## Abstract

The influence of atmospheric processes on deposition of inorganic nitrogen into the southern North Sea was investigated. The three dimensional mesoscale chemistry transport model MECTM (model system: 'Gas') was extended by a simple aerosol scheme (model system: 'Gas + einf. Aero') and by the sectional aerosol model SEMA (model system: 'Gas + SEMA'). The calculated aerosol ions are  $NO_3^-$ ,  $SO_4^{2-}$ ,  $Cl^-$ ,  $NH_4^+$ ,  $Na^+$ ,  $H^+$ . For the gas phase chemistry, 60 species (e.g.  $HNO_3$ , NO,  $NO_2$ ,  $O_3$ ,  $SO_2$ ) are used. Calculations with the three model systems for concentration and for deposition of species were performed for the period of June 16-20, 1998. The results of all model systems were compared with measurements of the campaign of the EU-project ANICE and measurements from the ETC/ACC- network.

The mean concentrations from the model systems agree well with observations for the species  $NH_3$ ,  $NO_3^-$  and  $NH_4^+$ . The concentrations calculated by all model systems for  $HNO_3$  were too high, but the model system 'Gas + SEMA' yielded the smallest deviation. The results of all model systems were compared with measurements of the ETC/ACC-network. In the beginning of the simulated period the modelled concentrations of  $O_3$  were too low and at the end too high. The simulated concentrations of NO,  $NO_2$ ,  $NH_3$ , and  $SO_2$  were within the allowed deviations. With the model systems 'Gas' and 'Gas + einf. Aero' maximum nitrogen depositions of  $3.3 \, mg \, m^{-2} \, day^{-1}$  were calculated whereas the model system 'Gas + SEMA' calculated a deposition of up to  $12.8 \, mg \, m^{-2} \, day^{-1}$ . Investigating the influences on the input of nitrogen in coastal waters led to the conclusion that the sedimentation velocity is a key factor for the amount of deposition.

Scenarios of changing emissions of  $NO_x$  and NMVOC were calculated with the model system 'Gas + einf. Aero'. The scenarios showed that changing the fraction of emission of  $NO_x$  and NMVOC considerably influenced the concentration of  $O_3$ . The spatial and temporal patterns of the concentrations of  $O_3$  and  $HNO_3$  and the deposition of nitrogen differ from the reference case. The scenarios show a nonlinear effect of changes in  $NO_x$  and NMVOC emissions on the concentration of  $O_3$  and  $HNO_3$ .