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Comment

## ***Interactive comment on “Biogenic emissions of NO<sub>x</sub> from recently wetted soils over West Africa observed during the AMMA 2006 campaign” by D. J. Stewart et al.***

### **Anonymous Referee #1**

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This paper presents aircraft NO<sub>x</sub> observations obtained over West Africa during July and August 2006, as part of the AMMA campaign. These observations show a pronounced increase in NO<sub>x</sub> following rain events, consistent with biogenic soil NO<sub>x</sub> emissions. The measurements are very interesting and the relationship between observed NO<sub>x</sub> concentrations and land surface temperature anomaly (a proxy for soil moisture) is striking in the three case studies. Beyond that, I find the interpretation of these measurements in terms of flux measurements and their extrapolation to be flawed. I suggest significant revisions to this paper.

Specific comments

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## 1. Flux calculation: time-dependent vs. steady-state

The authors do not justify the way that they have set up their flux calculation (section 3.4). They assume that at sunrise NO<sub>x</sub> concentrations are the background concentrations measured over dry soil. Then, between sunrise and the time of observations (~8-10 hours later), constant soil NO<sub>x</sub> emissions are applied. Are there no biogenic emissions at night? The observations were typically 18 hours-3 days after a rain event, so why assume that soil NO<sub>x</sub> production only occurred since sunrise? Why assume that the initial NO<sub>x</sub> concentrations at sunrise are at the values observed over dry soil? The dry soil measurements are also taken in the afternoon and that NO<sub>x</sub> is also subject the same loss processes as the NO<sub>x</sub> over wet soils.

Why not, more simply, assume that the NO<sub>x</sub> concentrations are at steady-state, such that  $F = H L C_{wet}$ . This eliminates the need to assume a time over which soil NO<sub>x</sub> emissions occur and also does not require any assumptions about background NO<sub>x</sub> concentrations. In practice this gives results that are very similar to the ones obtained with equation (3), especially for short lifetimes (for OH = 5e6 and 10e6 molec/cm<sup>3</sup>, the NO<sub>x</sub> lifetime is 4 and 2 hours, respectively using the values given by the authors). For OH = 0 and 1e6 molec/cm<sup>3</sup> the NO<sub>x</sub> lifetimes are much longer: 9 days and 21 hours. Actually, a 9 day lifetime for NO<sub>x</sub> is unrealistically long and there is no reason to believe that OH concentrations would be zero and that there would be no chemical loss. I suggest eliminating the case where OH = 0 from the paper.

Also, a more detailed discussion of the OH observations on board the aircraft would be useful. What range of values were observed under what conditions?

## 2. Flux calculation: dry versus wet

It appears that the authors assume that soils that have not been wetted recently do not emit NO<sub>x</sub>. Field measurements (for example summarized by Davidson et al., 1997) show that semi-arid soils emit little NO<sub>x</sub> during the dry season, but then after an initial pulsing following the first rains of the season, soil NO<sub>x</sub> emissions are lower but still

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significant during the remaining of the wet season. Indeed, Stewart et al. measure significant background NO<sub>x</sub> concentrations over dry regions (outside the wet regions), with values ranging from 210-400 ppt. If there are no other significant sources in this region (biomass burning or fossil fuel combustion) as suggested by low VOC and CO concentrations, then significant soil NO<sub>x</sub> emissions would be needed to maintain these high NO<sub>x</sub> concentrations, especially given the deeper boundary layer height.

In fact, assuming steady-state and the same OH levels as the authors, I find that the soil NO<sub>x</sub> emissions needed to maintain these observed NO<sub>x</sub> levels are about 25-60% lower than over wet soils. The factor of 3 difference in NO<sub>x</sub> concentrations is counterbalanced by the factor of 2 difference in boundary layer height. These fluxes cannot be neglected.

### 3. Significance of calculated flux (section 3)

The authors scale up their flux over wet patches to a 2.3 million km<sup>2</sup> over the Sahel. They assume that at any given time 11% of that region has seen recent rain and apply the following formula:

$$\text{Total N} = F \times A \times 0.11 \times t$$

For a 2-month period between July and August they find that these regions account for 0.01-0.05 TgN. They assume that the regions outside these recently wetted areas do not emit biogenic NO<sub>x</sub>. Again, this is not realistic (see section 2 above).

Using flux values inferred from NO<sub>x</sub> observations over dryer areas, I calculate that these dryer regions emit significant levels of NO<sub>x</sub>. Applying the following formula (assuming that these fluxes occur over the remaining 89% of the region):

$$\text{Total N (dry)} = F_{\text{dry}} \times A \times 0.89 \times t$$

the resulting NO<sub>x</sub> emissions range from 0.001 (OH=0) to 0.3 TgN!

Adding the two together, NO<sub>x</sub> emission over the Sahel range of 0.01 to 0.36 TgN. Compared to the top-down satellite estimates in Jaegle et al (1994) of 0.19 TgN, these

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values are not too different, and even exceed them by a factor of almost 2 at the high end of the OH concentrations assumed.

#### 4. Comparison to the Delon et al. paper in ACPD.

The authors briefly mention the Delon et al. paper which applies a mesoscale model and a neural network approach to modeling soil NO<sub>x</sub> emissions. Given that there is significant overlap between the 2 papers, it would be useful to see more of a discussion comparing the results of the Stewart et al. paper to the Delon et al. paper. For example how do the model calculated soil NO<sub>x</sub> emissions in Delon et al. compare to the simpler upscaling approach presented in the present paper?

#### 5. Other comments

+ page 16257, line 2. What is the detection limit and accuracy of the TECO instrument? The values are given for the UEA Noxy instrument but not for TECO.

+ Given that the main argument against an anthropogenic origin for NO<sub>x</sub> is the lack of correlation with CO, I suggest that the authors show the CO measurements on Figure 3.

In summary, the measurements are very interesting and provide new information on biogenic soil NO<sub>x</sub> emissions over regional scales. I commend the authors for attempting a simple approach in extrapolating their results to a larger region, but find their efforts flawed.

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