

Available online at www.sciencedirect.com

SciVerse ScienceDirect

journal homepage: www.elsevier.com/locate/watres

A novel methodology to quantify nitrous oxide emissions from full-scale wastewater treatment systems with surface aerators

Liu Ye^{a,b}, Bing-Jie Ni^a, Yingyu Law^{a,c}, Craig Byers^d, Zhiguo Yuan^{a,*}^a Advanced Water Management Centre, The University of Queensland, QLD, Australia^b School of Chemical Engineering, The University of Queensland, QLD, Australia^c Nanyang Technological University, SCELSE, Singapore 637551, Singapore^d Water Corporation, Perth, WA, Australia

ARTICLE INFO

Article history:

Received 10 July 2013

Received in revised form

14 September 2013

Accepted 17 September 2013

Available online 10 October 2013

Keywords:

Nitrous oxide

Emission factor

Greenhouse gases

Wastewater treatment

Surface aerators

Modelling

ABSTRACT

The quantification of nitrous oxide (N₂O) emissions from open-surface wastewater treatment systems with surface aerators is difficult as emissions from the surface aerator zone cannot be easily captured by floating hoods. In this study, we propose and demonstrate a novel methodology to estimate N₂O emissions from such systems through determination of the N₂O transfer coefficient ($k_L a$) induced by surface aerators based on oxygen balance for the entire system. The methodology is demonstrated through its application to a full-scale open oxidation ditch wastewater treatment plant with surface aerators. The estimated $k_L a$ profile based on a month-long measurement campaign for oxygen balance, intensive monitoring of dissolved N₂O profiles along the oxidation ditch over a period of four days, together with mathematical modelling, enabled to determine the N₂O emission factor from this treatment plant ($0.52 \pm 0.16\%$). Majority of the N₂O emission was found to occur in the surface aerator zone, which would be missed if the gas hood method was applied alone.

© 2013 Elsevier Ltd. All rights reserved.

1. Introduction

Nitrous oxide (N₂O) is a significant greenhouse gas, with a global warming potential approximately 300-fold stronger than that of carbon dioxide (IPCC, 2001). It also reacts with ozone in the stratosphere leading to ozone layer depletion (Portmann et al., 2012). N₂O can be produced and directly emitted from wastewater treatment systems (Ahn et al., 2010). N₂O emissions can contribute substantially to the carbon footprint of wastewater treatment plants (WWTPs). It is estimated that an emission factor as low as 0.5% would lead to a greenhouse gas emission that is comparable to that of the

indirect CO₂ emission due to energy consumption in conventional biological nutrient removal WWTPs (De Haas and Hartley, 2004). Therefore, reliable quantification of N₂O emissions from full-scale WWTPs is of importance.

Presently, the UN Climate Panel (IPCC) is using an over-simplified fixed emission factor as their current global climate tool to quantify N₂O emissions, without considering specific designs and operational conditions (IPCC, 2006). However, the N₂O emission data collected from WWTPs to date show a huge variation in the fraction of influent nitrogen load emitted as N₂O. By monitoring 12 WWTPs in the US, Ahn et al. (2010) reported emission factors in the range of 0.01–1.8%. Some other studies

* Corresponding author. Tel.: +61 7 33654374; fax: +61 7 33654726.
E-mail address: zhiguo@awmc.uq.edu.au (Z. Yuan).

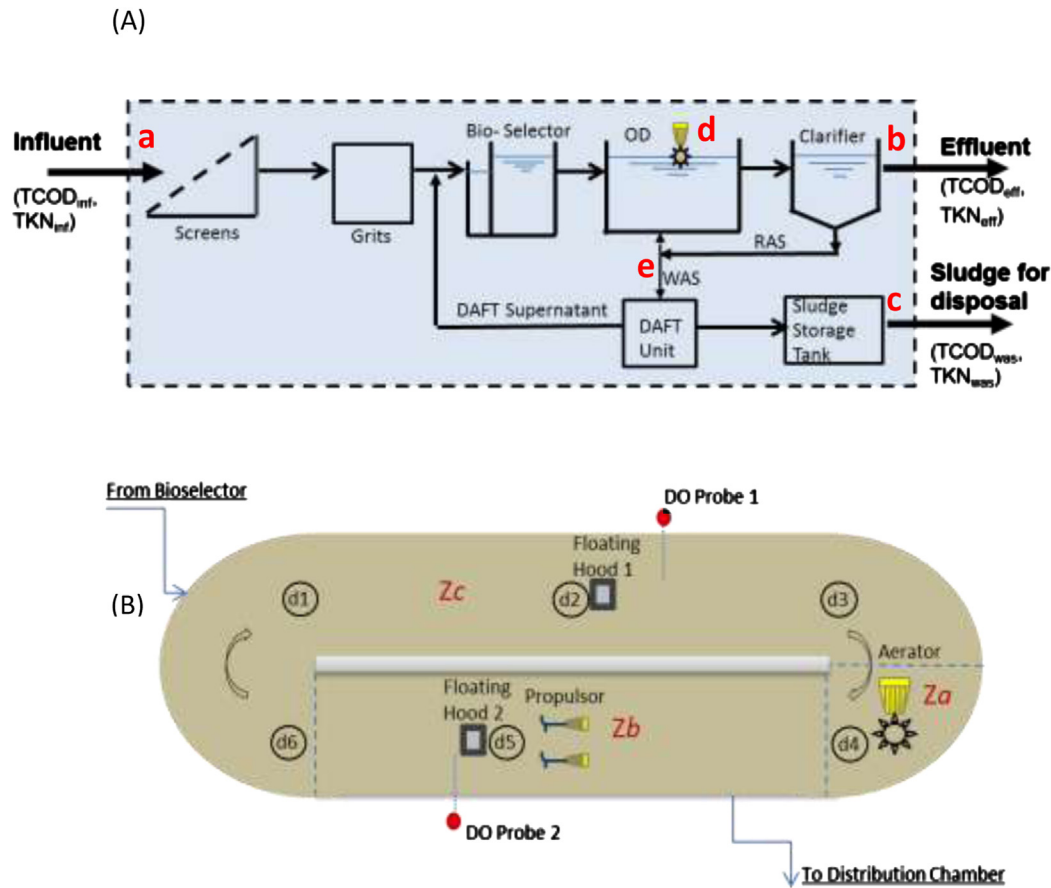


Fig. 1 – (A) A simplified flowchart of the WWTP studied with the long-term sampling/data collection locations (a-raw wastewater; b-effluent; c-disposed sludge; d-oxidation ditch; e-waste activated sludge) (B) Sampling locations in the oxidation ditch during intensive sampling period (d1-anoxic zone where the wastewater enters; d2-anoxic zone near floating hood 1; d3-anoxic zone before mixed liquor enters the aeration zone; d4-aeration zone with a surface aerator; d5-aerobic zone near the propulsor and floating hood 2; d6-anoxic zone, at the end of the ditch. The ditch is divided into three zones based on visual observation (Za – aerator zone, Zb – propulsor zone with noticeable turbulence on surface, Zc – non-turbulent zone).

revealed even higher (>10%) emission factors (Kampschreur et al., 2009; Foley et al., 2010). A high degree of temporal variability in N₂O emission has also been observed within the same WWTP (Ahn et al., 2010). These variabilities in measured emissions strongly contrast with the fixed emission factor method currently applied to estimate N₂O emissions from wastewater treatment (IPCC, 2001). Therefore, each WWTP needs to be assessed individually through either direct measurement of N₂O emission or reliable model-based estimation (Foley et al., 2011).

For the WWTPs that are fully covered, the N₂O emission rate can be calculated by measuring N₂O concentration in the gas phase, as well as the total gas flow rate. Using this method, Daelman et al. (2013) monitored N₂O emissions from a fully covered municipal WWTP in Netherlands for sixteen months, and observed that 2.8% incoming nitrogen was emitted as N₂O, which constituted three quarters of the carbon footprint of the plant.

However, most WWTPs are open-surface sludge systems. The floating chamber method is typically used for the quantification of N₂O in such systems (Law et al., 2012). N₂O flux is captured using a floating hood, a method adapted from gas

emission measurement from soils. Recently, the online, continuous monitoring of N₂O with floating hoods enabled relatively accurate quantification of N₂O emissions from some open-surface activated sludge systems (Kampschreur et al., 2008; Joss et al., 2009; Ahn et al., 2010; Desloover et al., 2011).

An alternative method to quantify N₂O emissions is based on mass transfer estimation. In WWTPs, N₂O is produced in the liquid phase during nitrogen removal, which may be transferred into gas phase when N₂O is over-saturated. The rate of emission can be estimated using the volumetric mass transfer coefficient ($k_L a$, d⁻¹) and the liquid phase N₂O concentration. Foley et al. (2010) estimated N₂O emissions from seven WWTPs in Australia with various process configurations using this method, with a reported N₂O emission factor from 0.6 to 25% of the denitrified nitrogen. However, $k_L a$ is related to many factors in a WWTP, including reactor geometry (particularly aerator immersion depth), aeration bubble size, diffuser layout and liquid viscosity, among others (Gillot et al., 2005). Therefore, the estimation of $k_L a$ for N₂O emission is very difficult in reality, especially considering the dynamic conditions and spatial variability in a wastewater treatment

Download English Version:

<https://daneshyari.com/en/article/6367172>

Download Persian Version:

<https://daneshyari.com/article/6367172>

[Daneshyari.com](https://daneshyari.com)