Optimization of working cathode position in sleeve-type bioelectrochemical system with inner chamber/outer chamber for azo dye treatment

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HIGHLIGHTS

• Working position in sleeve-type BES was evaluated and optimized for decolorization.
• Cathode in outer chamber performed better both at \( V_{\text{cathode}:V_{\text{anode}} = 1:1} \) and \( 3:1 \).
• Current and EIS analysis showed outer cathode benefited proton/electron transfer.
• Lower decolorization with increased \( V_{\text{cathode}} \) might be due to less substrate supply.
• It would be further improved through increasing substrate concentration at anode.

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ABSTRACT

In this study, the optimization of working cathode position in sleeve-type bioelectrochemical system (BES) was evaluated with inner/outer chamber for azo dye decolorization. Results showed that the working position in outer chamber performed better with decolorization efficiencies of 97.8 ± 2.1% (7 h) and 94.0 ± 2.3% (16 h) than that in inner chamber as the volume ratio \( V_{\text{cathode}:V_{\text{anode}} = 1:1} \) and 3:1, respectively. The current and electrochemical impedance spectroscopy (EIS) analysis indicated that the proton/electron transfer and anolyte diffusion could be improved using outer chamber as working position. The decolorization with increased volume ratio could be further improved through the strategy of increasing substrate concentration, which would provide enough electrons and decrease diffusion resistance, further improving the whole performance with increased outer cathode volume. It has the great potential in sleeve-type configuration application and would create more challenges for process optimization and maintenance.

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1. Introduction

Bioelectrochemical system (BES) as a novel and promising approach, is very suitable for treating wastewater rich in unsaturated bonds (such as azo dye, antibiotics, pentachlorophenol, nitrobenzene, and heavy metals) at cathode, being established world widely as an alternative wastewater treatment process (Huang et al., 2012; Kong et al., 2015a, 2013; Mu et al., 2009a, 2009b; Tao et al., 2014). It can catalyze the degradation of organic matter at anode to provide electrons and protons for the cathode reduction and to reduce overpotential (Wang and Ren, 2013).

Considering the separation between the oxidation and the reduction reactions, dual chamber structures are required as suitable BES configuration to increase the antitoxic feasibility when the reactions take place in the processing of contaminants from wastewaters, resulting in a possibly whole range of application (Huang et al., 2012; Kong et al., 2015a, 2013; Mu et al., 2009a, 2009b; Tao et al., 2014).

It has reported that the effectiveness of BES will be highly dependent on the design choice (Li et al., 2014; Wang and Ren, 2013; Wang et al., 2012). The favorable BES application in the future will be obtained in the BES reactor with the lowest internal resistance, thus considerable effort should be applied to reduce the internal resistance (Sleutels et al., 2012). Recent research has modified the configuration of BES to be sleeve-type that with large area and small distance between anode and cathode, reducing the electrons transfer resistance and mass transfer resistance.
its advantages have been demonstrated in some cases, for example desalination, decolorization and the treatment of chromium (Huang et al., 2010; Jacobson et al., 2011; Kong et al., 2015b, 2015c). These studies indeed provided the feasibility of the BES with sleeve-type configuration to treat wastewater, however, as a sustainable and environment-friendly technology, the practical application of BES is strongly limited by their configuration optimum and the assembled optimization for operation (Ge et al., 2014; Gil-Carrera et al., 2011; Ping and He, 2013; Raman and Lan, 2012; Uria et al., 2012), which could provide important theory for further scale-up. The emerging alternative application of BES for recalcitrant pollutants has not yet been evaluated in optimizing the state of working position, with inner chamber or outer chamber. The working position in inner chamber or outer chamber can help us, not only to gain better understanding of the BES configuration, but also enable us to influence reaction properties.

Thus, aiming to find the best arrangement in the sleeve-type BES reactor for azo dye treatment, this work evaluated and optimized the working cathode position of outer or inner chamber at $V_{\text{cathode}} : V_{\text{anode}}$ of 1:1. Further optimization was also investigated in the increased volume ratio of 3:1. This will provide a theoretical basis and reference for future practical application.

2. Methods

2.1. Configuration of sleeve-type BES

The BES reactor was constructed in sleeve-type configuration as previously described with an inner tube and an outer tube using Perspex (Kong et al., 2013). The inner tube with full of holes was identified as the inner chamber when cation exchange membrane (Ultrex CMI7000, Membranes International Inc., USA) was pasted. The space between the inner tube and outer tube was identified as the inner chamber when cation exchange membrane was placed into the outer chamber and anode in the inner chamber. While on the contrary, the working position of BES-2# and BES-4# was the inner chamber, with cathode in the inner chamber and anode in the outer chamber. The cathode chamber and anode chamber of BES-1# and BES-2# have the same volume with cathode and anode volume ratio of 1:1 ($V_{\text{cathode}} : V_{\text{anode}} = 1:1$), and the BES-3# and BES-4# with $V_{\text{cathode}} : V_{\text{anode}} = 3:1$ were constructed to evaluate the influence of increased cathode volume on azo dye treatment. Both anode and cathode electrode materials were carbon brush (ID 2 cm × L 8 cm) displaced to be surrounding deployment as described previously (Fig. 1) (Kong et al., 2014).

The experiments were performed with the applied voltage of 0.5 V supplied with a DC power supply. For the reason that the anode potential was sustained at −0.4 V or even lower with acetate as the substrate, only with the applied voltage of 0.5 V, the cathode potential could be kept at −0.9 V, which was feasible for AO7 reduction decolorization (Mu et al., 2009a; Zille et al., 2004).

2.2. Operation of experiments

Sodium acetate (NaAc) was the substrate in the anode chamber used as the sole electron donor, and azo dye (acid orange 7, AO7) was treated in the cathode chamber used as the sole electron accepter. In principle, the acetate could produce protons and electrons at anode that transferred to the cathode for the azo bond cleavage of azo dye. Anode and cathode were inoculated with anaerobic sludge obtained from wastewater treatment plant in Harbin, China. The anolyte was NaAc 1 g L$^{-1}$ (expect for the effect of substrate concentration), KCl 0.13 g L$^{-1}$, NH$_4$Cl 0.31 g L$^{-1}$, PBS 50 mM, trace element 1 mL L$^{-1}$ and Wolf’s vitamin 1 mL L$^{-1}$ (with conductivity of 7.0 ± 0.5 mS cm$^{-1}$). The catholyte was AO7 100 mg L$^{-1}$, KCl 0.13 g L$^{-1}$, NH$_4$Cl 0.31 g L$^{-1}$, PBS 50 mM, trace element 1 mL L$^{-1}$ and Wolf’s vitamin 1 mL L$^{-1}$ (with conductivity of 7.0 ± 0.5 mS cm$^{-1}$). The anolyte and catholyte were refilled with fresh medium each 24 h. All tests were conducted and repeated at least five times. Sleeve-type bioelectrochemical reactors with inner chamber or outer chamber as working chamber were established for decolorization of azo dye. BES-1# and BES-2# with $V_{\text{cathode}} : V_{\text{anode}} = 1:1$ were investigated to evaluate the effect of working cathode position on sleeve-type BES performance if the electrons supply or anolyte diffusion that affected the performance (Mu et al., 2009a; Zille et al., 2004).

Meanwhile, BES-3# and BES-4# with $V_{\text{cathode}} : V_{\text{anode}} = 3:1$ were also compared to study the effect of working position on sleeve-type BES performance if increased the volume ratio of working chamber. In order to identify if the electrons supply or anolyte diffusion that affected the performance with outer cathode in BES-3# at $V_{\text{cathode}} : V_{\text{anode}} = 3:1$ and to improve the performance for scale-up, experiments were performed with increased acetate concentration from 1000 to 3000 mg L$^{-1}$ in the inner anode chamber.

2.3. Analytical methods

The AO7 concentration was measured at $\lambda_{\text{max}} = 484$ nm by UV–vis scanning spectrophotometer (Shimadzu UV2550, Japan). Anode and cathode as well as the reference electrode (saturated calomel reference electrode, SCE, model-217, Shanghai Precise, Sci. Instru. Co., Ltd., China; 0.247 V vs. standard hydrogen electrode, SHE) were connected to a data acquisition unit (keithley 2700, Keithley Co., Ltd., USA) with external resistance of 20 Ω to record electrode potentials and current. Electrochemical impedance spectroscopy (EIS) was carried in a frequency range of 10$^{-2}$–0.01 Hz with a perturbation signal of 10 mV using electrochemical workstation (model-660D, CH Instruments Inc., USA).

3. Results and discussion

3.1. The optimization of working cathode position in sleeve-type BES with $V_{\text{cathode}} : V_{\text{anode}} = 1:1$

In order to investigate the decolorization performance with different working positions in the sleeve-type BES, two BES reactors...
were firstly performed and compared at the volume ratio $V_{\text{cathode}}:V_{\text{anode}}$ of 1:1 to ensure the comparability, including BES-1# with outer chamber as the working position (i.e. cathode electrodes were placed in the outer chamber) and BES-2# with inner chamber as the working position (i.e. cathode electrodes were placed in the inner chamber) (Fig. 1). They were analyzed in terms of decolorization, current and EIS to obtain the decolorization behavior of sleeve-type BES with different working positions and the reason leading to the different results.

### 3.1.1. Decolorization performance

For BES with $V_{\text{cathode}}:V_{\text{anode}}$ = 1:1, results showed that the working cathode position in the outer chamber took obvious advantages compared to that in the inner chamber. The decolorization efficiency in the BES-1# with outer cathode was 97.8 ± 2.1% at 7 h, which was higher than that in BES-2# with inner cathode (90.9 ± 3.4%), and it would take 12 h to reach the similar decolorization efficiency of 97.2 ± 2.8% (Fig. 2A). HPLC analysis indicated that sulfanilic acid (SA) and 1-amino-2-naphthol (AN) were the dominant products of the AO7 decolorization at the cathode, demonstrating the cleavage of the azo bond of azo dye in BES. The formation efficiencies of SA were higher than 90% as over 98% AO7 decolorized simultaneously (data not shown). In order to investigate the dynamic characteristics for AO7 decolorization in BES, the decolorization behavior with different working positions was also analyzed in terms of kinetic study by fitting AO7 concentrations as a function of time. Results exhibited that AO7 decolorization in the BES conformed to the first-order model, and the corresponding reaction rate constant $k$ (h$^{-1}$) was 0.644 h$^{-1}$ in the BES-1#, which was 1.6-fold higher than BES-2# with inner cathode (0.404 h$^{-1}$) (Fig. 2A), further indicating the better decolorization in the BES-1# with outer cathode. Thus, the outer working position was based on a kinetic mechanism that accelerated the reaction rate, which was a significant kinetic factor in the BES (Liang et al., 2013).

These different decolorization results between BES-1# and BES-2# showed that the working position in the wastewater treatment performance, and working position in the outer chamber was better than that in inner chamber at $V_{\text{cathode}}:V_{\text{anode}}$ = 1:1.

### 3.1.2. Current

Current is a key parameter to describe conversion rate (Zhao et al., 2009). It was observed that the BES-1# generated a peak current of 0.022 A, while the peak current of BES-2# was 0.012 A, which was 1.8-fold decrease (Fig. 2B). According to that current was related to the electron flows, which were inherent to the microbial metabolism that microorganisms transfer electrons from an electron donor to an electron acceptor (Zhao et al., 2009), it was expected that a higher current might be attributed to the efficient electrolyte transport and enhanced electron transfer between the microbial biofilm and the electrode. These current results strongly suggested that the BES-1# with outer working position enabled superior performance due to an increase in electron transfer with current generation, which would result in faster azo dye decolorization (Fig. 2A).

Moreover, the current of BES-1# reached the maximum at the beginning and then gradually decreased (Fig. 2B). It has reported that the peak current was observed due to the favorable oxidation of the electron donor (Zhao et al., 2009), thus, the faster substrate degradation at inner anode that provided enough electrons and protons for the AO7 reduction at cathode, resulting in the maximum current in a short time. Moreover, the maximum current cannot be sustained because of the rapidly depleted donor near the electrode (Zhao et al., 2009), leading to the gradual decrease as shown in Fig. 2B. While for the BES-2#, the substrate degradation of anode was in the outer chamber, which might be constricted by the anolyte diffusion, performing the gradual increase in current. Similar to BES-1#, the subsequent decrease might be resulting from the finish of substrate degradation and diminished AO7.

These results indicated that the working position in the sleeve-type BES with $V_{\text{cathode}}:V_{\text{anode}}$ = 1:1 could affect the current with wastewater in the outer chamber or inner chamber. The working position in the outer chamber performed better performance in electron transfer, probably due to the faster substrate degradation at inner anode and enough electrons and protons transferred to the cathode for decolorization.

### 3.1.3. EIS

EIS analysis is another principal method to examine the electrochemical behavior of BES configuration (He and Mansfeld, 2009). In order to further analyze the reason of improved BES performance, EIS of BES-1# and BES-2# had been compared and discussed in the total resistances (Fig. 2C) and their individual components (Fig. 2D). Fig. 2C showed that EIS results of BES-1# and BES-2# were different as operated with different working positions. The total resistance of BES-1# was 35.8 ± 1.9-fold lower than that of BES-2# (66.9 Ω). The advantage of BES-1# with this lower total resistance was due to the combination of lower anode resistance and cathode resistance. Results showed that the anode resistance of BES-1# was 28.2 Ω, 31.1% lower than that of BES-2# (40.9 Ω), while the cathode resistance of BES-1# was 7.5 Ω, 70.8% lower than that of BES-2# (26.0 Ω) (Fig. 2C). It can be seen that the sleeve-type BES with working position in the outer chamber could reduce anode resistance and cathode resistance of BES reactor, especially cathode resistance.

According to that resistance was the ability of circuit element to hinder the flow of electrons (Kashyap et al., 2014), the anode resistance and cathode resistance of BES-1# and BES-2# were analyzed in detail through the components of resistance, including ohmic resistance ($R_{\text{ohm}}$), charge transfer resistance ($R_{\text{ct}}$) and diffusion resistance ($R_{\text{diff}}$) to find the reason of difference performance with different working positions (Fig. 2D). Firstly, for the anode resistances in BES-1# and BES-2#, the lower anode resistance of BES-1# might be due to the lower $R_{\text{ct}}$ and $R_{\text{diff}}$ of anode compared with BES-2#. The $R_{\text{ct}}$ of anode in BES-1# and BES-2# was 9.9 and 15.5 Ω, respectively, while the corresponding $R_{\text{diff}}$ was 13.2 and 19.4 Ω (Fig. 2D). The lower charge transfer resistance and diffusion resistance of anode in BES-1# demonstrated the enhanced charge transfer at anode and anolyte diffusion in the anode chamber. The improved anolyte diffusion would accelerate the substrate degradation and promote the produce of electrons that provided to cathode, resulting in the improved decolorization (Fig. 2A).

Secondly, for the cathode resistances in BES-1# and BES-2#, it indicated that the different cathode resistances in BES-1# and BES-2# were caused by the interaction of $R_{\text{ohm}}$, $R_{\text{ct}}$, and $R_{\text{diff}}$. The $R_{\text{ohm}}$, $R_{\text{ct}}$, and $R_{\text{diff}}$ of cathode in BES-1# were 0.8, 2.9 and 3.9 Ω, respectively, while those in BES-2# were 7.5, 6.8 and 11.7 Ω (Fig. 2D). Thus, the better BES-1# with working position in the outer chamber might be associated with the better proton/electron transfer related to the substrate degradation at anode. With the inner anode in BES-1#, the substrate was concentrated degraded in the center, with protons transferred from the center to the surrounding CEM, which making the relative large surface area of proton transfer. Moreover, the concentrated substrate in the center might have decreased diffusion resistance, which was benefit to the substrate degradation at anode and would promote electron supply for the cathode, finally resulting in the good decolorization performance (Fig. 2A).

These results indicated that in the sleeve-type BES with $V_{\text{cathode}}:V_{\text{anode}}$ of 1:1, the working position in the outer chamber was better than the inner chamber, which could improve the charge transfer and substrate diffusion in the anode chamber, and probably
enhance the supply of electrons and protons for AO7 reduction at cathode.

3.2. The optimization of working cathode position in sleeve-type BES with V_{cathode}:V_{anode} = 3:1

The optimization of working positions (BES-3# with outer cathode chamber and BES-4# with inner cathode chamber) was also evaluated at increasing V_{cathode}:V_{anode} of 3:1 to study the effect of working positions in BES with increased working volume.

3.2.1. Decolorization performance

In the BES with increasing volume ratio of V_{cathode}:V_{anode} = 3:1, BES-3# with outer cathode exhibited the better decolorization (94.0 ± 2.3% at 16 h), followed by BES-4# with inner cathode (84.2 ± 4.1% at 16 h) (Fig. 3A). The reaction rate constant k for BES-3# was 0.227 h^{-1}, which was 1.9-fold higher than BES-4# with inner cathode (0.117 h^{-1}) (Fig. 3A). It can be seen that the BES with outer working position took obvious advantage of decolorization in comparison with BES with inner working position at V_{cathode}:V_{anode} = 3:1. Based on that the k of BES-1# was 1.6-fold higher than that of BES-2# (Fig. 2A) and the k of BES-3# was 1.9-fold higher than that of BES-4# (Fig. 3A), this increase of 1.6-fold to 1.9-fold indicated that the BES with outer working position could play more predominant priority as V_{cathode}:V_{anode} increased within a certain range.

In all the BES reactors, BES-1#, BES-2#, BES-3# and BES-4# removed 97.8 ± 2.1% (7 h), 97.2 ± 2.8% (12 h), 94.0 ± 2.3% (16 h), and 91.2 ± 3.6% (24 h), respectively. The time needed for 90% decolorization was 4, 6, 12, and 24 h, respectively (Figs. 2A, 3A, and Table 1). It can be seen that the BES-1# with outer working position (V_{cathode}:V_{anode} = 1:1) exhibited the best decolorization performance, followed by BES-2# with inner working position (V_{cathode}:V_{anode} = 1:1), BES-3# with outer working position (V_{cathode}:V_{anode} = 3:1) and then BES-4# with inner working position (V_{cathode}:V_{anode} = 3:1). These results indicated that the decolorization was dependent on the working position in the sleeve-type BES with inner chamber and outer chamber. It should be noted that along with the increased volume ratio, the decolorization was decreased. Take BES with outer working position for example, the reaction rate kinetics were 0.644 h^{-1} for BES-1# with V_{cathode}:V_{anode} = 1:1 and 0.227 h^{-1} for BES-3# with V_{cathode}:V_{anode} = 3:1 (Table 1). Considering that substrate degradation at anode would supply protons and electrons for decolorization at cathode, the less effective decolorization might be related to the substrate in a short supply at anode for more AO7 decolorization when the cathode volume increased.

3.2.2. Current

BES-3# generated a higher peak current of 0.016 A than BES-4# (0.008 A) (Fig. 3B). The results also demonstrated that the BES with outer working position exhibited better performance than that with inner one. The current of BES-3# and BES-4# was increased gradually from the maximum current and then decreased. The lower current of BES in the early period (before 3.5 h for BES-3# and 5.6 h for BES-4#) might be limited by the substrate degradation. If the substrate degradation feeding electrons for the cathode was sufficient at the beginning of the process, it would be expected to be a sharp current increase as the substrate addition (Wang et al., 2011). The substrate degradation at anode might be depending upon the concentration of substrate and the diffusion of substrate (Montpart et al., 2015; Velasquez-Orta et al., 2011; Wang et al., 2013). These results also showed that the improved decolorization and enhanced current were not only caused by the anodic reaction,
but by the reaction rate of the azo dye at cathode surface which decisively affected on the current.

3.2.3. EIS

The total resistance of the BES-3# with the outer working position (49.0 Ω) was also lower than the BES-4# with inner one (79.4 Ω) as the $V_{\text{cathode}}:V_{\text{anode}}$ increased to 3:1 (Fig. 3C). It was similar to the comparison of BES-1# and BES-2# in the volume ratio of 1:1, further indicating the improved BES performance with outer cathode in the sleeve-type configuration. This also pointed out that as the volume ratio of cathode and anode chamber increased in a certain range, the working electrode placed in the outer chamber was more favorable. Moreover, the anode resistances of BES-3# and BES-4# were 37.1 and 50.0 Ω, respectively, and the corresponding cathode resistances were 11.9 and 29.4 Ω (Fig. 3C). It indicated that both the anode resistance and cathode resistance in BES-3# were lower than those in BES-4#. Thus, the performance of BES-3# was improved by the combined enhancement of anode and cathode. It would be further testified in detail by the analysis of components of anode resistance and cathode resistance as showed in Fig. 3D.

Results indicated that the different resistances between the two reactors were mainly resulted from the different $R_{\text{ohm}}$ and $R_{\text{diff}}$. The $R_{\text{ohm}}$ of anode in BES-3# and BES-4# was 7.6 and 13.8 Ω, respectively, and the corresponding $R_{\text{diff}}$ was 20.1 and 25.9 Ω. Meanwhile, the $R_{\text{ohm}}$ of cathode in BES-3# and BES-4# was 0.5 and 7.2 Ω, respectively, and the corresponding $R_{\text{diff}}$ was 7.1 and 15.2 Ω (Fig. 3D). The data suggested that the BES with outer working position caused a remarkable decrease in the $R_{\text{ohm}}$ and $R_{\text{diff}}$, indicating a better performance could be obtained through the optimization of working position and the BES with outer cathode was a favorable way for sleeve-type BES application in wastewater treatment.

According to that the $R_{\text{ohm}}$ was related to the electron transfer resistance through the electrode material and proton transfer in solution or through the membrane (He and Mansfeld, 2009), the BES-3# and BES-4# might be with different electrode distance and proton transfer distance due to the different diameters of inner tube and outer tube at the same $V_{\text{cathode}}:V_{\text{anode}} = 3:1$ (Fig. 1), which resulting in the different $R_{\text{ohm}}$. The different diameters also led to the different electrolyte diffusion, which resulting in the different diffusion resistance (Fig. 3D).

In conclusion, compared with BES with inner working position, the substrate diffusion in the anode chamber of BES with outer...
working position could be improved, accelerating the transport of substrate between solution and anode and then improving the substrate degradation which supplied protons and electrons for decolorization at cathode. Thus, the BES performance with outer working position performance might be further optimized by increasing the substrate concentration at anode or the electrolyte diffusion.

3.3. Further improvement through increasing substrate concentration

The above results demonstrated that the best decolorization performance within 7 h was observed in the BES with outer working position (97.8 ± 2.1%) in comparison with inner one (90.9 ± 3.4%) at V_{cathode}:V_{anode} = 1:1 (Fig. 2A). As increasing the volume ratio to be V_{cathode}:V_{anode} = 3:1, the BES with outer working position (77.6 ± 1.3%) was also taken superiority to the BES with inner one (61.2 ± 1.9%) (Fig. 3A). As can be seen, BES performance was inhibited as increasing the working volume ratio, which needed to be further improved to construct the application of BES with sleeve-type configuration. The current and EIS analysis indicated that the electron transfer and diffusion of anode were the limiting factors that affected the performance of BES with the relative large cathode volume (Figs. 2 and 3). Meanwhile the reduction of decolorization as increasing the substrate concentration, which was inhibited as increasing the working volume ratio, which resulted in the decreased decolorization might be due to the limited supply of electrons that provided by substrate degradation at anode. Thus, much effort should be made in improving its performance through increasing the substrate concentration.

The decolorization as a function of time was well described by first-order kinetics with different substrate concentrations from 1000 to 3000 mg L⁻¹. The results of the impedance measurement can be presented in two common ways: Nyquist plot (Fig. 4C) and the Bode plot (Fig. 4D). Nyquist plot expresses the impedance with a real part (plotted on the X-axis) and an imaginary part (plotted on the Y-axis that is negative). The Bode plot, on the other hand, shows the information of impedance and frequency (He and Mansfeld, 2009). In this study, the Nyquist curves of the BES were typical comprised by the semicircle high frequency region and ray low frequency region. The impedance at the high frequency limit was the R_{ohm}, the diameter of the semicircle was R_{ct}, and the ray at low frequency region was R_{diff} (You et al., 2007). EIS was closely related to the processes of substrate degradation which supplied protons and electrons for decolorization at cathode, thus, much effort should be made in improving its performance through increasing the substrate concentration.

In this study, the Nyquist curves of the BES were typical comprised by the semicircle high frequency region and ray low frequency region. The impedance at the high frequency limit was the R_{ohm}, the diameter of the semicircle was R_{ct}, and the ray at low frequency region was R_{diff} (You et al., 2007). EIS was closely related to the processes of substrate degradation which supplied protons and electrons for decolorization at cathode.
related with the reactor configuration and operation parameters. It has been reported that $R_{\text{ohm}}$ was greatly dependent on the design and configuration of the system rather than the microbial population at anode, which was predominant the total internal resistances of the system or even higher than 50% (Sekar and Ramasamy, 2013; You et al., 2008). It has demonstrated that increasing the sieve area could decrease the internal resistance and ohmic resistance to be 27.5 and 14 $\Omega$, respectively (You et al., 2008). In the sleeve-type BES with three different operation conditions, the average ohmic resistances were about 7.9 $\Omega$, which were relative low and only accounted for 25.4% of the average internal resistance (Fig. 4C). These results could further demonstrate that reducing the electrode spacing and increasing surface area could be effective solutions to reduce ohmic resistance (Sekar and Ramasamy, 2013). In comparison with substrate concentration of 1000 mg L$^{-1}$, the higher concentration of 2000 mg L$^{-1}$ showed significantly better electron transfer performance, with $R_{\text{ct}}$ of 7.5 $\Omega$, about 65.3% lower than that of 1000 mg L$^{-1}$ (12.4 $\Omega$) (Fig. 4C). It indicated that the biofilm at anode had enhanced electrochemical activity that could benefit to the electron transfer from microorganisms to anode. From the Bode curves (Fig. 4D), it was clearly demonstrated the better electron transfer (He and Mansfeld, 2009). Moreover, the diffusion resistance for 1000, 2000, 3000 mg L$^{-1}$ was 20.1, 12.0 and 8.4 $\Omega$, respectively (Fig. 4C). It also exhibited more and more advantages in the anolyte diffusion with increasing substrate concentration. Therefore, with the substrate concentration of 2000 mg L$^{-1}$, BES with outer working position showed the best decolorization and reaction rate in the combined action of improving electron transfer and anolyte diffusion. It should be noted that EIS was also related with operation parameters. For example, the bacterial biofilm, mediators and electrolyte $pH$ played an important role in anode impedence, meanwhile, the choices of the substrate type, substrate diffusion, mediators, cathode material type and cathode binding agent were also significant for cathode impedance (Sekar and Ramasamy, 2013).

These results demonstrated that BES with outer working position at an increased volume ratio could be further improved through increasing substrate concentration in a certain range.

3.4. Potential and significance

The versatility of the dual chamber BES has notably expanded the range of application for the treatment of refractory pollutants, especially contaminants those were toxic to microorganisms at anode and inhibited the effectiveness of treatment (Huang et al., 2012; Kong et al., 2015a, 2013; Mu et al., 2009a, 2009b; Tao et al., 2014). In recent years, BESs including microbial fuel cells (MFCs) and microbial electrolysis cells (MECs), have been explored extensively to treat azo dye for their innovative features and environmental benefits. In an MFC, electrical energy can be extracted from the electrical circuit. In an MEC, electrical energy needs to be supplied to the electrical circuit by a DC power supply (Mu et al., 2009a). The use of MFC takes advantage of the microbial oxidation of organic matter to generate bioelectricity simultaneously with wastewater treatment. BES can reduce the azo dye, driven by microbial oxidation of organics at the anode. It has been reported that the sleeve-type dual-chamber BES with cathode as working electrode achieved very good results in Cr (VI) removal and azo dye decolorization (Huang et al., 2010; Kong et al., 2013), confirming the priority of this novel configuration compared with other configurations, such as rectangular configuration and tubular configuration (Kong et al., 2014). However, it still posed a challenge on the understanding of this system and the reaction to determine the performance of BES. Taking into account that the sieve-type configuration was composed by two tubes, including the inner chamber consisted by the inner tube, and the outer chamber consisted by the space between inner tube and outer tube, thus, the pollutants position (i.e. the working position) played an important part in the wastewater treatment. It caused the reason analysis of different effects had also become the main point in understanding and study the new configuration. Therefore, this study was performed to investigate the effect of working position in inner chamber/outer chamber on decolorization of azo dye.

Results indicated that the cathode in outer chamber performed better decolorization compared to that in inner chamber at the cathode and anode chamber volume ratio of 1:1. To further investigate the effect of amplified cathode volume ratio, BES reactors with $V_{\text{cathode}}$:$V_{\text{anode}}$ of 3:1 were also compared. It showed that the working position in outer chamber was also much better than inner chamber. EIS results demonstrated that when the cathode was placed in the outer chamber (BES-1# and BES-3#), the total resistance of BES was decreased, as a result of decreased anode resistance and cathode resistance (Figs. 2C and 3C). Moreover, the detailed analysis of components in anode resistance showed the improvement of charge transfer resistance and diffusion resistance in BES with outer working position (Figs. 2D and 3D). It should be noted that with the increased cathode volume ratio, the performance was inhibited, possibly for the inadequate electrons supply with the decreasing anode volume at the same substrate concentration. It could be further improved by increasing the substrate concentration in anode chamber, through reducing the charge transfer resistance and diffusion resistance (Fig. 4C), leading to increased decolorization rate (Fig. 4A).

This study not only investigated the optimization of working position in outer chamber or inner chamber, but also evaluated and optimized the working electrode position in increasing volume of cathode chamber. It has very significant significance in configuration optimization and will promote the application of sleeve-type BES in wastewater treatment. It is expected that the broad platform described here will stimulate new thoughts and ideas which can then develop into a new generation of BES application.

4. Conclusion

This study demonstrated that the outer working position in the sleeve-type BES took obvious priority in azo dye decolorization with the same volume ratio of cathode and anode (1:1 or 3:1). The performance with increased volume ratio could be further improved through increasing the substrate supply. It provided the basic theory for the optimization of BES with sleeve-type configuration and would have a great potential in the application.

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