

For The Reduction of Greenhouse Gas Emissions from Coal Mine Ventilation Air, Employ a Coal-Packed Methane Bio Filter

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Abstract

Methane emitted by coal mine ventilation air (MVA) is a significant greenhouse gas. A mitigation strategy is the oxidation of methane to carbon dioxide, which is approximately twenty-one times less effective at global warming than methane on a mass-basis [1]. The low non-combustible methane concentrations at high MVA flow rates call for a catalytic strategy of oxidation. A laboratory-scale coal-packed biofilter was designed and partially removed methane from humidified air at flow rates between 0.2 and 2.4 L min⁻¹ at 30°C with nutrient solution added every three days [2]. Methane oxidation was catalysed by a complex community of naturally-occurring microorganisms, with the most abundant member being identified by 16S rRNA gene sequence as belonging to the methanotrophic genus *Methylocystis*. Additional inoculation with a laboratory-grown culture of *Methylosinus sporium*, as investigated in a parallel run, only enhanced methane consumption during the initial 12 weeks [3]. The greatest level of methane removal of 27.2±0.66 g methane m⁻³ empty bed h⁻¹ was attained for the non-inoculated system, which was equivalent to removing 19.7±2.9% methane from an inlet concentration of 1% v/v at an inlet gas flow rate of 1.6 L min⁻¹ (2.4 min empty bed residence time). These results show that low-cost coal packing holds promising potential as a suitable growth surface and contains methanotrophic microorganisms for the catalytic oxidative removal of methane [4].

Keywords: Coal mine; Ventilation; Methane; Biofiltration

Introduction

To help explosions during underground coal mining, mine shafts are continuously voiced to adulterate methane released from the coal confluence to non-combustible attention(generally $\leq 1 \text{ v/v}$ since the lower ignitable limit of methane is 5 (v/v) in air) [5]. The mine ventilation air (MVA) is also released undressed into the atmosphere causing significant hothouse gas emigrations. Methane has an roughly twenty- one times advanced implicit impact on global warming than carbon dioxide(on a mass- base in a 100- time time frame) arising from its advanced molar immersion measure for infrared radiation and a longer hearthstone time in the atmosphere [6]. On a molecular base this equals a7.6- times advanced impact of methane. Worldwide emigrations of methane from coal mining are expansive, estimated to be over 329 million tonnes(Mt) carbon dioxide- fellow in 2005 with roughly 70 of these methane emigrations released as MVA [7]. Major challenges for MVA methane mitigation are the low methane attention limiting its utility as an energy source, the high inflow rates(50 – 500 m³s⁻¹) and the considerable variability of these parameters. colorful thermal technologies have been considered and while able of treating MVA, they attract high capital and operating costs and bear considerable safety measures. Biofiltration technology is a safer and less precious approach as it utilises microorganisms as biocatalysts to oxidise methane to carbon dioxide and biomass at ambient temperature. Methane- oxidising organisms (methanotrophs) do nowhere and laboriously grow in surroundings where both methane and either oxygen or indispensable electron acceptors are present(e.g. soils, lakes, ponds, tips, and coal mine spots). The biology of colorful methanotrophs has been lately reviewed. Over the once decade, methane biofiltration technology has garnered considerable attention for the treatment of effluent feasts generated during tip and beast husbandry operations, where nicely low gas inflow rates do [8]. still, only limited perceptivity are available regarding the eventuality of biofiltration for the junking of methane at the veritably low attention and high inflow rates of MVA (for a recent review see. Studies of methane junking with quilting accoutrements similar as polypropylene Raschig rings, a

batch study), glass tubes, mature compost, clay or pine dinghy revealed fairly slow conversion. Accordingly, large biofilter volumes may be needed for MVA operations. To accommodate large volume biofilters, affordable quilting accoutrements would be needed. At a mine point the most accessible and affordable quilting material may be coal [9]. Chakravorty and Forrester (1985) have reported that methanotrophic microorganisms were suitable to grow on the face of coal as a thin biofilm and oxidized methane in batch trials. To our current knowledge, coal has not been utilised preliminarily as a biofilter quilting material for microbial methane oxidation. The end of the present study was to design a laboratory- scale nonstop biofilter with coal as the quilting material for methane oxidation at 1(v/v) – a attention representative of MVA- in order to estimate whethernon-sterilised coal may serve as a suitable volition to other quilting accoutrements . Specific objects were to cover biofilter performance over a range of gas inflow rates and to probe the effect of inoculation with the methanotrophic bacterium, *Methylosinus sporium*. At the end of the trial, the composition of the mixed biofilm community was analysed to determine the dominant methanotrophic organism at that time [10].

Material and Methods

Nitrate mineral mariners(NMS) medium, coal quilting and feasts The nitrate mineral mariners(NMS) medium according to the German Resource Centre for Biological Material(DSMZ) contained

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per litre 1 g MgSO₄·7 H₂O; 0.2 g CaCl₂·6 H₂O; 0.004 g Fe(III) NH₄-EDTA; 1 g KNO₃; 0.272 g KH₂PO₄; 0.717 g Na₂HPO₄·12 H₂O. This medium was mixed with 1 ml L-1 methanol (as fresh carbon source to accelerate the slow growth of methanotrophs) and 0.5 ml L-1 trace element result. The trace element result contained (per L) 0.5 g Na₂-EDTA; 0.2 g FeSO₄·7 H₂O; 0.01 g ZnSO₄·7 H₂O; 0.003 g MnCl₂·4 H₂O; 0.03 g H₃BO₃; 0.02 g CoCl₂·6 H₂O; 0.001 g CaCl₂·2 H₂O; 0.002 g NiCl₂·6 H₂O; 0.003 g Na₂MoO₄·2 H₂O. The pH of the medium was acclimated to 6.8 with 2 M NaOH. It was also autoclaved by autoclaving for 15 minutes at 15 psi and 121 °C. Bituminous coal, for use as the biofilter quilting material, was kindly handed by BHP Billiton (www.bhpbilliton.com) from the Appin Colliery point in New South Wales, Australia. Coal characteristics are available in the supplementary section Data S1. The coal was base using a mortar and pestle and settled to gain pieces of 2 – 3 cm periphery. Methane with a purity of 99.95, argon with a purity of 99.996 and compressed air were obtained from Core Gas, Australia. Biofilter design and operation Two identical biofilters were designed and constructed from acrylic resin (Perspex). The two biofilters were packed with unsterilized coal to a bed height of 22 cm, equating to an empty bed volume of 3.89 L. The void volume within the coal bed was 0.39 L as determined by filling the airspace in the quilting with water. Biofilter 1 was invested with a 100 ml of pure *M. sporium* culture mixed with 900 mL of NMS. Biofilter 2 was drenched with 1 L of sterile NMS alone (i.e. no *M. sporium*). therefore both biofilters contained microbes that naturally passed on the coal quilting and biofilter 1 also contained *M. sporium*. The coal beds were left to soak in their separate media for 24 hours after which the redundant liquid was drained. A humidified methane and air gas sluce (1 (v/v) methane in air at 0.2 L/min) was also continuously passed through each biofilter. moisture was handed to help the coal bed from drying out. Sterile NMS medium (2 L) was added via the top of each biofilter every three days, the drain was opened after 10 min and the medium was also recirculated doubly and drained again so that only a liquid film remained on the shells to feed the microbes. Gas analysis and determination of biofilter performance Methane and carbon dioxide attention in the two biofilter bay and exit aqueducts were determined using a Shimadzu GC-8A gas chromatograph (GC) equipped with a thermal conductivity sensor. Separation was achieved using an Alltech Hayesep DB100/120 column with helium as the carrier gas. To assess whether methane junking began from microbial action or system leakage, one hour before analysis argon (1 (v/v)) was introduced to the gas sluce entering the biofilter as an inert internal standard. A cold trap was used to remove moisture from the gas sluce previous to injection into the GC. Analyses were performed in triplet every three days with attention determined relative to the internal argon standard grounded on peak areas. Biofilter performance was estimated on the base of empty bed volume to grease comparison with other studies. Methane bay cargo (IL), methane junking effectiveness (shaft), methane elimination capacity (EC), and carbon dioxide product rate (PCO₂) were calculated using the equations listed.

Discussion

The current study highlights the implicit operation of coal-packed biofilters for partial methane junking from coal MVA. adding the methane IL by raising inflow rates at a constant low methane attention representative of MVA increased the methane EC to an optimal position after which EC values dropped. analogous trends have been observed in other biofiltration systems where farther inflow rate increases saw EC values either be maintained or dropping. Several physical factors may affect the bioavailability of methane. Originally, the movement of methane motes within the gas phase plays a part. The Reynolds

number remained well below 10 for all tested inflow rates, which indicates laminar inflow [11]. Thus, the movement of methane motes within the gas towards the water face wasn't supported by turbulence, but governed by convection and proximity. Secondly, interfacial transfer from the gas to the water phase may be a limiting factor. As methane is inadequately answerable in water (0.022 g of methane/kg of water), the driving force for methane uptake is low. Accordingly, methane proximity into the liquid phase is slow. As the empty bed hearthstone time (EBRT) decreases with adding gas inflow rate, the time available for methane transfer across the gas/liquid interface is shorter and in turn, confining methane vacuity to the microorganisms and dwindling shaft. On the other hand, the adding gas inflow rate might increase the rate of proximity of methane from the gas phase into the liquid phase. This is because an adding IL replenishes used methane briskly and thus increases the methane attention grade across the interface. similar effect may have contributed to the adding EC with adding IL up to 139 g m⁻³ h⁻¹. Thirdly, the distribution of methane that has entered the liquid phase would be governed by methane proximity through the water and extracellular polymeric matrix of the microorganisms as well as by cellular methane uptake and oxidation. It's thus likely that an increase in the consistence of the microbial community will drop the vacuity of methane to the deepest subcaste of cells on the coal face [12]. At this stage still, it remains unclear which of the below-mentioned factors is the most limiting for the biofilter performance. The change of the oxidative capacity of the microbial community over time is unknown and may overlay the goods of changing ILs. It's known still, that the final return of the inflow rate to the original value revealed catalytic conditioning of the microbial community at the morning and end within the same order of magnitude. Biological factors that impact the oxidative capacity of the biofilter include biomass volume and thebio-catalytic exertion of the biomass. adding biomass volume as indicated by the carbon accumulation rate in Table III, will only be salutary until all coal face area is covered with microbial community up to a consistence that allows effective methane proximity to the cells. Maximising the quilting face area for microorganisms in order to increase the EC may be achieved by using lower coal pieces or high-porosity coal; still this could affect in increased back pressure and ultimately in blockage of the biofilter [13]. In addition to the volume of the microbial biomass, itsbio-catalytic exertion is important for biofilter performance. The natural methane oxidising exertion is determined by the community composition of microorganisms and their applicable enzymes as well as by thebio-available methane attention. The community analysis illustrated that in both biofilters a large proportion of the microbial diversity set up had no given capacity for methane oxidation and this may give an occasion for optimisation. The primary difference between biofilter 1 and 2 was the original inoculation of biofilter 1 with the methanotrophic *M. sporium*. This addition of *M. sporium* to the naturally being microbial community may have directly or laterally caused the original advanced EC and shaft values in biofilter 1 compared to biofilter 2. At the loftiest IL, performance of the biofilters came similar and the methanotroph *Methylocystis* sp. was dominant in both biofilters [14]. Interestingly, no *M. sporium* was detected. It isn't known at what point *Methylocystis* sp. began to dominate in the biofilters and when *M. sporium* faded in biofilter 1. Several strains of *Methylocystis* sp. are known for their high affinity for methane and the capability to oxidise methane indeed at low atmospheric attention. Reported methane monooxygenases of the rubric *Methylocystis* have lower Km values (3.2 – 4 µM) than those of the rubric *Methylosinus* (8.3 – 62 µM). Since the Km value signifies the methane attention at which the enzyme reaches half-minimal rate, *Methylocystis* sp. might be better acclimated to low methane attention than *M. sporium*, still

these values may vary at the strain position. While all K_m values are well below the solubility of methane in water (1375 μM), mass transfer limitations might have led to a much lower available methane attention in contact with the enzymes so that K_m values may play a part in biofilter performance [15].

Conclusion

The Dalong coal is composed of low rank coals with an R_o value of 0.57 to 0.59. The natural humidity content of coal samples is advanced with an average humidity value of 7.0. The effect of water on the gas adsorption of coal is substantially that water moles block the gas inflow channel, and the competition of adsorption between methane and water moles occupies part of the adsorption eventuality in the coal matrix. The adsorption test showed that the Langmuir volume of the Dalong coal sample is 4026 m^3/t , indicating a strong gas adsorption capability. The desorption test indicates that the gas desorption quality is vastly lesser in the first 30 min. still, the desorption volume increased gradationally in the last 80 to 120 min. When the gas pressure of adsorption equilibrium is 4.30 MPa, 10.1 mL/g methane was desorbed in 120 min. When the gas pressure dropped to 1.16 MPa, 4.6 mL/g gas was desorbed in 120 min. Trials of the impact of external humidity on gas desorption indicate that the gas desorption volumes and humidity content conform to a logarithmic relation. The advanced water content (lower than the critical achromatism of water content), the lower the accumulation desorption volume. After humidity correction, we can gain the outburst vaticination (coal slices desorption indicator) indicator critical value of long-honey coal. This indicator can give a theoretical base for the low rank coal mines to determine the critical value of the coal confluence outburst vaticination indicator.

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Conflict of Interest

None

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