such as inorganic polymer flocculants, biopolymer flocculants and natural coagulants.¹⁵ Many problems occur under enhanced coagulation. For example, preoxidation may damage the cell membrane, resulting in toxin release and increased taste and odor,¹⁶ and the addition of microsand will adversely affect the maintenance of the pump. Therefore, it is not only necessary but also meaningful to study the working mechanism, challenges and new discoveries associated with strengthening the coagulation process to minimize its disadvantages. This article summarizes the application of enhanced coagulation in water treatment, including the current advantages and disadvantages, and discusses the challenges associated with the development of enhanced coagulation technology in the future.

2. Ballasted reaction nuclei enhance coagulation

The process of ballasted reaction nuclei-enhanced coagulation mainly refers to improving the flocculation stage to strengthen its sedimentation characteristics and optimize the flocculation agent and hydraulic conditions in the flocculation stage to form flocs with good sedimentation performance.^{17–19}

2.1 Technical principle

Floc characteristics are important parameters for controlling the flocculation process and have a significant impact on subsequent sedimentation units and effluent water quality. Related research results showed²⁰ that flocs with large particle sizes and high densities have higher sedimentation speeds and better solid–liquid separation effects than those with small particle sizes and low densities and the turbidity of water after they sunk was correspondingly lower. Therefore, the formation of flocs with good sedimentation performance is an important way to improve the water quality of submerged water.^{12,21} Miyahara²² confirmed the higher fractal dimension and higher density of flocs with larger particle diameters using the Stokes formula. Many researchers^{17,18,23,24} believe that denser flocs have higher strength than less dense flocs, are not easily broken and are better for achieving solid–liquid separation. Ballasted nuclei-enhanced coagulation technology aims to increase the speed of floc formation and the density of flocs through ballasted nuclei, such as silica sand, magnetite sand, anthracite, recycled glass, *etc.*, so that the flocs can settle quickly.^{21,25} The ballast nuclei strengthening flocculation process keeps the core medium in suspension by mixing, and the mixing strength is controlled by the specific gravity and size of the nuclei medium.

Ballast nuclei with different specific gravities have the advantages of increasing the sedimentation speed of the shallow layer or increasing the particle removal rate because the specific gravity of the flocs is determined by the specific gravity of the nuclei.¹⁷ Different ballast nuclei have different characteristics. For example, the total suspended solid load of magnetite sand is twice that of silica sand, although the residual turbidity after sedimentation increases.²⁵

2.2 Microsand

The ballasted sand flocculation (BSF) technique is based on coagulation and flocculation with the injection of microsand by increasing the sedimentation rate²⁶ and overflow rate of the microsand to enhance coagulation. Depending on the type of wastewater, the amount of microsand added, and the amount of flocculant, there are significant differences in treatment effects. The ballasted sand flocculation technique has appeared to be an efficient method for the treatment of water and wastewater. The BSF trade named “Actiflo” was introduced in 1990.¹⁷ The BSF unit is shown in Fig. 1.²⁷ Before the first stage, the water is typically screened through a settling tank to remove large particulates. Then, the coagulant, microsand and polymer are injected and then mixed and stirred at a high rpm to fully coagulate. In the second stage, the maturation process, gentle mixing is performed at a low rpm to enhance the coagulation process. The third stage is clarification, wherein the mixed flocs and influent move downward in the unit. The clarified effluent is then either discharged into natural water bodies or directed to the next treatment process.

The BSF process improves the sedimentation speed and overflow rate, reduces the footprint of the system, and greatly reduces the cost of coagulation treatment. Many researchers have conducted in-depth research. For example, Desjardins¹⁷ conducted comparative experiments in three different water treatment plants, and the results showed that the amount of coagulant was the key parameter that controlled the response parameters. Plant no. 1 confirmed that the amount of coagulant was the key to clarification. The raw water of plant no. 2 was different, and the effect of coagulant dose on clarification was small. The results were consistent with the research by Edzwald.²⁸ Experiments were conducted at plant no. 3 in winter, and the quality of the raw water was very similar to that of the

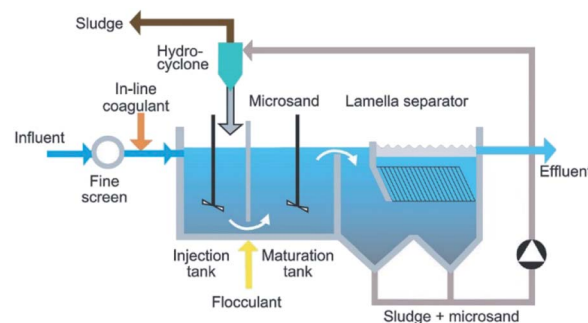


Fig. 1 Schematic of a ballasted sand flocculation process.²⁰



water in plant no. 1. Considering the cold water conditions, the hydraulic load rate was increased by 10 m h^{-1} compared to that of plant no. 1. The effects of coagulation time and precipitation time were investigated, and the results showed that as the reaction kinetics decreased with decreasing temperature, a longer contact time favored floc growth in cold water, excessive microsand (7 mg L^{-1}) caused erosion of the flocs, and a low microsand dose (1 mg L^{-1}) caused poor flocculation. James²⁹ found that the sedimentation velocity of discrete microsand particles and discrete ballast flocs fit the traditional sedimentation theory with reasonable accuracy. The key to increasing the sedimentation rate is to increase the size and roundness of the floc³⁰ and reduce its shape factor. The most important factors contributing to the high settling velocities of ballasted flocs seem to be the increased floc density. Zhu³¹ used alum and anionic polymers as the main coagulants and used silica sand as a carrier. The settleability of the silica sand was improved approximately five times, and the removal efficiency of suspended solids and BOD was 98% and 60%, respectively. Sumant Kumar²⁰ summarized the combined sewer overflow (CSO) treatment method. Depending on the type and characteristics of the water/wastewater, the amount of flocculant (cation/anionic polymer) ranged from 0.3 to 1 mg L^{-1} . The size was 40 – $150 \text{ }\mu\text{m}$, and the dose was 2 – 12 g L^{-1} . In most cases, the turbidity, total suspended solids (TSS) and TP removal rates were reported to be over 90%, while the BOD was over 60%. On this basis, researchers have conducted further research. SIELIECHI²³ proved that the root cause of the improved flocculation of sedimentation in ballast aggregates is related not only to the increase in the specific gravity of flocs but also to the increase in the specific gravity of the ballast. The sediment volume is relatively small in the CSO treatment, which proves the reorganization of flocs. Alan²⁶ proposed a new method to predict the settling conditions encountered by concentrated suspensions containing sand and mud floc mixtures. The results showed that by considering the total reflux effect of the two fractions, the best results can be provided. A frame model was used to predict hindered settlement conditions in various sand and mud mixtures. In addition, there are many literature data showing the improvement of the overall coagulation efficiency of microsand-enhanced flocculation.^{21,31,32}

The BSF process has been widely used in many countries, and the advantages of the BSF process are as follows:²⁰ (1) the floor area is reduced by 5–20 times, which reduces the cost of equipment installation and operation; (2) its adaptability is very high, the treatment effect does not change with the change in raw water conditions, and it can be applied to surface water, wastewater and industrial sewage; and (3) it only takes 20–30 min for the system to reach stable operations. The limitations of the system are as follows: (1) the chemical reagent dose is increased; (2) the microsand will adversely affect the operation and maintenance of the pump; and (3) there is a low removal rate during startup. The BSF process has proven to be a promising technology that can be used to clarify drinking water and reduce pollution loads before wastewater and COS treatment systems enter biological processes or natural water systems, and it has a small footprint and low cost. However, in

some applications, it may be better to replace microsand with a different core medium: (1) nuclei media with a higher density than that of microsand can increase the specific gravity and sedimentation speed of flocs and achieve a high rate clarification; (2) nuclei media with adsorption–desorption capabilities can simultaneously achieve a high rate of clarification and adsorption of pollutants;³³ (3) dolomite as a nuclei medium can increase the alkalinity of water while increasing the sedimentation speed;³⁴ and (4) low-density nuclei media can reduce the mixing strength, thereby limiting the shear force of sensitive flocs. Lapointe²⁵ evaluated the effects of 5 nuclei media (anthracite, recycled broken glass, conventional silica sand, garnet sand, and magnetite sand) on ballast flocculation. The results show that the choice of core media is determined by the sedimentation water quality and the highest applicable flow rate. The highest applicable flow rate can directly affect operating costs.

2.3 Magnetic technology

To achieve separation, magnetic separation technology affects the aggregation of particles through various external forces, such as magnetic gravity, van der Waals forces, an electric double layer, gravity, Brownian motion and fluid motion. Ferromagnetic materials have been widely used in magnetic separation processes^{35–37} and successfully applied in the separation and recovery of solid waste, nanotechnology, ultra-purification and biotechnology.^{38–41} CoMag technology is a new water treatment technology combining conventional coagulation and magnetic separation. On the basis of conventional coagulation, magnetic seeds are used as flocculated nodules and they work to combine flocs to form composite flocs. Under the action of an external magnetic field, precipitation is accelerated to improve the treatment efficiency.⁴² Magnetic separation is the core part of CoMag technology. The first patent for magnetic separation was proposed by William Fullarton in 1792 and applied to the beneficiation industry.⁴³ In the late 1960s, the Massachusetts Institute of Technology developed the first set of magnetic separation equipment and proposed an iron-free, high-gradient magnetic separation patent. In the late 1980s, Sweden began to apply magnetic disk separation to steel wastewater treatment, thus marking the beginning of magnetic technology in the field of water treatment.

In recent years, the CoMag process has been widely used in water treatment due to its advantages, such as a short solid–liquid separation time and good running effect. Li³⁸ used magnetite (Fe_3O_4) for magnetic seed flocculation combined with high-gradient superconducting magnetic separation (HGMS) to remove arsenic. Under the conditions of a polymeric ferric sulfate (PFS) dose as a flocculant of 108.75 mg L^{-1} and a seeding time of 15 min, the efficiency of removing arsenic reached 94.5%. Studies have shown that ultrafine magnetic particles have low separation efficiency and recovery due to low magnetic and collision efficiency issues. Many techniques induce ultrafine particles to form aggregates by increasing the effective particle size and mass.^{41,44,45} Mandel⁴⁶ used magnetic





Table 1 Application of ballast flocculation

Raw water	Ballast agent	Coagulation	Polymer	Ballast agent dose (g L^{-1})	Particle size	Coagulation dose (mg L^{-1})	pH	Stirring intensity (rpm)	Observation	Reference
Surface water	Microsand	$\text{Al}_2(\text{SO}_4)_3$	PASS-100	5	<160 mm	6.1–9.3	6		Filterability is more sensitive to coagulant and microsand dosage	17
Combined sewer overflow	Silica sand	Alum	A-3330	3	74–250 μm	9.7–17.8			Suspended solids and BOD removal efficiencies of >98% and >60%	31
Microalgae	Flocculated biomass	FeCl_3	PAA			50		200	Flocculation efficiency achieved approximately 90%	32
Synthetic water	Pozzolana particles	FeCl_3		0.5			6	250	Describes the turbulent viscous effect of floc compaction	23
Distillery wastewater	Activated carbon	FeCl_3		1.4		840	7	3–250	Precipitation rate increased by 8 times	18
Synthetic water As(v)	Magnetite	PFS	CPA	0.3		108.75		200	The removal efficiency of arsenic was more than 90.9%	38
Backside grinding wastewater	$\text{FeO} \cdot \text{Fe}_2\text{O}_3$	PAC		1.24	310–450 nm	10	5	30–100	PAC use was reduced sixfold	19
Synthetic water	MNPs	$\text{Al}_2(\text{SO}_4)_3$		0.1–5			7	20–100	Significant removal of dissolved organic matter (98.7%)	35

nanocomposite particles as the flocculation core of iron hydroxide and protected its surface with silicon dioxide to avoid its dissolution, which is conducive to recycling. Under the action of external magnetic fields, flocs adsorb and trap heavy metal ions. Shuang⁴⁷ prepared a new quaternized magnetic resin, NDMP, to study its adsorption behavior in dye wastewater. The results showed that NDMP has good adsorption, regeneration and separation and is an efficient adsorbent. Many researchers have also synthesized various types of magnetic particles for removing algae from lakes.^{48–50} The interaction of forces in the flocculation process is one of the important factors controlling the flocculation effect. Under the action of an external magnetic field, when a single particle approaches another particle, the external force on the particle changes (such as with gravity, van der Waals forces, magnetic attraction, etc.). The size of the external force depends on various parameters, such as the particle size, particle density, magnetic susceptibility and magnetic field. Therefore, a single study of the effects of individual forces is needed. Luo⁵¹ summarized the effects of various forces in magnetic flocculation and conducted in-depth discussions on the factors of magnetic flocculation stability, such as collision efficiency, flocculation frequency, flocculation potential energy, and flocculation kinetics.

Magnetic flocculation can be used for separation and has applications in improving the solid coagulation rate, producing clean water, increasing the amount of filtration and reducing the amount of flocculant added. Magnetic flocculation has many advantages, such as no chemical use, fast separation, low operating costs, relatively few ecological problems, space savings, etc.^{52,53} The applications of ballast flocculation are summarized in Table 1.

3. Chemical oxidation-enhanced coagulation

The study of chemical oxidation-enhanced coagulation is one of the hotspots in coagulation treatment research. The main goals of scholars are to reduce the toxicity of traditional oxidation and improve the efficiency of coagulation removal. Potassium permanganate preoxidation as a simple enhanced coagulation method that has been successfully applied in countries around the world.^{54–56} There are an increasing number of preoxidant coagulation agents, such as ferrate, ozone, permanganate complex agents, persulfate, Fenton reagent and sodium hypochlorite.^{57–60}

3.1 Technical principle

Although the mechanism of preoxidation for improving coagulation is different for different oxidants, the progress of preoxidation is usually aimed at changing the zeta potential by destroying the organic coating on the surface of the particles, which results in colloids and suspended particle aggregates, as shown in Fig. 2.⁶¹ For highly toxic and difficult-to-degrade wastewater, the combination of one or more oxidation methods is also an approach,^{62–64} and this is described in detail below.

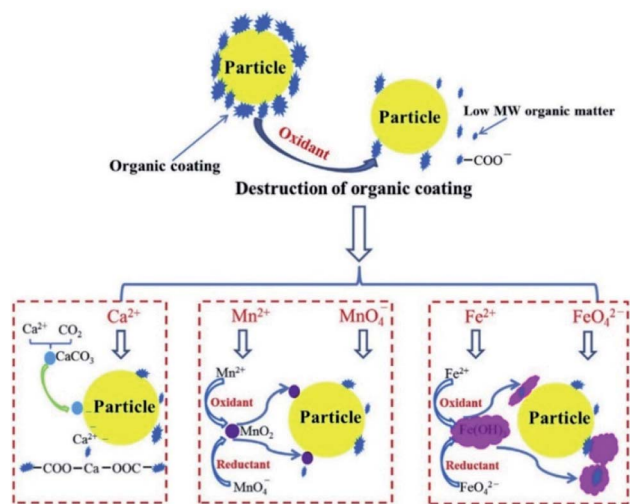


Fig. 2 Mechanisms of peroxidation to enhance coagulation.⁶¹

3.2 Chlorine and chlorine dioxide

Chlorine is a disinfectant. Chlorine oxidation of slightly contaminated water can effectively remove the color and taste of water. Chlorination has little use for enhancing coagulation, but chlorine pre-oxidation can effectively enhance coagulation and improve the removal of algae cells.^{65,66} Chlorine dioxide for potable water application, chlorine dioxide is commonly generated from sodium chlorite reacting with gaseous chlorine, hypochlorous acid, or hydrochloric acid. Chlorine dioxide pretreatment, like chlorine, has a positive effect on algae removal.⁶⁷ Lin⁶⁸ found that ClO_2 dosing had a strongest ability to lower algae viability and oxidize ionic and complexed soluble Mn. Shi⁶⁹ found that the optimal dose of 0.5 mg L^{-1} of Cl_2 could prevent the outbreak of *Pseudomonas aeruginosa*. The pre-oxidation of ClO_2 will destroy the cell walls of algae, leading to severe degradation of membrane glycoproteins or certain amino acids;⁷⁰ the reduction of cell integrity by ClO_2 pre-oxidation significantly improves the removal of algae by the coagulation precipitation method. However, the pre-oxidation of chlorine dioxide only has a limited effect when treating surface water that contains little or no algae. Alam⁷¹ used 0.6 and 1.0 mg L^{-1} of chlorine dioxide for pre-oxidation in the drinking water process and found no significant improvement in drinking water quality. In addition, NaOCl possesses the ability to penetrate the cell membrane of *Microcystis aeruginosa* causing the release of intracellular organic matter (IOM) and potassium ions (K^+) from the cells, thereby destroying cell chemosphere and enhancing the removal of the algae by coagulation–sedimentation.⁶⁵

3.3 Ozone

Ozone is a powerful disinfectant and oxidant, and it has been used in water treatment facilities in Europe since 1906.⁷² The main uses of ozone are to disinfect, decolorize, inactivate algae and reduce inorganic substances.⁷³ Advantages of ozone in water treatment include the following:⁷⁴ (a) ozone kills chlorine-

resistant viruses and spores; (b) it is scarcely affected by sewage pH and temperature; (c) it removes color, odor, and phenols from sewage, increases the dissolved oxygen content and improves water quality; (d) it can improve the biodegradability of pollutants; and (e) it is difficult to degrade and will not cause secondary pollution due to residues. Therefore, it has been widely studied and applied. Studies have shown that moderate amounts of ozone (low doses) enhance coagulation.⁷⁵ Several explanations have been proposed:^{10,76–79} (1) ozone oxidation can increase the concentration of oxygenated functional groups on solid surfaces (such as carboxylic acids), leading to complexation with cations (such as Ca^{2+} , Mg^{2+} , Al^{3+} , etc.) and formation of metal humic acid complex precipitates; (2) ozone can convert high-molecular-weight chemicals into low-molecular-weight chemicals, reducing space and electrostatic repulsion, and ozonation reduces the stabilization of organics that cover the surface of particles, which results in their desorption; (3) ozonation polymerizes metastable organics, which results in particle aggregation through bridging reactions; (4) ozonation changes the surface chemistry of organics and increases coagulation *via* charge neutralization; and (5) ozone oxidation ruptures organometallic complexes, resulting in the *in situ* production of coagulants.

Ozone peroxidation has been widely applied to enhance coagulation⁷⁵ and is often used to treat NOM, algae, heavy metal ions, humus and dye wastewater.^{77,80,81} NOM is adsorbed on the surface of minerals. Studies have shown that ozone can induce the accumulation of NOM. However, due to the composition of NOM and the complexity of the ozone chemical reaction, it is difficult to develop accurate models. A large number of studies have shown that preozonation has a more significant effect on the coagulation of particles in water of medium hardness than on the coagulation of particles in soft water.^{77,79} Chandrakanth⁷⁸ studied the mechanism of ozone removal of NOM, and the results show that ozone can cause particle instability only in the presence of calcium. A theory based on electrostatic (complexation/charge neutralization) and calcium oxalate precipitation was proposed: (a) the components of ozonated NOM can complex with calcium, and (b) the ozonation of NOM increases the number of coordination sites on the surface and the complexation of NOM with calcium is relatively easy. Many studies have shown the biggest influencing factors that affect the efficiency of ozonation coagulation: the characteristics of the raw water (hardness, pH, alkalinity, and type and concentration of NOM), and ozone dose and coagulation conditions (dose and type of coagulant).^{9,82,83} Francisco's⁷⁹ research showed that the effect of the efficiency of preozonation-enhanced coagulation seems to vary depending on the characteristics of the raw water, especially the calcium hardness and NOM fractionation. The ozone dose was $0.25\text{--}2.5 \text{ mg L}^{-1}$, and the THMFP removal rate was $5\text{--}25\%$. Smith⁹ developed a study of preozonation-enhanced electroflocculation for the treatment of municipal wastewater. The results showed that the technology could effectively reduce the total phosphorus and TSS from water sources, such as highly toxic textile wastewater and landfill leachate. Preozonation can be used in combination with other oxidation methods (Fenton, permanganate, etc.).⁶⁴ The



application of preozonation to enhance coagulation is very wide. Table 2 describes the application of ozonation in natural water and sewage treatment and the corresponding results.

In most cases, only low amounts of ozone can improve the coagulation effect and high doses of ozone may increase the turbidity of the final water. The optimal dose of ozone is always closely related to the charge density of the humus. The higher the charge density of humus is, the higher the required dose of ozonation.⁸⁴ Therefore, for some raw water, ozone can always find its optimal dose to enhance coagulation.

3.4 Permanganate

Permanganate shows high reactivity in oxidizing a wide variety of organic substances and inorganic substances. The main uses of permanganate are to control odor and taste, remove iron, manganese, and algae and act as a disinfectant.^{62,85,86} The principle of permanganate preoxidation to enhance coagulation is as follows. Permanganate is oxidized to produce manganese dioxide. Manganese dioxide is a strong adsorbent. It can increase the flocculation kinetics by increasing the particle concentration. It can also be adsorbed on the surface of other particles to enhance flocculation.^{87–89} Permanganate preoxidation can effectively remove heavy metals and organics in water.^{87,90,91}

Over the past three decades, permanganate preoxidation-enhanced coagulation has attracted increasing attention, especially in the field of surface water treatment.^{92–94} Cyanobacteria in surface water treatment can produce metabolites or

toxins that are difficult to remove by conventional water treatment processes. Therefore, preoxidation technologies are widely used, such as chlorine, ozone, chlorine dioxide, and permanganate. To control indicators, such as water taste, odor and toxin content,⁹⁵ and because chlorine gas and chlorine dioxide can produce disinfection byproducts (DBP), permanganate preoxidation technology is widely used. Ma⁹⁶ reported that permanganate can significantly enhance the coagulation of several surface waters. Studies show that the enhanced coagulation of permanganate is the formation of the strong adsorbent manganese dioxide, which adsorbs onto particle surfaces. Colthurst⁹⁷ found that manganese dioxide adsorbs humus only when Ca^{2+} is present, which may be due to the negative charge of both. Chen⁸⁷ studied the mechanism of permanganate preoxidation to enhance coagulation. With increasing calcium ion concentration, the sedimentation speed was significantly accelerated, and calcium ions promoted permanganate function by electrically neutralizing surface bridging. In addition to permanganate incorporation of algae flocs to increase sedimentation speed, the use of permanganate may also induce algal cells to release organic matter.¹⁰⁴ As shown in Fig. 3, SEM observations showed that EOM may enhance the combination of algae flocs and manganese dioxide. Studies have found that the flocculation efficiency of severely damaged algal cells is very low.¹⁰⁵ Takaara¹⁰⁶ reported that materials with too high of a molecular weight will adhere to the coagulant, causing its failure, which may also reduce the coagulation efficiency. In addition, permanganate is used at a common dose (0–2 mg L^{-1}); thus, there is little or no damage to algae. Wang⁹⁴

Table 2 Application of pre-ozone oxidation to treat water

Raw water	Coagulant	Dose (mg L^{-1})	Observation	Reference
Disperse dye-bath	FeSO_4	2300	Eliminated 95% color and 48% COD	98
Saline landfill leachates	$\text{Fe}_2(\text{SO}_4)_3$	49	Leachate treatment led to 100% removal of leachate color and 78% removal of COD	99
Textile wastewater	$\text{Al}_2(\text{SO}_4)_3$, FeSO_4	53.7–82.3	Almost complete color absorbance (over 98%) was removed in 20 min ozone contact time	80
Taihu Lake	Alum	0.45	THMFP removal by approximately 60%	100
Oilfield-produced water	PAC, PAM	5	PAC and PAM could be saved by approximately 10 mg L^{-1} and 5 mg L^{-1}	101
Waterworks	PAC	0.5	Improve THMFP coagulation removal by 14%	73
Municipal wastewater	PACl	0.8	DON, DOC, and UV254 removal of 71%, 66%, and 97%	10
<i>Chodatella</i> sp.	Alum	0–2.1	Increasing algae removal by approximately 50%	88
<i>Oscillatoria agardii</i>	$\text{Al}_2(\text{SO}_4)_3$	0–2	Increasing algae removal by 8%	102
Green algae mixture	FeCl_3	1	Increasing algae removal by approximately 20%	103



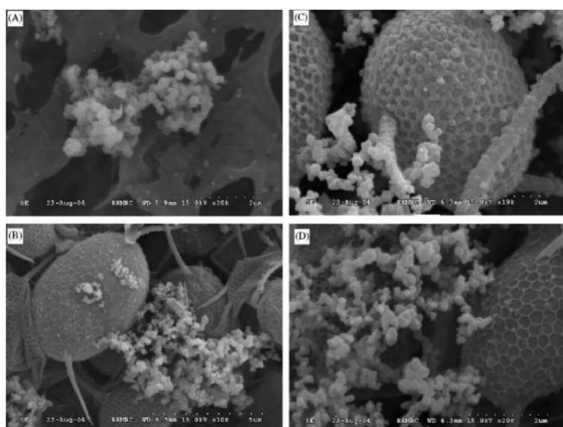


Fig. 3 SEM micrographs of algal cell surface morphology with and without preoxidation. (A) Preformed MnO_2 , (B) preformed MnO_2 adsorbed onto algal cell surface, (C) KMnO_4 preoxidized algae cell, and (D) EOM elimination due to trypsin dosing after KMnO_4 preoxidation.⁸⁷

studied the damage of the dose of permanganate to algal cells and observed the cell morphology at different potassium permanganate concentrations using stereoscopic scanning camera technology. The potassium permanganate concentrations were 1 mg L^{-1} , 2 mg L^{-1} and 6 mg L^{-1} . As shown in Fig. 4, an appropriate concentration of potassium permanganate (for example, 2 mg L^{-1}) promoted EOM secretion by cells. Chen⁸⁷ showed that EOM may improve the combination of algae flocs and manganese dioxide and achieve the purpose of enhanced coagulation. Many studies have also proven that the commonly used dose of high-acid salt oxidation ($0\text{--}2 \text{ mg L}^{-1}$) does little or no damage to algae.^{60,107–109} Excessive manganese will affect the quality of the effluent water. Usually, excess manganese dioxide is removed through a precipitation tank during the water treatment process, although it increases the burden on the filter and shortens the backwash cycle. The characteristics of the raw water need to be studied to ensure the complete conversion of the peracid salt into manganese dioxide. The disadvantages of permanganate preoxidation are overcome to enhance coagulation to the greatest extent.

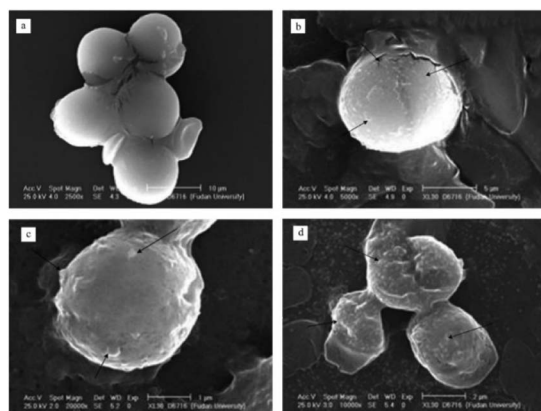


Fig. 4 Stereoscan photograph of algal cell influenced by different concentrations of KMnO_4 . (a) *Protococcus*; (b) KMnO_4 1 mg L^{-1} ; (c) KMnO_4 2 mg L^{-1} ; and (d) KMnO_4 6 mg L^{-1} .⁹⁴

3.5 Ferrate

Ferrate is a chemical reagent (coagulant and oxidant) with dual functions in drinking water treatment and has great potential in the treatment of water and wastewater.¹¹⁰ It has a strong redox potential across the entire pH range, from 2.2 V in acidic conditions to 0.7 V in alkaline conditions.^{110,111} During the reduction of ferrate, with the precipitation of ferric hydroxide, hydrolyzed substances with a high positive charge may be produced. These hydrolysates will lead to colloids, cause the particles to be electrically neutralized, and adsorb water molecules and fine particles, thus increasing the sedimentation speed of the coagulation.^{110,112,113} Ferrate can destabilize colloidal particles in one minute, while ferrous and iron salts need 30 min to achieve stable results.¹¹⁰ Ferrate is an emerging multifunctional oxidant, disinfectant and coagulant. Ferrate does not produce any mutagenic byproducts in water treatment. It is an ideal water and wastewater treatment material in the 21st century.^{111,114–116}

Research has shown that ferrate can effectively remove many organic and inorganic pollutants, such as nitrogenous organics, drugs, antibiotics, hydrogen sulfide, algae, persistent organics, *etc.*^{112,113,117–123} Ferrate can also remove various metals, such as Fe^{2+} , Fe^{3+} , Mn^{2+} , Cu^{2+} , Pb^{2+} , Cd^{2+} , Cr^{3+} , Hg^{2+} , *etc.*^{117,124} Jiang evaluated the application and preparation of ferrate, and the results showed that ferrate was superior to iron sulfate under low-dose conditions when treating colored water. Ferrate can effectively reduce the UV250 and turbidity of water. Dissolved organic carbon can also reduce the formation of trihalomethane (THMFP).^{119,125,126} In terms of wastewater treatment,¹²⁰ compared with the same doses of AS and FS, potassium ferrate showed a 50% increase in color removal rate and a 30% increase in COD, resulting in less sludge. Ma^{112,113} used ferrate preoxidation to improve coagulation and treat algae-containing water and surface water. Studies have found that ferrate preoxidation can enhance the removal and turbidity of algal cells at various oxidant doses. In terms of synthetic preparation, Stanford and Jiang¹²⁷ electrochemically synthesized potassium ferrate and showed that 1 mg L^{-1} potassium ferrate and 10 mg L^{-1} iron salt had the same treatment effect. Alsheyab and Jiang designed an electrochemical method for a Chinese factory. The reactor proved the operation concept and online application of the electrochemical generation of potassium ferrate, which is a step forward for the comprehensive application of potassium ferrate in water and wastewater. The decomposition rate of ferrate depends on the pH value, solution temperature, initial potassium ferrate concentration and coexisting ions. Before using ferrate, the characteristics and quality of raw water need to be analyzed.^{110,128}

4. Other enhanced coagulation methods

4.1 Ultrasound-enhanced coagulation

In recent years, ultrasound has proven to be an effective water treatment technology.^{129,130} Ultrasound irradiation produces cavitation, thereby causing reactive free radicals and shock



waves.^{131,132} Studies have shown that ultrasound combined with other technologies has better results than ultrasound alone.^{133–135} Ultrasound-enhanced coagulation is one of the current research hotspots, and the working principle is as follows:¹³⁶ (1) low-frequency (20–150 kHz) ultrasonic irradiation will increase molecular vibrations, improve the ion collision efficiency and make the flocs more compact and easy to precipitate; and (2) ultrasound irradiation can change the morphology and surface characteristics of ions, cause particle surface damage, significantly increase specific surface area, and increase particle aggregation. A large number of studies have shown that ultrasonic treatment can significantly increase the flocculating capacity of algae in the treatment of algae-containing water, reduce the amount of coagulant added, and reduce costs.¹³⁷

In algae-containing water, it is difficult to remove algal cells due to their special characteristics (negative surface potential), and algae metabolites tend to adsorb to colloidal particles. Zhang¹³³ reported the removal of algal cells from the water source in a water treatment plant by ultrasound-enhanced coagulation. Studies have shown that ultrasound irradiation applied for 5 seconds can increase the desiccation efficiency by 20%. Zhang¹³⁴ reported that the main theory involved the destruction during ultrasonic irradiation of gas vacuoles inside algal cells that acted as “nuclei” for acoustic cavitation and collapsed during the “bubble crushing” period, causing the settlement of cyanobacteria, whereas the ultrasound frequency had little impact on the removal effect. However, studies have shown that increasing ultrasound duration and power can negatively affect coagulation,¹³⁸ such as by causing microcystin release.¹³⁷ Therefore, the parameters of ultrasound-enhanced coagulation are very important, as shown in Table 3. Li¹³⁹ reported the effects of ultrasonic power, action time, coagulant dose and pH value on the removal of organic matter in low-temperature and low-turbidity water. The results showed that the optimal ultrasound power was 25 W, the optimal ultrasound treatment time was 9 min, the optimal PFS dose was 12 mg L⁻¹, the optimal pH was 7, and the COD removal rate was 87.2%. Fast¹⁴⁰ reported the effects of several different process

parameters, such as chemical dose, ultrasonic time, pH value, concentration factor, and flocculation time, on algae removal. A comparison between ultrasound-alum and ultrasound-chitosan flocculation of turbid algal waters as well as a comparison with conventional rapid mix and direct flocculation were conducted. Three processes were designed on the basis of single-factor experiments. The results showed that the removal rate of the ultrasonic-chitosan-enhanced coagulation process could reach more than 98%.

The ultrasonic enhanced coagulation process has attracted much attention in the treatment of algae-containing water, although the current research is still very unsystematic, and it is often limited to static ultrasonic experiments at a single frequency. It lacks multi-frequency, multi-intensity and multi-time comparisons. The actual application is still far away. In the future, it is necessary to further strengthen the research on the influence of the interaction of various parameters, clarify the mechanism of ultrasound enhancement, and lay the foundation for future applications.

4.2 Composite coagulants

4.2.1 Inorganic polymer flocculants. Aluminum salts and iron salts have been widely used as inorganic flocculants in water and wastewater treatment. The main products are aluminum sulfate, ferrous sulfate, iron sulfate, polyaluminum chloride (PAC) and PFS. The hydrolyzed polymerization products of these products have a positive charge, and through the adsorption and neutralization sweeping effect, they condense with the negatively charged colloidal pollutants in the water, although their ability to adsorb and bridge the colloidal pollutants in the water is weak. A higher dose is required to achieve a good flocculation effect.^{2,142} To further improve the flocculation effect of inorganic flocculants and reduce the amount of products, a large amount of research and application practice found that iron salt and aluminum salt are copolymerized under certain conditions or an inorganic component with flocculation or coagulation aid is added to the iron salt, aluminum salt or iron salt/aluminum salt solution and reacted. Through synergistic effects, inorganic–inorganic composite

Table 3 Application of ultrasound-enhanced coagulation

Raw water	Coagulant	Ultrasonic irradiation	Ultrasonic power	Observation	Reference
Algae-containing water	Chitosan	<60 s	100 W	Higher removal efficiencies (>97%)	140
Luan River	FeCl ₃	15 s	60 W	Improve algae coagulation removal by 12.4%	137
Low-temperature and low-turbidity water	PFS	9 min	25 W	COD removal rate will reach 87.2%	139
Algae-containing water	PAC	5 min	0.02 W mL ⁻¹	Turbidity removal rate of approximately 80% to 90%	141
Algae-containing water	PACl	30 s	<100 W	Dosage of PACl was reduced, and the sedimentation rate of flocs was increased	136
Algae-containing water	PAC	1–5 s	48 W	Pre-sonication reduced the coagulant dose by two-thirds	133
Algae-containing water	PAC	5 s	47.2 W cm ⁻²	Increased algae removal efficiency from 35% to 67%	134



flocculant products produce better flocculation effects.^{143–145} Because the composite flocculant can overcome many of the shortcomings of using a single flocculant, it can improve the flocculation performance while reducing the cost of water treatment. Therefore, the development and application of composite flocculants has become one of the issues of interest in the current water and wastewater treatment field, and it is also the main development direction of new, efficient and economical flocculants. An early composite flocculant was polyaluminum plus polyacrylamide¹⁴⁶ and then polyaluminum plus activated silicic acid was used^{147,148} in an attempt to combine an original coagulant and a new coagulant into one dose. With a more in-depth understanding of the principle of flocculants, it was gradually realized that the combination of flocculants is based on the principle of strengthening complementarity, combining flocculants with different flocculating properties, and improving the performance of a single coagulant. At present, there are two main types of composite coagulants: inorganic composite types and inorganic–organic composite types. For example, in polyaluminum chloride composite flocculants,¹⁴⁵ the active component is still aluminum hydrolysis and polymerization to form high-valent hydroxyl polymer ions. By adding an organic flocculant, the inorganic flocculant enhances its ability to adsorb to particles and bridge and neutralize electrical charges, thereby improving its coagulation ability. Inorganic polymer composite flocculants are mainly prepared by adding inorganic salt compounds,^{143,145} such as Al^{3+} , Fe^{3+} , Ca^{2+} , SO_4^{2-} , Cl^- , *etc.* According to the type of metal ions that the composite flocculant is associated with in flocculation, the inorganic polymer composite flocculant can be divided into the polyaluminum salt type, polyiron salt type and polyaluminum iron salt type.^{144,149–151} One or several kinds of them are added to form inorganic composite polymer flocculants, fully optimizing their respective advantages and achieving the purpose of efficient purification treatment. At present, a variety of inorganic–inorganic and inorganic–organic composite flocculant products have been developed, mainly including polyaluminum chloride sulfate (PACS), polyaluminum ferric chloride (PAFC), polyaluminum silicate sulfate

(PASiS), polyaluminum silicate chloride (PASiC), polyferric silicate sulfate (PFSiS), polyaluminum ferric silicate chloride (PAFSiC), PAC/PDM composite coagulants and PAC/PDMAAC composite coagulants.^{143,144,149,152–155} In recent years, many researchers^{156–158} have also enhanced coagulation by improving the performance of coagulant aids. Table 4 summarizes the application of composite flocculants in water treatment.

The research on composite flocculants has developed rapidly in recent years. Based on the research results of its performance and flocculation mechanism, a series of new and efficient inorganic–inorganic and inorganic–organic composite polymer flocculants have been developed. It is used in water and wastewater treatment and sludge dewatering, although the research, development and application of composite high-resolution flocculants are still in their infancy. At present, the research and development of composite flocculants is mainly focused on the development of new, efficient, nontoxic or low-toxicity composite flocculants. This work should be strengthened in the following ways.

(1) Further strengthen research on compound polymer flocculant ratio screening, preparation process technology, process parameters and supporting equipment to improve the flocculation effect and reduce the cost.

(2) Carry out in-depth and systematic research on the interaction between the components of the composite flocculant and strengthen the synergy between the component, thereby laying a theoretical foundation for the development of new flocculants.

(3) Carry out in-depth and systematic research on the performance of composite flocculants, the dynamic process of flocculation, and the physical characteristics of the formed flocs. The results of the research are helpful for clarifying the flocculation behavior and mechanism of composite polymer flocculants.

(4) Systematic research is performed on the application conditions and influencing factors of composite flocculants to clarify the applicable objects and lay the foundation for the promotion of new products.

Table 4 Application of composite flocculants in water treatment

Raw water	Coagulation	Materials	Dose (mg L ⁻¹)	Observation	Reference
Kaolinite suspensions	P(DADMAC-AM-VTMS)	DADMAC + AM + VTMS	0.1	<5 NTU (turbidity)	159
Kaolinite suspensions	PAFC	Coal gangue + HCl + CaCO ₃	0.05	<5 NTU (turbidity)	149
Kaolinite suspensions	PACS	Na ₂ CO ₃ + AlCl ₃ + Al ₂ (SO ₄) ₃	2	<4 NTU (turbidity)	143
Synthetic suspensions	PASiC	SiO ₂ + AlCl ₃ + NaOH	3	<1 NTU (turbidity)	144
Oil field wastewater	PAC–PDMDAAC	PDMDAAC + PAC	8	<100 (COD)	155
Landfill leachate	PFSiS	PFSiS + Fe ₂ (SO ₄) ₃	50	>90% (turbidity removal)	150
Lake Taihu waters	PAC–PDM	PAC + PDM	8.32	0.41 NUT (turbidity)	153
Oily wastewater	PFASS	Fly ash	140	>95% (turbidity removal)	151
Kaolinite suspensions	CMC-g-PAM	CMC + CAN + PAM	15	>80% (turbidity removal)	160
Synthetic suspensions	PTSS	PTS + PSiA	35	70% (UV254 removal)	161
Taihu Lake	PAC–PDMDAAC	PAC + PDMDAAC	10	<1.5 NTU (turbidity)	154
Cd(II) solution	APAM	PAM	2.87	93.65% (Cd(II) removal)	146
Synthetic textile wastewater	PAFC-starch-g-p(AM-DMDAAC)	PAFC + starch + AM + DMDAAC	0.2	>85% (dye removal)	162



4.2.2 Biopolymer flocculants. Natural polysaccharides derived from biomass feedstocks, marine resources, and microorganisms have been attracting considerable attention as benign and environmentally friendly substitutes for synthetic polymeric products. There are many types of biopolymer flocculants, such as lignin, cellulose, chitosan, xanthan gum, pectin, polysaccharides and proteins, *etc.*^{163,164} Among them, cellulose and chitosan are currently the most abundant biopolymers on Earth.¹⁶⁵ Biopolymer flocculants are derived from oceans, plants, and microorganisms. For example, chitosan mostly comes from seaweeds, crustacean shells and microorganisms, and cellulose, starch, *etc.* come from agricultural raw materials. Microorganisms, such as bacteria, yeast and algae, are important sources of biopolymer flocculants.¹⁶⁶ Biopolymers have many advantages because they are environmentally friendly, highly biochemical, nontoxic, *etc.* and can be used in agricultural wastewater,¹⁶⁷ oily wastewater,¹⁶⁸ heavy metal wastewater,¹⁶⁹ algal water, *etc.*^{170,171} This research field has received extensive attention. Turunen¹⁶⁷ used tannin and chitosan to treat agricultural wastewater. The tannin and chitosan coagulant performed best at a dose of 5–10 mL L⁻¹, with a turbidity removal rate of 80–98%. Henry¹⁶³ studied the removal of orthophosphate by a variable ternary system of alum, biopolymer and biomass. The removal rate of orthophosphate was 98%. Lichtfouse¹⁷² studied the modified chitosan treatment of phosphate solution, and the principle is shown in Fig. 5. This treatment could reduce the phosphate content by 97% under all pH values. Biopolymer flocculants usually exhibit low flocculation activity. Many researchers use chemical modification to improve their flocculation activity. There are two main forms:¹⁶⁶ (1) physical reactions, such as ionic interaction, polysaccharide–protein interaction, and (2) chemical reactions, such as graft copolymerization, crosslinking with aldehyde, esterification, etherification, amination, carboxyalkylation, hydroxyalkylation and other addition reactions, and

condensation. Peng¹⁷³ studied a series of amphoteric chitosan-based grafting flocculants (CM-chi)-*g*-PDMDAAC (denoted as CgPD), which were successfully synthesized by grafting diallyl dimethyl ammonium chloride (DMDAAC) onto carboxymethyl chitosan (CM-chi) with different grafting ratios. By carboxymethyl and grafting modification, dramatically increased water solubility of chitosan was obtained. Many biopolymer flocculants have been synthesized through chemical modification, for example, polysaccharide-*g*-(PDMA-*co*-AA),¹⁷⁴ AP-*g*-poly(AM-*co*-AA),¹⁷⁵ PHAP.¹⁷⁶ The main challenges faced by biopolymer flocculants are the complex synthesis process, monomer health problems, future production scale and consumption. Future research should focus on reducing production costs and developing large-scale production processes.

5. Conclusions

Because of their simple operation, high versatility, low energy consumption and high cost performance, coagulation and flocculation are still essential steps in the field of water treatment. This review summarized the application development of the enhanced coagulation process and outlined the effectiveness of the enhanced coagulation process from the perspective of different enhanced coagulation processes. Ballast reaction nuclei-enhanced coagulation has the potential to treat surface water, wastewater and overflow water from sewers. This method can increase the density of flocs, increase the sedimentation speed of flocs, and improve the fractionation of flocs. Preoxidation and ultrasound are widely used in the treatment of algae-containing waters. Preoxidation enhances coagulation by breaking the organic coating and changing the surface potential of particles. However, the mechanism of oxidative coagulation is completely different for different oxidants and colloids. Ultrasound increases the particle collision efficiency by increasing the molecular vibrations so that the flocs are denser and the sedimentation speed is increased. Composite flocculants were developed based on research on the performance and mechanism of different flocculants, and they represent an optimization and complementation of the action of two or more flocculants. The current research indicates that enhanced coagulation is based on the mechanism and dynamics of traditional coagulation. As long as the coagulation effect is substantially improved, it can be considered enhanced coagulation. In future research, a mathematical model and a stability model for enhanced coagulation should be established, flocculants should be developed, appropriate coagulation nodules should be selected, and different water qualities, preoxidation or ultrasonic methods can be considered to improve raw water flocculation characteristics. Enhanced coagulation is greatly affected by the process parameters. A method for evaluating coagulation and a coagulation process control strategy should be proposed and established to resolve issues at the intersection of coagulation and flocculation stages in the coagulation process during water treatment, as well as coagulation process and coagulant issues such as coordination, mixing conditions, and coagulation process control.

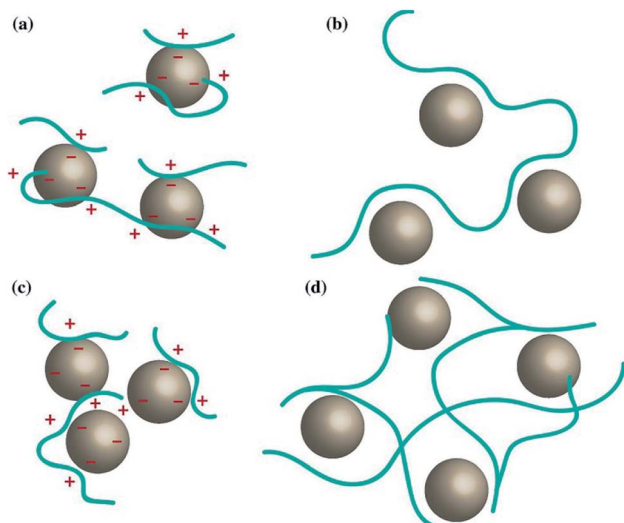


Fig. 5 Bioflocculation mechanism: (a) charge neutralization, (b) bridging, (c) electrostatic patching and (d) sweeping.



Conflicts of interest

Authors declare no competing interest.

Acknowledgements

This study was supported by the Northeast Petroleum University Youth Science Fund Project (2019QNL-30).

References

- J.-Q. Jiang, *Curr. Opin. Chem. Eng.*, 2015, **8**, 36–44.
- C. Y. Teh, P. M. Budiman, K. P. Y. Shak and T. Y. Wu, *Ind. Eng. Chem. Res.*, 2016, **55**, 4363–4389.
- J. K. Edzwald, *Water Sci. Technol.*, 1999, **9**, 63–70.
- S. Xue, Q. Zhao, L. Wei, X. Ma, Y. Wen and Z. Zhang, *Environ. Prog. Sustainable Energy*, 2015, **34**, 751–760.
- J. Yu, D. Wang, M. Yan, C. Ye, M. Yang and X. Ge, *Environ. Monit. Assess.*, 2007, **131**, 377–386.
- M. Yan, D. Wang, J. Qu, J. Ni and C. W. Chow, *Water Res.*, 2008, **42**, 2278–2286.
- M. Yan, D. Wang, S. You, J. Qu and H. Tang, *Water Res.*, 2006, **40**, 3621–3627.
- S. Ma, C. Liu, K. Yang and D. Lin, *Sci. Total Environ.*, 2012, **439**, 123–128.
- K. Nielson and D. W. Smith, *J. Environ. Eng. Sci.*, 2005, **4**, 65–76.
- B. Liu, L. Gu, Q. Li, G. Yu, C. Zhao and H. Zhai, *Environ. Technol.*, 2019, **40**, 2684–2694.
- R. D. Ambashta and M. Sillanpaa, *J. Hazard. Mater.*, 2010, **180**, 38–49.
- W. Yu, M. Liu and N. J. D. Graham, *ACS Sustainable Chem. Eng.*, 2019, **7**, 18566–18573.
- J. M. Younker and M. E. Walsh, *Water Res.*, 2016, **98**, 1–8.
- S. Kadouche, H. Lounici, K. Benaoumeur, N. Drouiche, M. Hadioui and P. Sharrock, *J. Polym. Environ.*, 2012, **20**, 848–857.
- S. Boulaadjoul, H. Zemmouri, Z. Bendjama and N. Drouiche, *Chemosphere*, 2018, **206**, 142–149.
- M. Ma, R. Liu, H. Liu and J. Qu, *Water Res.*, 2012, **46**, 73–81.
- C. Desjardins, B. Koudjonou and R. Desjardins, *Water Res.*, 2002, **30**, 744–754.
- M. Zhang, Y. Chen, L. Xie and Q. Zhou, *Chemosphere*, 2018, **211**, 1054–1064.
- T. J. Wan, S. M. Shen, S. H. Siao, C. F. Huang and C. Y. Cheng, *Water Res.*, 2011, **45**, 6301–6307.
- S. Kumar, N. C. Ghosh and A. A. Kazmi, *Environ. Technol. Rev.*, 2016, **5**, 57–67.
- M. Lapointe and B. Barbeau, *Water Res.*, 2016, **90**, 119–127.
- K. Miyahara, Y. Adachi, K. Nakaishi and M. Ohtsubo, *Colloids Surf.*, 2002, **196**, 87–91.
- J. Sieliechi, B. Lartiges, S. Skali-Lami, J. Kayem and R. Kamga, *Water Res.*, 2016, **105**, 361–369.
- A. J. Manning and D. H. Schoellhamer, *Mar. Geol.*, 2013, **345**, 266–280.
- M. Lapointe, C. Brosseau, Y. Comeau and B. Barbeau, *J. Environ. Eng.*, 2017, **143**(11), 04017071.
- A. Cuthbertson, P. Dong, S. King and P. Davies, *Coastal Engineering*, 2008, **55**, 1197–1208.
- V. Plum, C. P. Dahl, L. Bentsen, C. R. Petersen, L. Napstjert and N. B. Thomsen, *Water Sci. Technol.*, 1998, **37**, 269–275.
- J. E. Van Benschoten and J. K. Edzwald, *Water Res.*, 1990, **24**, 1527–1535.
- J. C. Young and F. G. Edwards, *Water Environ. Res.*, 2003, **75**, 263–272.
- W. Yu, J. Gregory, L. C. Campos and N. Graham, *Water Res.*, 2015, **68**, 119–126.
- W. Zhu, R. Seth and J. Lalman, *Environ. Technol.*, 2007, **28**, 761–770.
- K. V. Gorin, Y. E. Sergeeva, V. V. Butylin, A. V. Komova, V. M. Pojidaev, G. U. Badranova, A. A. Shapovalova, I. A. Konova and P. M. Gotovtsev, *Bioresour. Technol.*, 2015, **193**, 178–184.
- J. T. Mayo, C. Yavuz, S. Yean, L. Cong, H. Shipley, W. Yu, J. Falkner, A. Kan, M. Tomson and V. L. Colvin, *Sci. Technol. Adv. Mater.*, 2007, **8**, 71–75.
- L. Piirtola, B. Hultman, C. Andersson and Y. Lundeberg, *Water Res.*, 1999, **33**, 1799–1804.
- M. Kumari and S. K. Gupta, *J. Cleaner Prod.*, 2020, **244**, 118899.
- A. D. Ebner, J. A. Ritter, H. J. Ploehn, R. L. Kochen and J. D. Navratil, *Sep. Sci. Technol.*, 1999, **34**, 1277–1300.
- J. B. J. Janssen, A. P. Gelder and J. Perenboom, *J. Phys. D: Appl. Phys.*, 1990, **23**, 1447.
- Y. Li, J. Wang, Y. Zhao and Z. Luan, *Sep. Purif. Technol.*, 2010, **73**, 264–270.
- M. Omran, T. Fabritius, A. M. Elmahdy, N. A. Abdel-Khalek, M. El-Aref and A. E.-H. Elmanawi, *Sep. Purif. Technol.*, 2014, **136**, 223–232.
- J. Svoboda and T. Fujita, *Miner. Eng.*, 2003, **16**, 785–792.
- Y. Wang, E. Forssberg, R. J. Pugh and S. Åke Elming, *J. Dispersion Sci. Technol.*, 1995, **16**, 137–154.
- N. Karapinar, *Int. J. Miner. Process.*, 2003, **71**, 45–54.
- C. T. Yavuz, A. Prakash, J. T. Mayo and V. L. Colvin, *Chem. Eng. Sci.*, 2009, **64**, 2510–2521.
- T. Miettinen, J. Ralston and D. Fornasiero, *Miner. Eng.*, 2010, **23**, 420–437.
- R. Sivamohan, *Int. J. Miner. Process.*, 1990, **28**, 247–288.
- K. Mandel, F. Hutter, C. Gellermann and G. SEXTL, *Sep. Purif. Technol.*, 2013, **109**, 144–147.
- C. Shuang, P. Li, A. Li, Q. Zhou, M. Zhang and Y. Zhou, *Water Res.*, 2012, **46**, 4417–4426.
- S.-K. Wang, A. R. Stiles, C. Guo and C.-Z. Liu, *Algal Res.*, 2015, **9**, 178–185.
- Y. R. Hu, C. Guo, L. Xu, F. Wang, S. K. Wang, Z. Hu and C. Z. Liu, *Bioresour. Technol.*, 2014, **158**, 388–391.
- P. Y. Toh, B. W. Ng, C. H. Chong, A. L. Ahmad, J.-W. Yang, C. J. Chieh Derek and J. Lim, *RSC Adv.*, 2014, **4**, 4114–4121.
- L. Luo and A. V. Nguyen, *Sep. Purif. Technol.*, 2017, **172**, 85–99.
- M. Stolarski, C. Eichholz, B. Fuchs and H. Nirschl, *China Particuol.*, 2007, **5**, 145–150.
- E. Carissimi, J. D. Miller and J. Rubio, *Int. J. Miner. Process.*, 2007, **85**, 41–49.



- 54 H. T. Kemp, R. G. Fuller and R. S. Davidson, *J. AWWA*, 1966, **58**, 255–263.
- 55 J. C. Zhou, H. C. Zhao, K. H. Pan and Q. P. Xu, *J. Zhejiang Univ., Sci., B*, 2011, **12**, 156–162.
- 56 J. Gasperi, B. Laborie and V. Rocher, *Chem. Eng. J.*, 2012, **211–212**, 293–301.
- 57 X.-J. Ma and H.-L. Xia, *J. Hazard. Mater.*, 2009, **162**, 386–390.
- 58 S. Barredo-Damas, M. I. Iborra-Clar, A. Bes-Pia, M. I. Alcaina-Miranda, J. A. Mendoza-Roca and A. Iborra-Clar, *Desalination*, 2005, **182**, 267–274.
- 59 J. Zhou, J. Liu, Z. Zhao, W. Peng, F. Cui and Z. Liang, *Chem. Eng. J.*, 2020, **382**, 123012.
- 60 J. Naceradska, M. Pivokonsky, L. Pivokonska, M. Baresova, R. K. Henderson, A. Zamyadi and V. Janda, *Water Res.*, 2017, **114**, 42–49.
- 61 P. Xie, Y. Chen, J. Ma, X. Zhang, J. Zou and Z. Wang, *Chemosphere*, 2016, **155**, 550–563.
- 62 H. Liang, J. Y. Tian, W. J. He, H. D. Han, Z. L. Chen and G. B. Li, *J. Chem. Technol. Biotechnol.*, 2009, **84**, 1229–1233.
- 63 F. Wu, X. Li, Z. Tan, B. Xue, Y. Wang and X. Yan, Research on cellar rainwater of northwest region villages and small towns enhanced coagulation by potassium permanganate combined with powdered activated carbon, *2012 International Symposium on Geomatics for Integrated Water Resource Management, Lanzhou*, 2012, Lanzhou, pp. 1–5.
- 64 S. Meric, H. Selcuk and V. Belgiorno, *Water Res.*, 2005, **39**, 1147–1153.
- 65 M. Ma, R. Liu, H. Liu, J. Qu and W. Jefferson, *Sep. Purif. Technol.*, 2012, **86**, 19–25.
- 66 D. L. Sedlak and U. von Gunten, *Science*, 2011, **331**, 42–43.
- 67 D. Yan, Z. Sun, J. Wang, L. Wang, R. Pan, Q. Wu and X. Liu, *Water*, 2019, **11**, 2302.
- 68 J. L. Lin, L. C. Hua, Y. Wu and C. Huang, *Water Res.*, 2016, **89**, 261–269.
- 69 X. Shi, R. Bi, B. Yuan, X. Liao, Z. Zhou, F. Li and W. Sun, *Sci. Total Environ.*, 2019, **656**, 1063–1070.
- 70 S. Zhou, Y. Shao, N. Gao, L. Li, J. Deng, M. Zhu and S. Zhu, *Sci. Total Environ.*, 2014, **482–483**, 208–213.
- 71 M. Z. B. Alam, R. E. Cantwell, R. Hofmann, R. C. Andrews, J. L. Rand, G. A. Gagnon, M. VanderMarck, E. Moffat and S. A. Andrews, *J. Environ. Eng.*, 2008, **134**, 478–485.
- 72 S. D. Chang and P. C. Singer, *J. - Am. Water Works Assoc.*, 1991, **83**, 71–79.
- 73 Q. F. Liu, X. T. Zhou, Z. D. Li and Z. G. Zhan, *Adv. Mater. Res.*, 2012, **487**, 74–80.
- 74 M. G. Nagarkatti, *J. Environ. Qual.*, 1991, **20**, 881.
- 75 V. Camel and A. Bermond, *Water Res.*, 1998, **32**, 3208–3222.
- 76 F. Tumpa, J. Liang, L. Perez Estrada, M. Gamal El-Din and Y. Liu, *J. Environ. Manage.*, 2014, **146**, 535–542.
- 77 M. F. Rahman, S. Y. Jasim, E. K. Yanful, S. Ndongue and D. Borikar, *Ozone: Sci. Eng.*, 2010, **32**, 295–304.
- 78 M. S. Chandrakanth and G. L. Amy, *Environ. Sci. Technol.*, 1996, **30**, 431–443.
- 79 F. J. Rodríguez, L. A. Marcos, L. A. Núñez and M. García, *Ozone: Sci. Eng.*, 2012, **34**, 342–353.
- 80 H. Selcuk, *Dyes Pigm.*, 2005, **64**, 217–222.
- 81 L. Szpyrkowicz, C. Juzzolino and S. N. Kaul, *Water Res.*, 2001, **35**, 2129–2136.
- 82 M. Edwards and M. M. Benjamin, *J. - Am. Water Works Assoc.*, 1992, **84**, 63–72.
- 83 M. R. Jekel, *Ozone: Sci. Eng.*, 1994, **16**, 55–66.
- 84 M. R. Farvardin and A. G. Collins, *Water Res.*, 1989, **23**, 307–316.
- 85 M. S. Hobbs, R. S. Grippo, J. L. Farris, B. R. Griffin and L. L. Harding, *Environ. Toxicol. Chem.*, 2006, **25**, 3046–3052.
- 86 K. J. Ficek, *Opflow*, 1984, **10**, 1–7.
- 87 J. J. Chen and H. H. Yeh, *Water Res.*, 2005, **39**, 4420–4428.
- 88 J. J. Chen, H. H. Yeh and I. C. Tseng, *Chemosphere*, 2009, **74**, 840–846.
- 89 J. Guo, J. Ma and X. Shi, *Front. Environ. Sci. Eng. China*, 2007, **1**, 251–256.
- 90 X. Lu, X. Huangfu, X. Zhang, Y. Wang and J. Ma, *Water Res.*, 2014, **65**, 22–31.
- 91 X. Lu, X. Huangfu and J. Ma, *J. Hazard. Mater.*, 2014, **280**, 71–78.
- 92 Y. L. Song, B. Z. Dong, N. Y. Gao, Y. F. Yang and C. Chang, *Appl. Mech. Mater.*, 2012, **170–173**, 2360–2366.
- 93 J. Ma and G. Li, *Water Sci. Technol.*, 1993, **27**, 47–54.
- 94 L. Wang, J. Qiao, Y. Hu, L. Wang, L. Zhang, Q. Zhou and N. Gao, *J. Environ. Sci.*, 2013, **25**, 452–459.
- 95 J. L. Lin, L. C. Hua, S. K. Hung and C. Huang, *J. Environ. Sci.*, 2018, **63**, 147–155.
- 96 N. Graham, J. Ma and G. Li, *Aqua*, 1997, **46**, 1–10.
- 97 J. M. Colthurst and P. C. Singer, *J. - Am. Water Works Assoc.*, 1982, **74**, 78–83.
- 98 I. Arslan, *J. Hazard. Mater.*, 2001, **B85**, 229–241.
- 99 I. Monje-Ramirez and M. T. Orta de Velasquez, *Water Res.*, 2004, **38**, 2358–2366.
- 100 P. C. Chiang, E. E. Chang, P. C. Chang and C. P. Huang, *Sci. Total Environ.*, 2009, **407**, 5735–5742.
- 101 L. Cheng, X. Bi and Y. Ni, *Presented in part at the 2011 International Conference on Computer Distributed Control and Intelligent, Environmental Monitoring*, 2011.
- 102 J. Ma, J. Y. Fang, L. N. Wang, J. Guo and Z. L. Chen, *Water Sci. Technol.: Water Supply*, 2006, **6**, 145–152.
- 103 A. Montiel and B. Welte, *Water Sci. Technol.*, 1998, **37**, 65–73.
- 104 T. Gayathri, S. Kavitha, S. Adish Kumar, S. Kaliappan, I. T. Yeom and J. Rajesh Banu, *Ultrason. Sonochem.*, 2015, **22**, 333–340.
- 105 M. Drikas, C. W. K. Chow, J. House and M. D. Burch, *J. - Am. Water Works Assoc.*, 2001, **93**, 100–111.
- 106 T. Takaara, D. Sano, H. Konno and T. Omura, *Water Res.*, 2007, **41**, 1653–1658.
- 107 F. Qu, X. Du, B. Liu, J. He, N. Ren, G. Li and H. Liang, *Chem. Eng. J.*, 2015, **279**, 56–65.
- 108 E. C. Wert, M. M. Dong and F. L. Rosario-Ortiz, *Water Res.*, 2013, **47**, 3752–3761.
- 109 J. Fan, R. Daly, P. Hobson, L. Ho and J. Brookes, *Chemosphere*, 2013, **92**, 529–534.
- 110 J.-Q. Jiang and B. Lloyd, *Water Res.*, 2002, **36**, 1397–1408.
- 111 S. J. DeLuca, A. C. Chao and C. Smallwood, *J. Environ. Eng.*, 1983, **109**, 1159–1167.



Review

- 112 W. Liu and J. Ma, *Water Res.*, 2002, **36**, 871–878.
- 113 W. Liu and J. Ma, *Water Res.*, 2002, **36**, 4959–4962.
- 114 V. K. Sharma, R. Zboril and R. S. Varma, *Acc. Chem. Res.*, 2015, **48**, 182–191.
- 115 V. K. Sharma, *Adv. Environ. Res.*, 2002, **6**, 143–156.
- 116 V. K. Sharma, S. K. Mishra and N. Nesnas, *Environ. Sci. Technol.*, 2006, **40**, 7222–7227.
- 117 R. Bartzatt, M. Cano, L. Johnson and D. Nagel, *J. Toxicol. Environ. Health*, 1992, **35**, 205–210.
- 118 V. K. Sharma, J. O. Smith and F. J. Millero, *Environ. Sci. Technol.*, 1997, **31**, 2486–2491.
- 119 J.-Q. Jiang and S. Wang, *Environ. Eng. Sci.*, 2004, **20**, 627–633.
- 120 J. Q. Jiang, A. Panagouloupoulos, M. Bauer and P. Pearce, *J. Environ. Manage.*, 2006, **79**, 215–220.
- 121 V. K. Sharma, T. M. Triantis, M. G. Antoniou, X. He, M. Pelaez, C. Han, W. Song, K. E. O'Shea, A. A. de la Cruz, T. Kaloudis, A. Hiskia and D. D. Dionysiou, *Sep. Purif. Technol.*, 2012, **91**, 3–17.
- 122 K. Manoli, L. M. Morrison, M. W. Sumarah, G. Nakhla, A. K. Ray and V. K. Sharma, *Water Res.*, 2019, **148**, 272–280.
- 123 V. K. Sharma, *J. Environ. Sci. Health, Part A: Toxic/Hazard. Subst. Environ. Eng.*, 2010, **45**, 645–667.
- 124 R. K. R. Robinson, *Water Res.*, 1974, **8**, 543–547.
- 125 B. Lloyd, J.-Q. Jiang and L. Grigore, *Environ. Eng. Sci.*, 2004, **18**, 323–328.
- 126 J. Q. Jiang, S. Wang and A. Panagouloupoulos, *Chemosphere*, 2006, **63**, 212–219.
- 127 C. Stanford, J.-Q. Jiang and M. Alsheyab, *Water, Air, Soil Pollut.*, 2009, **209**, 483–488.
- 128 M. D. Johnson and K. D. Sharma, *Inorg. Chim. Acta*, 1999, **293**, 229–233.
- 129 D. Chen, *Handbook on application of ultrasound: Sonochemistry for sustainability*, 2012, p. 373, DOI: 10.1201/b11012-16.
- 130 J. H. Gibson, H. Hon, R. Farnood, I. G. Droppo and P. Seto, *Water Res.*, 2009, **43**, 2251–2259.
- 131 M. Furuta, M. Yamaguchi, T. Tsukamoto, B. Yim, C. Stavarache, K. Hasiba and Y. Maeda, *Ultrason. Sonochem.*, 2004, **11**, 57–60.
- 132 I. Tsukamoto, E. Constantinoiu, M. Furuta, R. Nishimura and Y. J. Maeda, *Ultrason. Sonochem.*, 2004, **11**, 167–172.
- 133 G. Zhang, B. Wang, P. Zhang, L. Wang and H. Wang, *J. Environ. Sci. Health, Part A: Toxic/Hazard. Subst. Environ. Eng.*, 2006, **41**, 1379–1390.
- 134 G. Zhang, P. Zhang and M. Fan, *Ultrason. Sonochem.*, 2009, **16**, 334–338.
- 135 R. Lall, R. Mutharasan, Y. T. Shah and P. Dhurjati, *Water Environ. Res.*, 2003, **75**, 171–179.
- 136 H.-X. Shi, J.-H. Qu, H.-J. Liu, Y.-L. Mu, K.-T. Xiao and L. Wang, *J. Water Supply: Res. Technol.-AQUA*, 2008, **57**, 101–108.
- 137 L. Heng, N. Jun, H. Wen-jie and L. Guibai, *Desalination*, 2009, **239**, 191–197.
- 138 P. Rajasekhar, L. H. Fan, T. Nguyen and F. A. Roddick, *Water Res.*, 2012, **46**, 1473–1481.
- 139 J. W. Li, X. Li, X. H. Xu and D. B. Wang, *Adv. Mater. Res.*, 2011, **356–360**, 291–297.
- 140 S. A. Fast, B. Kokabian and V. G. Gude, *Chem. Eng. J.*, 2014, **244**, 403–410.
- 141 Y. Li, X. Shi, Z. Zhang and Y. Peng, *Ultrason. Sonochem.*, 2019, **55**, 232–242.
- 142 D. Ghernaout and B. Ghernaout, *Desalin. Water Treat.*, 2012, **44**, 15–28.
- 143 B. Y. Gao, Q. Y. Yue and B. J. Wang, *J. Environ. Sci. Health, Part A: Toxic/Hazard. Subst. Environ. Eng.*, 2003, **38**, 897–907.
- 144 B. Y. Gao, Q. Y. Yue, B. J. Wang and Y. B. Chu, *Colloids Surf., A*, 2003, **229**, 121–127.
- 145 B. Y. Gao, G. Abbt-Braun and F. H. Frimmel, *Acta Hydrochim. Hydrobiol.*, 2006, **34**, 491–497.
- 146 C. Zhao, S. Shao, Y. Zhou, Y. Yang, Y. Shao, L. Zhang, Y. Zhou, L. Xie and L. Luo, *Sci. Total Environ.*, 2018, **645**, 267–276.
- 147 J. Ma, R. N. Wang, X. Y. Wang, H. Zhang, B. Zhu, L. L. Lian and D. W. Lou, *J. Environ. Chem. Eng.*, 2019, **7**, 103049.
- 148 A. Arnold-Smith, R. Christie and C. Jolicoeur, *Chemical Water and Wastewater Treatment II*, Springer, 1992, pp. 203–219.
- 149 B. Gao, Q. Yue and J. Miao, *Water Sci. Technol.*, 2003, **47**, 127–132.
- 150 P. A. Moussas and A. I. Zouboulis, *Sep. Purif. Technol.*, 2008, **63**, 475–483.
- 151 T. Sun, C.-h. Sun, G.-l. Zhu, X.-j. Miao, C.-c. Wu, S.-b. Lv and W.-j. Li, *Desalination*, 2011, **268**, 270–275.
- 152 M. A. Aboulhassan, S. Souabi, A. Yaacoubi and M. Baudu, *J. Hazard. Mater.*, 2006, **138**, 40–45.
- 153 Y. J. Zhang, X. L. Zhao, X. X. Li, C. Liu and L. L. Zhu, *Water Sci. Technol.*, 2010, **62**, 330–339.
- 154 X. Li, Y. Zhang, X. Zhao, N. Gao and T. Fu, *Sep. Purif. Technol.*, 2015, **147**, 125–131.
- 155 B.-Y. Gao, Y. Wang and Q.-Y. Yue, *Acta Hydrochim. Hydrobiol.*, 2005, **33**, 365–371.
- 156 Y. X. Zhao, B. Y. Gao, Y. Wang, H. K. Shon, X. W. Bo and Q. Y. Yue, *Chem. Eng. J.*, 2012, **183**, 387–394.
- 157 S. Zhao, B. Gao, Y. Wang and Z. Yang, *Colloids Surf., A*, 2013, **417**, 161–169.
- 158 X. Huang, X. Bo, Y. Zhao, B. Gao, Y. Wang, S. Sun, Q. Yue and Q. Li, *Bioresour. Technol.*, 2014, **165**, 116–121.
- 159 H.-Z. Zhao, Z.-K. Luan, B.-Y. Gao and Q.-Y. Yue, *J. Appl. Polym. Sci.*, 2002, **84**, 335–342.
- 160 Z. Yang, B. Yuan, X. Huang, J. Zhou, J. Cai, H. Yang, A. Li and R. Cheng, *Water Res.*, 2012, **46**, 107–114.
- 161 X. Huang, B. Gao, Y. Wang, Q. Yue, Q. Li and Y. Zhang, *Chem. Eng. J.*, 2014, **245**, 173–179.
- 162 L. Zhou, H. Zhou and X. Yang, *Sep. Purif. Technol.*, 2019, **210**, 93–99.
- 163 H. K. Agbovi, L. D. Wilson and L. G. Tabil, *Ind. Eng. Chem. Res.*, 2017, **56**, 37–46.
- 164 S. Bhalkaran and L. D. Wilson, *Int. J. Mol. Sci.*, 2016, **17**, 1662.
- 165 P. Kanmani, J. Aravind, M. Kamaraj, P. Sureshbabu and S. Karthikeyan, *Bioresour. Technol.*, 2017, **242**, 295–303.



- 166 H. Salehizadeh, N. Yan and R. Farnood, *Biotechnol. Adv.*, 2018, **36**, 92–119.
- 167 J. Turunen, A. Karppinen and R. Ihme, *SN Appl. Sci.*, 2019, **1**, 210.
- 168 N. Qiao, M. Gao, X. Zhang, Y. Du, X. Fan, L. Wang, N. Liu and D. Yu, *Appl. Microbiol. Biotechnol.*, 2019, **103**, 2821–2831.
- 169 A. S. Ayangbenro, O. O. Babalola and O. S. Aremu, *Chemosphere*, 2019, **231**, 113–120.
- 170 H. K. Agbovi and L. D. Wilson, *Carbohydr. Polym.*, 2018, **189**, 360–370.
- 171 J. Pérez-Calderón, M. V. Santos and N. Zaritzky, *J. Environ. Chem. Eng.*, 2018, **6**, 3808–3818.
- 172 E. Lichtfouse, N. Morin-Crini, M. Fourmentin, H. Zemmouri, I. O. do Carmo Nascimento, L. M. Queiroz, M. Y. M. Tadza, L. A. Picos-Corrales, H. Pei, L. D. Wilson and G. Crini, *Environ. Chem. Lett.*, 2019, **17**, 1603–1621.
- 173 B. Peng, Z. Yao, X. Wang, M. Crombeen, D. G. Sweeney and K. C. Tam, *Green Energy & Environment*, 2020, **5**, 37–49.
- 174 H. Kolya, D. Sasmal and T. Tripathy, *J. Polym. Environ.*, 2017, **25**, 408–418.
- 175 D. Sasmal, J. Maity, H. Kolya and T. Tripathy, *Int. J. Biol. Macromol.*, 2017, **97**, 585–597.
- 176 D. Sasmal, H. Kolya and T. Tripathy, *Int. J. Biol. Macromol.*, 2016, **91**, 934–945.

