

Nitrogen circulation in a Mediterranean holm oak forest, La Castanya, Montseny, northeastern Spain

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Abstract

Bulk deposition, wet-only deposition, throughfall and dry deposition inferred from washing foliage and surrogate surfaces were used to describe inorganic N inputs to a forested catchment in the Montseny Mountains (La Castanya, Catalonia, Spain). Bulk inputs of inorganic N were moderate, with a mean of $5.7 \text{ kg N ha}^{-1} \text{ yr}^{-1}$, ranging between 4 and $10 \text{ kg N ha}^{-1} \text{ yr}^{-1}$ for the period 1983 to 2000. Dry deposition fluxes estimated from washing branches added about $9 \text{ kg N ha}^{-1} \text{ yr}^{-1}$ to wet inputs and the total atmospheric deposition was estimated in $15 \text{ kg N ha}^{-1} \text{ yr}^{-1}$. Despite this substantial input flux, nearly all the inorganic nitrogen was retained within the forest ecosystem: NH_4^+ and NO_3^- concentrations decreased dramatically as water crossed the canopy and the soil profile. In the stream, at baseflow conditions, NH_4^+ and NO_3^- concentrations were always below the analytical detection limit ($< 2 \mu\text{eq L}^{-1}$). Only briefly during peak flows did NO_3^- concentrations increase up to $100 \mu\text{eq L}^{-1}$. Averaged over 10 years (1984–1994), the export of N at the catchment outlet was $0.05 \text{ kg N ha}^{-1} \text{ yr}^{-1}$. This indicates a very tight N cycling allowing for an increase of N availability in these undisturbed forest ecosystems.

Keywords: bulk deposition, throughfall, dry deposition, soil water, nitrate, ammonium, Spain, Montseny, La Castanya

Introduction

Aggrading forests receiving moderate N inputs retain most of the deposited N within the ecosystem, with only a minor fraction being lost to streams (Vitousek, 1977). However, in recent years the deposition of nitrogen has increased in many regions of Europe and North America because of enhanced agricultural and industrial activities (Galloway 1995; Vitousek *et al.*, 1997). The combined effect of high N inputs and high nitrification (which depends on site specific characteristics, such as soil depth, soil temperature, soil moisture, and high pH) may lead to nitrogen saturation in terrestrial ecosystems (Aber *et al.*, 1989, 1998; Agren and Bosatta, 1988; Nihlgard, 1985). Nitrogen saturation has been found for forests receiving large loads of N, in the range $20\text{--}100 \text{ kg N ha}^{-1} \text{ yr}^{-1}$. In particular, these elevated inputs have been described for the Netherlands (Van Breemen *et al.*, 1987; Wyears *et al.*, 1992), Great Britain (Emmet *et al.*, 1993), Scandinavia (Andersen *et al.*, 1993) and the United States (Aber *et al.*, 1989, 1998; Bytnerowicz and Fenn, 1996; Fenn *et al.*, 1996). A symptom of N

saturation in forests is the increase of NO_3^- leached to the streams.

In Spain, increasing N emissions have occurred because of recently increased fossil fuel consumption and enhanced livestock production: from 1983 to 1998, NO_x emissions increased by 17% and NH_3 emissions increased by 31% (EMEP, 2000). Nitrate concentrations in rainwater also increased significantly during this period although the trend was statistically non-significant when considering NO_3^- -N deposition due to lower rainfall in later years (Avila and Rodà, 2002).

In this paper, the results obtained since 1983 for a study of a holm oak forested catchment of La Castanya, Montseny Mountains (Catalonia) are used to:

- (1) describe the inorganic N deposition rates (wet and dry).
- (2) describe the changes occurring in NO_3^- and NH_4^+ concentrations as water moves through the ecosystem.
- (3) discuss the effects of the atmospheric inputs on the N cycle in these holm oak forests.

Study site

The Montseny Mountains (41° 46' N, 2° 21' E; 40 km NNE from Barcelona) cover an area of about 400 km², and reach their highest altitude at 1707 m.a.s.l. They are mostly forested (holm oak, pines and beech), with heathlands and grasslands in the upper parts and arable crops in the lower reaches. The bedrock consists mostly of metamorphic phyllites and schists. The climate is montane Mediterranean. Mean annual precipitation was 901 mm yr⁻¹ and ranged between 640 and 1660 mm yr⁻¹ in the period August 1983–July 2000 (hydrological years being defined from 1 August to 31 July). Mean annual temperature ranged from 10 to 14°C.

The sampling site, named La Castanya, was at La Castanya valley (altitude 700 m.a.s.l.), amidst extensive holm oak forests in the Montseny massif. The main topographic characteristics and the stand structure of this forest are summarised in Table 1. A station for air pollution monitoring from the Departament de Medi Ambient de la Generalitat de Catalunya was used to compare bulk and wet collecting methods. This station (named PA) is at the foot of the Montseny Mountains, at a linear distance of 6 km from La Castanya in the surroundings of the town of Santa Maria de Palautodera (200 m.a.s.l., 5000 inhabitants).

Table 1. Site characteristics (mean ± SE, n=4) of La Castanya holm oak at Montseny. Stand and tree data are for stems having a diameter at breast height >5 cm.

Variable	
Altitude (m)	731 ±4.3
Slope (°)	37.5 ±0.6
Stand density (stems ha ⁻¹)	2127 ±16.2
Basal area (m ² ha ⁻¹)	26.4 ±2.0
Tree height (m)	5.9 ±0.3
Canopy depth (m)	3.1 ±0.2
Tree cover (%)	128 ±0.2

Material and methods

At La Castanya, bulk deposition was sampled in the holm oak forest in a clearing of sufficient diameter to avoid interference by surrounding trees. Bulk deposition was collected weekly from four (until 1993) and two (from 1993 to 2000) replicate bulk collectors consisting of a polyethylene funnel connected by tygon tubing to a 10-l polyethylene bottle. At PA, precipitation was collected

weekly with two bulk collectors and one wet-only collector (MCV trademark).

Throughfall was sampled weekly in four replicate plots inside the holm oak forest of La Castanya. Throughfall plots were at a distance of 100–200 m from the bulk deposition clearing. The plots were circular, of 7-m radius. Each plot contained eight throughfall collectors, consisting of a 10-cm diameter funnel connected to a 2-l polyethylene bottle. The sampling bottles, both for throughfall and bulk deposition, were kept in the dark to minimise biological transformations of the sample. Stemflow was also sampled using rings around the tree stems, with ten trees sampled in one of the plots (Rodrigo, 1998). As stemflow N fluxes were very low compared to throughfall, these results are not discussed here.

Dry deposition rates were calculated from an experiment of parallel branch- and plate-washings during rainless periods. Eight extruding branches were chosen and six metacrylate plates of 20 × 20 cm and 0.2 cm thickness were placed at the top of dominant trees near the throughfall plots. On five rain-free periods both these branches and plates were washed *in situ* with distilled deionised water after exposure times ranging between 72 to 187 h since the last rainfall, and dry deposition fluxes per unit leaf or plate area (µg cm⁻² h⁻¹) for NO₃⁻-N and NH₄⁺-N were calculated. The resulting values were then extrapolated to estimate dry deposition fluxes to the canopy at annual scale, assuming a leaf area index (LAI) of five for the La Castanya holm oak site (Sabaté *et al.*, 1999).

At La Castanya, the movement of dissolved NO₃⁻ and NH₄⁺ through the soil profile was studied in a 3-m long trench, with two zero tension through-flow collectors installed at different depths to collect: (1) the forest floor solution (0–5 cm-depth), and (2) the deep flow at the bedrock contact (160 cm-depth). In addition, the soil solution was sampled with ceramic-cup tension lysimeters evacuated at –65 kPa. Eight lysimeters were used, four at each depth (20 cm and 40 cm depth; Avila *et al.*, 1995).

Streamwater samples were collected manually weekly, but more frequent sampling was programmed during stormflows using an automatic collector (Manning S-4040). Sampling frequency was at fixed time intervals, with the frequency adjusted to cover the whole hydrograph.

Study periods differed among sites and among the processes studied. The longest record was for rainwater chemistry, for which 17 years of data were available (August 1983 to July 2000). Throughfall data were collected from 6 June 1995 to 25 June 1996. The washing experiment of branches and plates was conducted on May–June 1996. Streamwater chemistry and discharge measurements from the stream (named TM9) that flows through a 5.9 ha

catchment within the holm oak forest extended from August 1983 to July 1994. Storm events were sampled from August 1983 until July 1988. Lysimeters were emptied weekly from November 1983 to June 1986. Through-flow solutions from the trench were collected weekly from May 1985 to May 1988.

All water samples (bulk deposition, wet deposition, throughfall, leaf and plate washes, soil solution and streamwater) were taken to the laboratory on the day of sampling, where pH, conductivity and alkalinity were analysed within 24–48h. Samples were then filtered with 0.45- μm Millipore filters, and the filtrate was deep-frozen for later analyses of major anions (SO_4^{2-} , NO_3^- and Cl^-) and cations (Na^+ , K^+ , Ca^{2+} , Mg^{2+} , NH_4^+). Nitrate was analysed by ion chromatography and NH_4^+ by flow injection analysis until 1996 and by ion chromatography thereafter. All ion analyses included synthetic samples of known concentrations within the runs to check for replicability and accuracy. Data were further screened for analytical quality by the ratios sum of cations / sum of anions (on an equivalent basis) and calculated conductivity / measured conductivity. Since 1991, the laboratory participated in a joint laboratory intercomparison of analytical results coordinated by ISPRA (AQUACON Project, Mosello *et al.*, 1998), with very satisfactory results.

Results and discussion

WET AND DRY DEPOSITION

Mean rainwater composition at La Castanya was not acidic: the volume-weighted mean (VWM) alkalinity for the period 1983–2000 was 14.3 $\mu\text{eq l}^{-1}$. Calcium (VWM concentration = 52.5 $\mu\text{eq l}^{-1}$) was the main cation counterpart to acidic anions (VWM concentration for SO_4^{2-} and NO_3^- = 41.2 and

21.6 $\mu\text{eq l}^{-1}$, respectively; Avila and Rodà, 2002). In northeastern Spain, calcium carbonate can be incorporated into the atmosphere from local edaphic, urban and industrial sources, but wet deposition of African dust is very important, providing on average 46% of annual Ca deposition (Avila, 1996).

Annual mean concentrations of inorganic N in rainwater were moderate, those of NH_4^+ being slightly higher than NO_3^- (Table 2). As expected, NH_4^+ concentrations were higher at PA, the lowland site closer to cultivated fields and farms with animals (Table 2). At PA, the comparison of bulk and wet collecting methods showed very small differences statistically (paired *t*-test) non-significant for all major elements, except for NH_4^+ which was 5% higher in wet collectors (Table 2).

Averaging over the period 1983–2000, the N inputs in bulk deposition at La Castanya were 2.9 $\text{kg ha}^{-1} \text{yr}^{-1}$ for NH_4^+ -N and 2.7 $\text{kg ha}^{-1} \text{yr}^{-1}$ for NO_3^- -N resulting in a wet input of 5.6 $\text{kg N ha}^{-1} \text{yr}^{-1}$. Forest canopies are very effective surfaces for dry deposition, but dry deposition fluxes are difficult to measure and dry inputs are not often taken into account. For N, various studies in north-American deciduous forests have shown the predominant role of deposition of HNO_3 vapour and coarse nitrate aerosols which produce a dry input flux that can be even higher than that of wet N deposition (Lovett and Lindberg, 1986, 1993).

Dry deposition of NO_3^- -N and NH_4^+ -N at La Castanya was higher on branches than on metacrylate plates, probably because the foliage is more efficient in capturing airborne substances than horizontal plates due to differences in both the surface area of catch and the degree of air turbulence. Therefore, it was considered that branch washes gave a better estimate of dry-deposited N than metacrylate plates (Rodrigo and Avila, 2002). The fluxes estimated from washing branches amounted to 9.2 $\text{kg ha}^{-1} \text{yr}^{-1}$, about 60% higher

Table 2. Volume-weighted mean concentrations in bulk or wet-only precipitation and deposition of inorganic N at Montseny at two study sites: La Castanya (LC) and Santa Maria de Palautordera (PA).

Site	Length of study period (yr)	Precipitation (mm yr^{-1})	Concentration ($\mu\text{eq l}^{-1}$)		Deposition ($\text{kg N ha}^{-1}\text{yr}^{-1}$)	
			NH_4^+	NO_3^-	NH_4^+ -N	NO_3^- -N
LC, bulk	17 ^a	901	22.8	21.6	2.88	2.73
PA, bulk	4 ^b	740	44.7	32.4	4.63	3.36
PA, wet	4 ^b	705	47.0	31.7	4.64	3.13

^a From January 1983 to September 2000.

^b From June 1995 to June 1999.

than the estimated bulk deposition flux. The total deposition flux of inorganic N was estimated as 14.8 kg ha⁻¹ yr⁻¹. The same methodology was used at a nearby holm oak site Riera de Sant Pere (7 km linear distance from La Castanya and 1.5 km from PA, 530 m.a.s.l.) with a higher exposure to traffic and industrial pollution. At this site, total N deposition was estimated to be 21.9 kg ha⁻¹ yr⁻¹, with 67% of the total N amount being as dry deposition (Rodà *et al.*, 2002). For the forests within the Integrated Forest Study in eastern United States, N dry deposition (HNO₃ and NO₃⁻ and NH₄⁺ in fine and coarse particles) ranged between 39% to 59% of total deposition, and total N deposition varied between 4.7 and 27 kg ha⁻¹ yr⁻¹ (Lovett and Lindberg, 1993).

At sites with moderate N inputs, canopies intercept and retain much of the incident N input fluxes because of stomatal or cuticular uptake or retention by phyllosphere organisms (Parker, 1983; Pucket, 1990; Lovett and Lindberg, 1993). That was the case at La Castanya: net throughfall fluxes for NO₃⁻-N and NH₄⁺-N were negative indicating net retention at the canopy level despite the non-negligible total deposition fluxes. On the other hand, at the more pollutant-exposed site of RP, net throughfall fluxes shifted to positive, although very small, values (0.3 and 1.2 kg ha⁻¹ yr⁻¹ for NO₃⁻ and NH₄⁺ respectively).

Nitrogen leaching

The changes in the chemistry as water crosses the ecosystem are shown in Fig. 1. To highlight the changes for N compounds, the changes in their concentrations are displayed in Table 3. At La Castanya, concentrations of NO₃⁻ and NH₄⁺ in throughfall were similar to those in rainwater, throughfall fluxes being lower (and hence net fluxes negative) due to a 20% water interception at the canopy

Table 3. Water chemistry profile for NH₄⁺ and NO₃⁻ at La Castanya (Montseny)

Variable	Water flux mm yr ⁻¹	NH ₄ ⁺ µeq l ⁻¹	NO ₃ ⁻ µeq l ⁻¹
Bulk Precipitation	891 ^a	22.8	21.6
Throughfall	565 ^b	27.1	22.4
Forest Floor Throughflow	—	7.0	249
Soil Solution	—	<2	4.0
Deep Flow	—	<2	40.0
Streamwater	311 ^a	<2	1.3

^a From August 1984 to July 1996

^b From December 1978 to December 1980

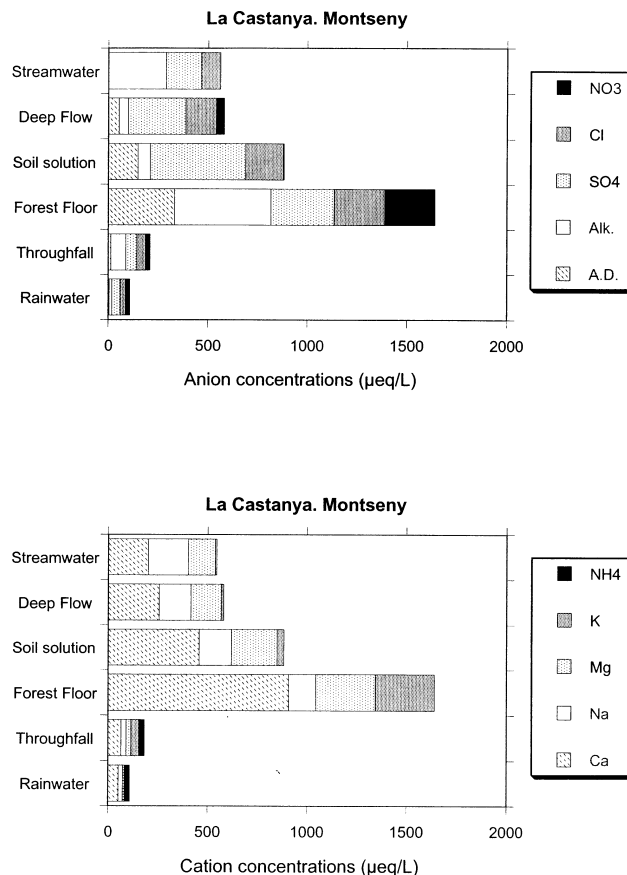


Fig 1. Water chemistry profiles for the holm oak forest at La Castanya (Montseny). a) anions, b) cations. A.D = anion deficit (sum of cations - sum of anions).

(Rodrigo, 1998). At the forest floor, mean volume-weighted NO₃⁻ was 250 µeq l⁻¹, while NH₄⁺ concentration was only 7 µeq l⁻¹. Despite the low NH₄⁺ concentration, H-layer throughflow was the only sampled soil water type that consistently showed detectable NH₄⁺.

Changes occurring along the profile across the ecosystem indicate a decreasing trend in inorganic N concentrations, although the pattern is different for the two forms of N. Ammonium disappeared when water entered the soil at the forest floor level never to be detected again, either in the soil solution or in the streamwater. For NO₃⁻, maximum concentrations were found at the forest floor level, decreased to 4 µeq l⁻¹ in soil solution and increased again in the deep subsurface flow to 40 µeq l⁻¹ (Fig.1). The high NO₃⁻ concentrations in the H horizon correspond with the site in the soil profile of highest nitrification rates (Serrasolses *et*

al., 1999). In the soil solution, NO_3^- and NH_4^+ concentrations decreased, probably due to plant uptake and microbial immobilization. Deep through-flow presented intermediate NO_3^- concentrations between the water collected below the forest floor and that collected from tension lysimeters at 20 and 40 cm depth. This indicates that free flowing water at the contact with the bedrock is a mixture of macropore water flowing from the forest floor and pre-event displaced soil solution from the mineral soil (Avila *et al.*, 1995). The behaviour of other ions and, in particular, that of ionic ratios to Cl^- taken as a tracer scarcely affected by exchange processes or biological uptake, gives support to this interpretation (Melià *et al.*, 1999).

The inorganic N concentrations in streamwater were very low: as mentioned for NH_4^+ , they were always under the analytical detection limit. For NO_3^- , the same applied for baseflow samples. However, NO_3^- concentrations increased during high flows. Nitrate concentration maxima usually occurred in advance of peak flow; therefore, NO_3^- presented usually clockwise hysteretic loops when plotted against discharge (Avila, 1988). Such behaviour can result either

from an initial flushing of a limited ion supply, or from changing the proportional contribution of different flowpaths during the events.

A distinction has to be made between two types of hydrograph responses: those corresponding to dry antecedent conditions and those to wet antecedent conditions. Hydrograph characteristics and the variation of NO_3^- concentrations were very different between both types. An example of an event under dry antecedent conditions is depicted in Fig. 2 and with wet antecedent conditions in Fig. 3. For dry antecedent conditions, the peak flow was reached quickly and the recession returned to pre-storm levels in a scale of hours (Fig. 2). This implied small runoff, the percentage of precipitation that resulted in quick flow ranging between 0.5% and 8.7% for the different storms (Table 4). For storm flow in humid antecedent conditions, the time to reach the peak flow increased and high flow was maintained for days (Fig 3): the storm runoff was higher, and percentages of quick flow with respect to precipitation ranged between 18% and 58% for the different storm events sampled (Table 4).

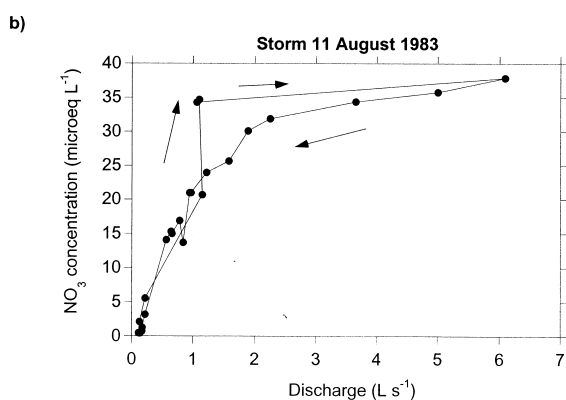
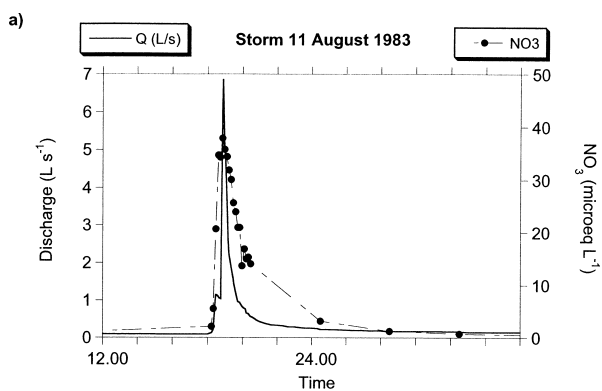


Fig 2. Dry antecedent conditions event in 11 August 1983. a) Hydrograph and variation of NO_3^- concentrations in TM9 streamwaters, and b) NO_3^- hysteresis with discharge.

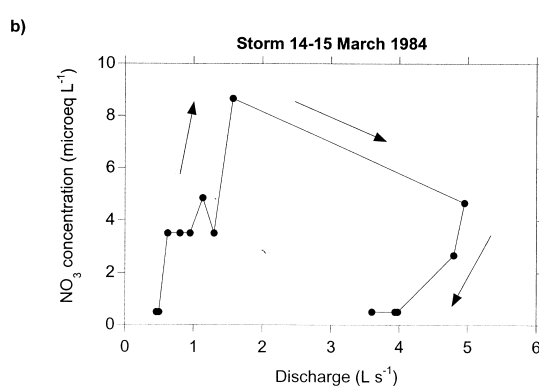
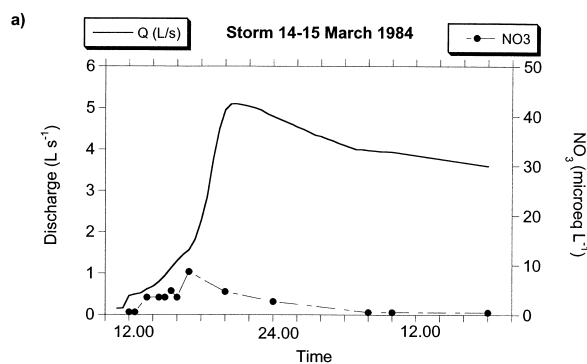


Fig 3. Wet antecedent conditions event in 14-15 March 1984. a) Hydrograph and variation of NO_3^- concentrations in TM9 streamwaters, and b) NO_3^- hysteresis with discharge.

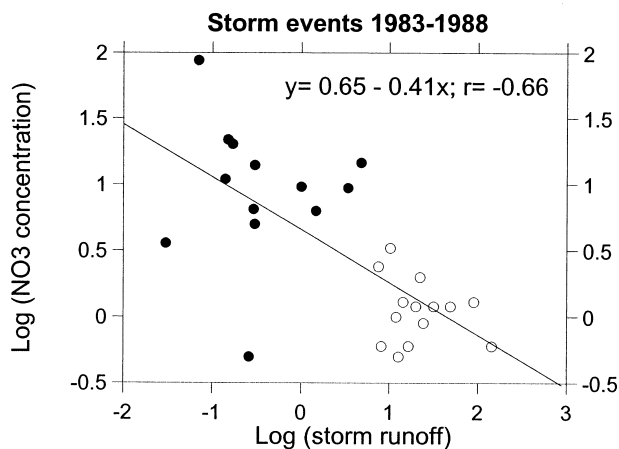


Fig 4. Log-log relationship between storm flow volume-weighted mean NO_3^- concentration and storm runoff. Storms are distinguished according to the humidity antecedent conditions at the TM9 catchment (La Castanya, Montseny). Filled dots correspond to storms with dry antecedent conditions; blank dots correspond to storms with wet antecedent conditions.

Table 4. Hydrograph characteristics for stormflows separated according to humidity antecedent conditions at the TM9 catchment in La Castanya valley (Montseny). Arithmetic means (in brackets, standard error) are given for the storms recorded from August 1983 to July 1988.

Variable	Dry antecedent conditions n=13	Wet antecedent conditions n=14
Peakflow ($\text{l s}^{-1} \text{ km}^{-1}$)	54.2 (17.0)	137.8 (42.2)
Runoff (mm)	0.95 (0.41)	30.7 (9.52)
Runoff/rainfall	0.02 (0.006)	0.44 (0.06)
Time to peakflow (hour)	4.96 (2.18)	53.5 (113)

Solute behaviour mirrored these differences in hydrological response. The changes in NO_3^- concentrations reflected the various water pathways through the ecosystem compartments and the different water residence times. In stormflow associated with dry antecedent conditions, NO_3^- reached brief peaks of high concentration at or near the peak of flow (Fig. 2), probably because of the contribution of Hortonian partial overland flow with the chemical characteristics of the H layer forest floor (Table 3). In wet antecedent conditions, a dominant contribution of subsurface flow (with chemical characteristics of soil water, or deep flow, Table 3) would attest to the lower concentrations during these storm events (Fig. 3). Corresponding with these

differences in water pathways and solute content, when representing, for each storm event, the volume-weighted mean NO_3^- concentration against the storm runoff, an inverse log-log relationship appeared (Fig. 4). Storms occurring with dry antecedent conditions presented low storm runoff and high volume-weighted mean NO_3^- concentrations. Storms in wet antecedent conditions produced higher drainage but, because of subsurface major pathways, they had very low volume-weighted mean NO_3^- concentrations.

At baseflow, with groundwater as the major component of streamflow, NO_3^- concentrations were very low throughout the year. This contrasted with the seasonality reported for other Mediterranean streams in Catalonian forests (Butturini and Sabater, this issue) and for northern deciduous forest ecosystems, with higher NO_3^- levels during the inactive winter season (Likens and Bormann, 1995). The fact that at La Castanya NO_3^- showed high concentrations ($>50 \mu\text{eq l}^{-1}$) only during very short times at peak flows of dry antecedent conditions testifies to the tight N cycle in this holm oak undisturbed ecosystem. Averaged over 10 years (1984–1994), the export of N at the catchment outlet was only $0.05 \text{ kg N ha}^{-1} \text{ yr}^{-1}$, a low figure when compared to the export at Riera Major ($0.66 \text{ kg N ha}^{-1} \text{ yr}^{-1}$, Butturini and Sabater, 2002) a stream draining similar holm oak forests but with N-fixing *Alnus glutinosa* on the streamsides.

Biological demand for N is a major process for the tightly closed N cycle at La Castanya forest. This was demonstrated by a root-trenching experiment where water and nutrient uptake by tree roots was prevented by building trenched plots. Six months after trenching, NO_3^- in the soil solution increased to concentrations between 2000 and $2500 \mu\text{eq l}^{-1}$ (Bonilla and Rodà, 1990), while at the control plots, NO_3^- concentration remained around $10 \mu\text{eq l}^{-1}$.

In a comparative study of N budgets for 65 forested catchments and plots in North and Central Europe submitted to a wide range of total N deposition, Dise and Wright (1995) proposed a N deposition threshold of $10 \text{ kg N ha}^{-1} \text{ yr}^{-1}$ in throughfall fluxes; thereafter significant NO_3^- leaching would start to occur from forests. The present study at the holm oak forest of La Castanya indicates that, although the estimated total N input is not negligible, canopy processes and biological uptake retain most of the input within the ecosystem.

Conclusion

Biological demand is a major controlling process for a tight N cycle at the holm oak forest of La Castanya (Montseny). The atmospherically deposited N is almost completely retained within the ecosystem, possibly because holm oak

forests at Montseny were harvested until around 1950 and are now in an aggrading phase. In contrast, high N deposition poses a problem of excess N in soils and streamwaters at more mature forests in North America or central and North Europe subjected over many years to high N loads.

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References

- Aber, J.D., Nadelhoffer, K.J., Steudler, P. and Melillo, J.M., 1989. Nitrogen saturation in northern forest ecosystems. *BioSci.*, **39**, 378–386.
- Aber, J.D., McDowell, W., Nadelhoffer, K., Magill, A., Berntson, G., Kamakea, M., McNulty, S., Currie, W., Rustad, L. and Fernandez, I., 1998. Nitrogen saturation in temperate forest ecosystems. *BioSci.*, **48**, 921–934.
- Agren, G.I. and Bosatta, E., 1988. Nitrogen saturation of terrestrial ecosystems. *Environ. Pollut.*, **54**, 185–198.
- Andersen, H.V., Hovmand, M.F., Hummelshoj, P. and Jensen, N.O., 1993. Measurement of ammonia flux to a spruce stand in Denmark. *Atmos. Environ.*, **27**, 189–202.
- Avila, A., 1988. Balanç d'aigua i nutrients en una conca d'alzinar del Montseny. *Estudis i Monografies*, 13. Diputació de Barcelona. Barcelona. 219 pp.
- Avila, A., 1996. Time trends in the precipitation chemistry at a montane site in north-eastern Spain for the period 1983-1994. *Atmos. Environ.*, **30**, 1363–1373.
- Avila, A., Bonilla, D., Rodà, F., Piñol, J. and Neal, C., 1995. Soilwater chemistry in a holm oak (*Quercus ilex*) forest: inferences on biogeochemical processes for a montane-Mediterranean area. *J. Hydrol.*, **166**, 15–35.
- Avila, A. and Rodà F., 2002. Assessing decadal changes in rainwater alkalinity at a rural Mediterranean site in the Montseny Mountains (NE Spain). *Atmospheric Environment*, **36**, 2881–2890.
- Bonilla, D. and Rodà, F., 1990. Nitrogen cycling responses to disturbance: trenching experiments in an evergreen oak forest. In: *Nutrient cycling in terrestrial ecosystems*. A.F. Harrison, P. Ineson and O.W. Heal (Eds.) Elsevier, London, 179–189.
- Butturini, A. and Sabater, F. 2002. Nitrogen concentrations in a small Mediterranean stream: I. Nitrate. II. Ammonium. *Hydrol. Earth Syst. Sci.*, 539–550.
- Bytnerowicz, A. and Fenn, M.E., 1996. Nitrogen deposition in California forests: a review. *Environ. Pollut.*, **92**, 127–146.
- Dise, N. B. and Wright, R. F., 1995. Nitrogen leaching from European forests in relation to nitrogen deposition. *Forest Ecol. Manage.*, **71**, 153–161.
- Emmett, B.A., Reynolds, B., Stevens, P.A., Norris, D.A., Hughes, S., Görres, J. and Lubrecht, I., 1993. Nitrate leaching from afforested Welsh catchment: interaction between stand age and nitrogen deposition. *Ambio*, **22**, 386–394.
- EMEP 2000 from <http://www.emep.int/>
- Fenn, M.E., Poth, M.A. and Johnson, D.W., 1996. Evidence for nitrogen saturation in the San Bernardino mountains in southern California. *Forest Ecol. Manage.*, **82**, 211–230.
- Galloway, J.N., 1995. Acid deposition: perspectives in time and space. *Water Air Soil Pollut.*, **85**, 15–24.
- Likens, G.E. and Bormann, F.H., 1995. *Biogeochemistry of a forested ecosystem*. 2nd Edition. Springer Verlag. New York. 159 pp.
- Lovett, G.M. and Lindberg, S.E., 1986. Dry deposition of nitrate to a deciduous forest. *Biogeochem.*, **2**, 137–148.
- Lovett, G.M. and Lindberg, S.E., 1993. Atmospheric deposition and canopy interactions of nitrogen in forests. *Can. J. Forest Res.*, **23**, 1603–1616.
- Melià, N., Bellot, J. and Vallejo, V.R., 1999. Soil water chemistry. In: *Ecology of Mediterranean Evergreen Oak Forests*. F. Rodà, J. Retana, C. Gracia, and J. Bellot (Eds.). Springer, Berlin, 237–251.
- Mosello, R., Bianchi, M., Geiss, H., Marchetto, A., Morselli, L., Muntau, H., Tartari, G.A., Serrini, G. and Serrini Lanza, G., 1998. *Italian Network for the Chemistry of Atmospheric Deposition. Intercomparison 1/98*. Consiglio Nazionale delle Ricerche. Istituto Italiano di Idrobiologia. Verbania Pallanza. 81 pp.
- Nihlgard, B., 1985. The ammonium hypothesis- An additional explanation for the forest dieback in Europe. *Ambio*, **14**, 2–8.
- Parker, G.G., 1983. Throughfall and stemflow in the forest nutrient cycle. *Adv. Ecol. Res.*, **13**, 57–133.
- Pucket, L.J. 1990. Estimates of ion sources in deciduous and coniferous throughfall. *Atmos. Environ.*, **24**, 545–555.
- Rodà, F., Avila, A. and Rodrigo, A., in press. Nitrogen deposition in Mediterranean forests. *Environ. Pollut.*
- Rodrigo, A., 1998. Deposició atmosfèrica en dos alzinars (*Quercus ilex*) del Montseny sotmesos a una exposició contrastada de contaminants de l'àrea barcelonina i vallesana. Ph Dissertation. Universitat Autònoma de Barcelona. 330 pp.
- Rodrigo, A. and Avila, A., 2002. Dry depositon to the forest canopy and surrogate surfaces in two Mediterranean holm oak forests in Montseny (N E Spain). *Water Air Soil Pollut.*, **136**, 269–288.
- Sabaté, S., Sala, A. and Gracia, C.A., 1999. Leaf traits and canopy organization. In: *Ecology of Mediterranean Evergreen Oak Forests*. F. Rodà, J. Retana, C. Gracia, and J. Bellot (Eds.). Springer, Berlin, 121–133.
- Serrasolses, I., Diego, V. and Bonilla, D., 1999. Soil nitrogen dynamics. In: *Ecology of Mediterranean Evergreen Oak Forests*. F. Rodà, J. Retana, C. Gracia, and J. Bellot (Eds.). Springer, Berlin, 223–235.
- Van Breemen, N., Mulder, J. and Van Grinsven, J.J.M., 1987. Impacts of acid atmospheric deposition on woodland soils in the Netherlands. II. Nitrogen transformations. *Soil Sci. Amer. J.*, **51**, 1634–1640.
- Vitousek, P.M., 1977. The regulation of element concentration in mountain streams in the northeastern United States. *Ecol. Monogr.*, **47**, 65–87.
- Vitousek, P. M., Aber, J.D., Howarth, R.W., Likens, G.E., Matson, P.A., Schindler, D.W., Schlesinger, W.H. and Tilman, D.G., 1997. Human alteration of the global nitrogen cycle: sources and consequences. *Ecol. Appl.*, **7**, 737–750.
- Wyears, G.P., Vermuelen, A.T. and Slanina, J., 1992. Measurement of dry deposition of ammonia on a forest. *Environ. Pollut.*, **75**, 25–28.

