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Core–shell architecture based on bio-sourced porous carbon: the shape formation mechanism at the solid/liquid interface layer†

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The overall goal of this work was to activate agri-food wastes by microbial action, which makes it possible to produce bio-digestate and energy (methane). The resulting bio-digestate could be transformed to porous carbon (PC), which was used for the preparation of core–shell particles with alginate (bio-polymer) and a calcium ion layer. Furthermore, surface charge measurements showed electrostatic attractions occurring between the alginate, calcium (Ca²⁺) ions and the PC, hence leading to the formation of core (PC)–shell (alginate–calcium ions) particles. However, in the absence of calcium ions, no electrostatic attractions were observed between the PC and the alginate. In the dried state (using scanning electronic microscopy analysis (SEM)) and in the hydrate state (using numerical microscopy), the designed core–shell architecture was confirmed. Transmission electron microscopy (TEM) shows that the PC particles were graphitic and porous. In addition, both Raman spectroscopy (RS) and X-ray photoelectron spectroscopy (XPS) showed the presence of several chemical functions, in particular hydroxyl (–O–H) and carboxylic groups (–COO–H). In aqueous media, the results showed that the PC was negatively charged and its surface charge and particle size were found to be very sensitive to the variation in pH. Finally, the core–shell particles were used as an adsorbent for the removal of methylene blue (MB), crystal violet (CV) and congo red (CR) molecules from wastewater. The overall data indicated efficient dye removal, without the occurrence of the solid/liquid separation problem.

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1. Introduction

Over the last few years, human activities have allowed the increase in toxic industrial effluents, irrespective of their sources. These contaminants result in various diseases and disorders in the ecosystem,^{1–3} as they are easily accumulated in living organisms and dispersed into aquatic systems. As an example, in Morocco, agri-food industries are a major consumer of water and the largest producers of pollution, and this is characterized by their higher COD and microorganism content.⁴ Therefore, it is necessary to remove these contaminants in order to prevent hazardous health effects in human beings. Many techniques including adsorption and anaerobic digestion have been used to degrade these pollutants.^{5–18} However, these techniques are generally limited when applied separately, requiring coupling and/or hybrid processes.⁵

To better understand the complexity of agri-food waste treatment based on single techniques, the anaerobic digestion process (AD) was taken as an example. In the AD process, the degradation of agri-food wastes to bio-methane (75%) was performed using microorganisms under certain temperature and pressure conditions, and this technique was highly recommended for these types of waste in terms of energy production.⁴ Nevertheless, the resulting bio-digestate remains the big problem of the AD process. In fact, the production of methane gas from agri-food wastes does not mean that the problem has been solved, but unexpectedly, the bio-product could also be toxic to the environmental media. Recently, researchers around the world have found ways to reduce this problem by converting the digestate into carbonaceous materials (*e.g.* Biochar, activated carbon and char).^{19–22} Indeed, they considered that the physicochemical properties of the bio-digestate could become suitable for the preparation of functional carbonaceous materials after microbial action using the AD process. Then, these carbonaceous materials were used for environmental remediation (in most cases the carbonaceous materials have been used as adsorbents).²³ In fact, adsorption using porous materials is crucial for applications relating to the environment due to their large specific surface area and physicochemical properties.²⁴

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The two main problems related to carbonaceous materials produced by digestate are their regeneration and the solid/liquid phase separation. So, the objective of the present work was to use the raw bio-digestate of agri-food organic waste to produce porous carbon, which was then used as a basic matrix to prepare core-shell particles based on the bio-polymer alginate and Ca^{2+} ions. These core-shell particles finally allowed both problems relating to material regeneration and phase separation to be solved.

During the preparation and utilization of the prepared core-shell particles, several chemical interactions occurred between several functional groups of the various components (carbon, alginate and Ca^{2+}) that are involved. The study of these interactions will be fruitful for a good understanding of the solid/liquid surface reactions, and for elucidating the core-shell mechanism formation as well as the mechanism of the removal of the pollutants from wastewater, especially as very few articles in the literature deal with these interactions in more detail. For these reasons, powerful surface characterizations and interface techniques were used.

The experimental section of the present work began by the preparation of the PC from the bio-digestate derived from agri-food organic waste. Then, the physicochemical, structural and morphological properties of this material were evaluated on two levels: in the hydrated and dried states. After determining these properties, the PC was then used to prepare core-shell particles with the use of alginate and calcium. The procedure used is described and the mechanism of formation of the particles is discussed based on the characterization techniques in the dispersed medium. Finally, the efficiencies of the final particles to eliminate toxic pollutants and to solve the problem of the solid/liquid phase separation have been evaluated and discussed.

2. Experimental section

2.1. Chemical reagents and preparation of aqueous solutions

The reagents used in this work were: NaOH (Sigma-Aldrich, >95%), HCl (Sigma-Aldrich, >37%), methylene blue (Sigma-Aldrich, dye content >82%), H_2SO_4 (Sigma-Aldrich, 99.99%), alginate (Sigma-Aldrich, primary grade), distilled water (laboratory grade), CaCl_2 (Sigma-Aldrich, >99%), and cetyltrimethyl ammonium chloride (CTACl) (Sigma-Aldrich, 25 wt% in H_2O). The aqueous dispersions used in this work were prepared and placed in glass vials and kept for potential use. The PC solution (0.12 g L^{-1}) was prepared using 6 mg of PC mixed with 50 mL of distilled water and then ultrasonicated for 10 min. Adjustment of the pH was performed by adding small amounts of NaOH (0.1 M) and HCl (0.1 M) aqueous solutions. The alginate solution (1 g L^{-1}) was elaborated using 1 g of alginate mixed with 1 L of distilled water and then stirred for 6 h. CaCl_2 (0.1 M) and CTACl solutions ($6.25 \times 10^{-4} \text{ M}$) were prepared from dilution of 1 M stock solutions.

2.2. Preparation of PC

The experimental protocol that was followed to prepare the PC can be counted in several stages. Initially, four agri-food organic waste materials were methanized in a mesophilic system at $38 \pm$

1°C for 72 h, followed by 13 days of stabilization. Then, the residue (bio-digestate) was dried at 100°C for 24 hours and washed with tap water several times. After this, the filtrate was chemically activated using sulfuric acid, H_2SO_4 ($\text{pH} = 2$), with 12 hours of stirring. The final product was filtered, washed with water and ethanol, and finally dried at 100°C for 24 hours. In the last step (thermal activation), the filtrate was placed in a furnace at a heating rate of $5^\circ \text{C min}^{-1}$ under nitrogen for 3 h. In the present work, eight treatment temperatures ranging from 200 to 600°C were investigated. The optimum temperature was chosen to be at 350°C . Indeed, in both temperature treatments at 200 and 250°C , a small amount of leaching of the PC was observed during the adsorption of methylene blue (MB) as a model. For the temperature treatment performed at 300°C , the prepared material was thermally stable and the removal of MB reached a maximum value of 100% with an initial concentration of 50 mg L^{-1} (Fig. 1).

2.3. Design of the core (PC)-shell (alginate-calcium) particles: PC@alginate

The core (PC)-shell (alginate-calcium) particles used in this work were prepared as follows: 1 g of alginate powder was dissolved in 100 mL of distilled water with 6 hours of stirring, then, 1 g of PC was added and the mixture was stirred for 2 hours followed by 30 min of sonication (Sonimasse 150 TS, 20 kHz, sonicating power output of 160 W). In addition, a solution of CaCl_2 with a known concentration was placed in a crystallizer and kept under low stirring (30 rpm). The preparation of PC@alginate was ensured by mixing both solutions using a pump with a speed of 20 mL h^{-1} . Indeed, when each droplet of the PC@alginate solution was brought into contact with the Ca^{2+} ions, a capsule membrane was instantly formed with the carboxylic groups of the alginate, and the PC was immobilized inside. Finally, the capsules were washed with distilled water and stored in aqueous medium (Fig. 2). The core-shell particles were labelled as PC@alginate.

2.4. Solid/liquid interface characterization

The curve representing the surface potential variation as a function of the amount of the cetyltrimethyl ammonium chloride (CTACl) surfactant added, mainly shows the interaction that occurs between the negatively charged PC surface sites

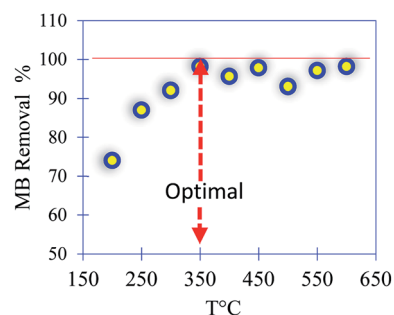


Fig. 1 MB removal efficiency versus the temperature of treatment of the PC.



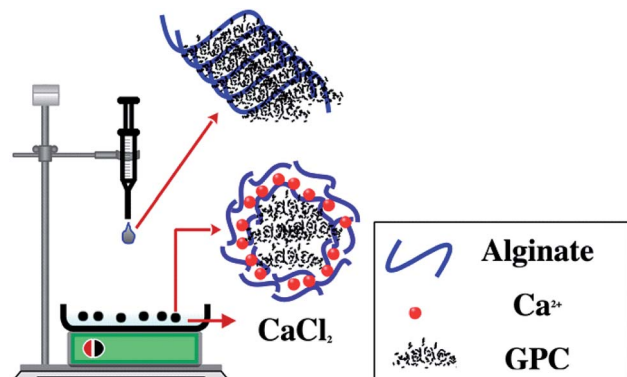


Fig. 2 Experimental protocol for (carbonaceous) core-shell (alginate-calcium ions) particle preparation.

and the positively charged CTACl head groups. The resulting titration curve of the PC by the CTACl surfactant finally reflects the surface electrical properties of the PC. The Muteck PCD 02 apparatus was used for monitoring the titration curve of the PC by the CTACl surfactant. This apparatus allowing the PC surface potential measurements, contained a cylindrical chamber and a piston made of poly(tetrafluoroethylene).²⁵ Thus, 10 mL of the PC aqueous suspension (0.12 g L^{-1} , dispersed under ultrasound for 10 min) was first placed in the space (0.5 mm) between the cylindrical chamber wall and the piston. Thereafter, at each minute, a given amount of the aqueous CTACl solution was added and the potential was measured between the two electrodes placed inside the cylindrical chamber. Since the PCD 02 apparatus allows measurements of relative potential values, to avoid data disparity, the obtained surface potential was then normalized to its maximum magnitude. The measurements of the particle sizes of the PC in aqueous dispersion as a function of pH variation were performed by Dynamic Light Scattering (DLS), using Coulter Model N4S (Coultronics) apparatus that operates with a 4 mW helium-neon laser (wavelength = 632.8 nm) and an optical system that detects at a 90° angle the light scattered by the sample, as described elsewhere.^{26,27} The measurements of the PC zeta potentials were carried out at ambient temperature with a Zetaphoremeter II model Z3000 purchased from SEPHY, as described elsewhere.²⁵

2.5. Characterization techniques in the dried state

Transmission Electron Microscopy (TEM) analyzes were done using a Philips CM 200, operating at an accelerating voltage of 20–200 kV and with a magnification ranging from $\times 10\,000$ to $\times 200\,000$. The PC was analyzed using a scanning electron microscope FEI, Model Quanta 400. The equipment used to perform the various Raman analyzes was a spectrometer Horiba model LabRAM BX40, using a CCD detector, a laser line of 532 nm and a $100\times$ objective lens with a laser power of 1.6 mW. Moreover, X-ray Photoelectron Spectroscopy (XPS) was carried out using a Thermo Fisher Scientific ESCALAB 250Xi X-ray photoelectron spectroscopy system equipped with an Al $K\alpha$ X-ray source = 1486.7 eV to study the surface chemical compositions of the synthesized PC samples. Confocal microscopy (Axio

Imager M2 LSM 700 ordered by the ZEN 2012 software) was used to confirm the designed core-shell adsorbent. X-ray diffraction patterns were collected on a PANalytical MPD X'Pert Pro diffractometer, operating with Cu K_α radiation ($K_\alpha = 0.15418 \text{ nm}$), equipped with an X'Celerator real-time multiple strip detector (active length = $2.12^\circ 2\theta$). The powder patterns were collected at 295 K in the range $5^\circ < 2\theta < 90^\circ$, with step = $0.017^\circ 2\theta$, and time/step = 220 s.

2.6. MB molecule removal experiments

The removal tests of dyes molecules from aqueous solution using the PC@alginate particles were conducted using a batch system. The effects of pH on the adsorption of dyes onto PC@alginate were investigated using 40 mL (10 mg L^{-1}) at different pHs (3–12). The pH of the solution was adjusted by adding small amounts of HCl (0.1 M) and NaOH (0.1 M) aqueous solutions to various PC@alginate dispersions. The effects of the initial dye concentration ($5\text{--}40 \text{ mg L}^{-1}$) were evaluated using 25 mg of PC@alginate and 40 mL aqueous solution of the dyes. Finally, the mixture was easily separated using a previously cleaned spatula, and the equilibrium concentration of MB was analyzed by a UV-vis spectrophotometer. The removal efficiency and adsorption capacity at equilibrium were calculated as follows:

$$\% \text{ removal} = \frac{(C_0 - C_e)}{C_0} \times 100 \quad (1)$$

$$\% \text{ removal} = (C_0 - C_e) \times \frac{m}{V} \quad (2)$$

where C_0 and C_e are the initial and equilibrium concentrations, respectively, of the dyes in mg L^{-1} , m (g) is the amount of adsorbent and V (L) is the volume of the solution.

In order to investigate the adsorbent reusability, eight cycles of adsorption-desorption were carried out using 25 mg of PC@alginate loaded with dyes, with 20 mL of HCl (0.1 M) and NaOH (0.1 M) with 30 min of agitation. To fit the isotherm data, Langmuir and Freundlich equations were used (eqn (3) and (4)).^{2,28–31}

$$\frac{C_e}{Q_e} = \frac{1}{k_L Q_{\max}} + \frac{C_e}{Q_{\max}} \quad (3)$$

$$\ln(Q_e) = \frac{1}{n} \ln(C_e) + \ln(k_F) \quad (4)$$

where C_e and Q_e are the concentration and amount at equilibrium, k_L is a direct measure of the intensity of the adsorption process, Q_{\max} is the maximum adsorption capacity, k_F is the adsorption capacity, and n reflects the adsorption intensity ($1/n = 0$ irreversible adsorption; $1/n > 1$ unfavorable adsorption, $0 < 1/n < 1$ favorable adsorption).

3. Results and discussion

3.1. Characterization of prepared material in the dried state

3.1.1. Scanning electron microscopy (SEM) and transmission electron microscopy (TEM). To acquire the



morphological characteristics of the digestate and PC material, SEM analysis was performed (Fig. 3). Before treatment, the digestate surface is quite compact and it does not have many pores. When the surface is chemically attacked by sulfuric acid and then heat-treated, a change in the morphological characteristics is observed. After treatment, the morphological properties are modified. Indeed, the surface becomes very smooth and macroporous. The MEB resolution is still not able to permit observation of smaller pores. Therefore, TEM analysis was performed to assess the presence of meso- and micro-pores.

Micrographs of typical areas of PC at 20 nm are shown in Fig. 4a and b. As can be observed in these figures, the PC has a disordered microporous structure, and is made up of arranged curved graphene layers.^{32,33} Furthermore, numerous graphene sheets with different dimensions and shapes are clearly observed, that are at 5 Å spacings, which allow the creation of micropores.^{33,34} However, the PC exhibits a much lower packing density of carbon sheets in some regions (more compact structure) and not in others.³⁵ Finally, we can see in Fig. 4a and b that the TEM images confirm the amorphous structure of our material, with graphite layers being observed as is already reported in some works.³⁶ From Fig. 4c, we can observe the characteristic planes of amorphous carbon around 20–25 2θ and the region of 40–45 2θ degree.^{37,38} The XRD analysis shows an amorphous phase of the PC adsorbent by the presence of characteristic peaks of graphite (around 26 2θ) and activated carbon, and this confirms the chemical carbonaceous structure of our prepared sample.

3.1.2. Elemental analysis. Elemental analyzes of the digestate, and of the PC, were performed by the EDS technique, as presented in Fig. S1.† These analyzes showed the presence of some impurities such as Na, Al, Si, S, K, Ca, Fe, Mg and Cl. The presence of these elements has been explained by the use of some chemical reagents during the process of preparation of the PC (H_2SO_4 for example) and/or the origin of raw material (agri-food wastes), which is characterized by heterogeneous chemical composition. The EDS analysis, on the other hand,

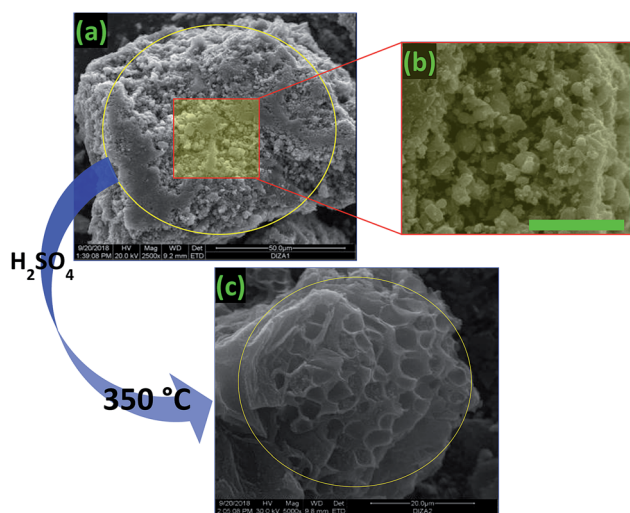


Fig. 3 (a) and (b) SEM images of the raw digestate, and (c) an SEM image of the prepared PC.

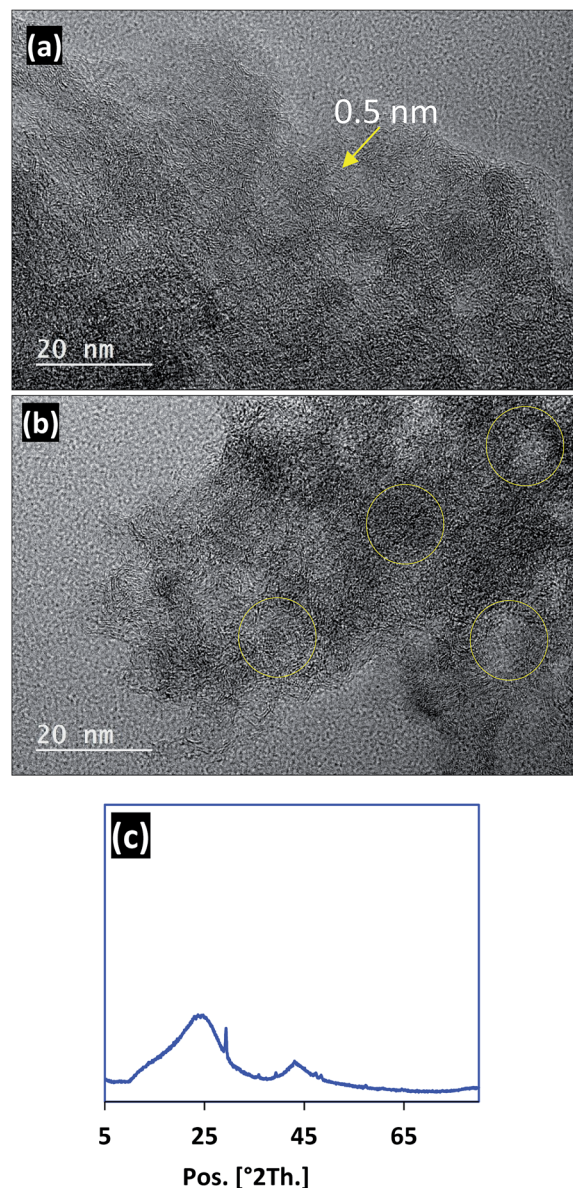


Fig. 4 (a) and (b) TEM images of the prepared PC and (c) the XRD spectrum of the prepared PC.

indicated an increase in the carbon content and a decrease in the oxygen content after chemical and thermal treatments. Table S1† shows that the atomic percentage of carbon increased by 45.26%, while a decrease by 30.56% was observed for the oxygen concentration after the preparation of PC. The present analysis shows that the PC consists mainly of carbon and oxygen, and this is consistent with other techniques.

3.1.3. Functional group characterization. To characterize the oxygen-containing functional groups on the surface of the PC sample, XPS analysis was performed. From Fig. 5a and b, the surface of the PC was made up mainly of sp^2 -bonded carbon, and C–OH and C=O bonds. The sp^2 -bonded carbon atoms were detected at 284.99 eV, C–OH broad peaks were confirmed at 286.2 eV and C=O bonds were observed at 288.45 eV.^{2,29,39} The deconvolution of the O1s band shows three crests at 530, 532



and 533 eV, which can be attributed to the O atoms in phenolic and ether compounds, the carbonyl oxygen atom in anhydrides and lactones, and the oxygen atom in carboxylic acid groups, respectively.^{40,41} Nevertheless, impurities were detected in the PC structure related to the starting biomass and the reagents used for the preparation of the PC.

As a tool to investigate the functionalization and graphitization of the surface of the PC, Raman analysis was used (Fig. 5d). The results showed that the PC sample presents two broad bands at 1355.22 and 1604.89 cm^{-1} , noted D (disorders) and G (graphitic), respectively.² The G band is related to the stretching vibrations of sp^2 carbon atoms in the long carbon chains in graphite or the carbon ring. The D band, however, is assigned to the deflection of the crystalline structure and disorder arrangement, which gives low symmetry and defects.^{42,43} In addition, the relative intensity between I_G/I_D was used to calculate the average crystallite size " L_a ", which shows the structural disorder degree and hence informs us about the size of the graphitic domains of the prepared PC.^{44,45} The same results have been found in reported works aimed at the preparation of carbon materials.^{46,47}

3.2. Characterization of prepared material in the hydrate state

3.2.1. Surface charge measurements of PC particles. The surface charge of the PC was studied as a function of pH (Fig. 6) using cetyltrimethyl ammonium chloride (CTACl) as a titrant. After pH adjustment using small amounts of HCl (0.1 M) and NaOH (0.1 M) aqueous solutions, 10 mL of the PC suspension (0.12 g L^{-1}) was placed 0.5 mm between the cylinder and the

piston. Before analysis, the solutions were dispersed under ultrasound for 10 minutes. The titration was carried out by the addition of variable amounts of aqueous solution of CTACl ($6.25 \times 10^{-6} \text{ M}$) at determined periods using a micropipette. At the natural pH (pH = 8.3), the potential was initially negative, which reflects the fact that the PC particles are negatively charged due to the ionization of the $-\text{COOH}$ and $-\text{OH}$ groups present on the surface (already confirmed by XPS and Raman analyzes) (Fig. 5). When the pH > 6.5, and after addition of CTACl, the negative charges of the PC particles are progressively compensated by CTA^+ ions to the point where the particles become neutralized (SIP = 0 mV) (Fig. 6a). The number of moles of CTACl that made it possible to reach the zero charges correspond to the equivalent point. Therefore, the values of the equivalent points increase as the pH increases ($10^{-7} \text{ mol g}^{-1}$ for pH = 6.2, $1.75 \times 10^{-7} \text{ mol g}^{-1}$ for pH = 8.32, $2.25 \times 10^{-7} \text{ mol g}^{-1}$ for pH = 9.13 and $2.75 \times 10^{-7} \text{ mol g}^{-1}$ for pH = 10.07). Beyond the equivalent points, the excess of cationic surfactant reverses the charge of the PC particles and the surface potential becomes positive. As the surface of the PC is rich in carboxylic and hydroxyl groups, the variation in pH

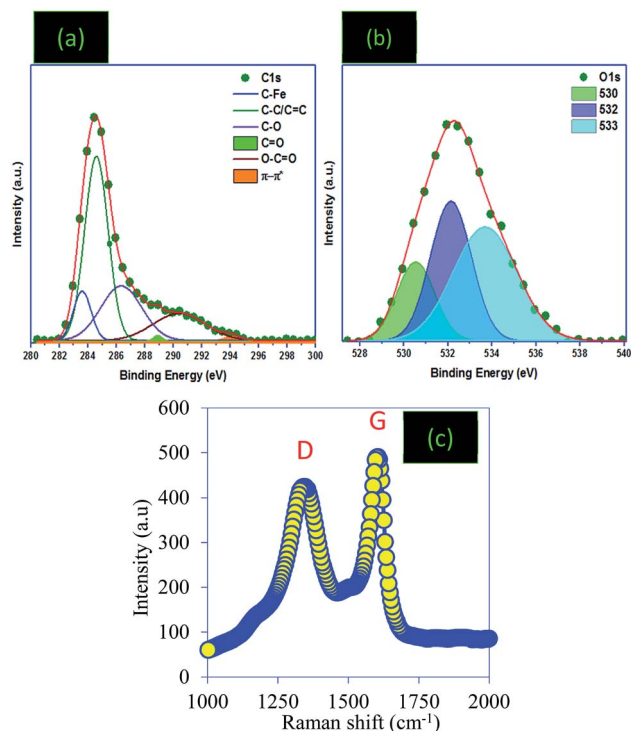


Fig. 5 (a) and (b) XPS and (c) Raman analysis of the PC material.

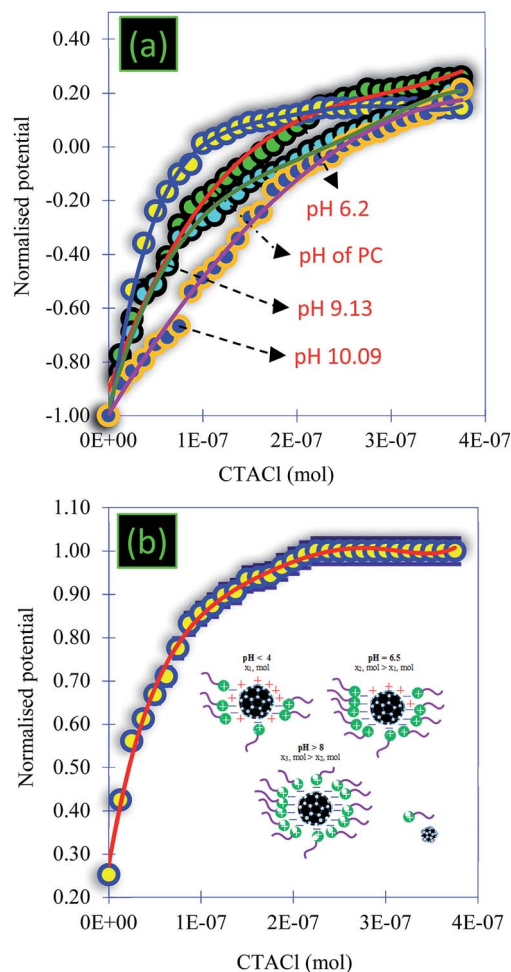


Fig. 6 (a) Titration of the PC solution at different pHs using CTACl ($6.25 \times 10^{-4} \text{ M}$) and (b) titration of the PC solution at acidic pH using CTACl ($6.25 \times 10^{-4} \text{ M}$).



modifies the charge of these groups and makes the surface very polarizable, which favors more electrostatic attraction with CTA⁺ cationic molecules. At pH < 4, the measured values indicate a positive charge due to the number of H⁺ ions added (the potential starting point is +50 mV) (Fig. 6b). After the addition of the CTA⁺ molecules, no reverse charge was observed. This analysis shows that the PC sample contains negative surface charges that are very sensitive to the aqueous phase pH changes.

3.2.2. Effect of pH and temperature on the size and zeta potential of the PC particles. Fig. 7 shows the variation in the size of the PC particles in aqueous medium as a function of pH. The pH was varied between 3 and 10 to evaluate the particle size. At pH = 3.4, the particle diameter of the PC was found to be 433 nm, and when the pH changed from 3.4 to 6.1, the size of the PC particles decreased from 433 nm to 336 nm. The size variation in the PC at these pH values can be explained by the decrease in the attraction forces as the pH increases (Fig. 6b confirms the presence of negatively charged particles on the surface, even when the pH was <4). In the range of pH 7 to 8, an increase in the diameter of the PC particles by 78 nm was observed, which can be explained by the electrical instability in this pH zone (pH near the PZC). At pH > 8, the surface becomes negative and the particle size decreased again to 307 nm. In fact, at these pH values, the carboxylic and phenolic groups were negatively charged, which leads to repulsive electrical forces occurring between the PC particles and hence avoidance of their agglomeration. This analysis is evidence that the particle size in

dispersed media is dependent on the aqueous phase pH, as well as evidence for the particles having smaller sizes in the liquid phase compared to those observed by SEM in the dry state.

Note that the PZC is an important parameter for characterizing the pH of a solid surface and ensuring colloidal stability.^{48,49} By definition, the PZC refers to the pH value at which the magnitude of the negative and the positive surface charges are equal. In this work, the surface charge of the PC was characterized using zeta-potential measurements as a function of pH in the range of pH 3–11 (Fig. 7b). As shown in Fig. 7b, the isoelectric points were located at pH 6.9, indicating that the surface charge of the PC has a positively charged surface below this value, while it has a negatively charged surface above it. The PC carbon powder showed positive zeta-potential values below about pH 3, indicating the coexistence of positively charged groups at the surface of the particles with less negatively charged oxygen groups (–COOH and –OH), and this coincides with other work.^{50,51} As we have already shown by surface charge detector measurements, the PZC confirmed that the surface of PC at the natural pH was negatively charged due to the introduction of oxygen-containing functional groups. We note that the variation in the electrical surface potential of the PC particles as a function of pH can be useful for the control of the desorption and the regeneration of PC.⁴⁸

The variation in the size of the particles with the temperature was studied and the temperature control was provided by a thermocouple immersed in the measuring cell (Fig. 7c). The

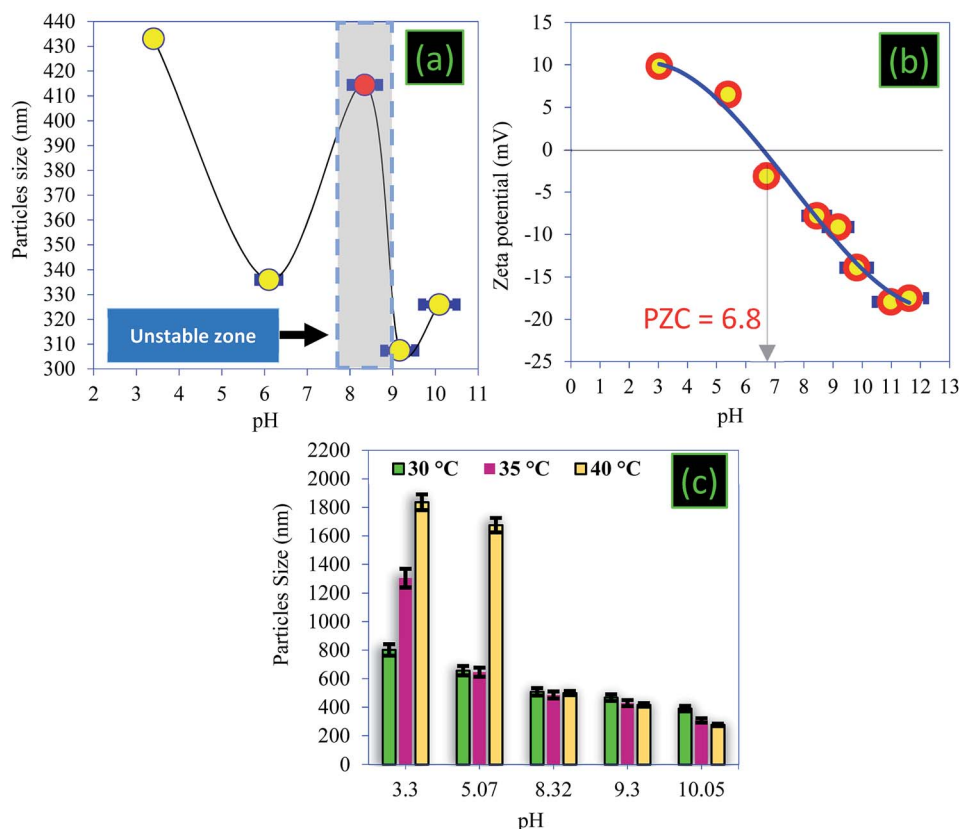


Fig. 7 (a) Particle mean size as a function of pH, (b) zeta potential of the PC, and (c) particle mean size as a function of temperature.



influence of temperature on the particle size of PC can be summarized in two main points:

- When the $\text{pH} < \text{PZC}$, an increase in temperature favors an increase in particle size, and this increase is very high where the $\text{pH} < 3$.

- When the $\text{pH} > \text{PZC}$, an increase in temperature favors a decrease in particle size up to a value equal to 276 nm, recorded at 40 °C.

The peak observed at 20 °C (Fig. 7a) gradually disappears when the temperature increases. Indeed, the sizes become larger in acidic medium, which prevents the influence of this unstable zone on the particle size. These results show that the highest yields of removal of BM and CV dyes will be recorded at alkaline pH and at higher temperatures, which is not the case in acidic conditions. As a result, the particle size decreases, resulting in an increase of the surface area of the PC.

3.3. Chemical mechanism of the formation of the core-shell particles

Despite the high performance of PC to remove pollutants (due to a developed porous structure and the presence of surface functions), the problem of separation of both liquid and solid phases is still apparent. To solve this problem, PC has been used as a matrix for preparing core-shell particles that are used to remove toxic dyes from aqueous solutions. Before running the adsorption experiments using core-shell particles, the formation process of these particles was investigated using the measurements of the surface charges (Fig. 8 and 9). Thus, the chemical mechanism of core-shell particle formation has been studied following successive titrations:

- Titration 1 (Fig. 8a): titration of alginate solution (1 g L⁻¹) with CaCl₂ (0.1 M). This titration showed a strong electrostatic interaction occurring between the Ca²⁺ ions and the carboxylic groups of the alginate. Indeed, the reaction at the beginning started with a potential equal to -510 mV and, after the addition of 6×10^{-7} mol of Ca²⁺, the surface potential decreased to 0 mV (Fig. 8a and 9).

- Titration 2 (Fig. 8a): titration of the PC solution (0.12 g L⁻¹) at pH = 8.4 by CaCl₂ (0.1 M). The main objective of this titration was to verify that the surface of the PC reacts weakly with the Ca²⁺ molecules. The analysis of Fig. 8a shows a weak interaction occurring between the PC and the Ca²⁺ ions after the addition of the same amount of CaCl₂ as in titration 1. The comparison between titrations 1 and 2 (Fig. 8a) allowed an exponential curve to be seen in the case of alginate/Ca²⁺ and the opposite for the PC/Ca²⁺ system. Indeed, in both cases we started with a normalized potential (ISP) equal to -1 and after the addition of 6×10^{-7} mol of Ca²⁺, the saturation plateau (ISP = 0 mV) was detected in the case of the alginate/Ca²⁺ system, whilst in the PC/Ca²⁺ system, a plateau of the ISP around 0.9 mV was reached following the addition of large amounts of CaCl₂.

- Titration 3 (Fig. 8b): titration of the PC solution (0.12 g L⁻¹) at pH = 8.4 with alginate (1 g L⁻¹). This experiment showed that the surface of the PC does not interact with that of alginate. At the beginning of the reaction, the surface of the PC indicates a negative potential equal to -156 mV, which remains constant

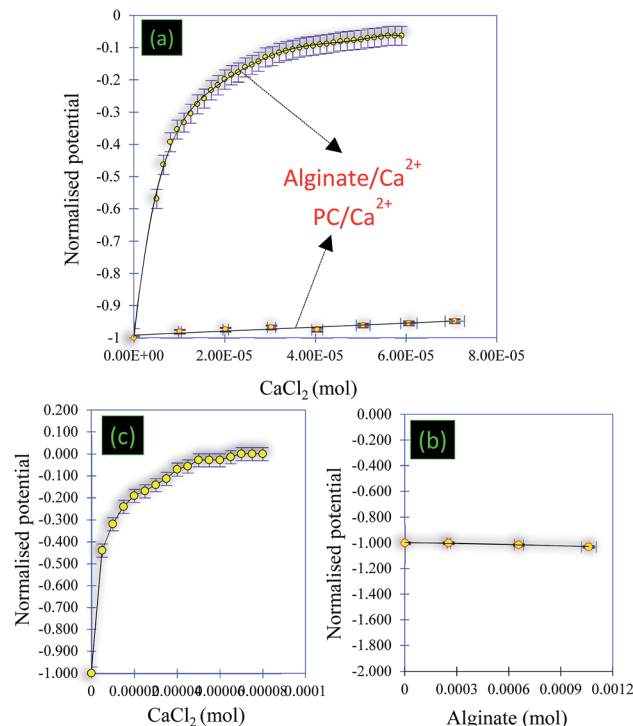


Fig. 8 (a) Titration of the PC solution (0.12 g L⁻¹) using CaCl₂ (0.1 M) and titration of the alginate solution (1 g L⁻¹) using CaCl₂ (0.1 M). (b) Titration of the PC solution (pH = 8.4) (0.12 g L⁻¹) using alginate (1 g L⁻¹) and (c) titration of the PC@alginate mixture using CaCl₂ (0.1 M).

after the addition of the alginate molecules (Fig. 8b and 9). The repulsive forces between the two negatively charged surfaces explained these results. Fig. 8b confirms these results and shows a linear form of the ISP variation as a function of the amount of alginate added.

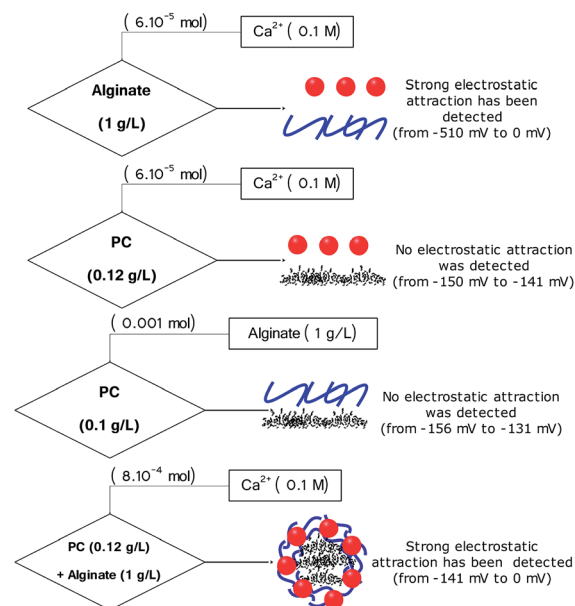


Fig. 9 Chemical mechanism of core-shell particle formation.



• Titration 4 (Fig. 8c): titration of a mixed solution of PC (0.12 g L^{-1}) and alginate (1 g L^{-1}) with CaCl_2 (0.1 M). Fig. 8c shows a strong electrostatic interaction (probably between Ca^{2+} and alginate) and the surface charge saturation is held after the addition of $8 \times 10^{-5} \text{ mol CaCl}_2$ (0.1 M).

In addition, Fig. 9 shows the performed titrations used in this work to describe the mechanism of core-shell particle formation. As a conclusion, the surface charge measurements showed electrostatic attraction and/or repulsion forces occurring between the alginate ions, the calcium ions (Ca^{2+}) and the PC, thus leading to the formation of core(carbon)-shell (alginate-calcium ions) particles. However, in the absence of calcium ions, no electrostatic attraction was observed between the PC and the alginate. In addition, the form of the core-shell particles was confirmed by numerical microscopy and the dimensions have been calculated (Fig. S2†).

3.4. PC surface charge interaction with MB, CV and CR

PC surface charge interactions with different dyes were investigated and CTACL was used as a reference. From Fig. 10, we can conclude that the CR molecule was weakly interfaced with PC compared with MB and CV. The number of moles of CR necessary to saturate the surface of the PC was equal $2.5 \times 10^{-10} \text{ mol}$ and, in the case of MB and CV, this value was higher by 60 and 100 times, respectively. These results can be explained by the characteristic surface of PC, which is more sensitive to cationic dyes than to anionic dyes. In addition, this analysis showed that the CV dye was adsorbed more than the MB dye on the surface of the PC by 0.6 times. This difference in the number of moles can be explained by the tail of the MB molecule, as well as the functional groups of CV, which can be more attractive with the $-\text{COO}-\text{H}$ and $-\text{OH}$ groups of PC, especially with the $\text{N}-\text{H}$ and SO_3^- groups. This analysis allows the characteristic interactions between the molecules and the surface of the PC to be seen. These results, and those of the characterizations, confirm that the mechanism of adsorption refers to the formation of hydrogen bonding and the electrostatic attraction of the dyes with C-O groups. In addition, the fixation of the

pollutant molecule inside the adsorbent pores of PC plays a very important role in the dye adsorption mechanism.^{2,52-54}

3.5. Adsorption studies

The effect of pH on the MB solution was carried out under the following conditions: 25 mg of adsorbent mass, a contact time of adsorbent-adsorbate of 100 min, 40 mL of dye solutions, and an initial dye concentration of 10 mg L^{-1} at room temperature.

Fig. S3a† shows the curve of the removal efficiency measured at different pHs. It can be seen that the pH of the solution has a critical impact on dye removal. Therefore, when the pH of the solution was very acidic, the removal of MB and CV was very low, whereas it was the opposite when the pH of the cationic dye solution was higher than 8. In fact, the availability of negative active sites in PC and the high surface area at this range of pH explained the higher adsorption capacity of these dyes compared to that of CR.

These results are in agreement with those cited in the above sections and are summarized as the following:

- From Fig. 7a, we demonstrated that the carboxylic and phenolic groups were charged negatively, which leads to a repulsion of the electrical forces between PC particles and avoids agglomeration, which allowed the recording of the lowest values of PC particle sizes (307 nm).

- From Fig. 6a, the titration of the PC solution at different pHs using CTACL ($6.25 \times 10^{-4} \text{ M}$) showed that the values of the equivalent point increase as the pH increases. Therefore, a high amount of carboxylic and hydroxyl functionalization groups characterized the surface of the PC and a variation in the pH allowed the modification of the charge of these groups, which favored more electrostatic attractions with MB and CV.

- From Fig. 7b, in alkaline pH ($\text{pH} > \text{pH}_{\text{PZC}}$), the surface of the PC@alginate was negatively charged, which is in favor of the electrostatic attraction forces that occur between the PC and the cationic dyes.⁵⁵

The adsorption isotherms of MB, CV and CR onto PC@alginate have been calculated using both Langmuir and Freundlich models.⁵⁶⁻⁶⁰ The overall data indicate efficient dye removal in the case of MB and CV (>96%), without the occurrence of the solid/liquid separation problem. As a result of the adsorption isotherms (Fig. S3b† and Table 1), the correlation coefficients showed a good fit in the case of the Langmuir model and the values of R_L showed a favorable adsorption, indicating the achievement of a homogeneously adsorbed monolayer of MB, CV and CR onto PC@alginate.

The reusability and the recovery ability of the adsorbents were considered as crucial economic factors when assessing the cost-effectiveness of the treatment process.^{61,62} The results of regeneration revealed that the core-shell particles showed high reusability efficiencies. Indeed, the regeneration test successfully eluted 91.28% (for MB) and 92.74% (for CV) of the dyes loaded onto PC@alginate after 5 cycles of sorption-desorption (Fig. 11a). In the case of CR, the reusability efficiencies decreased by 5.93 after 5 cycles using NaOH (0.1 M) (Fig. 11a). In all case systems (MB, CV and CR adsorption), a total recovery of the adsorbent particles was observed (Fig. 11b).

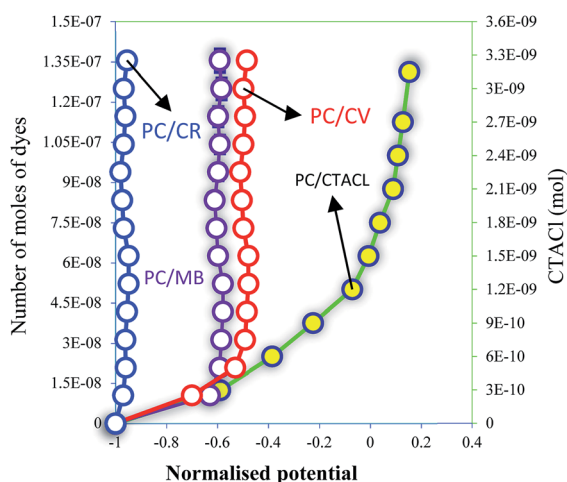


Fig. 10 PC surface charge interactions with MB, CV and CR.



Table 1 Characteristic kinetic adsorption parameters of MB, CV and CR adsorption

Langmuir				
Dye	Q_{\max} (mg g ⁻¹)	k_L (L mg ⁻¹)	R^2	R_L
MB	25.906	1.550	0.9978	0.11–0.01
CV	30.959	1.318	0.9930	0.13–0.01
CR	9.0334	0.422	0.9888	0.32–0.05
Freundlich				
Dye	k_F (mg ¹⁻ⁿ L ⁿ g ⁻¹)	n	R^2	k_F (mg ¹⁻ⁿ L ⁿ g ⁻¹)
MB	8.545	2.758	0.3613	8.545
CV	7.252	1.910	0.4299	7.252
CR	1.549	1.822	0.8858	1.549

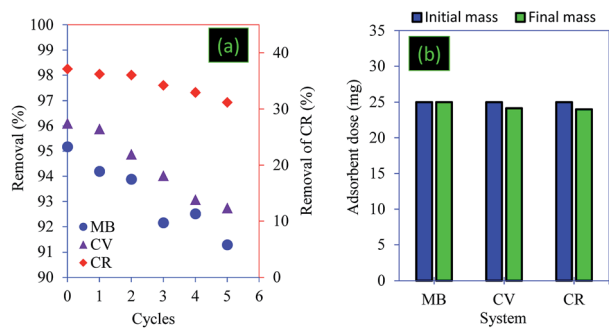


Fig. 11 (a) Reusability and (b) recovery ability of the PC@alginate particles.

4. Conclusions

In this work, agri-food organic waste has been transformed into core-shell particles. These particles allow for the elimination of toxic dyes as well as solve the problem relating to the separation of liquid and solid phases after the adsorption process. The main results of the work are:

- The bio-digestate has been transformed into PC with a large specific surface area.
- The characterization of PC shows the structure of activated carbon with the presence of carboxylic and hydroxyl groups.
- In a dispersed medium, the PC sample was very sensitive to the variation in pH. Indeed, the pH variation makes it possible to reduce the particle size and to increase the electrostatic attraction with the PC. These results were confirmed using characterization techniques in aqueous medium and the surfactant molecule CTACl as a probe. The particle size of PC was very sensitive to temperature variation and the sizes are very small when the medium becomes basic at higher temperatures.
- The mechanism of formation of the core-shell is well evidenced using several titrations and the final form is observed by two-level 2D and 3D analysis using numerical microscopy.
- Core-shell particles were used to remove toxic dyes. Experimental results showed that cationic dyes are more absorbed than anionic ones. According to the Langmuir model,

the adsorption capacities were: 25.90 mg g⁻¹ (for MB), 30.95 mg g⁻¹ (for CV) and 9.03 mg g⁻¹ (for CR).

- Core-shell particles are easy to regenerate. The results showed a decrease of 3.89, 3.31, and 5.93% for MB, CV and CR, respectively.

- The core-shell particles are easy to recover. The results showed that the adsorbent mass loss after adsorption does not exceed 3%.

Conflicts of interest

The authors declare that they have no conflicts of interest.

Author contributions

All authors have given approval to the final version of the manuscript.

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Notes and references

- 1 M. Alimohammady, M. Jahangiri, F. Kiani and H. Tahermansouri, *New J. Chem.*, 2017, **41**, 8905–8919.
- 2 M. Zbair, Z. Anfar, H. A. Ahsaine, N. El Alem and M. Ezahri, *J. Environ. Manage.*, 2018, **206**, 383–397.
- 3 M. Zbair, H. Ait Ahsaine and Z. Anfar, *J. Cleaner Prod.*, 2018, **202**, 571–581.
- 4 S. Lhanafi, Z. Anfar, B. Chebli, M. Benafqir, R. El Haouti, Y. Azougarh, M. Abbaz and N. El Alem, *Sustainable Environ. Res.*, 2018, **28**, 389–395.
- 5 Y. Ozay, E. K. Ünşar, Z. Işık, F. Yılmaz, N. Dizge, N. A. Perendeci, M. A. Mazmanci and M. Yalvac, *J. Cleaner Prod.*, 2018, **196**, 42–50.
- 6 R. Su, Q. Li, Y. Chen, B. Gao, Q. Yue and W. Zhou, *J. Cleaner Prod.*, 2019, **209**, 20–29.
- 7 S. Liu, X. Chen, W. Ai and C. Wei, *J. Cleaner Prod.*, 2019, **212**, 1062–1071.
- 8 S. Wong, H. H. Tumari, N. Ngadi, N. B. Mohamed, O. Hassan, R. Mat and N. A. Saidina Amin, *J. Cleaner Prod.*, 2019, **206**, 394–406.
- 9 E. Serrano León, J. A. Perales Vargas-Machuca, E. Lara Corona, Z. Arbib, F. Rogalla and M. Fernández Boizán, *J. Cleaner Prod.*, 2018, **198**, 931–939.
- 10 R. M. Novais, A. P. F. Caetano, M. P. Seabra, J. A. Labrincha and R. C. Pullar, *J. Cleaner Prod.*, 2018, **197**, 1137–1147.
- 11 T. L. da Silva, A. C. da Silva, M. G. A. Vieira, M. L. Gimenes and M. G. C. da Silva, *J. Cleaner Prod.*, 2016, **137**, 1470–1478.



- 12 O. Kuczman, M. V. D. Gueri, S. N. M. De Souza, W. N. Schirmer, H. J. Alves, D. Secco, W. G. Buratto, C. B. Ribeiro and F. B. Hernandez, *J. Cleaner Prod.*, 2018, **196**, 382–389.
- 13 A. Kumar and H. M. Jena, *J. Cleaner Prod.*, 2016, **137**, 1246–1259.
- 14 R. Bedoić, L. Čuček, B. Čosić, D. Krajnc, G. Smoljanić, Z. Kravanja, D. Ljubas, T. Pukšec and N. Duić, *J. Cleaner Prod.*, 2019, **213**, 700–709.
- 15 M. Moretti, M. Van Dael, R. Malina and S. Van Passel, *J. Cleaner Prod.*, 2018, **171**, 954–961.
- 16 A. Pfluger, J. Coontz, V. Zhiteneva, T. Gulliver, L. Cherry, L. Cavanaugh and L. Figueroa, *J. Cleaner Prod.*, 2019, **206**, 97–107.
- 17 A. Ahmad, A. H. Bhat and A. Buang, *J. Cleaner Prod.*, 2018, **171**, 1361–1375.
- 18 R. Kamaraj and S. Vasudevan, *New J. Chem.*, 2016, **40**, 2249–2258.
- 19 M. Inyang, B. Gao, Y. Yao, Y. Xue, A. R. Zimmerman, P. Pullammanappallil and X. Cao, *Bioresour. Technol.*, 2012, **110**, 50–56.
- 20 M. Inyang, B. Gao, A. Zimmerman, M. Zhang and H. Chen, *Chem. Eng. J.*, 2014, **236**, 39–46.
- 21 Y. Yao, B. Gao, M. Inyang, A. R. Zimmerman, X. Cao, P. Pullammanappallil and L. Yang, *Bioresour. Technol.*, 2011, **102**, 6273–6278.
- 22 M. Inyang, B. Gao, W. Ding, P. Pullammanappallil, A. R. Zimmerman and X. Cao, *Sep. Sci. Technol.*, 2011, **46**, 1950–1956.
- 23 M. Inyang, B. Gao, P. Pullammanappallil, W. Ding and A. R. Zimmerman, *Bioresour. Technol.*, 2010, **101**, 8868–8872.
- 24 B. Coasne, *New J. Chem.*, 2016, **40**, 4078–4094.
- 25 A. Jada, H. Debih and M. Khodja, *J. Pet. Sci. Eng.*, 2006, **52**, 305–316.
- 26 A. Jada, H. Ridaoui, L. Vidal and J. B. Donnet, *Colloids Surf., A*, 2014, **458**, 187–194.
- 27 H. Ridaoui, A. Jada, L. Vidal and J. B. Donnet, *Colloids Surf., A*, 2006, **278**, 149–159.
- 28 Z. Anfar, M. Zbair, H. A. Ahsaine, M. Ezahri and N. El Alem, *Fullerenes, Nanotubes, Carbon Nanostruct.*, 2018, **26**, 389–397.
- 29 H. A. Ahsaine, M. Zbair, Z. Anfar, Y. Naciri, R. El, N. El Alem and M. Ezahri, *Mater. Today Chem.*, 2018, **8**, 121–132.
- 30 A. Elouahli, M. Zbair, Z. Anfar, H. A. Ahsaine, H. Khallok, R. Chourak and Z. Hatim, *Surf. Interfaces*, 2018, **13**, 139–147.
- 31 Z. Anfar, R. El Haouti, S. Lhanafi, M. Benafqir, Y. Azougarh and N. El Alem, *J. Environ. Chem. Eng.*, 2017, **5**, 5857–5867.
- 32 P. J. F. Harris, Z. Liu and K. Suenaga, *J. Phys.: Condens. Matter*, 2008, **20**, 1–5.
- 33 M. E. Casco, M. Martínez-Escandell, K. Kaneko, J. Silvestre-Albero and F. Rodríguez-Reinoso, *Carbon*, 2015, **93**, 11–21.
- 34 Y. Huang, F. S. Cannon, J. K. Watson, B. Reznik and J. P. Mathews, *Carbon*, 2015, **83**, 1–14.
- 35 C. Bommier, R. Xu, W. Wang, X. Wang, D. Wen, J. Lu and X. Ji, *Nano Energy*, 2015, **13**, 709–717.
- 36 J. Liu, N. Sun, C. Sun, H. Liu, C. Snape, K. Li, W. Wei and Y. Sun, *Carbon*, 2015, **94**, 243–255.
- 37 B. Chang, D. Guan, Y. Tian, Z. Yang and X. Dong, *J. Hazard. Mater.*, 2013, **262**, 256–264.
- 38 B. Chang, H. Yin, X. Zhang, S. Zhang and B. Yang, *Chem. Eng. J.*, 2017, **312**, 191–203.
- 39 B. Chang, Y. Guo, Y. Li, H. Yin, S. Zhang, B. Yang and X. Dong, *J. Mater. Chem. A*, 2015, **3**, 9565–9577.
- 40 Y. Haldorai, W. Voit and J.-J. Shim, *Electrochim. Acta*, 2014, **120**, 65–72.
- 41 Z. Anfar, A. Amedlous, A. Ait El Fakir, H. Ait Ahsaine, M. Zbair, S. Lhanafi, R. El Haouti, A. Jada and N. El Alem, *ACS Omega*, 2019, **4**, 9434–9445.
- 42 S. Vasudevan and J. Lakshmi, *RSC Adv.*, 2012, **2**, 5234–5242.
- 43 W. Song, B. Gao, X. Xu, F. Wang, N. Xue, S. Sun, W. Song and R. Jia, *J. Hazard. Mater.*, 2016, **304**, 280–290.
- 44 F. Tuinstra and J. L. Koenig, *J. Chem. Phys.*, 1969, **31**, 646–650.
- 45 A. C. Ferrari and J. Robertson, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2013, **10**, 26–31.
- 46 A. Lazzarini, A. Piovano, R. Pellegrini, G. Agostini, S. Rudić, C. Lamberti and E. Groppo, *Phys. Procedia*, 2016, **85**, 20–26.
- 47 B. Chang, S. Zhang, L. Sun, H. Yin and B. Yang, *RSC Adv.*, 2016, **6**, 71360–71369.
- 48 H. W. Kwak, Y. Hong, M. E. Lee and H. J. Jin, *Int. J. Biol. Macromol.*, 2018, **120**, 906–914.
- 49 G. K. Poongavanam and V. Ramalingam, *Int. J. Therm. Sci.*, 2019, **136**, 15–32.
- 50 L. Liang, M. Zhou, W. Yang and L. Jiang, *Chem. Eng. J.*, 2018, **352**, 673–681.
- 51 J. Sun, X. Liu, F. Zhang, J. Zhou, J. Wu, A. Alsaedi, T. Hayat and J. Li, *Colloids Surf., A*, 2019, **563**, 22–30.
- 52 Z. Anfar, A. Amedlous, A. A. El Fakir, M. Zbair, H. Ait Ahsaine, A. Jada and N. El Alem, *Chemosphere*, 2019, **236**, 124351.
- 53 Z. Anfar, H. Ait Ahsaine, M. Zbair, A. Amedlous, A. Ait El Fakir, A. Jada and N. El Alem, *Crit. Rev. Environ. Sci. Technol.*, 2019, 1–42.
- 54 Z. Anfar, M. Zbair, H. A. Ahsaine, Y. Abdellaoui, A. A. El Fakir, E. H. Amaterz, A. Jada and N. El Alem, *ChemistrySelect*, 2019, **4**, 4981–4994.
- 55 V. Rocher, A. Bee, J. M. Siaugue and V. Cabuil, *J. Hazard. Mater.*, 2010, **178**, 434–439.
- 56 M. Şaban Tanyildizi, *Chem. Eng. J.*, 2011, **168**, 1234–1240.
- 57 M. Laabd, H. Ait Ahsaine, A. El Jaouhari, B. Bakiz, M. Bazzaoui, M. Ezahri, A. Albourine and A. Benlhachemi, *J. Environ. Chem. Eng.*, 2016, **4**, 3096–3105.
- 58 H. A. Ahsaine, M. Zbair and R. El Haouti, *Desalin. Water Treat.*, 2017, **85**, 330–338.
- 59 I. Langmuir, *J. Am. Chem. Soc.*, 1916, **38**, 2221–2295.
- 60 H. Freundlich, *Z. Phys. Chem.*, 1907, **57**, 385–470.
- 61 A. Benhouria, M. A. Islam, H. Zaghouane-Boudiaf, M. Boutahala and B. H. Hameed, *Chem. Eng. J.*, 2015, **270**, 621–630.
- 62 K. W. Jung, B. H. Choi, M. J. Hwang, T. U. Jeong and K. H. Ahn, *Bioresour. Technol.*, 2016, **219**, 185–195.

