

Removal of hard COD, nitrogenous compounds and phenols from a high-strength coal gasification wastewater stream

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ABSTRACT

The objective of this study was to identify the factors affecting the suspended and fixed biomass in the removal of hard COD, nitrogenous compounds and phenols from a coal gasification wastewater (CGWW) stream using a hybrid fixed-film bioreactor (H-FFBR) process under real-time plant operational conditions and actual wastewater composition. The composition of the influent and effluent was studied to determine which compounds were not removed by hydrolysis (bacterial activity) and how this correlated to the suspended and fixed biomass activity, COD and phenol removal. A H-FFBR with 12 g-m⁻³ attached biomass and 440 mg-l⁻¹ suspended biomass achieved 78% phenol removal and 49% COD removal but insignificant removal of nitrogenous compounds. During the operation period, fixed biomass concentration was higher than the suspended biomass. Parameters such as pH, phenols, alkalinity, metal ions, conductivity, total dissolved solids and aeration rate affected the fixed biofilm properties such as adhesion, thickness and structure. It can be concluded that the composition of the effluent had a direct effect on the fixed biomass properties and thus a direct effect on the removal of phenols, COD and nitrogenous compounds in the wastewater.

Keywords: ammonia stripping, AnoxKaldness, coal gasification, fixed-biofilm reactor, hydantoins, phenosolvan, *Pseudomonas*, refinery effluent, thiocyanate

INTRODUCTION

Coal gasification wastewater (CGWW) is generated in coal gas purification which poses numerous environmental problems in many countries. The chemical composition of coal gasification wastewater is complex and varies from one plant to another. The generated wastewater can contain ammonia, cyanide, thiocyanate, monohydric phenols, dihydric phenols, polycyclic hydroxyl compounds, monocyclic n-aromatics, polycyclic aromatics and aliphatic acids. Of these compounds the phenolic components, primarily, methyl-phenol, and phenols, constitute between 60 and 80% of the organic content (Li et al., 2003).

Physico-chemical methods for the extraction of the phenolic content have been realised in order to reduce the toxicity of the waste stream before conventional treatment. Such treatments as the Phenosolvan process for phenol recovery have been investigated. The Phenosolvan process employs a liquid-liquid extraction process where the incoming gas liquor is filtered through a gravel bed and then contacted with the solvent di-isopropyl ether (DIPE) (Beychok, 1974) or butyl ethanoate (Anastasi, 1980) in multistage mixer-settlers (Beychok, 1974; Bryant et al., 1988). The phenol-rich solvent (extract) is distilled to recover lean solvent for reuse and then stripped to remove and recover residual solvent (Beychok, 1974). The dephenolised liquor (raffinate) is gas-stripped to remove and recover solvent (Beychok 1974; Bryant et al., 1988). The liquor is then steam-stripped to remove acid gases (hydrogen sulphide and carbon dioxide), followed by steam-stripping to remove ammonia (Beychok, 1974).

As can be seen from above, the physico-chemical extraction method is complex, is difficult to optimise, and leaves

room for error. Due to the apparent difficulty and cost of utilising such a complex process, it has become necessary to investigate cheaper and more robust methods for treatment of the CGWW. One such possibility is the use of hybrid and homogeneous fixed-film reactor systems to take advantage of the mixed culture communities capable of metabolising a range of recalcitrant organics in biofilm environments (Stoodley et al., 1999).

Biological treatment has been widely used only to treat the wastewater following pre-treatment by processes of ammonia stripping and phenol solvent extractions to reduce the concentration of toxic compounds (Li et al., 2011). As part of the conventional treatment process, biofilm processes are widely utilised to eliminate the organic carbon and nitrogen content of the wastewater. In biofilm processes, bacteria are grown on suspended carrier particles, fluidised on flocs, or fixed on immobile structures within the reactor. Such processes are aimed to extend the cell retention time (sludge age) of the system indefinitely (Li et al., 2003).

In the biofilm system, the cells are held together by microbial secretions of various polysaccharides that act as a gel-like matrix surrounding the cells (Nelson et al., 1996). This encapsulated structure results in the formation of concentration gradients for all relevant dissolved compounds due to microbial activity and diffusion. The biofilm-associated micro-organisms detach from the surface of the biofilm to populate a planktonic phase that contributes to the removal of organic carbon and nitrogen. The attached biofilm and suspended biomass make the fixed-film bioreactor (FFBR) a hybrid reactor (H-FFBR) rather than an exclusively attached or suspended phase system (Goode, 2010).

MATERIALS AND METHODS

Composite samples (1 000 ml) of the influent stream and effluent stream (clarifier overflow) were collected at a rate of 50 ml-h⁻¹, and pH, suspended solids (SS), chemical oxygen demand (COD),

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ammonium-nitrogen (NH₄-N), nitrate-nitrogen (NO₃-N), ortho-phosphate (o-PO₄), total dissolved solids, conductivity and total phenols were measured using Standard Methods (APHA, 2012). Each aeration zone was linked to a data-collection-system (DCS) and parameters such as temperature, dissolved oxygen (DO), recycling rate, feed rate, de-sludging rate, hydraulic retention time, pH and hydraulic sludge age were controlled and optimised automatically. The oxygen uptake rate (OUR) of the fixed biomass solids was measured on-site using a HACH model HQ40d multimeter with a LDO 101 probe (Hou et al., 2014). A BOD sample bottle (300 ml) was filled with 150 ml effluent and topped up with 150 ml AnoxKaldnes K1 carrier media taken from the respective zones, to simulate a 50% media fill fraction (by volume). The suspended biomass OUR was measured by the same procedure; however, no carrier medium was added (Blanc et al., 2010). The dissolved oxygen (DO) was measured every minute for a period of 10 min. The OUR rates at 36°C ± 1 were calculated by linear regression obtained from the plot of DO concentration versus time (Qiqi et al., 2012). Fixed biomass solids on the carrier were measured by first rinsing with tap water (3 x), then drying duplicate samples, consisting of 100 carriers each, at 105°C for 24 h. The dried carriers were weighed and the solids removed by mixing carriers with 150 ml 0.25 N NaOH for 24 h. The alkali wash was repeated followed by sonication for 90 min. The carriers were then rinsed well with tap water and dried for 24 h at 105°C and re-weighed. The difference in mass was used to determine the biomass on the carriers (Goode, 2010). Scanning electron microscopy (SEM) and energy dispersive X-ray analysis (EDX), conducted by the University of KwaZulu-Natal (Westville, South Africa), were used to determine the structure, thickness and composition of the fixed biomass. Gas chromatography-mass spectrometry (GC-MS) organic fingerprinting was performed by ERWAT (Kempton Park, South Africa).

Pilot plant design and operating parameters

The H-FFBR (1 000 l) consisted of 3 aeration compartments (Zone 1, Zone 2, Zone 3) (Van Leeuwen, 1989) containing acclimatised fixed and suspended biomass (DiMassimo and Bundgaard, 2011; Boltz et al., 2009). The CGWW was diluted to 33% to reduce the toxicity towards the biomass and to maintain an influent COD concentration around 2 000 mg·l⁻¹ (Li et al., 2011; Zhou et al., 2014). Dilution was 1 part CGWW to 2 parts recycled sludge (Luthy et al., 1980). The reactor was operated at 36°C ± 1 (Merlo et al., 2011) and monitored for a 6-month period. The pH was adjusted with 20% (m/v) sulphuric acid (H₂SO₄) or 10% (m/v) sodium hydroxide (NaOH) to maintain a pH range between pH 6.5 and pH 7.5. Plastic biofilm carrier filling fraction (based on volume) for each aeration zone was approximately 70%, 50% and 30%, respectively. A filling ratio of ≤ 70% allowed the plastic media to move freely, minimising dead or unused space in the reactor (Ratcliffe et al., 2006; Zafarzadeh et al., 2011). A 70% filling (Zone 1) of biofilm carrier media (AnoxKaldnes K1) corresponded to an effective biofilm growth area equal to 350 m²·m⁻³; therefore, a 50% fill would equal 250 m²·m⁻³ and a 30% fill would equal 150 m²·m⁻³ for Zone 2 and Zone 3, respectively (Ratcliffe et al., 2006; Ayun et al., 2008). Jing et al. (2009) reported that a carrier fill of between 30% and 50% was suitable to treat a coking plant wastewater. Qiqi et al. (2012) reported minimal difference in performance between a 33% and 66% carrier filling; however, at a filling of 70% the attached growth density was 5 to 13 times higher and responded more strongly to influent COD than that of activated sludge floc found in suspended activated sludge systems.

A defoamer suitable for biological processes was used during excessive foaming periods in the respective aeration zones. In general, Zone 3 tended to foam the most over the 6-month test period. Bryant et al. (1985) reported that foaming tended to be high in aeration zones having the highest COD removal rates; when steady operating states were not being achieved; when biodegradable compounds were not degraded biologically during an upset; and/or when total dissolved solids (TDS) were high in the influent stream.

RESULTS AND DISCUSSION

Composite samples of the CGWW feed and effluent streams were collected and the main organic compounds in each stream were identified using GC-MS. The compounds identified in the CGWW were phenols, straight-chain carboxylic acids, aromatic carboxylic acids, ketones, diesel range organics (DRO), hydroquinones, indoles and hydantoin. The effluent stream consisted mainly of hydantoin, traces of carboxylic acids, traces of diesel range organics and phenols, thus indicating that the biological process was effective in removing carboxylic acids, monohydric phenols and hydroquinones, but not, however, polyhydric phenols, diesel range organics and hydantoin (5,5-dimethyl-hydantoin and 5-methyl-5-ethyl-hydantoin). The composition of the CGWW differs with different coal supplies, gasification processes and conditions. Beychok (1974) reported the presence of 75% to 85% monohydric phenols (phenols, cresols), 30% to 35% dihydric phenols (catechol, resorcinol and hydroquinones), organic acids (naphthenic acid), neutral oils and organic bases (pyridine) in gas liquor. Li et al. (2003) reported 27.6% methyl-phenol, 11.4% quinolinones, 10% dimethyl-phenol, 10% indole, 9.5% phenol, 6.7% methyl-quinolinone, and 5% isoquinolinone. Lignite coals (low-ranking coal) are a source of high amounts of aromatic compounds with higher oxygen content than higher-ranking coals (Kapusta and Stanczyk, 2011). Low-rank coal contains large amounts of polar groups such as -OH, -COO, -O, -N, and -S, and exhibit mostly hydrophilic behaviour (Molva, 2004).

Hydantoin is non-volatile, highly polar heterocyclic compounds, documented as being poorly biodegradable or non-biodegradable and which are not removed by solvent extraction with DIPE or steam stripping (Turner et al., 1985). Pavlovich and Luthy (1988) reported that 5,5-dimethyl-hydantoin (DMH) was slightly more soluble in aqueous media than 5-methyl-5-ethyl-hydantoin and both could form complexes with transition metals at pH 8.5.

Hydantoin is specific to slagging fixed-bed gasification and typical concentrations of 630 mg·l⁻¹ have been reported by Strain and Turner (1985), and of 1 700 mg·l⁻¹ to 2 600 mg·l⁻¹ by Pavlovich and Luthy (1980), after pre-treatment of the liquor by solvent and gas stripping. Cyanide combines with carbon dioxide, ammonia and a ketone during the gas-quenching process to form hydantoin at pH 8.5 (Diehl et al., 1985). If the ketone is 2-butanone (MEK) then 5-methyl-5-ethyl-hydantoin (EMH) is formed; if it is 4-methyl-2-pentanone (MIBK) then 5,5-dimethyl-hydantoin (DMH) is formed (Turner et al., 1985). Ketones and hydantoin are present in low-rank (lignite) coal (Olson et al., 1985).

Removal of phenols

The removal of total phenols in the H-FFBR ranged between 62% and 93% with an average of 78%; thus 22% of the total phenols were soluble, but recalcitrant (Fig. 1). The removal of

phenols is affected by factors such as biodegradation kinetics, substrate inhibition effect, pH, temperature, biomass concentration, microbial community and their metabolic potential, and nutrient concentration (Lepik and Tenno, 2011). Biodegradation depends on the structures of the phenolic compounds and the number of substituents on the aromatic nucleus. Biodegradation is also affected by the positions of the methyl groups of methylphenols. The *p*-substituted phenols are more readily biodegradable than the *m*- or the *o*-substituted phenols since they are weaker electron donors (Adabju, 2013).

Marrot et al. (2006) reported the optimum pH for phenol degradation to be between 6.5 and 7.5, Crutescu et al. (2008) reported a degradation rate of between 95% and 99% at pH between 6.8 and 7.5, Chakraborty et al. (2010) reported the highest phenol degradation rate (83%) at pH 7.0 (30°C); however, the rate of degradation at 35°C would still be considerable but less than that at 30°C (<83%). Arutchelvan et al. (2006) reported that temperature (34°C) and pH (8.0) influenced the rate of phenol degradation to a maximum of 1 750 mg·ℓ⁻¹. Lim et al. (2013) reported a pH between 7.3 and 7.8 for phenol degradation. A healthy biomass (acclimatised to wastewater constituents) requires a pH between 6.5 and 8.5 (Merlo et al., 2011).

In this study, phenol removal efficiencies of 62–93% were measured in the operation of the H-FFBR treating CGWW with the incoming pH controlled to pH 6.5–7.9 using acid or alkali, and a temperature of 36°C ± 1. These results are higher than a phenol reduction of 83% at pH 7 and 30°C as published by Chakraborty et al. (2010) and lower than 95–99% as reported by Crutescu (2008) for a pH range of 6.8–7.5.

Removal of soluble COD

The removal of soluble COD ranged between 37% and 62% with an average of 49% (Fig. 2). Turner et al. (1985) reported that the presence of 5,5-dimethyl-hydantoin and related compounds in the condensate waters accounted for 1% to 6% of the COD, phenols 59% to 76% of the COD, and dihydroxy benzenes 0.02% to 9.5% of the COD. Galil et al. (1988) reported COD removal of 40% to 60% when biologically treating similar coal condensate (solvent-extracted and ammonia-stripped), which is in line with the COD removal efficiencies obtained in this study.

The H-FFBR was not effective in removing all the hydantoin, aromatic carboxylic acids and diesel range organics (DRO). These results correlate with reports that biological oxidation provided good removal of many organic compounds, but that aromatic hydrocarbons containing aliphatic substitutions and certain polycyclic aromatic hydrocarbons (PAH) were only partially removed.

Wang et al. (1999) reported that phenolic compounds were the main organic components of coal gasification wastewater and made up more than 60% of the total organic carbon. Merlo et al. (2011) reported that 13% of COD in petroleum refinery effluent was non-biodegradable. Moretti and Neufeld (1989) reported that most of the COD was due to the presence of DMH and EMH, of which 1% to 6% was due to EMH, 59% to 76% due to DMH, and 38% to 85% due to phenols.

Mirhossaini et al. (2010) reported that the amount of COD in the influent stream resulted in competition between heterotrophs and autotrophs and therefore defined the total biomass and biofilm composition as well as oxygen diffusion into the biofilm. The total biomass population was thus heterotrophic since the COD removal rate (Fig. 2) was higher than ammonium-nitrogen removal (Mazumder, 2010). Dürr et al. (2006) reported that heterotrophs belonging to the genera

Arthrobacter, *Pseudomonas*, *Bacillus* and *Flavobacterium* were able to catalyse the hydrolysis of hydantoin in a ring-opening step by enzymes classified as hydantoinases.

Fixed biomass versus suspended biomass activity

The average fixed biomass activity was 62.58 (mg O₂·ℓ⁻¹·h⁻¹), 33.80 (mg O₂·ℓ⁻¹·h⁻¹) and 33.71 (mg O₂·ℓ⁻¹·h⁻¹) for the respective aeration zones. The suspended biomass activity was 48.47 (mg O₂·ℓ⁻¹·h⁻¹), 37.48 (mg O₂·ℓ⁻¹·h⁻¹) and 41.31 (mg O₂·ℓ⁻¹·h⁻¹) for the respective aeration zones. The suspended biomass activity was higher than the fixed biomass activity in Zone 2 and Zone 3 (Fig. 3). The biofilm thickness for Zone 1 ranged between 1.79 μm and 213 μm with an average of 51.45 μm. The biofilm

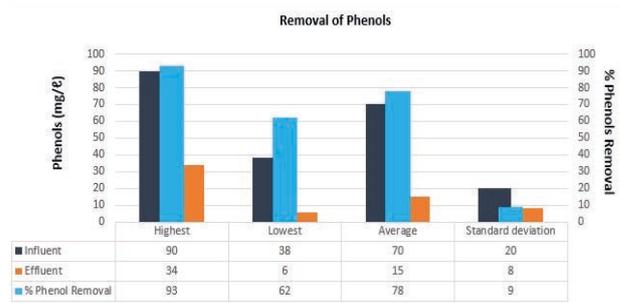


Figure 1

Data distribution of 132 samples tested for % phenol removal in the operation of the hybrid fixed-film bioreactor (H-FFBR)

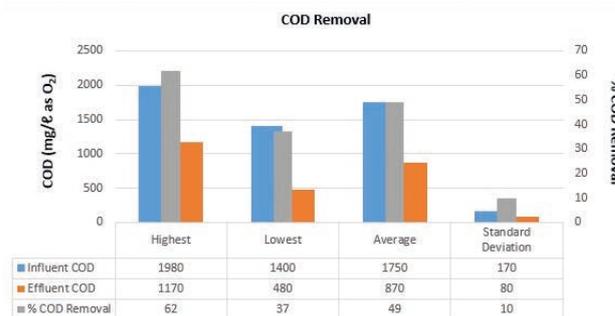


Figure 2

Data distribution of 132 samples tested for % COD removal in the operation of the hybrid fixed-film bioreactor (H-FFBR)

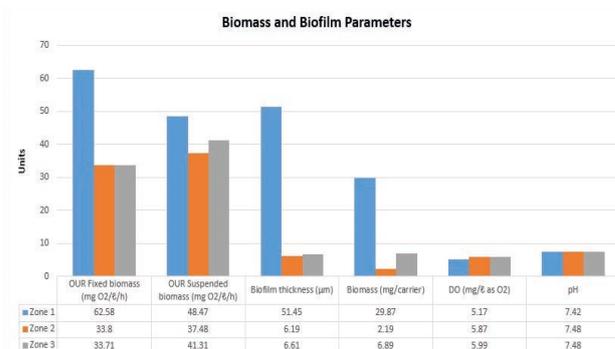


Figure 3

Differences between biomass and biofilm parameters in the respective aeration zones in the operation of the hybrid fixed-film bioreactor (H-FFBR)

thickness for Zone 2 ranged between 0.47 μm and 13.61 μm with an average of 6.19 μm . The biofilm thickness for Zone 3 ranged between 1.45 μm and 24.0 μm with an average of 6.61 μm . There was no direct correlation between fixed biomass thickness and biomass activity. However, there was a correlation between aeration rate and biofilm thickness (Fig. 3). Biofilm thickness is normally between 5 μm and 40 μm and rarely more than 100 μm to 200 μm (Ward and King, 2012). Boltz et al. (2009) reported biofilm thickness in aerobic H-FFBR reactors to be in the range of 50 μm to 200 μm . Asiedu (2001) reported the biofilm thickness should be less than 100 μm to allow for sufficient substrate diffusion into the biofilm.

Higher aeration rates increased the mixing energy (roll pattern) of the plastic media, resulting in higher shear forces across the biofilm layer (Sen and Randall, 2008), which influenced the biofilm thickness, microbial population and rate of growth. The mixing energy provided by the aeration is critical for sloughing of biomass and the formation of a thin biofilm yielding higher kinetic rates than previously measured in thick biofilm (Hubbell et al., 2006). Higher aeration rates increase the biological activity of the fixed biomass; however, extreme turbulence detaches biofilm from the carrier (Asiedu, 2003). Thin biofilms increase available surface area and reduce variability of solids discharge due to sloughing (Goode, 2010). Sen and Randall (2008) reported that the rate of sloughing and the biofilm yield for heterotrophs and nitrifiers in each cell are dependent on (i) shear forces imparted by mixing and roll pattern, (ii) design of biofilm carrier particle (media), and (iii) substrate and electron acceptor level in each cell. The biofilm thickness is a function of the substrate concentration in the bulk liquid, average mixed liquor volatile suspended solids (MLVSS) of the biofilm and shear forces (Sen and Randall, 2008).

Biofilm structure and properties

EDX analyses of the fixed biofilm layers indicated the presence of carbon, oxygen, sodium, aluminium, silicon, phosphorus, sulphur, chloride, potassium, calcium, nitrogen, copper and iron for all three aeration zones (Fig 4, 5 and 6). Aluminium, silicon, calcium and iron were found to be highest in Zone 1; nitrogen, oxygen, sodium, magnesium, sulphur, chloride and potassium highest in Zone 2, and carbon, phosphorus and copper highest in Zone 3 (Table 3). The same inorganic elements were reported by Revanuru and Mishra (2011), who reported that the silicon and aluminium increased biofilm resistance to toxic shocks and the presence of iron and sulphur contributed to the aggregation of biomass. Potassium, magnesium, sodium, calcium, iron and chlorides are macronutrients required for membrane stabilisation. Micronutrients such as zinc, manganese, molybdenum, selenium, copper, cobalt, nickel, vanadium and tungsten have been reported to allow proper function of enzymes. Hu et al. (2013) reported that calcium and phosphorus absorb and accumulate in the biofilm as insoluble compounds rendering the biofilm highly resistant to detachment. Todar (2006) reported that sulphur was required for protein synthesis, phosphorus for nucleic acids and metabolic cofactors, and nitrogen for proteins, some sugars and nucleic acids.

Goode (2010) reported that the extracellular polymeric substance (EPS) acts as an ion exchange resin where divalent bridges can be formed or broken due to relative concentrations of monovalent and divalent cations competing for negative sites. The presence of divalent cations (Ca^{2+} , Mg^{2+} , Fe^{2+}) influences biofilm formation directly through its effect on electrostatic interactions, and indirectly via physiologically-dependent

attachment processes by acting as cellular cations and enzyme cofactors. Calcium ions also impact the mechanical properties of the biofilm by acting as cross-linkers and making the extracellular polymeric substances more proteinaceous (Song and Leff, 2006).

In general, biofilms become thicker (lower detachment rate), denser and more mechanically stable (mature biofilm) when exposed to increasing concentrations of divalent cations.

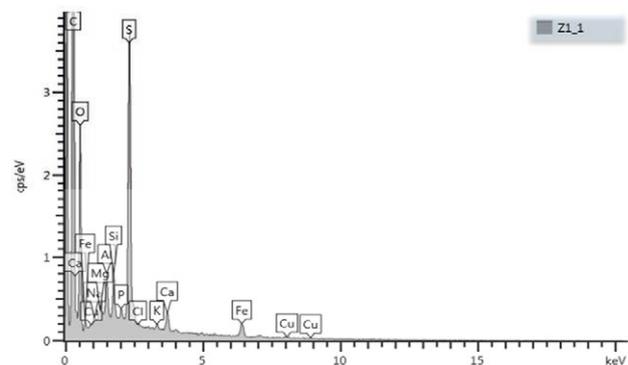


Figure 4

Energy dispersive X-ray analysis (EDX) of the fixed biofilm in aeration Zone 1 (Z1_1) in the operation of the hybrid fixed-film bioreactor (H-FFBR)

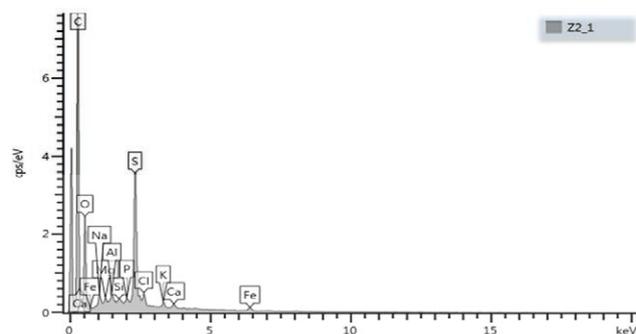


Figure 5

Energy dispersive X-ray analysis (EDX) of the fixed biofilm in aeration Zone 2 (Z2_1) in the operation of the hybrid fixed-film bioreactor (H-FFBR)

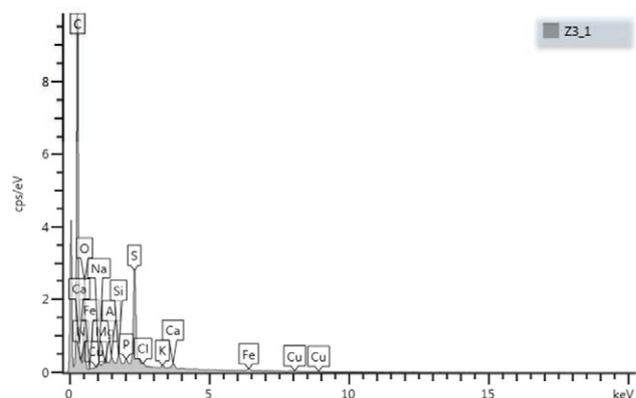


Figure 6

Energy dispersive X-ray analysis (EDX) of fixed biofilm in aeration Zone 3 (Z3_1) in the operation of the hybrid fixed-film bioreactor (H-FFBR)

Table 1 Energy dispersive X-ray analyses of the fixed biofilm			
Element	Zone 1	Zone 2	Zone 3
Carbon	67.74	66.22	68.38
Nitrogen	< 0.1	0.88	0.62
Oxygen	24.66	25.70	25.01
Sodium	0.17	0.47	0.22
Magnesium	0.10	0.19	0.01
Aluminium	0.42	0.24	0.22
Silicon	0.41	0.10	0.22
Phosphorus	0.24	0.21	0.28
Sulphur	4.79	4.86	4.23
Chloride	0.10	0.37	0.16
Potassium	0.12	0.21	0.15
Calcium	0.36	0.22	0.22
Iron	0.85	0.55	0.40
Copper	0.13	0	0.20

Unit: % by weight

However, elevated calcium levels could lead to the accumulation of inorganic salts and thus reduce the effective surface area of the biofilm (Goode, 2010). DMH is a weak acid and complexes with metals in the anionic form, thus affecting the equilibrium chemistry and speciation of calcium (Pavlovich and Luthy, 1988).

Scanning electron microscopy (SEM) analyses indicated the fixed biofilm morphology to be relatively uniform and dense for Zone 1 (Fig. 7), 'rippling' for Zone 2 (Fig. 8) and 'porous' for Zone 3 (Fig. 9). Lembre et al. (2012) reported that the bending of biofilm structures in the same direction of the shear force was due to the viscoelasticity response of a mixed culture biofilm. Goode (2010) reported that a porous structure was found to allow greater diffusion of dissolved solids compared to a homogenous biofilm. Ward and King (2012) reported that biofilms are initially homogenous, and then become heterogeneous complex structures when mature.

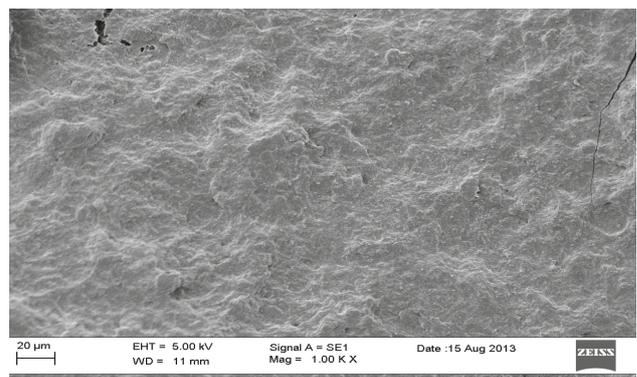


Figure 8
Scanning electron microscopy (SEM) of the fixed biofilm in Aeration Zone 2

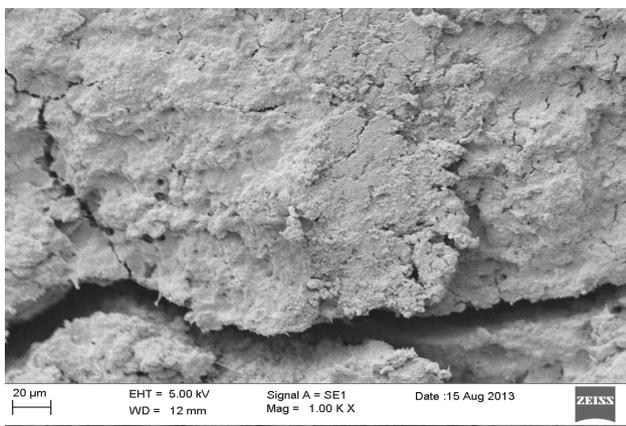


Figure 7
Scanning electron microscopy (SEM) of the fixed biofilm in Aeration Zone 1

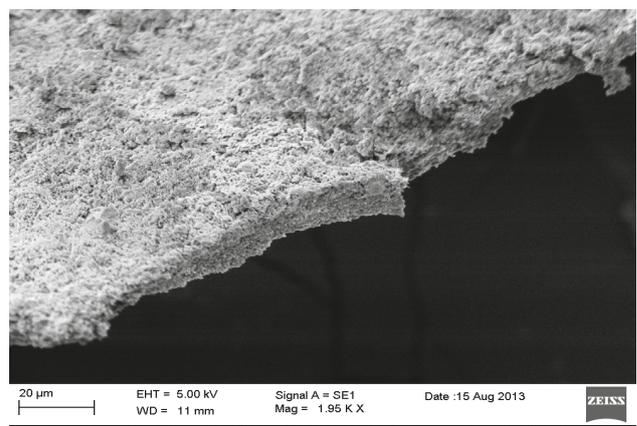


Figure 9
Scanning electron microscopy (SEM) of the fixed biofilm in Aeration Zone 3

Electrolyte concentration (osmolarity, conductivity) and nutrient concentration also affect biofilm morphology (Song and Leff, 2006). Biofilms are thicker and denser at higher nutrient concentration and the morphology of an established biofilm changes as the carbon concentration changes (Song and Leff, 2006). Increasing biofilm thickness and reducing diffusion of oxygen into the deeper regions of the biofilm would lead to better retention of denitrifiers (Goode, 2010).

Commercially available bio-augmentation products specifically designed for refinery wastewater were tested in the laboratory (results not included). Results indicated a 30% improvement in the removal of soluble COD. However, this needs to be confirmed on the pilot plant under similar operating conditions.

CONCLUSIONS

The results show that a hybrid fixed-film bioreactor (H-FFBR) can be successfully used for the removal of chemical oxygen demand (COD) and total phenols from coal gasification wastewater (CGWW). The attached biomass activity was higher than the suspended biomass activity. Thus, the removal of COD and total phenols was performed mainly by the attached biomass. Both populations were heterotrophic since there was no removal of nitrogenous compounds. The H-FFBR can thus be scaled up for the removal of COD and total phenols from CGWW. However, from an environmental perspective, bio-augmentation should be investigated further for enhancing the removal of COD, total phenols and nitrogenous compounds.

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