

Nitrous Oxide (N₂O) emissions in a continuum plot-groundwater-wetland-river in a farming catchment area



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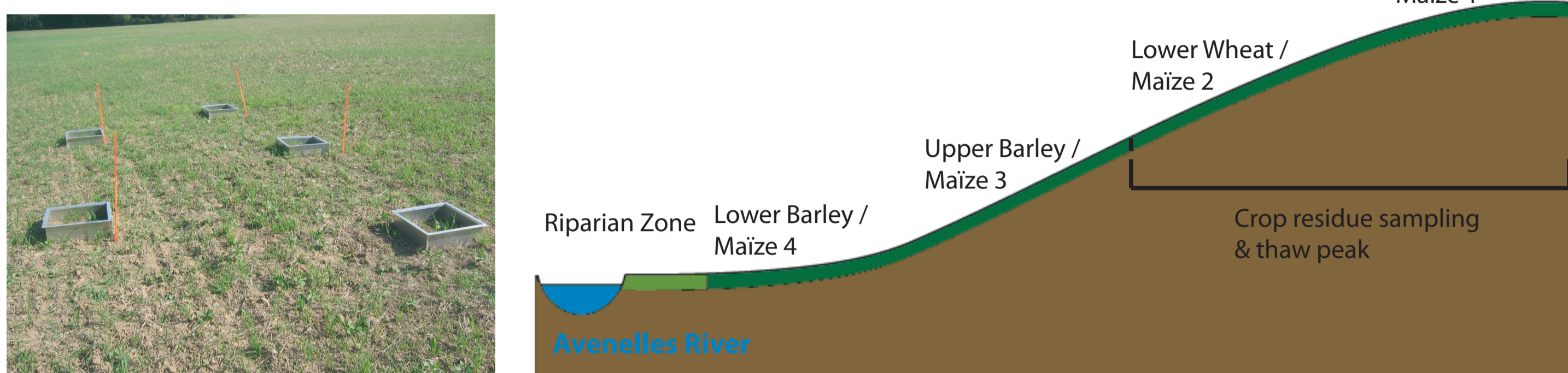
1 Scope of the study

Nitrous oxide (N₂O) has an important role in the atmosphere; it is a greenhouse gas with a 296 times larger global warming potential than carbon dioxide over a 100 year time period (IPCC, 2001). Soil contribution of global emission of N₂O is 70% (Conrad and Smith 1995) with a major contribution of agricultural soils, responsible for 67% of anthropogenic N₂O emissions (UNFCCC, 2003).

The purpose of this study was to analyse environmental factors (texture and organic matter content of soil, crop management and fertilizer application) in a continuum from an agricultural plateau to the riparian zone, in order to quantify emissions of N₂O (IPCC 2006).

3 N₂O emissions by soils

Material & Methods



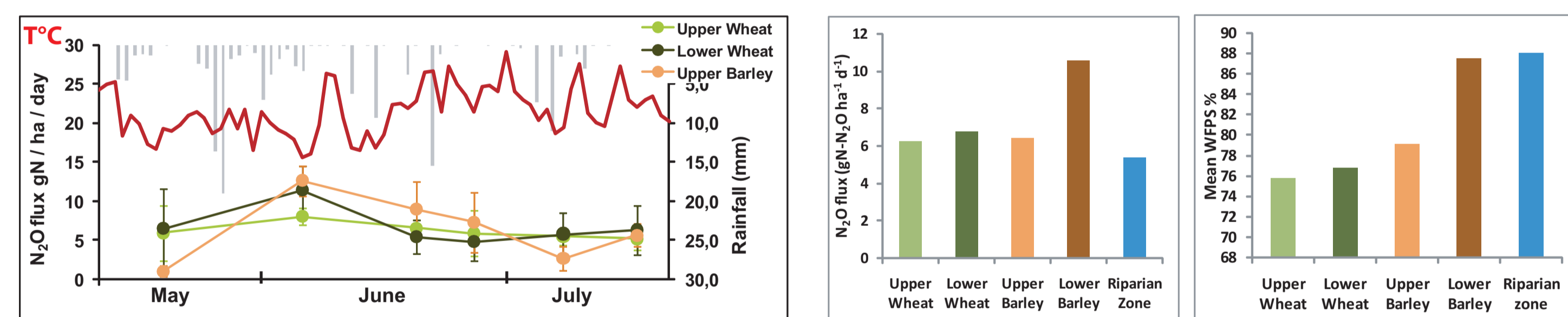
Nitrous oxide (N₂O) gas emission is measured with the "closed chamber technique" with 5 sampling plots at each site.

N₂O concentrations in the gas sample is analyzed using a gas chromatograph (GC) coupled with electron capture detector (ECD).

N₂O fluxes are determined by measuring N₂O concentration increases in the chamber headspaces and calculating the slope of linear regression of N₂O concentrations in function of time.

Soil and air temperature are measured around each chamber as well as soil moisture and organic matter content.

SPRING - SUMMER 2008 : Wheat and Barley



Mean N₂O emissions (a) and mean soil humidity (WFPS) (b)

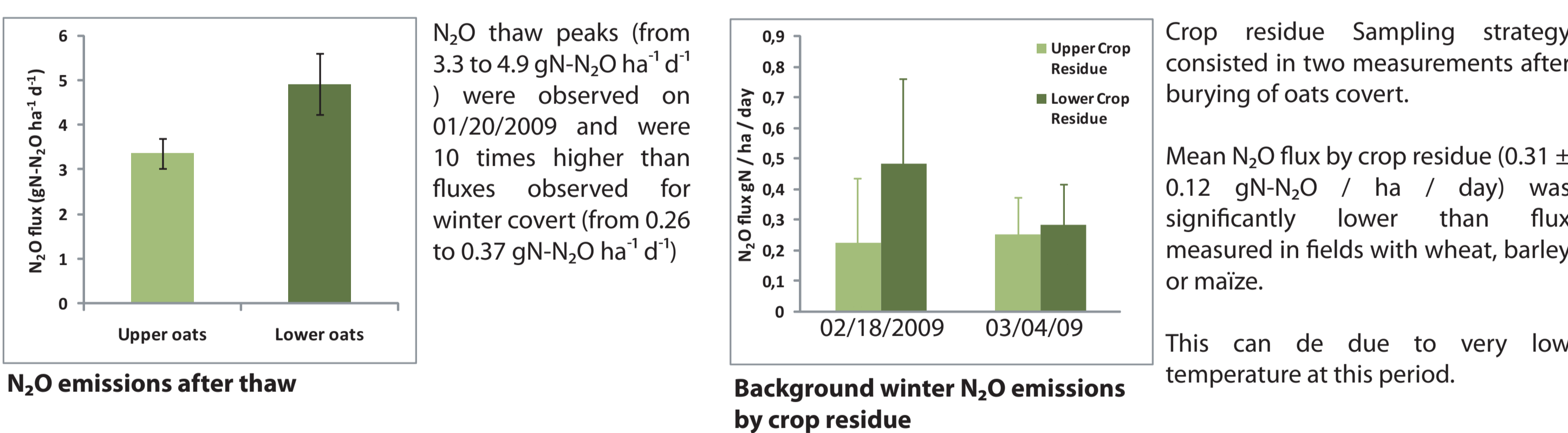
Mean N₂O emissions, from 6 to 10 gN-N₂O ha⁻¹ day⁻¹ for croplands are well in the range of those found in other regions of the world (Bouwman, 1996; Garnier et al., 2009).

Mean N₂O emissions show a distinct landscape pattern. The mean fluxes of the footslope level (LB) are significantly higher than the fluxes in shoulder position (UW) or midslope position (LW and UB). The same relationship with landscape position is observed with WFPS.

Topographic factor, to which water content is closely linked, seems to be a major controlling factor of N₂O emissions at the microscale level.

N₂O emissions during sampling campaign. Plots in slope (a) and backslope (b)

WINTER 2008-2009 : Thaw peak & N₂O emissions by crop residue



N₂O emissions after thaw

N₂O thaw peaks (from 3.3 to 4.9 gN-N₂O ha⁻¹ d⁻¹) were observed on 01/20/2009 and were 10 times higher than fluxes observed for winter covert (from 0.26 to 0.37 gN-N₂O ha⁻¹ d⁻¹)

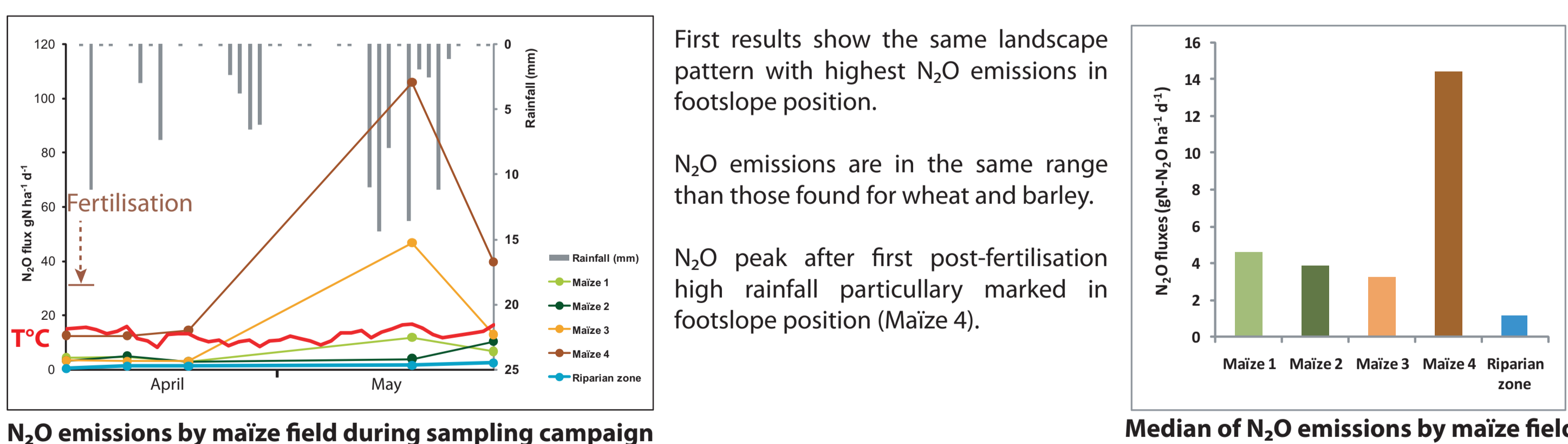
Background winter N₂O emissions by crop residue

Crop residue Sampling strategy consisted in two measurements after burying of oats cover.

Mean N₂O flux by crop residue (0.31 ± 0.12 gN-N₂O / ha / day) was significantly lower than flux measured in fields with wheat, barley or maize.

This can be due to very low temperature at this period.

SPRING - SUMMER 2008 : Maize (in progress)



N₂O emissions by maize field during sampling campaign

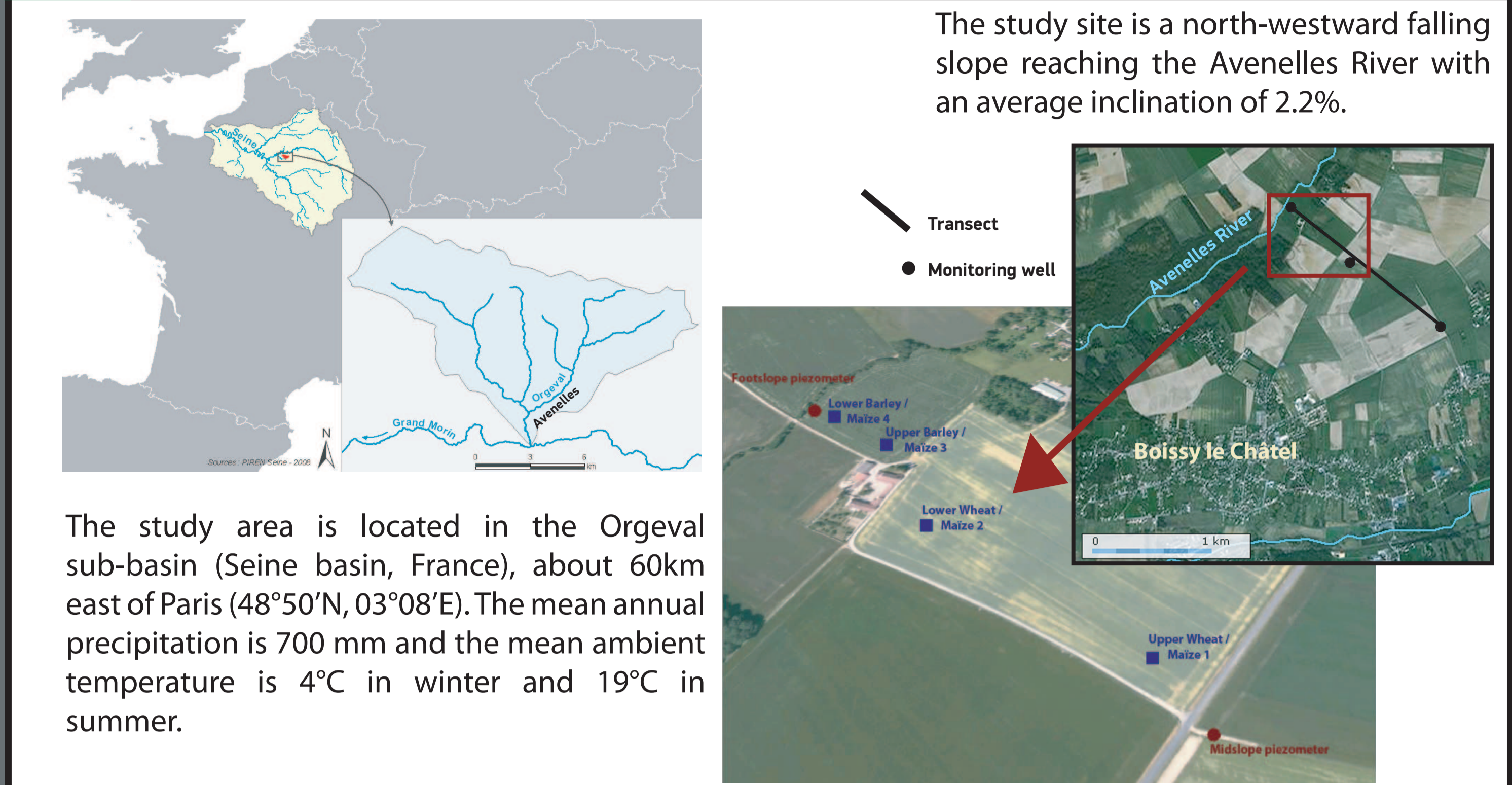
First results show the same landscape pattern with highest N₂O emissions in footslope position.

N₂O emissions are in the same range than those found for wheat and barley.

N₂O peak after first post-fertilisation high rainfall particularly marked in footslope position (Maize 4).

Median of N₂O emissions by maize field

2 Study site : The Orgeval sub-basin



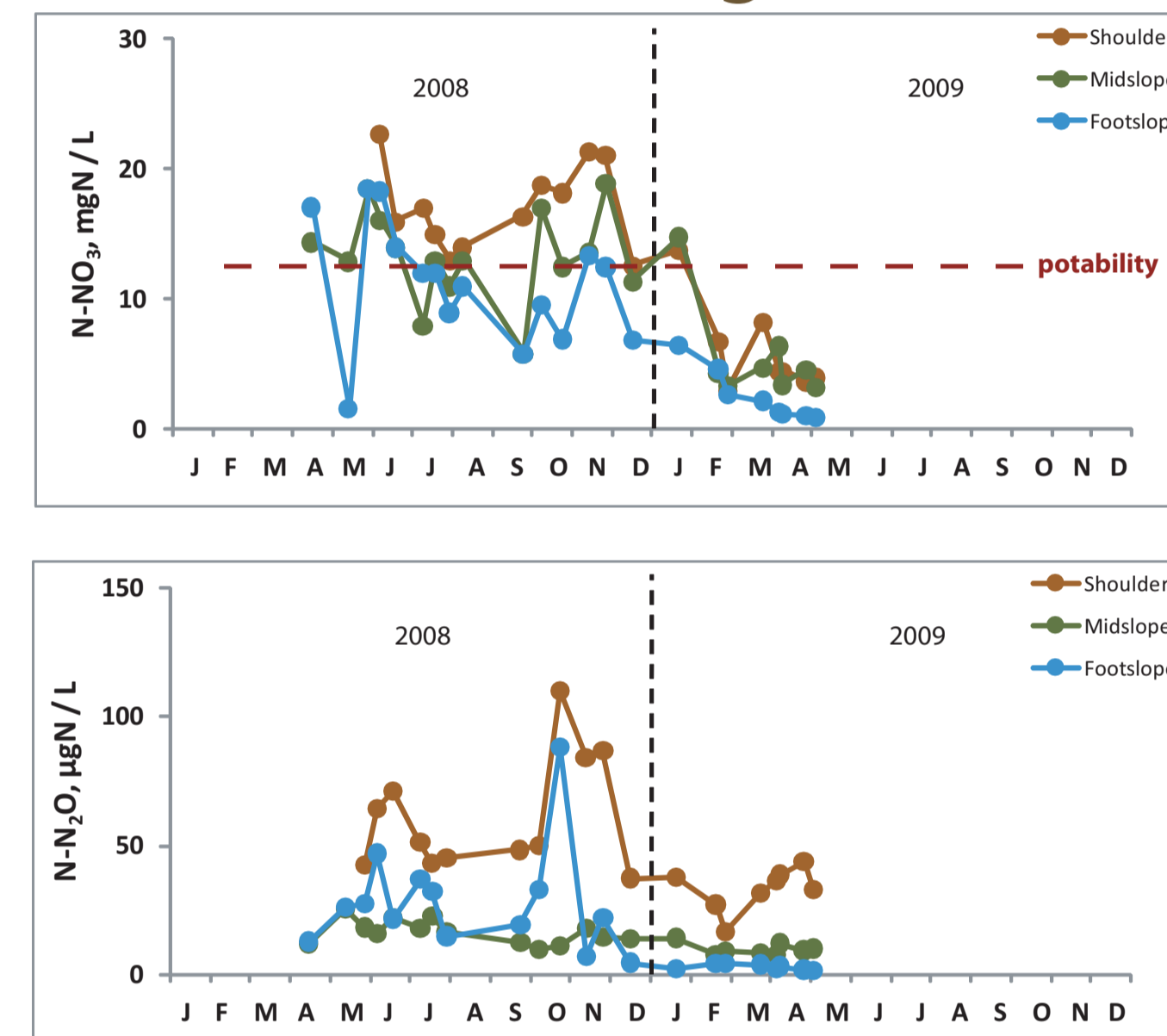
The study area is located in the Orgeval sub-basin (Seine basin, France), about 60km east of Paris (48°50'N, 03°08'E). The mean annual precipitation is 700 mm and the mean ambient temperature is 4°C in winter and 19°C in summer.

4 NO₃ and N₂O in groundwater and rivers

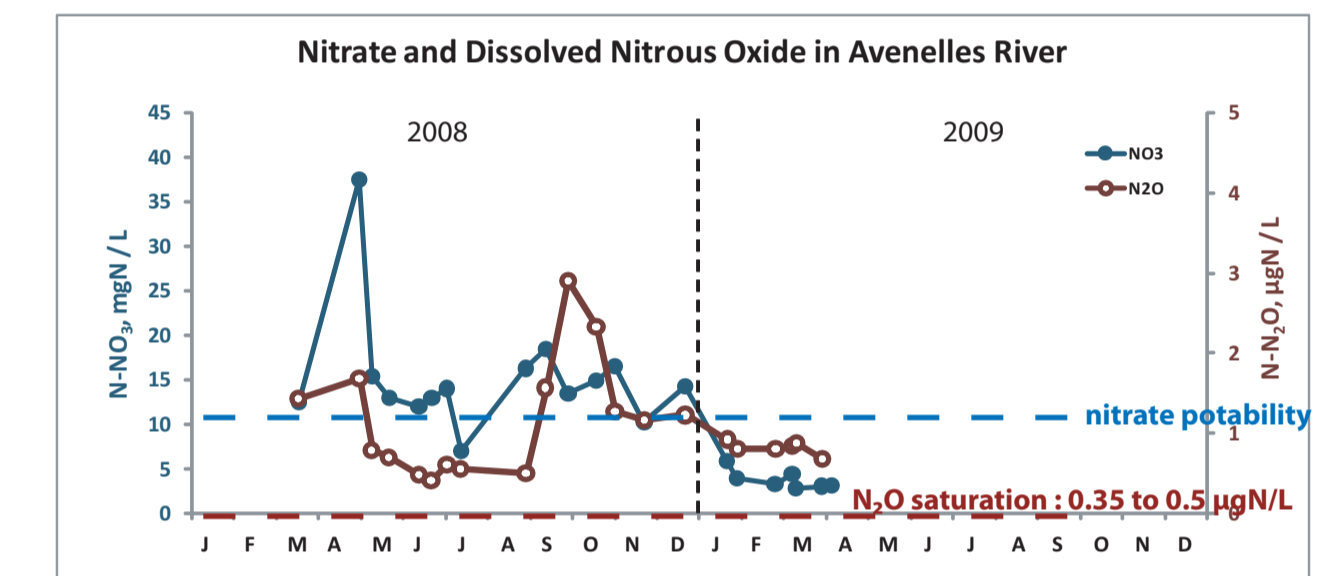
Dissolved N₂O and nitrate in the groundwater and rivers are sampled with a submersible pump. Dissolved nitrous oxide is analyzed, following desorption, with a gas chromatograph (PERICROM ST200) with an electron capture detector (ECD).

Nitrate is measured on filtered water (GF/F 0.7µm of porosity) with an autoanalyser (Quatro), after cadmium reduction into nitrite with the sulphanimide method.

Nitrate and N₂O in groundwaters



Nitrate and N₂O in rivers



Dissolved N₂O in groundwater and rivers always above saturation with very higher concentrations in groundwater.

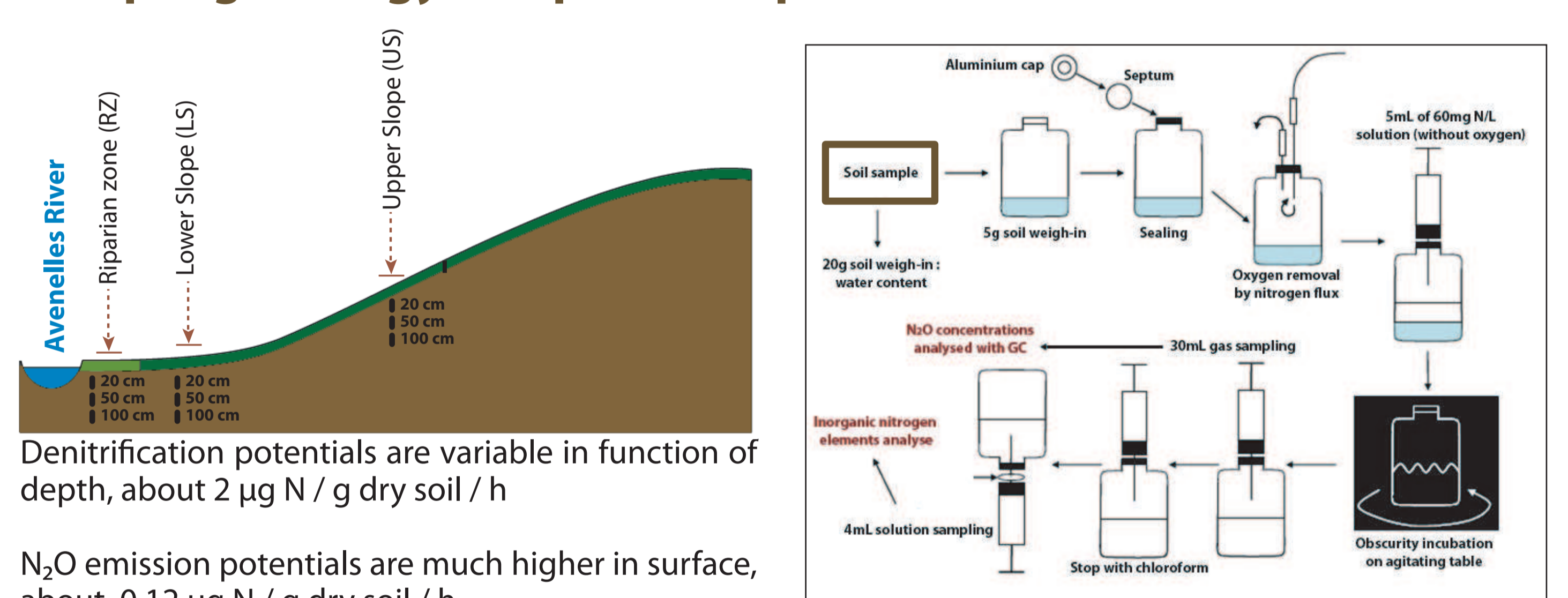
N₂O concentration gradient along the slope with highest concentrations in upper position (more marked for N₂O).

In 2009, nitrate concentrations is in the same order in groundwater and rivers.

Two key periods for highest concentrations of nitrate and dissolved N₂O : spring (2008 & 2009) after fertilisation and fall (after first high post-summer rainfalls)

5 Soil potential denitrification : a lab study

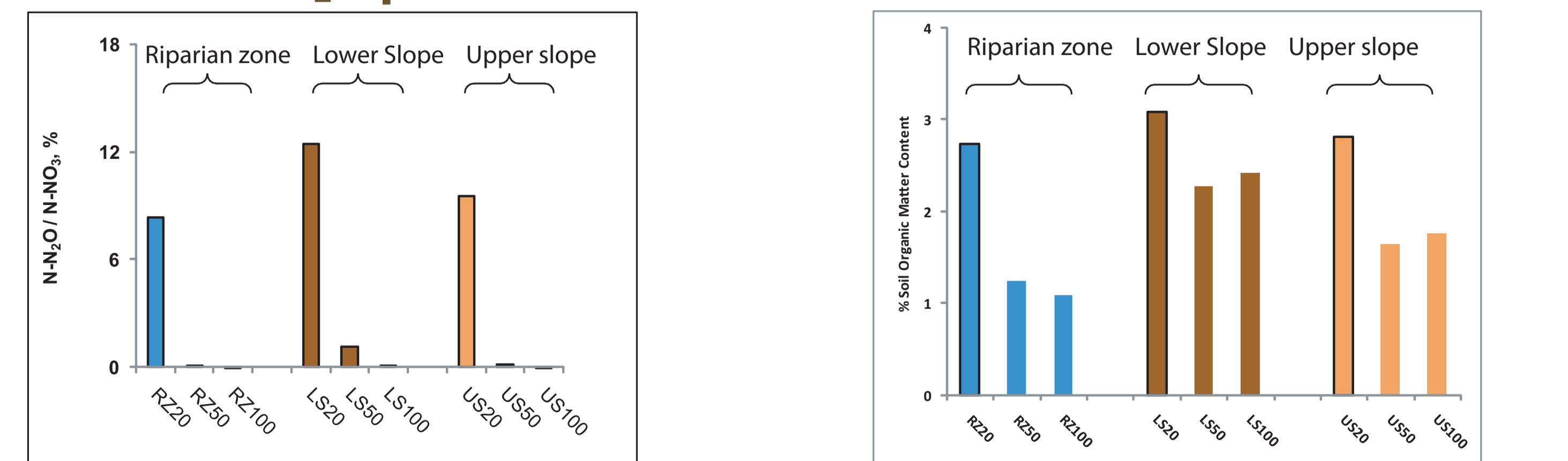
Sampling strategy & Experiment protocol



Denitrification potentials are variable in function of depth, about 2 µg N / g dry soil / h

N₂O emission potentials are much higher in surface, about 0.12 µg N / g dry soil / h

Results : % N₂O production



N₂O emission potentials about 10% of surface denitrification potential...

... in relation with a higher organic matter content

Conclusion

This study has permitted to show that landscape position affects N₂O emissions as well as soil characteristics like WFPS.

Values from 6 to 10 gN-N₂O ha⁻¹ day⁻¹ for croplands are well in the range of those found in other regions of the world (Bouwman, 1996; Garnier et al., 2009). Laboratory experiments show that N₂O production are potentially higher at the surface, so that besides being emitted N₂O could reach the groundwaters by infiltration. During the water transfer from aquifer to surface water, N₂O could be further reduced or emitted.

N₂O concentrations in groundwaters are clearly higher than concentrations found in rivers and huge compared with that of the saturation, with a clear gradient along the slope. Soils appear to be a real source of N₂O for the river water.

References

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