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Denitrification of groundwater using a biodegradable polymer as a carbon source: long-term performance and microbial diversity

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Nitrate pollution in groundwater is a worldwide problem. This paper reports on the denitrifying performance of using the biodegradable polymer polybutylene succinate (PBS) as a biofilm medium and carbon source to remove nitrate from groundwater *via* a packed bed bioreactor which was operated continuously for nearly 2 years. Results showed that the effluent nitrate concentration reached 3.3–8.8 mg L⁻¹ and 88–97% of nitrate removal was achieved. The denitrification rate range was 0.25–0.35 g N per L per d at 20–29 °C and decreased to 0.12 g N per L per d at 10–18 °C. According to microelectrode analysis, the nitrate consumption rate (1069 ± 103 μmol cm⁻¹ h⁻¹) was much higher than the ammonium production rate (74 ± 7 μmol cm⁻¹ h⁻¹), which proved that denitrification plays the major role in the system. A low level of DOC (1.7 ± 0.6 mg L⁻¹) and ammonium (0.5 ± 0.3 mg L⁻¹) was observed in the effluent, which was beneficial for practical application. The consumption rate of PBS was 2.75 ± 0.72 g PBS/g NO₃-N_{removed}. In the attached biofilm, *Proteobacteria*, *Betaproteobacteria*, *Burkholderiales* and *Comamonadaceae* were the major phyla (75.6%), classes (59.8%), orders (42.3%) and families (42.2%) in each level. In the top 20 genera accounting for 25% of total sequences, 9 genera including *Simplicispira*, *Comamonadaceae*, *Hydrogenophaga* and *Rhodocyclaceae* were affiliated with denitrifying groups with an abundance of 16%, whereas the bacteria belonging to the other 11 genera including *Veillonellaceae*, *Propionivibrio* and *Bdellovibrio* were reported to have the function of degradation and acidification of organic substance and might serve for degrading PBS in the system. The PBS solid-phase denitrification is promising for removing nitrate from groundwater.

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Introduction

Nitrate is naturally present in the environment as part of the nitrogen cycle. However, anthropogenic inputs, such as the agricultural application of nitrogenous fertilizers and wastewater emission, have greatly increased the nitrate loads in receiving waters, leading to potential risks to the shallow and deep groundwater aquifers.^{1–3} Drinking nitrate-contaminated groundwater has the risks of methaemoglobinaemia for infants and stomach cancer.⁴ The maximum permissible level of nitrate in drinking water was set as 10 mg N per L by the U.S. Environmental Protection Agency⁵ and in the standard for drinking water quality of China (GB 5749-2006) to reduce the risks to human health.

Compared to the methods commonly used for nitrate removal such as ion-exchange, distillation, reverse osmosis and electro dialysis, heterotrophic biological denitrification is a cost-

effective and practical on large-scale technology.⁶ The denitrifying microbes utilize organic substrates such as methanol, glucose and acetic acid that serve as electron donor and energy source, and convert nitrate into nitrogen gas under anoxic conditions.^{7,8}

In recent years, solid-phase denitrification, which uses water insoluble solid materials as carbon substrates for denitrification and biofilm media, has been developed to eliminate nitrate from groundwater *in situ* or *ex situ*,^{9–11} aquaculture effluent,^{12,13} and biologically treated effluent.¹⁴ There exist two kinds of solid carbon source: the natural plant materials such as woodchips, straws and reeds^{15,16} and the synthetic biodegradable polymers such as polycaprolactone (PCL), polybutylene succinate (PBS), polylactic acid and polyhydroxyalkanoate.^{17–22}

Healy *et al.*¹¹ investigated nitrate removal from groundwater by solid-phase denitrification using woodchips, pine needles, barley straw and cardboard, respectively as carbon source and media. The nitrate removal efficiency reached 67–89% at steady-state operation and the bioreactor filled with woodchips demonstrated less adverse effects such as dissolved organic carbon (DOC) release. Gibert *et al.*²³ built a permeable reactive barrier filled with softwood for *in situ* groundwater remediation. More than 98% of nitrate removal was obtained with the

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denitrification rate of 0.067 mg N per L per d per $g_{\text{substrate}}$. Wu *et al.*²⁴ reported that the nitrate removal reached 95% in a solid-phase denitrification system using PBS as carbon source to treat drinking water at 25 °C. Boley *et al.*²⁵ studied the treatment of wastewater from a recirculating aquaculture system with eels by a fluidized bed reactor filled with PCL for denitrification carbon source. The effluent nitrate concentration remained at low levels of 3–10 mg N per L, whereas a nitrate accumulation of 200 mg N per L was observed in the control system without denitrification treatment.

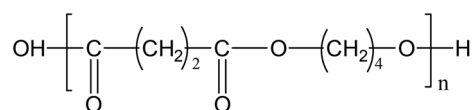
Solid-phase denitrification using biopolymers as carbon source is especially attractive in groundwater treatment due to the good and stable denitrifying performance and effluent quality such as low levels of DOC and color release.²⁶ The high production cost of the biopolymers is the major factor to limit its application. Among the biopolymers available, PBS denitrification is drawing increasingly attention owing to the good biodegradability and economics.^{24,27} In addition, most studies related to biopolymer solid-phase denitrification are operated in batch mode or continuously for a short time (3–8 months). For commercial application, the denitrifying performance and degradation properties of biopolymers during a long-term run should be studied.

In the present study, a PBS-packed bioreactor was established to eliminate nitrate in groundwater. The long-term performance of the bioreactor for nearly 2 years was evaluated in terms of nitrate removal, denitrification rate, and the quality of the treated groundwater. The characteristics of PBS degradation and the biofilm attached as well as the bacterial community were analyzed. The results of this study could enrich our understanding on solid-phase denitrification and provide a reference for its practical application in the management of groundwater nitrate pollution.

Materials and methods

PBS media

The granules of PBS used are cylindrical with an average diameter of 3.0 mm. The density and specific surface are 1.24 kg L⁻¹ and 0.826 m² g⁻¹, respectively. The chemical structure of PBS is as follows:



Experimental set-up and procedures

A packed bed reactor consisting of a cylindrical Plexiglas vessel (working volume 3.0 L) was established (Fig. 1). The packing ratio of PBS granules was 56.3% (v/v). A sieve was inserted into the reactor to prevent washout of PBS granules. Influent was continuously pumped into the bottom of reactor and effluent was discharged from overflow. The denitrification reactor was seeded with anoxic sludge collected from a municipal wastewater treatment plant with the concentration of 1500 mg total suspended

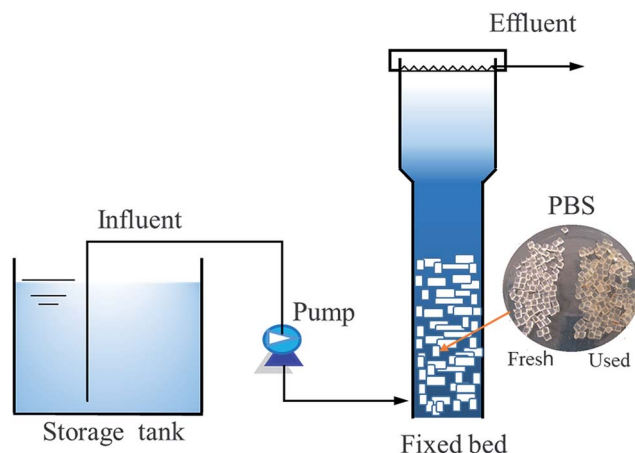


Fig. 1 The schematic diagram of the experimental set-up.

solids (TSS) per L. The initial hydraulic retention time (HRT) was 41 h and temperature was 20 °C. During the long-term operation of nearly 2 years, the temperature fluctuated in the range of 29–10 °C with the seasons (Fig. 2). The HRT was decreased gradually to 4.5 h and then varied in 5.6–13 h, depending on the performance of the reactor at different temperatures. Samples were periodically taken throughout the experiment to evaluate NO₃-N, NO₂-N and NH₄-N and DOC content in the effluent. Biofilm samples were taken at about 500 days to do DNA extraction, PCR amplification and pyrosequencing.

Influent characteristics

The groundwater was exploited from Changping district, Beijing. The content of NO₃-N, NO₂-N and NH₄-N and DOC were 10–13 mg L⁻¹, less than 0.09 mg L⁻¹, 0.17 mg L⁻¹ and 0.3 mg L⁻¹, respectively. The following metal ions (mg L⁻¹) were detected: Ca²⁺, 69; Mg²⁺, 41; Na⁺, 13; K⁺, 2.1; Mn²⁺, 0.2. The groundwater was fed into the PBS bioreactor by spiking with NaNO₃ and KH₂PO₄ to reach the influent NO₃-N concentration

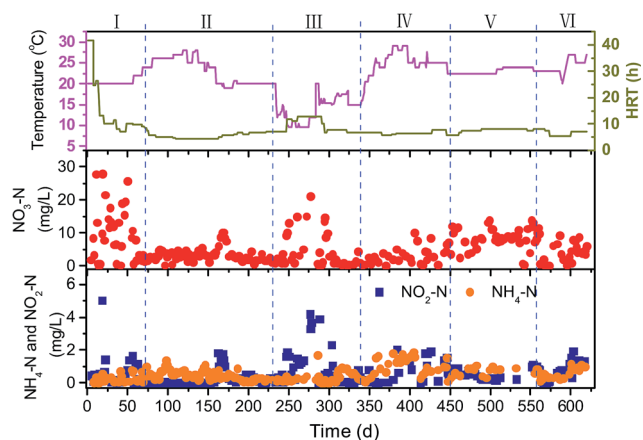


Fig. 2 Variation of temperature, HRT and effluent nitrogen concentrations with time.



of $68 \pm 2 \text{ mg L}^{-1}$ and TP levels of $1.1 \pm 0.2 \text{ mg L}^{-1}$, respectively. The pH value of the influent was 7.8 ± 0.1 .

Batch assays

After a certain period (1–2 months), batch tests were conducted to determine the specific denitrification rate (SDNR) of the biofilm attached to PBS media as described in previous studies.²⁸ The temperature was kept at $28.5 \text{ }^\circ\text{C}$. The SDNR was determined using linear regressions fitted to plots of decreasing nitrate concentration vs. time divided by the initial volatile suspended solid (VSS) content (mg N/g VSS per d). The changes in the weight per piece of PBS granule and the biofilm concentration (g VSS per L) were also evaluated.

Analytic methods

$\text{NH}_4\text{-N}$, $\text{NO}_2\text{-N}$ and $\text{NO}_3\text{-N}$ were measured according to Chinese SEPA Standard Methods.²⁹ DOC was determined by a multi N/C 2100 model TOC/TN analyzer (Analytic Jena, Germany). Three-dimensional excitation-emission matrix (3D-EEM) fluorescence spectroscopy was applied to analyze the characterization of dissolved organic matter in the effluent using a spectrometry (F-7000 fluorescence spectrophotometer, Hitachi Japan).

The molecular weight (MW) distribution of organic matters in the effluent was evaluated by a HPLC system (LC-20A, Shimadzu, Japan) equipped with a UV detector and an SEC column (TSK-GEL G3000PWXL). Sodium polystyrene sulphonate (3–30 kDa) and acetone (58 Da) were used as the standard solutions for M_w calibration curve construction.

The functional groups of PBS granules were characterized using one NTS FT-IR spectrometer (PerkinElmer Spectrum). The structure of the PBS granules and biofilm attached was observed by a JSM- 6700F scanning electron microscopy (SEM, JEOL, Japan).

Biofilm adhered to the PBS granules was collected by ultrasonic (40 kHz, 5 min). The content of TSS and VSS of the biofilm samples were evaluated using gravimetric method.²⁹ The extracellular polymer substances (EPS) were analyzed by the formaldehyde-NaOH extraction method.³⁰ The content of carbon, oxygen, nitrogen and hydrogen element was quantified by an Vario EL element analyzer (Elementar, Germany).

For analysis of bacterial communities in the biofilm samples, after DNA extraction and purification, PCR amplification, amplicon quantitation and pooling, the pyrosequencing was conducted using a 454 Life Sciences Genome Sequencer FLX instrument (Roche, NJ, USA) by Majorbio Bio-pharm Biotechnology Company (Shanghai, China). Analysis of sequencing data were performed using Mothur software.

The liquid-membrane microelectrodes sensitive to nitrate anions and ammonium ions were prepared, respectively to detect the profiles of nitrate and ammonium within biofilm. Briefly, the nitrate and ammonium microelectrodes with a tip diameter of $5 \text{ }\mu\text{m}$ and $15 \text{ }\mu\text{m}$, respectively were moved from the surface to the inner part of biofilm by a motor-driven micromanipulator. The output signals were transformed and recorded by computer using SensorTrace Pro 2.0 software (Unisense, Denmark). The

nitrate consumption rate and ammonium production rate were calculated using Fick's second law of diffusion.³¹ The applied molecular diffusion coefficients of nitrate and ammonium were 1.1×10^{-5} and $1.38 \times 10^{-5} \text{ cm}^2 \text{ s}^{-1}$, respectively.³²

Results and discussion

Performance of PBS-packed bed bioreactor

Fig. 2 shows the variation of operational parameters and the overall performance of the PBS-packed bed reactor with respect to the concentrations of effluent $\text{NO}_3\text{-N}$, $\text{NH}_4\text{-N}$ and $\text{NO}_2\text{-N}$ during the long-term operation of 620 d. The operational conditions and denitrification rate in the different phases were summarized in Table 1.

The effluent $\text{NO}_3\text{-N}$ concentration decreased on the third day from start-up and then HRT was reduced gradually. After 2 months of operation (Phase 1), the reactor was started up successfully and $\text{NO}_3\text{-N}$ removal reached 90%. Thereafter, the reactor was operated at temperatures of $20\text{--}28 \text{ }^\circ\text{C}$ and HRT of 5–7 h (Phase 2), the effluent nitrate maintained at low levels of around 3.0 mg L^{-1} with $\text{NO}_3\text{-N}$ removal efficiency of higher than 95%. The denitrification rate reached the highest level of $0.35 \text{ g N per L per d}$. As the operation temperature dropped to $10\text{--}18 \text{ }^\circ\text{C}$ (Phase 3), the nitrate removal efficiency decreased to 87.7% even extending HRT to 12–13 h. A nitrite accumulation was also found. The denitrification rate decreased by 65% compared to that in phase 2. When the temperature of the reactor rose to more than $20 \text{ }^\circ\text{C}$ (Phase 4), the nitrate removal efficiency and denitrification rate recovered to more than 95% and $0.25 \text{ g N per L per d}$, respectively. After 1.5 years of experiment (Phase 5), the concentration of effluent $\text{NO}_3\text{-N}$ increased remarkably, which might be attributed to the great degradation of PBS carbon source and the increased biofilm density which limited the mass transfer of both nitrate and substrate. Therefore, 200 g of fresh PBS granules, which equals to 15% of the initial amount, was replenished to the reactor (Phase 6). The effluent nitrate decreased rapidly and 94% of nitrate removal was obtained after PBS addition. It is obvious that renewal of the fresh carbon source after some period of time is necessary for the continuous operation.

Only a small increase in effluent DOC of $1.7 \pm 0.6 \text{ mg L}^{-1}$ was observed during the long-term operation, which is much lower than that using PCL of $1.7\text{--}5.2 \text{ mg L}^{-1}$ (ref. 9) and poly-3-hydroxybutyrate-co-hydroxyvalerate (PHBV) of $14.9 \pm 5.5 \text{ mg L}^{-1}$ (ref. 22) as carbon source for denitrification in treating the similar groundwater. This is promising for practical applications because the burden of post treatment to polish the effluent is reduced. The MW distributions of the DOC in the effluent revealed four fractions, 3.31, 2.32, 1.27 and dominant 0.83 kDa. The calculated M_w and M_n were 1.55 and 1.14 kDa, respectively. The effluent mainly contained soluble microbial products-like substances from 3D-EEM spectrum (spectrum not shown). The pH values and TP concentration in the effluent were 8.0 ± 0.1 and $0.5 \pm 0.2 \text{ mg L}^{-1}$, respectively.

It is reported that temperature is one of the important factors controlling solid-phase denitrification, in addition to HRT, influent nitrate concentration and pH, dissolved oxygen (DO),



Table 1 The operational parameters and performance of the PBS-packed reactor in the different phases

	Phase 1	Phase 2	Phase 3	Phase 4	Phase 5	Phase 6
Conditions						
Time (d)	0–70	71–231	232–335	336–449	450–560	561–620
Temperature (°C)	20–22	20–28	10–18	20–29	23–24	23–27
HRT (h)	41–10	5–7	12–13	6–8	6–8	6–8
Effluent						
NO ₃ -N (mg L ⁻¹)	10.7 ± 9.1	3.0 ± 1.5	8.8 ± 6.9	2.2 ± 1.9	8.4 ± 3.2	3.6 ± 2.8
NO ₂ -N (mg L ⁻¹)	1.3 ± 2.2	0.1 ± 0.1	1.8 ± 1.2	0.5 ± 0.5	0.57 ± 0.27	0.79 ± 0.61
NH ₄ -N (mg L ⁻¹)	0.3 ± 0.3	0.5 ± 0.3	0.2 ± 0.1	0.8 ± 0.3	0.72 ± 0.23	0.62 ± 0.33
Nitrate removal (%)	89.4 ± 8.6	96.1 ± 2.6	87.7 ± 11.5	96.9 ± 2.8	86.5 ± 5.0	94.2 ± 4.2
Denitrification rate (g N per L per d)	0.16 ± 0.02	0.35 ± 0.03	0.12 ± 0.02	0.25 ± 0.04	0.17 ± 0.03	0.22 ± 0.04

etc.^{26,33} The DO concentration detected in the influent was around 1.5 mg L⁻¹, which was consumed quickly through PBS-packed bioreactor, leading to low DO levels in the effluent (less than 0.3 mg L⁻¹) that was appropriate for denitrification. The denitrification rate in the PBS-packed reactor was lower by 52–66% at temperatures of 10–18 °C than that at 20–29 °C. The solid carbon source could not be utilized directly by denitrifying bacteria but needs to be hydrolyzed into soluble and micro-molecular substrates. Both the hydrolysis efficiency of PBS and the activities of the denitrifying microbes declined with decreasing temperature,³⁴ leading to a reduction in denitrification rate. Our previous studies showed that the denitrification rate decreased by around 50% with a 5 °C reduction in PCL-packed denitrification reactor.⁹ Cameron and Schipper³⁵ documented that the NO₃-N removal rate was raised by 1.7 times when temperature was increased by 10 °C using the softwoods-packed denitrification bed.

Nitrite is an intermediate of denitrification process. As shown in Fig. 2 and Table 1, NO₂-N accumulation was observed in the effluent during the initial stage (Phase 1, around 1.3 mg L⁻¹) and the operation at the lower temperatures (Phase 3, around 1.8 mg L⁻¹) owing to the incomplete denitrification. In the most period, the NO₂-N concentration in the effluent were less than 0.9 mg L⁻¹. The concentration of leached ammonium in the effluent was at low levels (0.5 ± 0.3 mg L⁻¹) throughout the total operation. This indicates that heterotrophic denitrification of reducing nitrate to nitrogen gas was predominant in the system using PBS as carbon source. It is known that there is another pathway, dissimilatory nitrate reduction to ammonium (DNRA), which competes with denitrification for nitrate.³⁶ Microelectrode data indicated that ammonium production was detected at the biofilm-media interface and NH₄-N concentration decreased toward the bulk surface, while NO₃-N concentration decreased gradually from the biofilm-bulk interface toward the biofilm bottom (Fig. 3). The calculated nitrate consumption rate (1069 ± 103 μmol cm⁻¹ h⁻¹) was much higher than the ammonium production rate (74 ± 7 μmol cm⁻¹ h⁻¹), which proved that denitrification plays the major role in the system.

Characterization of PBS degradation and biofilm attachment

From batch test, the weight of each piece of PBS granule declined almost linearly over the course of time (Fig. 4). The

degradation rate of PBS was around 0.04 mg d⁻¹. If the useful life of the solid carbon source is 50% of its initial weight, the PBS granules (with the initial weight of around 0.0289 g per piece) would last around 1 year, after that period refreshing some carbon source seems to be needed.

From SEM observation, the fresh PBS granule has relatively smooth surface with some protuberances (Fig. 5a), while the used PBS has concave cavities induced by microbial corrosion (Fig. 5b). According to FT-IR spectra (Fig. 6), PBS exhibits the characteristic adsorption at 2961 cm⁻¹, 2944 cm⁻¹ and 2850 cm⁻¹ due to CH₂ stretching vibration, 1713 cm⁻¹ assigned to carbonyl group, 1174 cm⁻¹ and 955 cm⁻¹ due to C–O stretching vibration and 1044 cm⁻¹ assigned to O[CH₂]₄O group. After PBS was used as carbon source, a broad peak in 3000–3750 cm⁻¹ region assigned to hydroxyl group has become obviously, indicating that there existed small fragments from PBS hydrolysis. The shape and position of the characteristic adsorption of the used PBS were similar to that of the fresh PBS, whereas the peak intensities declined greatly due to biodegradation.

When acting as carriers, PBS favors the attachment of biofilm which was composed of cocci and rod shape bacteria with some filamentous bacteria (Fig. 5c). The amount of biofilm

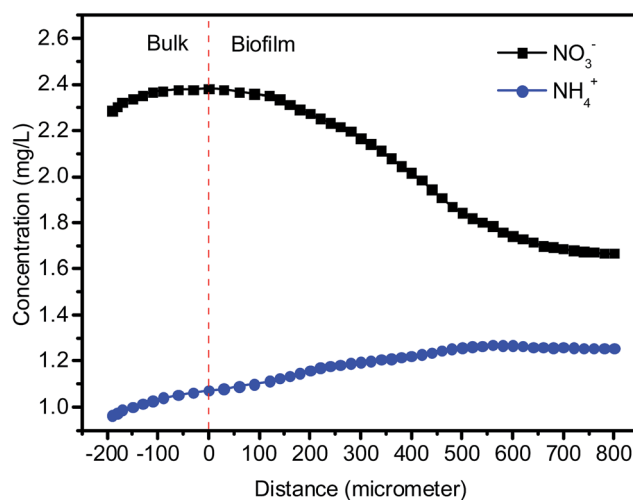


Fig. 3 Micro-profiles of nitrate and ammonium concentrations.



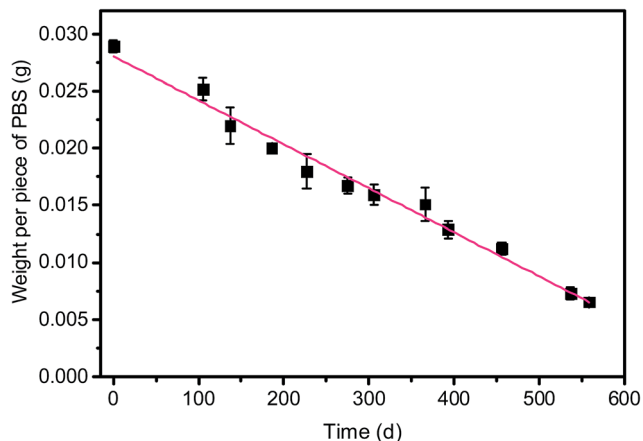


Fig. 4 Changes in weight per piece of PBS with operation time.

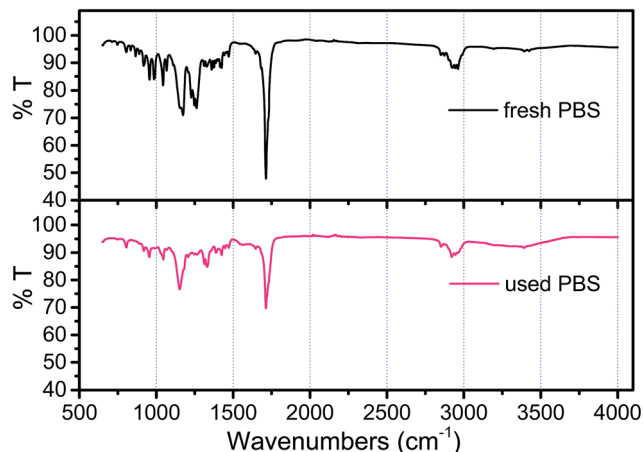
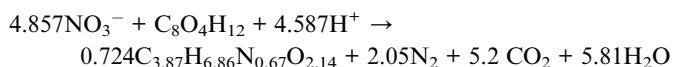


Fig. 6 FT-IR spectrum of the fresh and used PBS.

attached to PBS media increased gradually with the prolonged operating time and reached around 16 g VSS per L after 1 year of operation. With the increase in the biofilm density the mass transfer of both nitrate and substrate might be limited. Consequently, the SDNR of biofilm attached initially decreased remarkably and then stabilized at around 0.03 mg N/g VSS per d (Fig. 7). It is noted that the VSS/TSS ratio of biofilm reached more than 90%, indicating the high activity of biofilm attached to the biodegradable media. Similar results were found in a denitrification reactor using PCL as carbon source.⁹ In the EPS composition, the content of protein (129 ± 12 mg per g VSS) was much higher than that of polysaccharides (41 ± 5 mg per g VSS), which might be owing to the excretion of cytosolic proteins or cell lysis.³⁷

When PBS was utilized as carbon source for denitrification, the reaction of PBS monomer with nitrate leads to the production of carbon dioxide, nitrogen gas and biomass. The elemental composition of biomass was $C_{3.87}H_{6.86}N_{0.67}O_{2.14}$ from the elemental analysis of biofilm. Assuming a yield 0.45 g biomass/g PBS,³⁸ the equation of denitrification reaction can be given as:



The theoretical amount of PBS consumed in removing a gram of NO_3^-N was 2.68 g PBS/g NO_3^-N . The practical PBS consumption rate (2.75 ± 0.72 g PBS/g NO_3^-N) was slightly higher than the theoretical value. Similar results were also found by other researchers who utilized PCL and poly-3-hydroxybutyric acid (PHB) as denitrifying carbon source.^{13,25} This difference might be due to the residual oxygen consumption on carbon source. In the presence of dissolved oxygen, a portion of carbon source was consumed by aerobic respiration instead of denitrification.²⁵

The unit price of the domestic PBS products in China ranges 18–25 CNY/kg. The calculated denitrification cost using PBS as carbon source was 49–68 CNY/kg NO_3^-N , which are higher than using methanol.²² If the economic PBS polymers were available with the development of bioplastic industry, PBS solid-phase denitrification might be competitive in practical application.

Microbial community analysis

The numbers of effective sequence tags obtained from pyrosequencing were 13 950. Fig. 8 shows the bacterial community and abundance identified at the phylum, class, order and family levels in biofilm attached. At the phylum level, *Proteobacteria* was predominant with relative abundance of 75.6%, followed by

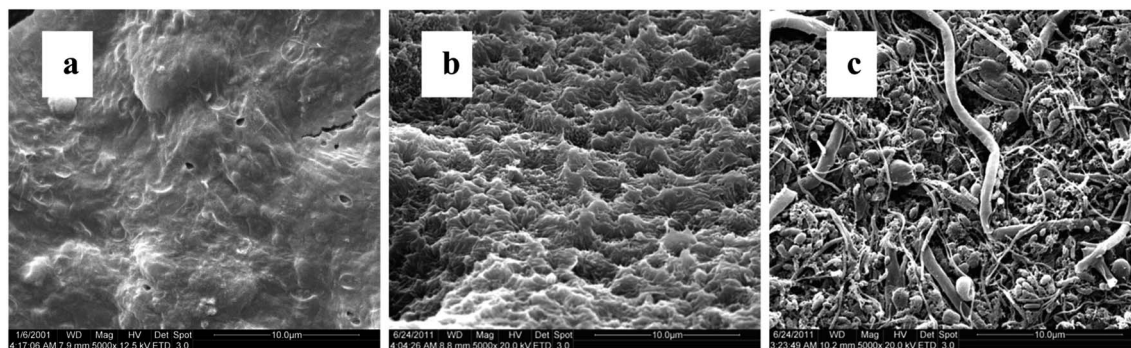


Fig. 5 SEM images of PBS granule, fresh (a) and used (b) and biofilm attached (c).



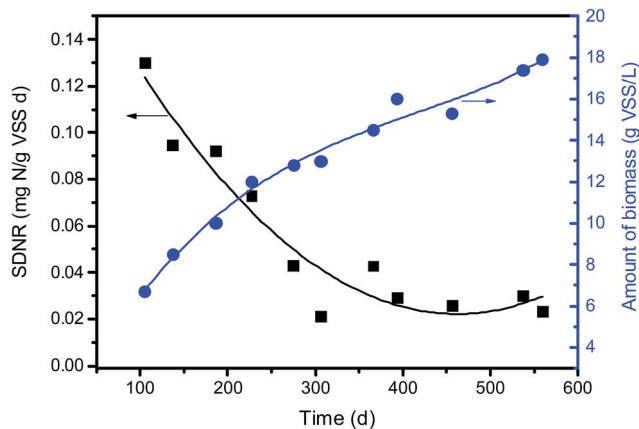


Fig. 7 Changes in SDNR and the amount of attached biomass with time.

Bacteroidetes (5.62%), *Firmicutes* (4.32%), *Chlorobi* (4.14%) and *Spirochaetes* (0.8%). *Betaproteobacteria*, *Burkholderiales* and *Comamonadaceae* were the major classes (59.8%), orders (42.3%) and families (42.2%) in each level, respectively. Hiraishi and Khan³⁸ have documented that most of the PHA-degrading denitrifying bacteria were affiliated to *Betaproteobacteria* and family *Comamonadaceae*. Khan *et al.*³⁹ have isolated 23 strains of PHBV-degrading denitrifying bacteria from different activated sludge. 16S ribosomal DNA sequence analysis demonstrated that 20 of the strains belonged to the family *Comamonadaceae*.

At the genus level, the identified bacteria of the major OTUs, the relative abundance and function reported were shown in Table 2. It is worthwhile to note that there were 71% of sequences unclassified in the biofilm samples. In the top 20 OTUs accounting for 25% of the total sequence, 9 OTUs

involving *Simplicispira*, *Comamonadaceae*, *Hydrogenophaga* and *Rhodocyclaceae* belonged to denitrifying groups with an abundance of 16%. All the genera of identified denitrifiers affiliated with phylum *Proteobacteria*. The abundance of genus *Simplicispira* was the highest. It was reported bacteria belonging to *Simplicispira* are able to reduce nitrate to nitrite and have been isolated from activated sludge in treating wastewater for inorganic nutrient removal.⁴⁰ Ruan *et al.*⁴¹ also reported that the denitrifiers *Azoarcus* and *Simplicispira* were dominant in a recirculating aquaculture system filled with PBS as the biofilm carrier and carbon source for simultaneous removal of ammonium and nitrate. In addition, genus *Desulfovibrio* was found in the pyrosequencing analysis (0.6%), which was reported to be able to reduce nitrate to ammonium.⁴² This is the reason why ammonium was detected in the effluent.

The denitrifiers using the solid carbon source were found to be different from that using the soluble carbon substrate. The denitrifying bacteria in the genera *Diaphorobacter*, *Simplicispira* and *Comamonas* which were found in the present study and *Acidovorax* belonging to family *Comamonadaceae* were commonly dominant in the solid-phase denitrification system.^{12,43,44} The genera *Pseudomonas*, *Paracoccus*, *Alcaligenes* and *Bacillus* are the important groups of denitrifiers in the traditional denitrification system using soluble carbon source such as methanol and ethanol.^{45–47}

As shown in Table 2, the bacteria including *Veillonellaceae*, *Propionivibrio* and *Bdellovibrio*, belonging to other 11 genera with an abundance of 9% were reported to have the function of degradation and acidification of organics. In the present study, they might serve for degrading PBS. The denitrifying microbes are not capable of utilizing the macromolecular PBS directly. In the solid-phase denitrification system there are microbes which secrete some hydrolases such as protease and lipases to

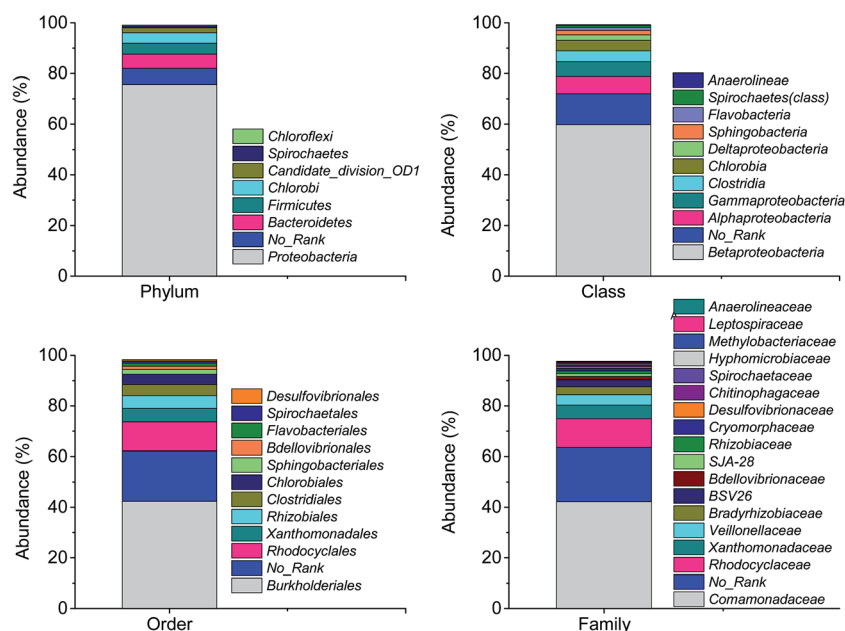


Fig. 8 The bacterial community and abundance at phylum, class, order and family levels in biofilm attached.



Table 2 Bacterial identification of the major OTUs at genus level, the relative abundance, the phylogenetic affiliation and function^a

OTU	Abundance (%)	Genus	Phylum/family	Function	Ref.
1	5.48	<i>Simplicispira</i> [★]	<i>Proteobacteria/Comamonadaceae</i>	Reduce nitrate to nitrite	40
2	4.14	<i>Veillonellaceae</i> uncultured	<i>Firmicutes/Veillonellaceae</i>	Fermentation of sugars, carboxylic acids and alcohols	49
3	3.86	<i>Comamonadaceae</i> uncultured [★]	<i>Proteobacteria/Comamonadaceae</i>	Denitrification	39
4	2.95	<i>Hydrogenophaga</i> [★]	<i>Proteobacteria/Comamonadaceae</i>	PHBV/PBS-degrading denitrifiers	24 and 39
5	1.88	<i>Propionivibrio</i>	<i>Proteobacteria/Rhodocyclaceae</i>	Fermentative acidogenic bacteria	50
6	1.25	<i>Rhodocyclaceae</i> uncultured [★]	<i>Proteobacteria/Rhodocyclaceae</i>	Reduce nitrate to nitrogen gas	51
7	1.08	<i>Bdellovibrio</i>	<i>Proteobacteria/Bdellovibrionaceae</i>	Degrade macromolecules of their prey	52
8	0.95	<i>Rhizobium</i> [★]	<i>Proteobacteria/Rhizobiaceae</i>	Anaerobic denitrification	53
9	0.75	<i>Owenweeksia</i>	<i>Bacteroidetes/Cryomorphaceae</i>	Aerobic oxidation metabolism	54
10	0.59	<i>Desulfovibrio</i> [★]	<i>Proteobacteria/Desulfovibrionaceae</i>	Reduce nitrate to ammonium	42
11	0.39	<i>Azospira</i> [★]	<i>Proteobacteria/Rhodocyclaceae</i>	Denitrifier isolated from activated sludge	55
13	0.36	<i>Meganema</i>	<i>Proteobacteria/Methylobacteriaceae</i>	Filamentous bacteria isolated from bulking activated sludge	56
14	0.35	<i>Spirochaetaceae</i> uncultured	<i>Spirochaetes/Spirochaetaceae</i>	Chemo-organotrophic, utilize carbohydrates or amino acids as carbon and energy sources	57
15	0.29	<i>Blastochloris</i>	<i>Proteobacteria/Hyphomicrobiaceae</i>	Phototrophs and nitrogen-fixers	58
16	0.26	<i>Ideonella</i>	<i>Proteobacteria/Comamonadaceae</i>	Enable to grow anaerobically by reducing chlorate to chloride	59
17	0.24	<i>Anaerolineaceae</i> uncultured	<i>Chloroflexi/Anaerolineaceae</i>	Degrade carbohydrate	60
18	0.22	<i>Leptospiraceae</i> uncultured	<i>Spirochaete/Leptospiraceae</i>	Free-living saprophyte found in fresh water	61
19	0.12	<i>Diaphorobacter</i> [★]	<i>Proteobacteria/Comamonadaceae</i>	PHB-degrading denitrifying	62
20	0.1	<i>Thiothrix</i> [★]	<i>Proteobacteria/Thiotrichaceae</i>	Reduce nitrate to nitrite and/or further to N ₂ O	63

^a The bacteria with ★ represent denitrifiers.

decompose PBS to small soluble monomers.^{17,26} The degraded products were then utilized by denitrifying bacteria to achieve biological denitrification. It should be noted that some species of genus *Diaphorobacter* which was detected in the PBS solid-phase denitrification system, were reported to be characterized with both degrading PHBV and denitrification.⁴⁸ In addition, no methanogen microbes were observed in the current work. It is reported that in the solid-phase denitrification process, the organic solid substances might be degraded by anaerobic digestion, leading to the production of CO₂ and CH₄.²⁶ In such cases, the methanogens might develop in the inner part of the carriers, which compete with denitrifiers for carbon source.

Conclusions

The long-term performance of a PBS-packed bed bioreactor for nearly 2 years demonstrated that the system is promising to remove nitrate from groundwater. The effluent nitrate concentration reached 3.3–8.8 mg L⁻¹ and 88–97% of nitrate removal was achieved. The denitrification rate at temperatures of 10–18 °C decreased by 52–66% compared to that at 20–29 °C. Only a small increase of DOC (1.7 ± 0.6 mg L⁻¹) and ammonium (0.5 ± 0.3 mg L⁻¹) was observed in effluent which reduce the burden of post treatment. The practical PBS consumption rate (2.75 ± 0.72 g PBS/g NO₃-N) was slightly higher than the theoretical value. Renewal of the fresh carbon source after certain period was necessary for the continuous operation. Bacteria belonging

to genera *Simplicispira*, *Veillonellaceae* uncultured, *Comamonadaceae* uncultured and *Hydrogenophaga* were highly enriched in PBS biofilm. In the top 20 OTUs accounting for 25% of the total sequence, 9 OTUs belonged to denitrifying groups with an abundance of 16%. Some members belonging to other 11 genera were reported to be capable of degradation and acidification of the organic substances.

Conflicts of interest

There are no conflicts to declare.

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