

# Nitrous oxide emissions from acid sulphate soil at high and low groundwater level in a lysimeter experiment

Asko Simojoki, Seija Virtanen and Markku Yli-Halla Department of Food and Environmental Science Environmental Soil Science University of Helsinki, Finland





## Related research presented in 7IASSC

- <u>Virtanen, S.</u>, Simojoki, A. & Yli-Halla, M. Aluminium and iron concentrations in pore water of an acid sulfate soil

   soil response to waterlogging.
- <u>Epie, K.E.</u>, Virtanen, S., Lammi, A., Simojoki, A. & Stoddard, F. Growth of reed canary grass in high moisture acid sulphate soils.
- <u>Simek, M.,</u> Virtanen, S., Simojoki, A., Krištůfek, V. & Yli-Halla, M. Microbial community in boreal acid sulfate soil: vertical distribution, activity assessment, and potential for greenhouse gas emissions.



#### Background

- 1) Various possible reactions for N<sub>2</sub>O production in wet conditions in soils rich in organic matter
- 2) Potential for high microbial denitrifying enzyme acitivity (DEA) in AS soils
- 3) Large pools of C and N in AS soils

A risk for high N<sub>2</sub>O fluxes from AS soils. The risk may even be increased, if water table is kept high to prevent the oxidation of sulfides.



## Reactions producing N<sub>2</sub>O emissions from AS soils

In agricultural soils the full range of redox reactions for N fertiliser species are possible. This includes: nitrification:

$NH_4^+ + 2O_2^- \rightarrow NO_3^- + H_2O + 2H^+$	
$NH_4^+ + O_2 \rightarrow 0.5 N_2O + 1.5H_2O + H^+$	[2]
denitrification:	
$NO_3^{-} + 1.25c + H^{+} \rightarrow 0.5N_2(g) + 1.75H_2O + 1.25CO_2(g)$	[3]
$NO_3^- + 0.75\{CH_20\} + H^+ \rightarrow NO(g) + 1.25H_2O + 0.75CO_2(g)$	[4]
$NO_3^- + \{CH_20\} + H^+ \rightarrow 0.5 N_2O(g) + 1.5 H_2O + CO_2(g)$	[5]
and reduction:	
$2NO_3^- + 2\{CH_2O\} + 8H^+ \rightarrow N_2 + 6H_2O + CO_2(g)$	[6]

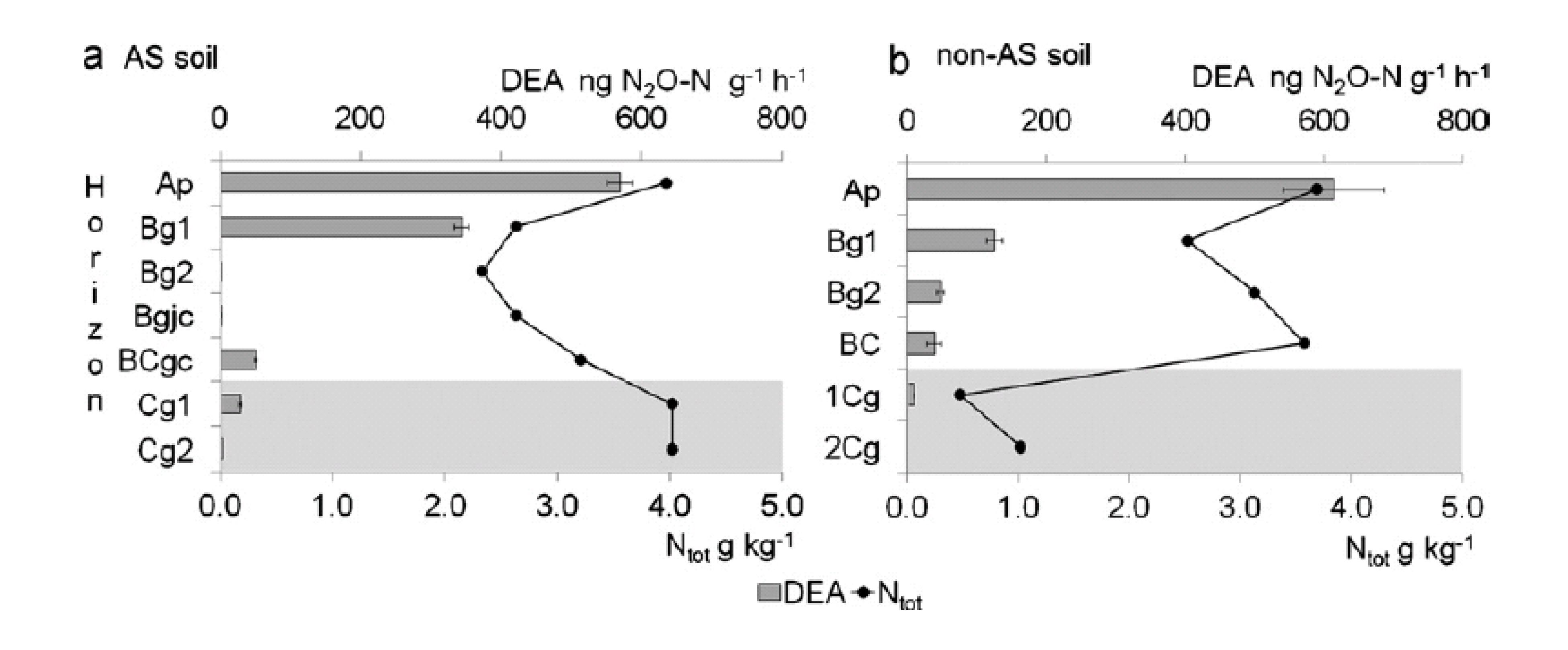
From eqns [3] to [6], acidic soils with high organic carbon content, favour denitrification, releasing N<sub>2</sub>, NO and N<sub>2</sub>O and possibly CO<sub>2</sub>, provided soil water conditions are favourable. In acid sulfate soils, nitrate can also oxidise underlying sulfides producing further denitrification:

$$NO_3^- + 0.7S^- + .03H^+ \rightarrow 0.5N_2(g) + 0.7SO_4^{2-} + 0.1H_2O$$
 [7]  
 $NO_3^- + 0.4S^- + 0.6H^+ \rightarrow NO(g) + 0.4SO_4^{2-} + 0.3H_2O$  [8]  
 $NO_3^- + 0.6S^- + 0.4H^+ \rightarrow 0.5N_2O(g) + 0.6SO_4^{2-} + 0.2H_2O$  [9]

Again reactions [7] to [9] seem favoured by acidic conditions. In acid sulphate soils conditions appear optimum for the conversion of fertiliser N to gaseous N provided there is sufficient soil N, organic matter and water.



# High denitrifying enzyme activity in AS soils



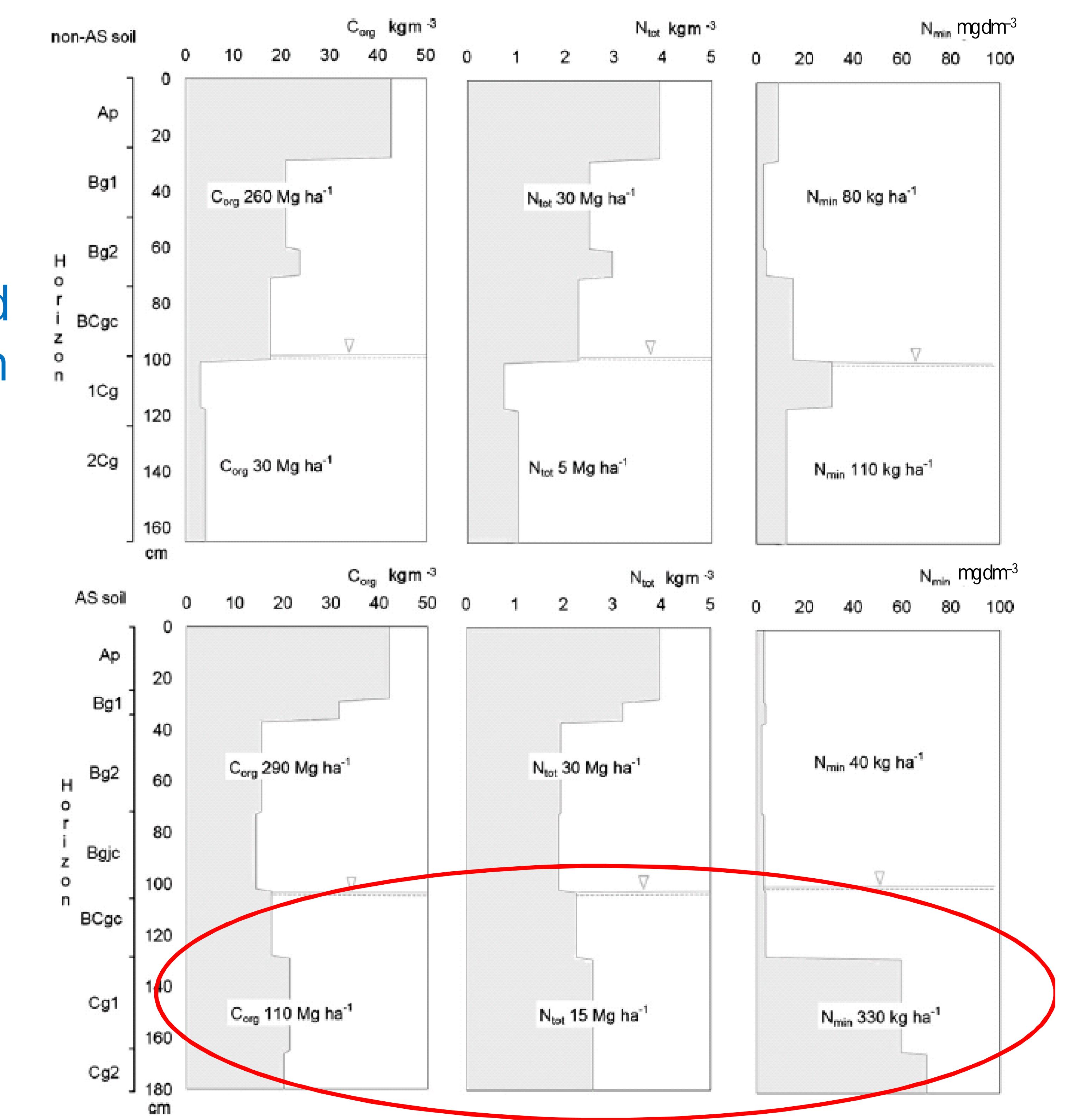
Evidence of rich microbial communities in the subsoil of a boreal acid sulphate soil conducive to greenhouse gas emissions

Miloslav Šimek<sup>b,c</sup>, Seija Virtanen<sup>a,\*</sup>, Václav Krištůfek<sup>b</sup>, Asko Simojoki<sup>a</sup>, Markku Yli-Halla<sup>a</sup>

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Total C, total N and mineral N stocks in the soil profiles of non-AS and AS soils.



\$imek et al. 2011. AEE 140:113-122



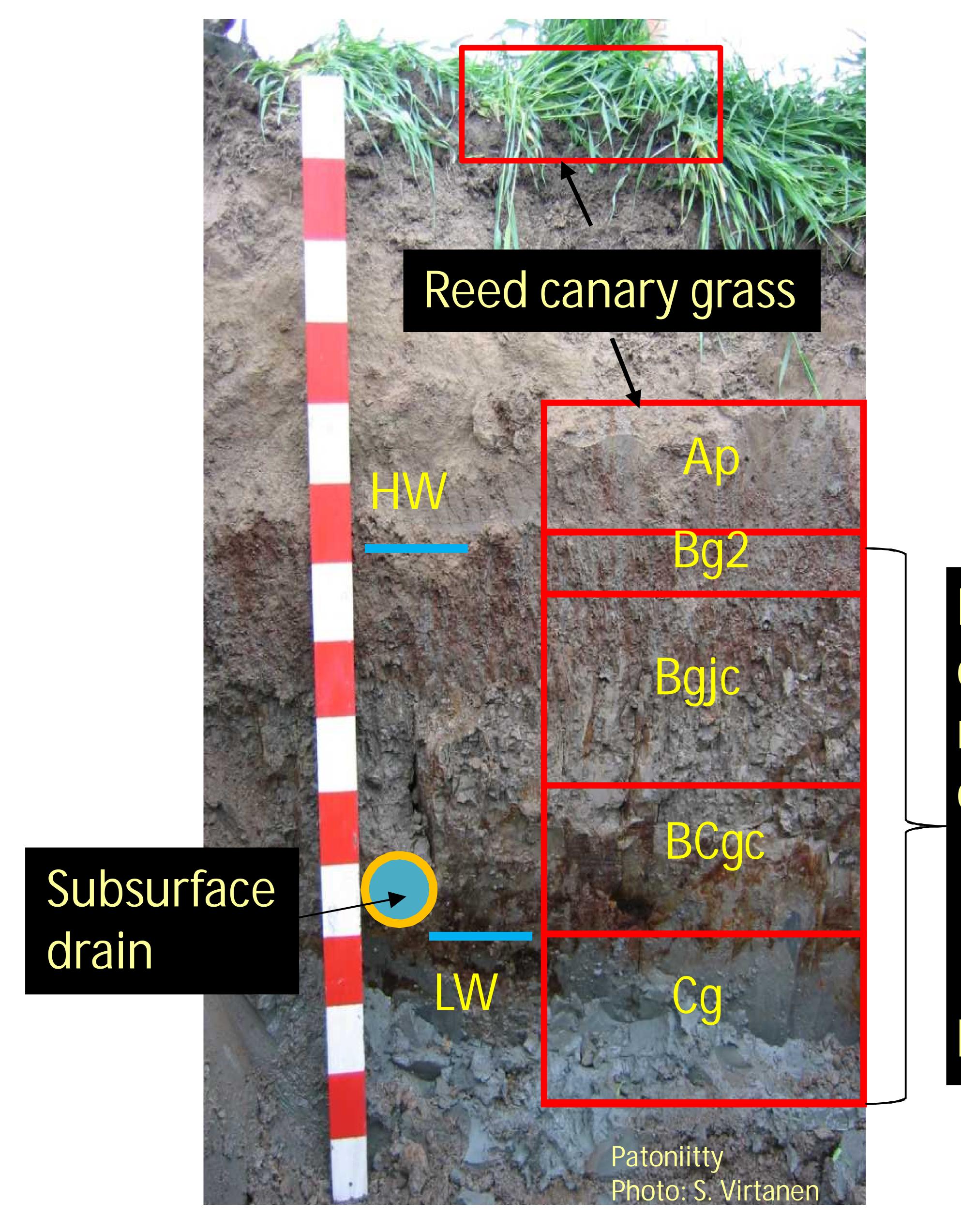
#### The aim of study

Do high groundwater levels increase N<sub>2</sub>O emissions from acid sulphate soils?

Are N<sub>2</sub>O emissions similar from cropped and uncropped acid sulphate soil?



#### The pedon and the characteristics of soil



Horizon		рH	$BS^1$	Total
	Clay	Fresh		S
	%		%	$g kg^{-1}$
Ap	33	6.4	73	0.91
Bg2	48	4.4	25	0.94
Bgjc	61	3.8	19	4.21
BCgc	57	4.2	27	4.57
Cg	58	6.5	76	14.87

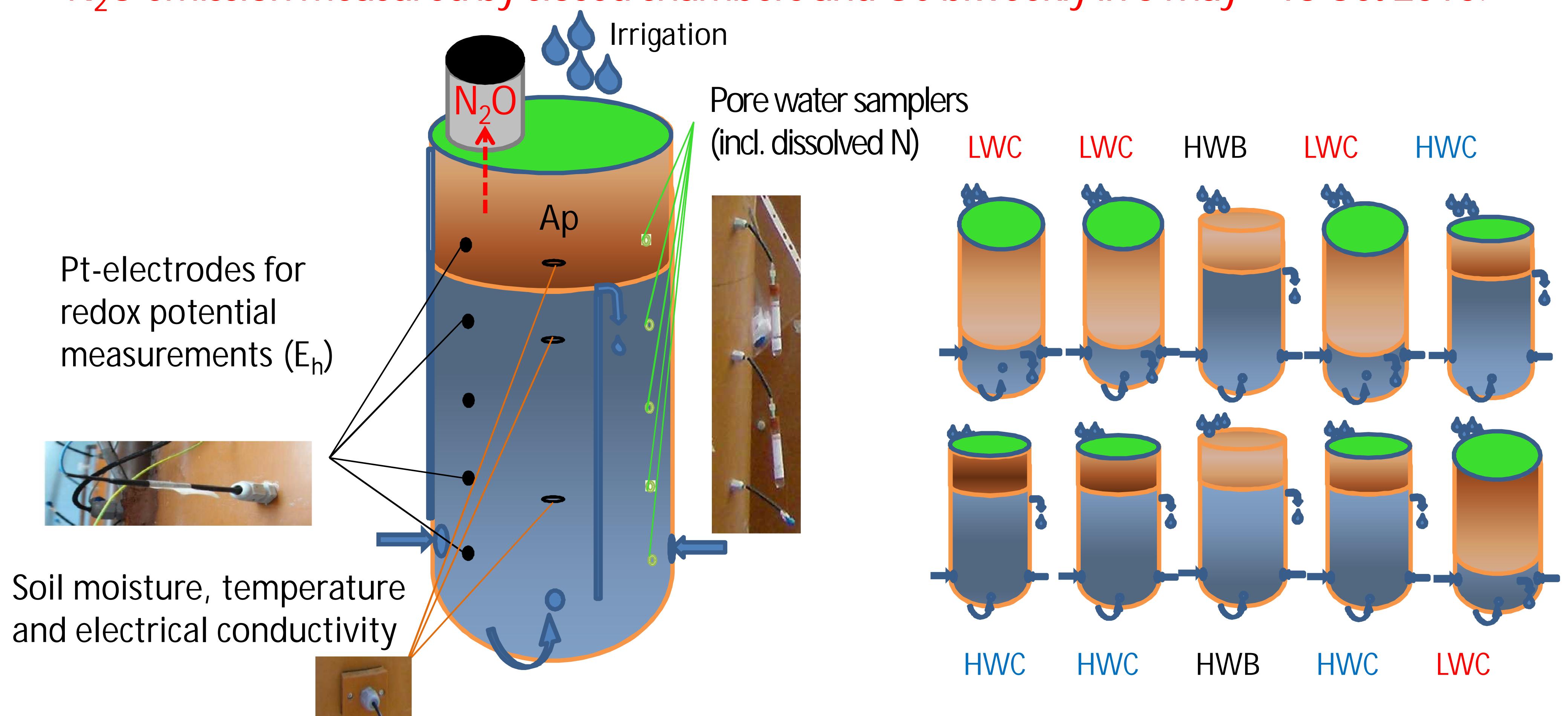
<sup>1</sup>(Na<sup>+</sup>, K<sup>+</sup>, Ca<sup>2+</sup>, Mg<sup>2+</sup>)/CEC\*100

Sulfic Cryaquept



#### The setup of experiment and measurements

- Ground water table: high (20 cm depth) or low (70 cm depth).
- Irrigation: according to local long-term precipitation + two 2-day periods with +25 mm/day.
- N<sub>2</sub>O emission measured by closed chambers and GC biweekly in 5 May 13 Oct 2010.



Eh, **\theta**, T and EC were logged continuously. pH, Al, Fe and DOC in pore water were determined biweekly or monthly.

LWC low water table, cropped HWC high water table, cropped HWB high water table, bare soil

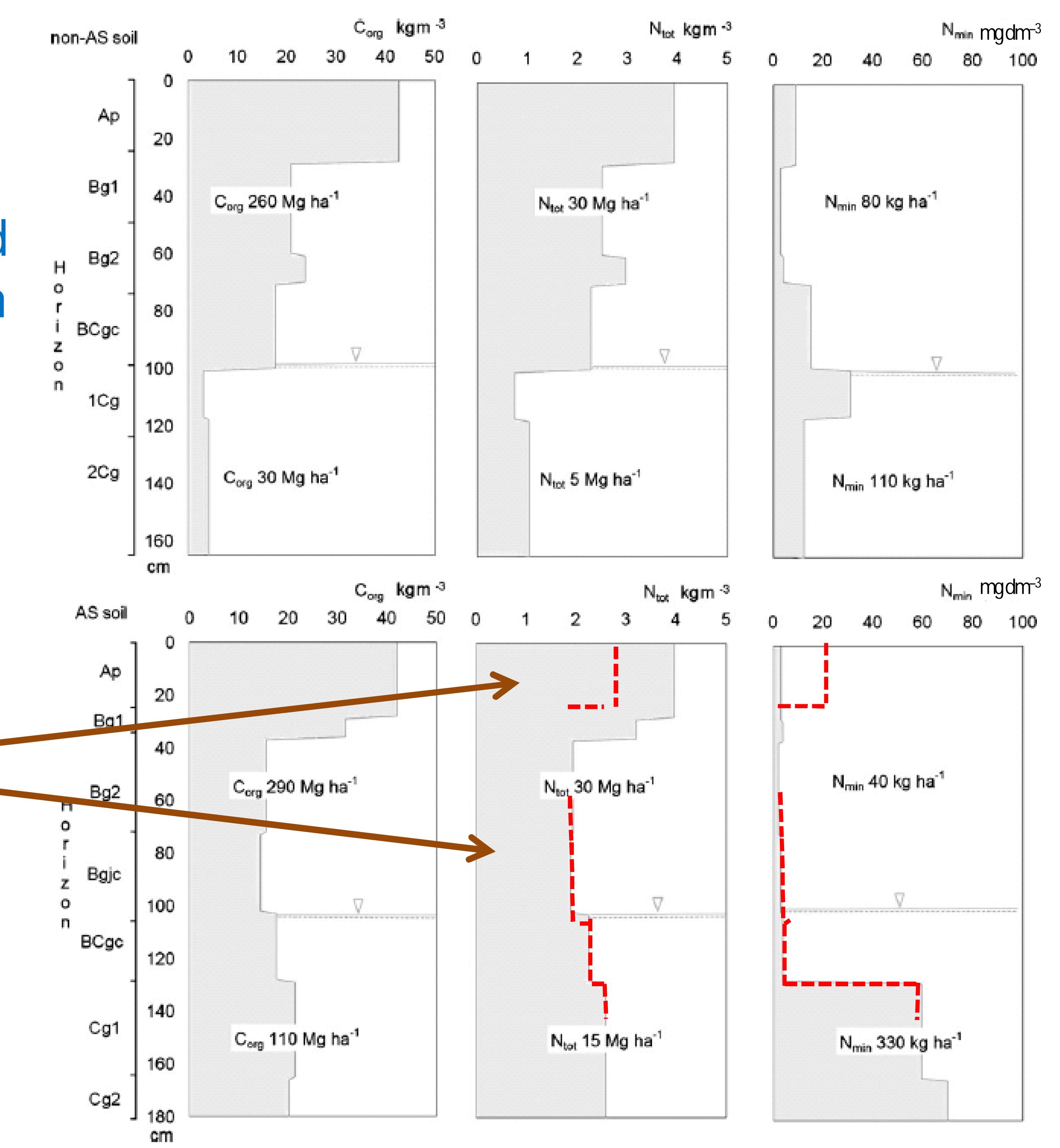


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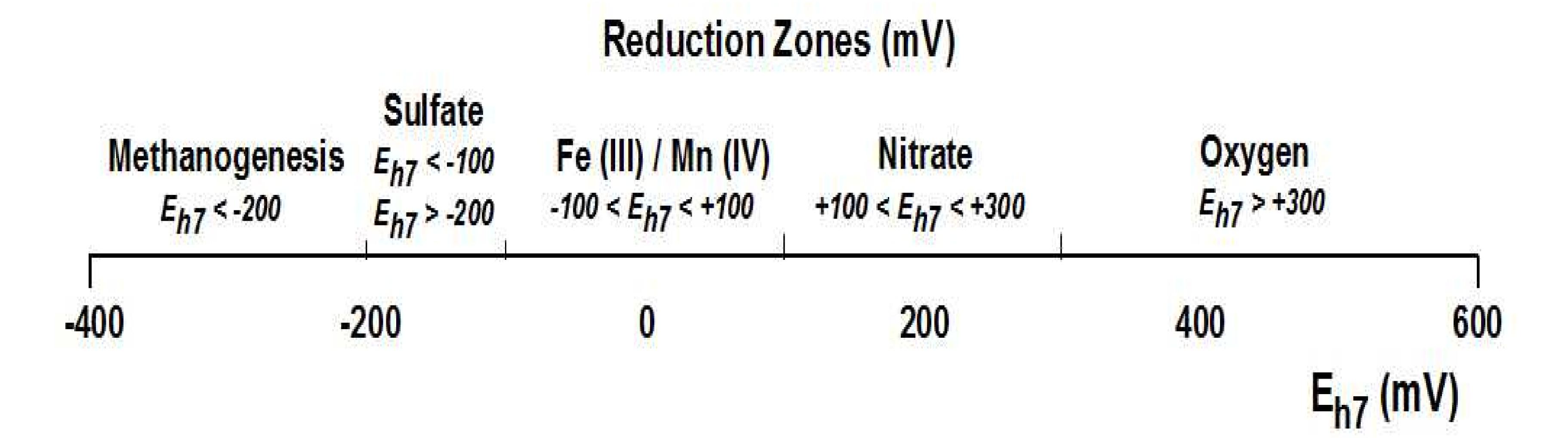
The soil layers taken into the Jusimeters.

N fertilization in early May corresponded to N 90 kg/ha/yr





# Ranges of reduction potentials for various compounds



The reduction zones adapted from Reddy et al. (2000).



Soil N<sub>2</sub>O emissions, dissolved N and water content in different experiment treatments in 2010

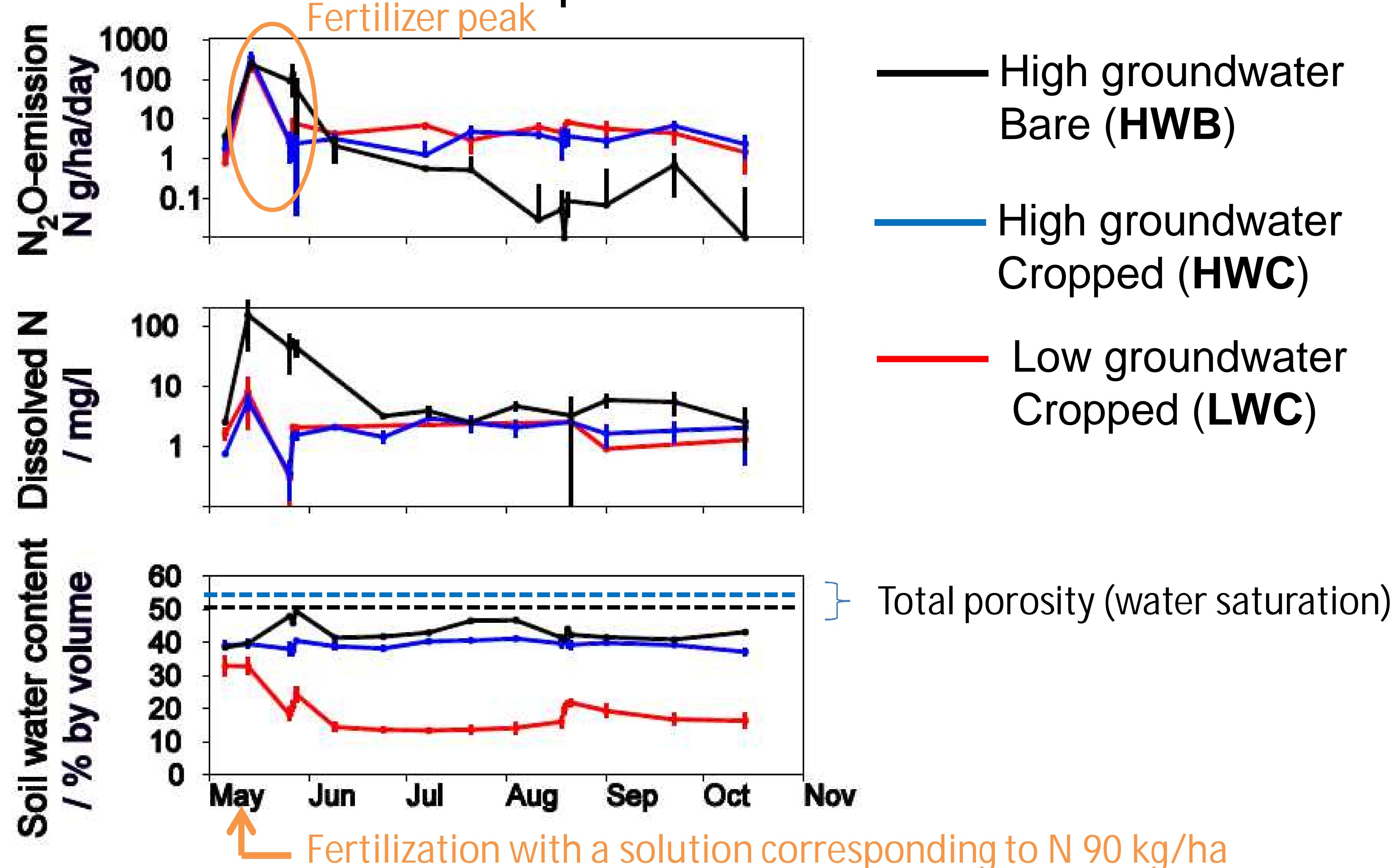


Fig 1. The emission of  $N_2O$  from soil, the concentration of dissolved N in pore water (10 cm depth), and the water content (15 cm depth) at different experimental treatments (mean  $\pm$  standard error, n = 2-4).



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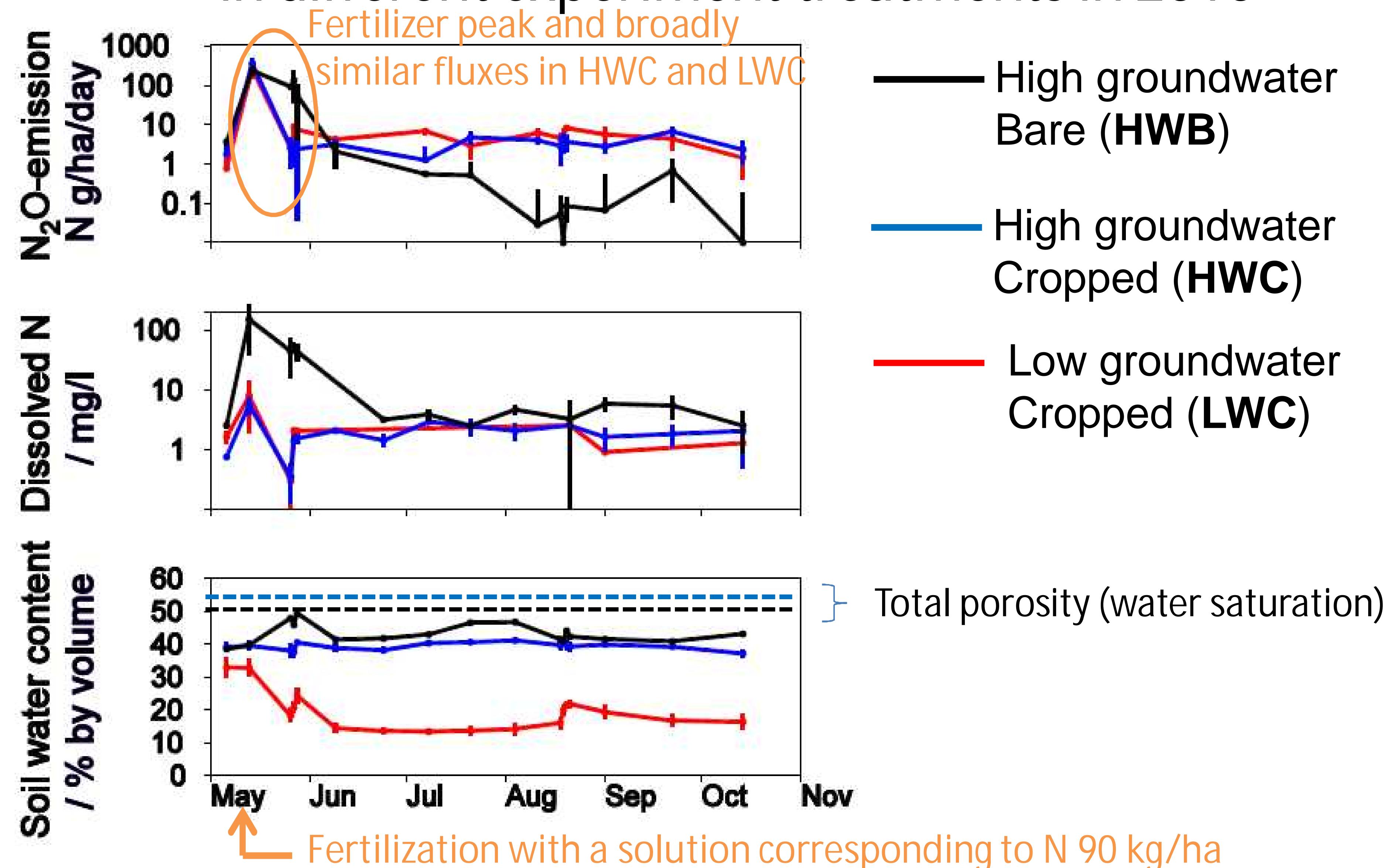


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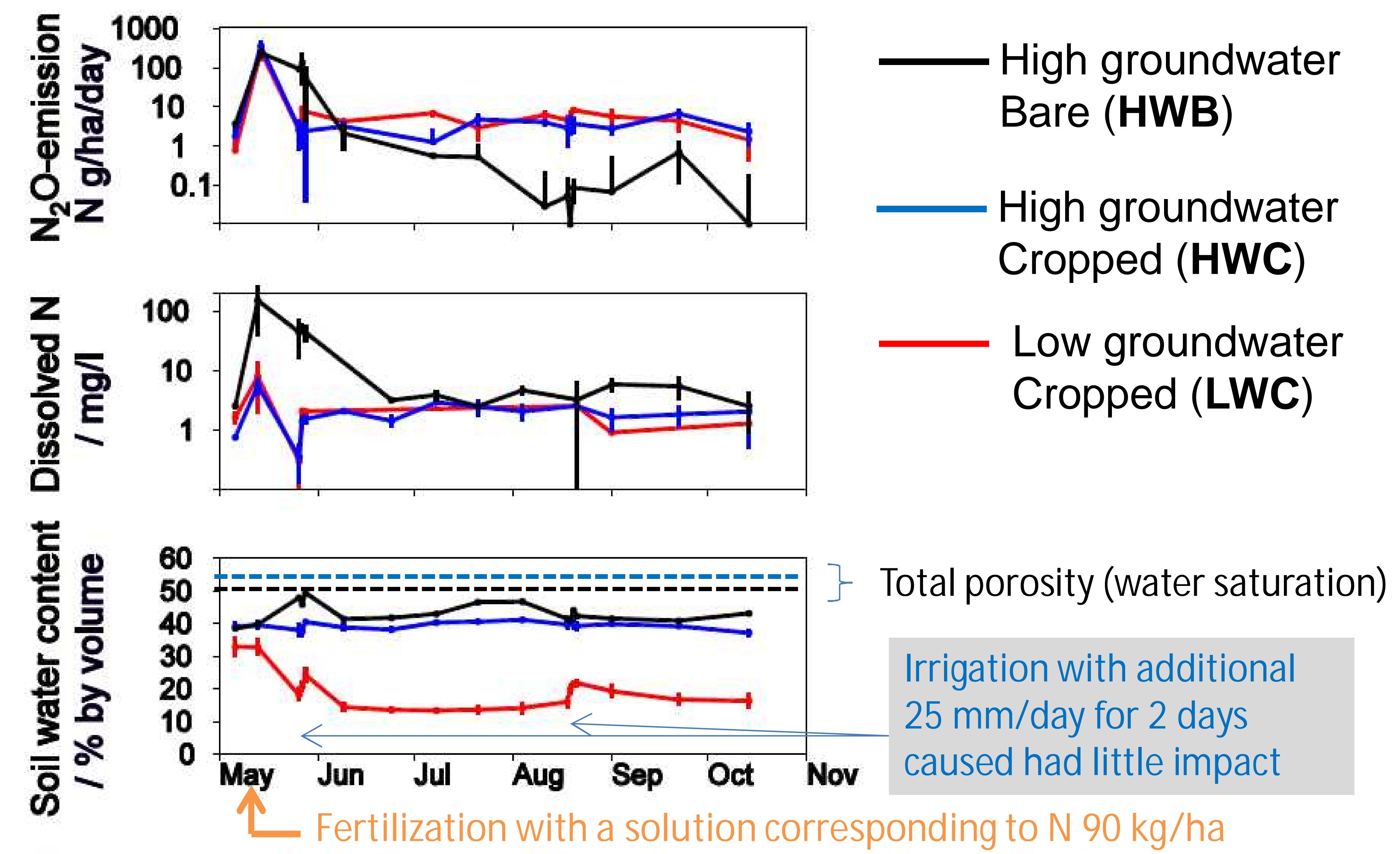


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#### Discussion

- In HWC, topsoil water content was always about 40%
- ⇒ max 10-15% air space, and WFPS (water-filled pore space) >70%.
- ⇒ at least temporarily deficient aeration in soil
- ⇒ high N<sub>2</sub>O emissions after fertilization
- In LWC, topsoil water content was usually about 15% (40% air) and WFPS < 30%
- ⇒ oxygen deficiency and large denitrification in topsoil unlikely
- $\Rightarrow$  production of N<sub>2</sub>O in deeper soil horizons?
- Immediately after the fertilization, however, topsoil water contents in the LWC were still >30% (WFPS about 60%)
- ⇒ significant denitrification and large N<sub>2</sub>O fluxes possible

In HWB, topsoil water contents were still >40% (<10 % air) and WFPS >80%

⇒ significant denitrification and large N<sub>2</sub>O fluxes likely



#### Conclusions

Nitrogen fertilization causes large N<sub>2</sub>O emissions from AS soils immediately after the fertilization as in other soil types.

Plant nutrient uptake decreases efficiently the dissolved N in pore water and the N<sub>2</sub>O emissions from soil to the atmosphere.

Cumulative N<sub>2</sub>O emissions (corresponding to about N 4 kg/ha during 161 days) are somewhat higher compared with most mineral soils.

This experiment does not support the view that raising the groundwater level to prevent sulphide oxidation would significantly increase N<sub>2</sub>O emissions from AS soils to the atmosphere.



### Literature

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