

INTRODUCTION

The sustainability of any cellulosic bioenergy feedstock production system is in part a function of its ability to decrease net greenhouse gas emissions into the atmosphere compared to the combustion of fossil fuels. Nitrous oxide emissions can be a significant component of the total greenhouse gas budget for any bioenergy feedstock production system. Recent estimates suggest that the feedstock production phase of cellulosic bioenergy production may emit 60% of the total GHG's and that emissions of nitrous oxide (N_2O) are responsible for 36% of this number. Nitrous oxide is a potent GHG that is emitted from soils containing inorganic N (NO₃ and NH₄) (Fig. 2). Therefore N_2O emissions are generally proportional to the rate of N fertilizer applications. However, the rate of N_2O emission may also be influenced by meteorological conditions, soil texture, soil organic C, microbial activity, and crop type. Few studies have evaluated N_2O emissions from forage sorghum and switchgrass grown for cellulosic bioenergy and no research has been conducted to evaluate this important component of the production system in Oklahoma.

MATERIALS AND METHODS

- Nitrous oxide emissions were measured in small plots (9.1 m by) 9.1 m) of forage sorghum and switchgrass located at Stillwater, OK.
- Sorghum and switchgrass plots were treated with urea ammonium nitrate (UAN) (28-0-0) fertilizer at rates of 0, 84, 168, and 252 kg N ha⁻¹ at 4 leaf stage and green up, respectively.
- Nitrous oxide emissions were measured for all sorghum N rates and the 84 kg N ha⁻¹ rate for switchgrass using a vented chamber technique (Mosier et al., 1991).
- Base anchors (38.1 cm by 12.7 cm) were forced into the soil of each plot.
- + A vented chamber lid was placed on the base anchor such that an air tight seal was formed, only allowing air exchange through the vent tube. The air tight seal was formed by placing water in the trough into which the lid was placed (Fig. 1).
- 20 ml air samples were collected from a septum in the chamber at 0, 15, 30 and 45 minutes after the chamber lids were placed on the base anchors (Fig. 1).
- Air samples were stored in evacuated vials and analyzed using a gas chromatograph with an electron capture detector.
- Chamber headspace N_2O concentrations were used to calculate fluxes using linear regression between concentration and time. +These flux measurements were made at 24, 48, 72, 96, 120, 144,
- and 168 hours after application and then weekly for the remainder of the growing season.
- Total growing season emissions were estimated with linear extrapolation between sampling periods.





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Figure 1: Parallel oriented chamber with sampling vials and syringe primed for sampling.

NITROUS OXIDE EMISSIONS AS A FUNCTION OF N APPLICATION RATE TO BIOFUEL FEEDSTOCKS

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Annual N Rate (kg/ha) Figure 3: The relationship between the cumulative N₂O emitted during the 19 month observation period and the annual fertilizer N rate applied.

250



