

## Using the N2/Ar-Method to check modelled diffuse NO<sub>3</sub><sup>-</sup> emissions from soils into the groundwater of Lower Saxony (Germany)

Lisa Krienen (1), Heinrich Höper (2), Wolfram Eschenbach (1), Reinhard Well (1), and Jörg Elbracht (2) (1) Thünen-Institut für Agrarklimaschutz, (2) Landesamt für Bergbau, Energie und Geologie

Diffuse  $NO_3^-$  emissions derived from agricultural N surpluses are the main cause of  $NO_3^-$  pollution of aquifers and open water bodies. Denitrification is the key process for the attenuation of these anthropogenic  $NO_3^-$  concentrations in soils and groundwater. Since the greenhouse gas N2O is an obligate intermediate of denitrification this process is also a major regulator of N2O emissions from soils and indirect N2O fluxes from aquifers and open water bodies which result from  $NO_3^-$ -leaching. Up to now the denitrification potential of soils and the potential  $NO_3^-$  concentration in the groundwater recharge are modelled from agricultural N-surpluses, water balances (GROWA) and soil properties (DENUZ) (Wendland et al. 2009) (LBEG 2008).

In this study we compare modelled NO<sub>3</sub><sup>-</sup> emissions (pot-NO<sub>3</sub><sup>-</sup>) (DENUZ) to the groundwater recharge with the calculated initial NO<sub>3</sub><sup>-</sup> concentrations in the groundwater at time of groundwater recharge (NO<sub>3</sub><sup>-</sup>t0) (N2/Armethod (Weymann et al. 2008)). NO<sub>3</sub><sup>-</sup>t0 can be calculated from the measurement of dissolved gases N2, N2O, Ar and NO<sub>3</sub><sup>-</sup> concentrations in groundwater samples.

We analysed groundwater samples from 534 groundwater monitoring wells throughout Lower-Saxony (Germany). Median  $NO_3^-$  and  $NO_3^-t0$  concentrations were 0,4 and 29 mg  $NO_3^-t^-1$  respectively, showing that considerable proportions of the anthropogenic N-surplus is denitrified within the saturated zone.

First results showed a good agreement between measure and modelled  $NO_3^-$  emissions for areas of coastal marshes in the North of Lower-Saxony (predominantly Fluvisols). Medians of measured and modelled  $NO_3^-$  emissions are 12,5 mg  $NO_3^-$ t0 l<sup>-1</sup> and 0,3 mg pot- $NO_3^-$  l<sup>-1</sup> (mean values 20 mg l<sup>-1</sup>  $NO_3^-$  pot and 9,3 mg l<sup>-1</sup>  $NO_3^-$ t0), respectively. Compared to the coastal marshes and in accordance with modelled pot- $NO_3^-$  concentrations our measurements show small-scale spatial heterogeneities of  $NO_3^-$ t0 concentrations in soil regions where the dominant parent material of soils are glacio fluviatile and moraine deposits (predominantly Podzols, Cambisols and Gleysols) in Lower-Saxony. In these regions the median of measured  $NO_3^-$ t0 concentrations was between 29 and 38 mg  $NO_3^-$  l<sup>-1</sup> and on average 25 to 30 mg  $NO_3^-$  l<sup>-1</sup> below the modelled  $NO_3^-$  l<sup>-1</sup> concentrations. To further compare the modelled  $NO_3^-$  emissions (pot- $NO_3^-$ ) with results of the N2/Ar-method ( $NO_3^-$ t0 values) we select groundwater monitoring wells with more homogenous soil properties, soil water residence times in their catchment areas.

We expect that further analysis will help to validate existing denitrification models.  $NO_3^-t0$  values might then be used as a lower boundary condition if denitrification in soils is modelled.

References:

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