

1 Long-term nitrogen addition decreases carbon leaching in a  
2 **nitrogen-rich forest ecosystem**

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4 Xiankai Lu<sup>1</sup>, Frank S. Gilliam<sup>2</sup>, Guirui Yu<sup>3</sup>, Linghao Li<sup>4</sup>, Qinggong Mao<sup>1</sup>, Hao Chen<sup>1</sup>,  
5 Jiangming Mo<sup>1\*</sup>

6

7 <sup>1</sup>Key Laboratory of Vegetation Restoration and Management of Degraded Ecosystems,  
8 South China Botanical Garden, Chinese Academy of Sciences, Guangzhou 510650,  
9 China;

10 <sup>2</sup>Department of Biological Sciences, Marshall University, Huntington, West Virginia,  
11 25755-2510, USA;

12 <sup>3</sup>Institute of Geographical Sciences and Natural Resources Research, Chinese  
13 Academy of Sciences, Beijing 100101, China;

14 <sup>4</sup>State Key Laboratory of Vegetation Environmental Change, Institute of Botany,  
15 Chinese Academy of Sciences, Xiangshan, Beijing, 100093, China.

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17 **\*Corresponding author:** Jiangming Mo; Tel: +86758-2621187; Fax:  
18 +86758-2623242; E-mail: [mojm@scib.ac.cn](mailto:mojm@scib.ac.cn)

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26 **Abstract**

27 Dissolved organic carbon (DOC) plays a critical role in the carbon (C) cycle of forest  
28 soils, and has been recently connected with global increases in nitrogen (N) deposition.

29 Most studies on effects of elevated N deposition on DOC have been carried out in  
30 N-limited temperate regions, with far fewer data available from N-rich ecosystems,

31 especially in the context of chronically elevated N deposition. Furthermore,

32 mechanisms for excess N-induced changes of DOC dynamics have been suggested to

33 be different between the two kinds of ecosystems, because of the different ecosystem

34 N status. The purpose of this study was to experimentally examine how long-term N

35 addition affects DOC dynamics below the primary rooting zones (the upper 20 cm

36 soils) in typically N-rich lowland tropical forests. We have a primary assumption that

37 long-term continuous N addition minimally affects DOC concentrations and effluxes

38 in N-rich tropical forests. Experimental N addition was administered at the following

39 levels: 0, 50, 100 and 150 kg N ha<sup>-1</sup> yr<sup>-1</sup>, respectively. Results showed that seven years

40 of N addition significantly decreased DOC concentrations in soil solution, and

41 chemo-physical controls (solution acidity change and soil sorption) rather than

42 biological controls may mainly account for the decreases, in contrast to other forests.

43 We further found that N addition greatly decreased annual DOC effluxes from the

44 primary rooting zone and increased water-extractable DOC in soils. Our results

45 suggest that long-term N deposition could increase soil C sequestration in the upper

46 soils by decreasing DOC efflux from that layer in N-rich ecosystems, a novel

47 mechanism for continued accumulation of soil C in old-growth forests.

48

49 **Key words:** Nitrogen deposition; Nitrogen saturation; N-rich; DOC efflux; Carbon

50 cycle; Carbon sequestration; Soil solution; Tropical forest; Acidification

51 **1 Introduction**

52 Terrestrial ecosystem carbon (C) cycling and storage are a global concern in the  
53 context of increasing atmospheric deposition of N in the biosphere, especially in  
54 recent decades (Schindler and Bayley, 1993; Nadelhoffer *et al.*, 1999; Galloway *et al.*,  
55 2004; LeBauer and Treseder, 2008; Hyvönen *et al.*, 2008). Although it is generally  
56 known that N deposition can significantly alter terrestrial ecosystem C cycle, most  
57 studies on the responses of ecosystem C cycling to N enrichment focused on net  
58 primary productivity (NPP), net ecosystem productivity, net ecosystem CO<sub>2</sub> exchange,  
59 and labile pools of C (LeBauer and Treseder, 2008; Hyvönen *et al.*, 2008; de Vries *et*  
60 *al.*, 2009; Liu and Greaver, 2010; Thomas *et al.*, 2010). In contrast, effects of N on  
61 dissolved organic C (DOC) have received less attention, likely because these effluxes  
62 are small relative to the C fluxes associated with primary productivity or heterotrophic  
63 respiration in terrestrial systems (Kalbitz *et al.*, 2000; Neff and Asner, 2001). However,  
64 the dynamics of DOC are receiving increased attention, considering their essential  
65 links in the bio-, hydro- and pedosphere (Kalbitz *et al.*, 2000) and their central  
66 importance in soil-forming processes and carbon sequestration via DOC mobilization  
67 and transport for both temperate and tropical soils (McDowell, 1998; Monteith *et al.*,  
68 2007; Cusack *et al.*, 2010; Liu & Greaver, 2010; Kindler *et al.*, 2011).

69 Forest soils play a key role in the global C cycle (Lal, 2005). To explore the  
70 importance of DOC effluxes under elevated N deposition in forest ecosystems,  
71 ecologists have conducted such studies by the methods of simulating N deposition or  
72 using natural N deposition gradients (Evans *et al.*, 2008; Sleutel *et al.*, 2009). Until  
73 now, these studies are limited to DOC dynamics (e.g., concentrations or effluxes), and  
74 have not been linked to the possible C sequestration induced by N deposition in  
75 ecosystems. Meanwhile, these studies are mostly focused in temperate regions,

76 especially in North American and Europe, where ecosystems commonly belong to  
77 glaciated landscapes and are N-limited under natural conditions (e.g. Vitousek and  
78 Howarth, 1991; Aber et al., 1998, 2003; Magill et al., 2004). These studies often find  
79 that DOC concentration in soil solution increases with elevated N deposition (Yano et  
80 al., 2000; McDowell et al., 2004; Pregitzer et al., 2004; Adams et al., 2005; Findlay,  
81 2005; Sleutel et al., 2009; Rappe-George et al., 2012).

82 Tropical forest ecosystems, which store approximately 13% of global soil C,  
83 contribute greatly to the global C cycle; thus, even relatively small fluctuations in C  
84 cycling can have global consequences (Post et al., 1982; Phillips et al., 1998; Findlay,  
85 2005; Townsend et al., 2011). In contrast to their temperate and boreal counterparts,  
86 many lowland tropical forests are typically N-rich ecosystems as compared with P  
87 availability, with high soil N availability, rapid rates of N cycling, and the lack of N  
88 limitation to NPP (Vitousek & Sanford, 1986; Matson et al., 1999; Hedin et al. 2009;  
89 Wright et al., 2011; Brookshire et al., 2012). However, our understanding of how N  
90 additions control DOC dynamics in these N-rich ecosystems remains far from  
91 complete.

92 The purpose of this study was to examine the effects of how long-term (7 yr)  
93 experimental addition of N affects DOC dynamics in the N-rich tropical forests. In  
94 2002, we established long-term N deposition research plots in typical N-rich lowland  
95 tropical mature forests of Southern China (Mo et al., 2006. 2008; Fang et al., 2009; Lu  
96 et al., 2010), where atmospheric N deposition rates are commonly  $> 19 \text{ kg N ha}^{-1} \text{ yr}^{-1}$ ,  
97 and are expected to increase greatly in the future due to the rapid development of  
98 agricultural and industrial activities (Zhou and Yan, 2001; Galloway et al., 2004; Lü  
99 and Tian, 2007; Liu et al., 2011). Because soil solution chemistry can be considered as  
100 a sensitive indicator of biogeochemical processes within forest stands, responding

101 quickly to disturbances or stresses such as excess N (e.g. McDowell et al., 2004;  
102 Pregitzer et al., 2004; Michel et al., 2006; Gilliam and Adams, 2011), we mainly  
103 focused our study on the response of soil solution chemistry to N addition. Earlier  
104 measurements in these forests have indicated no changes in DOC dynamics in  
105 response to short-term (1 to 2 years) N deposition treatment (Fang et al., 2009). In the  
106 present study, we expected to find that long-term continuous N addition to N-rich  
107 tropical forests has minimal effect on DOC concentrations and effluxes, because  
108 highly weathered tropical soils commonly have high levels of N availability and rapid  
109 N cycling (Martinelli et al. 1999; Vitousek and Sanford, 1986; Fang et al., 2009). At  
110 the same time, we assumed that mechanisms for N-addition induced changes of DOC  
111 dynamics may be different from those of N-limited temperate forests, because of the  
112 different N status.

113

## 114 **2 Materials and methods**

### 115 **2.1 Study Site**

116 We carried out our work in the Dinghushan Biosphere Reserve (DBR). This site is  
117 part of the UNESCO/MAB network and is within the Guangdong Province of  
118 southern China (112°10' E, 23°10' N). The DBR extends approximately 1,200 ha  
119 within the subtropical/tropical moist forest life zone. It was established in 1950 for the  
120 protection of remnant of undisturbed monsoon evergreen broadleaf forests in the  
121 lower subtropics, being the first National Natural Reserve in China in 1956. The  
122 monsoon climate of this site averages 1927 mm precipitation per year with  
123 approximately 75% occurring between March and August, and 6% between December  
124 and February (Huang & Fan, 1982). Relative humidity averages 80% throughout the  
125 year. Mean annual temperature is 21.0 C, ranging from mean coldest in January (12.6

126 C) and hottest in July (28.0 C). Currently, the region surrounding DBR experiences  
127 high rates of atmospheric N deposition (21-38 kg N ha<sup>-1</sup> yr<sup>-1</sup> as inorganic N in bulk  
128 precipitation) (Huang et al., 1994; Zhou and Yan, 2001; Fang et al., 2008). In  
129 2004—2005 wet N deposition averaged ~33 kg N ha<sup>-1</sup> yr<sup>-1</sup> (Fang et al. 2008).

130 We established the research site at DBR in 2002 between 250 and 300 m above  
131 sea level. According to <sup>14</sup>C measurement of forest soils, forest stands have been  
132 protected from direct human disturbance for > 400 years (Shen et al., 1999). These  
133 support a rich assemblage of plant species, most of which are evergreen tree species  
134 native to the tropics and subtropics. These include *Castanopsis chinensis* Hance,  
135 *Schima superba* Chardn. & Champ., *Cryptocarya chinensis* (Hance) Hemsl.,  
136 *Cryptocarya concinna* Hance, *Machilus chinensis* (Champ. Ex Benth.) Hemsl., and  
137 *Syzygium rehderianum* Merr. & Perry (Cao et al., 2002). Canopy closure is typically  
138 above 95% (Lu et al., 2010). Soils are oxisols (lateritic red earths) formed from  
139 sandstone approximately 30 cm to 70 cm in depth.

140

## 141 **2.2 Experimental treatments**

142 The experiments involving N amendments were established in July 2003 (Mo et al.,  
143 2006), with four N addition rates used: Control (0 N added), Low-N (50 kg N ha<sup>-1</sup>  
144 yr<sup>-1</sup>), Medium-N (100 kg N ha<sup>-1</sup> yr<sup>-1</sup>) and High-N (150 kg N ha<sup>-1</sup> yr<sup>-1</sup>), which were  
145 based on the present atmospheric N deposition rate and the expected increase in the  
146 future due to the rapid development of agricultural and industrial activities (Galloway  
147 et al., 2004; Lü and Tian, 2007). Considering that any effects of chronic low level N  
148 addition are likely to be similar in direction, if not magnitude, to the short-term effects  
149 of high rates of N addition (e.g. Báez et al. 2007; Clark & Tilman 2008; Lu e al.,  
150 2010), results from our present concentration gradients could be as a prediction for the

151 future changes. A 10 m wide buffer strip surrounded each of 12 10-m x 20-m plots,  
152 with plots and treatments replicated in triplicate and randomly located. A  
153 hand-applied NH<sub>4</sub>NO<sub>3</sub> solution was added each month to the forest floor of each plot  
154 as 12 equal, monthly applications per year. Fertilizer was weighed and mixed with  
155 20 L of deionized water (equivalent of 0.1 mm rainfall), with solution added via  
156 backpack sprayer below the canopy. Two passes were made across each plot to ensure  
157 an even distribution of fertilizer. Control plots received an equivalent volume of  
158 deionized H<sub>2</sub>O.

159

### 160 **2.3 Field water sampling and laboratory analysis**

161 Precipitation and air temperature were monitored in an open area adjacent to the study  
162 plots. The data used in this study were from the weather station in the reserve  
163 (Appendix 1).

164 Soil solution was collected at a 20 cm depth, a depth which represents the primary  
165 rooting zone, and containing >70% of fine root biomass and 68% of total root  
166 biomass (Liao et al., 1993; Wen et al., 1999). Soil solution was sampled with two  
167 replicate zero tension tray lysimeters (755 cm<sup>2</sup> per tray) per plot, which were installed  
168 in April/May 2003 (i.e., 3-4 months prior to our experiment). Each lysimeter was  
169 connected with Tygon tubing to a 10-L bottle.

170 Soil solution samples were taken after each rain event (particularly after heavy  
171 rainstorms) from July 2009 to June 2010. Soil solution volume was recorded and  
172 composited within a plot on data of collection. All collectors were washed with  
173 deionized H<sub>2</sub>O immediately following each collection.

174 Within 24 to 48 hr of field collection, soil solution samples were filtered through  
175 0.45 mm micron filters in the laboratory, then stored in plastic bottles at 4°C until

176 chemical analysis, which included  $\text{NH}_4^+$ -N,  $\text{NO}_3^-$ -N, DOC, and pH. A Shimadzu  
177 TOC-VCSH Total Organic Carbon analyzer was used to determine DOC, with  
178 samples combusted at 680°C via platinum catalyst and  $\text{CO}_2$  determined with a  
179 non-dispersive infrared (NDIR) detector. Samples were analyzed for dissolved  
180 inorganic nitrogen ( $\text{NH}_4^+$ -N and  $\text{NO}_3^-$ -N) using a Lachat QC8000 Flow Injection  
181 Analyzer.

182

#### 183 **2.4 Field soil sampling and laboratory analysis**

184 Samples of mineral soil were collected in August 2009 with a 5-cm diameter corer at  
185 0–10 and 10–20 cm depths. From 0–10 cm, cores were taken beneath the loose litter  
186 layer (O<sub>i</sub>) and comprised Oe and Oa horizon plus mineral soil to a total depth of 10  
187 cm. Following this, the corer was driven to a 20 cm depth for sample collection.  
188 Sampling in each plot took place in three randomly-selected locations

189 In the laboratory, roots and stones were removed by sieving soil to pass a 2-mm  
190 screen; sieved soils mixed thoroughly by hand. For water-extractable DOC (WDOC)  
191 measurements, one 10 g sub-sample from each sample was extracted with 50 ml of  
192 deionized H<sub>2</sub>O for 30 min and filtered through 0.45  $\mu\text{m}$  cellulose–acetate filters, as  
193 modified from (Hagedorn et al., 2002). Water-extractable DOC was determined with a  
194 Shimadzu TOC analyzer as previously described. Other subsamples were air-dried  
195 and used to measure pH (soil:water = 1:2.5) and nutrient content. Total C (total soil  
196 organic C) was measured via titration with  $\text{Fe}^{2+}$  solution following dichromate  
197 oxidation (Liu et al., 1996). Total N was determined by determination of  $\text{NH}_4^+$   
198 following semi-micro Kjeldahl digestion (Liu et al., 1996). Exchangeable Fe and Al  
199 were extracted with 0.1mol/L BaCl<sub>2</sub> (50:1, solution:soil). Subsamples of soil were  
200 oven-dried at 105°C to a constant weight (at least 24 hr) to allow reporting soil results  
201 on an oven-dry basis.

202 **2.5 Field litterfall sampling**

203 Two 1-m x 1-m litter traps with a 1-mm mesh size were placed randomly in each plot  
204 at an approximate 0.5-m height above ground surface. Traps were emptied each  
205 month during the year, with litterfall separated into three components: leaves, small  
206 woody material (branches and bark), and miscellaneous (mainly reproductive parts).

207

208 **2.6 Data analyses**

209 Monthly and annual C effluxes from the primary rooting zone for each plot were  
210 calculated by multiplying DOC concentrations of soil leachate by the recorded water  
211 volume for each sample collection and then summed appropriately. We calculated  
212 mean values per month for  $\text{NH}_4^+$ -N,  $\text{NO}_3^-$ -N, DOC and pH in water samples for  
213 further analysis. Effects of N treatments on soil solution chemistry ( $\text{NH}_4^+$ -N,  $\text{NO}_3^-$ -N,  
214 DOC and pH) and litterfall during the study period were assessed with repeated  
215 measure analysis of variance (ANOVA). One-way ANOVA with Tukey's honestly  
216 significantly different (Tukey's HSD) test was used to test N treatment effects on  
217 concentrations of  $\text{NH}_4^+$ -N and  $\text{NO}_3^-$ -N, pH, and annual DOC effluxes for the whole  
218 study period. One-way ANOVA with Tukey's HSD test was also employed to identify  
219 N treatment effects on soil properties (soil pH, concentrations of total C and N, C/N  
220 ratios, and extractable Fe and Al) and WDOC. Extractable Fe and Al pools were  
221 estimated by multiplying extractable concentrations by soil bulk density, which were  
222 taken from the Dinghushan research station. We conducted the planned contrast  
223 analysis to test for differences between Control plots and N-treatment plots.

224 We also used general linear models to analyze relationships between DOC  
225 concentrations and pH for soil solution sampled at 20 cm depth in all plots during the  
226 study period. Linear regression analysis was also used to examine the relationship

227 between mean DOC of cm soil solution at 20 cm depth and extractable Fe and Al  
228 pools in the upper 20 cm soil, respectively. All analyses were conducted using SPSS  
229 14.0 for Windows® (SPSS, Chicago, IL, USA), with significant differences set with  $P$   
230  $< 0.05$ , unless otherwise stated.

231

### 232 **3 Results**

233 During the study period (July 2009 to June 2010), the total precipitation was 1992  
234 mm, most falling during the March to August wet season (Appendix. 1). Mean  
235 monthly precipitation in wet season (245 mm) nearly three times that of dry season  
236 (88 mm). Mean monthly temperature was 22.2 °C. Total wet N deposition was 34.4  
237 kg N ha<sup>-1</sup>, with 18.2 kg ha<sup>-1</sup> dissolved inorganic N (7.7 kg ha<sup>-1</sup> NO<sub>3</sub><sup>-</sup>-N and 10.5 kg  
238 ha<sup>-1</sup> NH<sub>4</sub><sup>+</sup>-N, respectively) and 16.2 kg ha<sup>-1</sup> dissolved organic N, respectively.

239

#### 240 **3.1 DOC concentration and effluxes**

241 The repeated measures ANOVA revealed that N additions significantly decreased the  
242 DOC concentrations and DOC effluxes at 20 cm depth over the study period ( $df=3$ ,  
243  $F=21.4$ ,  $P=0.001$ ;  $df=3$ ,  $F=6.8$ ,  $P=0.02$ , respectively) (Figure 1a and c). There were  
244 also significant interaction effects between treatment and time (months) on DOC  
245 concentrations and effluxes ( $df=33$ ,  $F=3.6$ ,  $P<0.001$ ;  $df=33$ ,  $F=2.1$ ,  $P=0.006$ ,  
246 respectively). For DOC concentrations, the decreased trends were more pronounced in  
247 Medium-N and High-N plots than that of Low-N plots, and relative measures showed  
248 they decreased by 15%, 28% and 31% in the Low-N, Medium-N and High-N plots,  
249 respectively, relative to that of the Control plots over the whole year (Figure 1b). For  
250 DOC effluxes, they decreased by 44%, 34% and 18% in the Low-N, Medium-N, and  
251 High-N plots, respectively (Figure 1d).

252 Mean DOC concentrations in the Medium-N and High-N treatments were  
253 significantly lower than that of the Controls ( $P<0.05$ ; Table 1). Further analysis  
254 showed that N additions decreased annual DOC effluxes at 20 cm, especially in the  
255 Low-N and Medium-N plots, where the decreases were significant ( $P<0.05$ ; Table 1).  
256 Planned contrast analysis showed that there were significant N-treatment effects for  
257 both mean DOC concentrations and annual DOC effluxes. The annual DOC effluxes  
258 were 99.6, 63.6, 61.0, and 79.1 kg C ha<sup>-1</sup> yr<sup>-1</sup> in the Control, Low-N, Medium-N and  
259 High-N plots, respectively.

260

### 261 **3.2 NO<sub>3</sub><sup>-</sup>-N, NH<sub>4</sub><sup>+</sup>-N and pH in soil solution**

262 The concentrations of NO<sub>3</sub><sup>-</sup>-N in N-treatment plots were generally higher than that of  
263 the Controls (Fig. 2a), and the mean concentrations of NO<sub>3</sub><sup>-</sup>-N across the whole study  
264 period increased, but not significant, in the N-treatment plots ( $P=0.099$ ; Table 1),  
265 although repeated measures ANOVA revealed that N additions did not significantly  
266 increase concentrations of NO<sub>3</sub><sup>-</sup>-N ( $df=3$ ,  $F=2.4$ ,  $P=0.16$ ).

267 Concentrations of NH<sub>4</sub><sup>+</sup>-N (commonly less than 1 mg N L<sup>-1</sup> as mean values for  
268 the whole period) were much lower than those of NO<sub>3</sub><sup>-</sup>-N at all plots (Figure 2b).  
269 There were no significant responses to N treatments across all plots and sampling  
270 times. This is further confirmed by the result of repeated measures ANOVA ( $df=3$ ,  
271  $F=1.4$ ,  $P=0.34$ ). In addition, the mean concentrations of NH<sub>4</sub><sup>+</sup>-N across the whole  
272 study period also showed no significant differences between N treatments and  
273 Controls (Table 1).

274 Repeated measures ANOVA showed that N additions significantly decreased soil  
275 solution pH it at 20 cm depth ( $df=3$ ,  $F=42$ ,  $P<0.001$ ). Similarly, across the whole

276 study period, N treatments changed significantly ( $P=0.001$ ) the mean values of soil  
277 solution pH, especially in the High-N plots (Table 1).

278

279 **3.3 Soil chemistry and litterfall**

280 Total soil N and C and extractable Al and Fe showed increasing trends with elevated  
281 N addition (Table 2). Total N increased by 8%, 12%, 17% in the Low-N, Medium-N,  
282 and High-N treatment plots, respectively, compared to the Controls; total C increased  
283 by 10%, 13%, 17% in the Low-N, Medium-N, and High-N treatment plots,  
284 respectively, compared to Controls. High-N treatments also showed marginally  
285 significant effects on total C ( $P=0.08$ ), and significant effects on extractable Fe  
286 ( $P=0.03$ ). Soil pH values decreased with increasing N treatment levels, especially in  
287 the Medium-N and High-N plots ( $P<0.1$ ). Further analysis showed that there were  
288 significantly linear relationships between treatment levels and pH across all plots  
289 ( $R^2=0.54$ ,  $N=12$ ,  $P=0.006$ ). There was no significant difference among treatments for  
290 soil C/N ratios. For WDOC, N additions increased their contents in this upper 0-20  
291 cm soil, where the increases were significant under N treatments compared to the  
292 Controls ( $P=0.032$ ; Table 2). Annual litterfall was not significantly different among  
293 treatments (Figure 3).

294

295 **3.4 Relationships between DOC, extractable Fe and Al, and pH**

296 Linear model analysis showed that DOC concentrations were significantly and  
297 positively correlated with pH ( $R^2=0.4$ ;  $N=144$ ;  $P<0.001$ ) in soil solution across all  
298 sampling data (Figure 4). Meanwhile, extractable Al pool was not significantly  
299 correlated with mean DOC concentration in the soil solution at 20cm soils ( $R^2=0.004$ ,  
300  $N=12$ ,  $P=0.84$ ; Figure 5a ), but extractable Fe pool exhibited significant and negative

301 correlations with DOC ( $R^2=0.42$ ,  $N=12$ ,  $P=0.023$ ) (Figure 5b).

302

303 **4 Discussions**

304 **4.1 Effects of N addition on DOC leaching**

305 Earlier measurements **in the year 2005** at our site showed that N addition had no  
306 significant effects on soil solution DOC concentrations **below the primary rooting**  
307 **zone** (Fang et al., 2009). Current results, however, indicate that N treatments  
308 significantly decreased DOC concentrations in soil solution **from this layer**,  
309 suggesting that responses of DOC dynamics to N addition may be time-dependent in  
310 N-rich tropical forests. This rejects our initial hypothesis, and also contrasts with other  
311 studies in primarily N-limited ecosystems. In N-limited forests, increased N  
312 availability generally results in more DOC production and subsequent leaching  
313 (Pregitzer et al., 2004; Findlay, 2005; Adams et al., 2005; Sleutel et al., 2009). Smemo  
314 et al. (2007) reported that deposition added as  $\text{NaNO}_3$  significantly increased soil  
315 solution DOC concentration and export from four different northern hardwood forests  
316 in the Great Lakes region. McDowell et al. (1998, 2004) found greater concentrations  
317 of DOC following the addition of  $50\text{-}150 \text{ kg N ha}^{-1} \text{ yr}^{-1}$  as  $\text{NH}_4\text{NO}_3$ . Sleutel et al.  
318 (2009) provide additional evidence on higher concentrations of DOC in boreal forests  
319 of Belgium under historic high N deposition.

320 **Biological mechanisms** (balance between processes that produce and consume  
321 **DOC**) are often suggested to explain changes of DOC concentrations in leachate with  
322 elevated N addition in typically N-limited ecosystems (Neff and Asner, 2001; Knorr et  
323 al., 2005; Zak et al., 2008; Evans et al., 2008). Pregitzer et al. (2004) suggested that  
324 increases in DOC **were** primarily biologically driven, resulting from changes in  
325 production of organic substrates and processing within soil food webs. Bragazza et al.

326 (2006) indicated that the increased release of DOC from litter peat was a consequence  
327 of enhanced enzymatic activity (e.g., phenol oxidase). In these studies, litterfall is the  
328 major source of DOC to the forest floor and thus deeper soil horizons with elevated N  
329 deposition (Currie et al., 1996; Magill and Aber, 2000; Park et al., 2002; Sleutel et al.,  
330 2009). Gundersen et al. (1998) showed a significant correlation between DOC  
331 concentration beneath the forest floor (Oa horizon) and litterfall amount. In a  
332 meta-analysis from multiple terrestrial ecosystems, Liu & Greaver (2010) found that  
333 N addition increased soil DOC concentration by an average of 18%, although soil  
334 respiration was not altered, suggesting C leaching loss may increase in N-limited  
335 ecosystems.

336 In our N-rich forest, however, there were no significant effects of N treatments on  
337 litterfall production (Mo et al., 2008; Figure 3); at the same time, there was no  
338 significant difference between N treatments in DOC dynamics in surface runoff (data  
339 not shown), suggesting that litterfall inputs may play a minor role in DOC production  
340 and subsequent fluxes into deeper soils under N treatments. Further studies showed  
341 that N addition significantly inhibited litter decomposition and decreased soil  
342 respiration in this forest (Mo et al., 2006, 2008). In addition, C mineralization, which  
343 is the conversion from the organic C form to inorganic compound as a result of  
344 decomposition reactions (Carter and Gregorich, 2008), is suggested to result in high  
345 absolute loss of DOC (Chantigny et al., 1999; Huang and Schoenau, 1998; Sjöberg et  
346 al., 2003). An incubation experiment this forest soils, however, showed that N  
347 addition decreased organic C mineralization (Ouyang et al., 2008), indicating that  
348 elevated N inputs may contribute to soil DOC accretion in the deeper soil by  
349 decreasing DOC decomposition/consumption Our findings support this suggestion in  
350 that N addition greatly increased water-extractable DOC (Table 2), a finding also

351 supported by other studies (e.g. Hagedorn et al., 2002; Sinsabaugh et al., 2004; Gallo  
352 et al., 2005). Increases in extractable DOC under experimental N additions are  
353 generally suggested to increase regional and global DOC effluxes from terrestrial  
354 ecosystems to aquatic ecosystems (Pregitzer et al., 2004; Findlay, 2005; Mo et al.,  
355 2008; Chapin et al., 2009). However, our results showed that N treatments decreased  
356 DOC concentrations in leachate solutions. Therefore, biological control mechanisms  
357 are unlikely responsible for declines in DOC in soil leachate of this N-rich tropical  
358 forest.

359 Indeed, as suggested by Neff and Asner (2001), physical controls may also play an  
360 important role in dominating DOC transformations in soils. Here, we propose that  
361 both changes in solution acidity and soil sorption dynamics play a dominant role in  
362 regulating DOC losses from N-rich ecosystems.

363 For example, acidity of soil solution may regulate the patterns of DOC responses.  
364 It has been recognized that the increase of soil solution pH (or acid neutralizing  
365 capacity) would lead to the net positive changes in DOC concentration by increasing  
366 DOC solubility in soil (Monteith et al., 2007; Evans et al., 2008). This was  
367 demonstrated by Evans et al. (2008) while reviewing field N addition experiments in  
368 Europe and North American. It has been widely accepted that high N deposition could  
369 accelerate soil acidification and have the potential to change the acidity of soil  
370 solution (Aber et al., 1989; Vitousek et al., 1997; Bowman et al., 2008; Van den Berg  
371 et al., 2008). In this study, we found that N treatments significantly decreased soil  
372 solution pH below the dominant rooting zone. Further analysis showed that there was  
373 a significant and positive relationship between soil solution pH and DOC  
374 concentration.

375 It should be noted that the effect of pH on DOC dynamics may be confounded

376 with other mechanism related to soil properties, for example, sesquioxides in the  
377 mineral soil (Moore et al., 1992; Guggenberger, 1994; Michalzik et al., 2001). Thus,  
378 we suggest an alternative mechanism for our observations. Soils containing high  
379 concentrations of extractable Fe or/and Al exhibit the capacity to adsorb DOC as  
380 water percolates down through the soil profiles thereby decreasing DOC  
381 concentrations (Boudot et al., 1989; Guggenberger, 1994; Kaiser and Guggenberger,  
382 2000; Sleutel et al., 2009). Corre et al. (2010) suggested that sorption by hydrous Al  
383 oxides could be an important reason for N -induced decreases of soil solution DOC. In  
384 our study, however, the extractable Al did not vary significantly among Controls and  
385 N-treatment plots, and there were no significant relationships between the extractable  
386 Al pool and mean DOC concentration (Table 2, Figure 4a). By contrast, N addition  
387 significantly increased extractable Fe. Also, there was a significant negative  
388 relationship between extractable Fe pool and mean DOC concentration at 20 cm soil  
389 solution (Figure 4b). Accordingly, it is possible that extractable Fe, rather than  
390 extractable Al, may play a key role in DOC adsorption in N-treatment plots after  
391 long-term N inputs. A better understand how Fe and Al oxides control DOC dynamics  
392 in tropical forests merits further study.

393

#### 394 **4.2 Effects of N addition on annual DOC effluxes**

395 Annual DOC effluxes below the 20-cm rooting zones in our study ranged from  
396 60-100 kg C ha<sup>-1</sup> yr<sup>-1</sup>, well within the range (30-139 kg C ha<sup>-1</sup> yr<sup>-1</sup>) reported for  
397 tropical forests by Aitkenhead and McDowell (2000). Our results demonstrated that  
398 long-term N addition decreased annual DOC effluxes from the primary rooting zones,  
399 especially in the Low-N and Medium-N plots. Considering that there were no  
400 significant differences for annual water effluxes among treatments (data now shown),

401 **DOC concentration may dominate DOC effluxes, and lower DOC concentration led to**  
402 **the decreased DOC fluxes under N-treatments.** The decreases in annual DOC effluxes  
403 indicated that soils may accumulate much more DOC with elevated N addition,  
404 consistent with the significant increase of water-extractable DOC at 0-20 cm soil layer  
405 in N-treatment plots (Table 2), and suggesting that elevated N deposition might  
406 enhance soil C sequestration by decreasing DOC effluxes in N-rich forests.

407 Zhou et al. (2006) found that this old-growth forest could accumulate soil C  
408 (0–20cm depth) at about  $610 \text{ kg C ha}^{-1} \text{ yr}^{-1}$  over the last two decades, but concluded  
409 that the reason for this accumulation was unclear. Our results showed that  
410 N-induced net C sequestration (via reduced DOC efflux, calculated by the difference  
411 between N-treatment plots and the Controls) was about  $36, 39, 21 \text{ kg C ha}^{-1} \text{ yr}^{-1}$  in the  
412 Low-N, Medium-N, and High-N plots respectively, with a mean value of  $32 \text{ kg C ha}^{-1} \text{ yr}^{-1}$   
413 in N treatment plots, relative to controls (Table 1). In fact, we have found that N  
414 treatments significantly increased soil total C after long-term N addition (Table 2).  
415 Therefore, such decreases in annual DOC effluxes may explain, in part, this  
416 accumulation of soil C observed by Zhou et al. (2006) considering the high N  
417 deposition during the past decades in this region.

418

#### 419 **4.3 Implications**

420 We have studied effects of long-term N additions on DOC dynamics of soil solution  
421 in N-rich lowland tropical forests under a warm and humid climate. Our results  
422 showed that long-term N additions significantly decreased DOC concentrations in soil  
423 solution of deeper soils. It was suggested that chemo-physical controls (solution  
424 acidity change and soil sorption) rather than biological controls could play a dominant  
425 role in regulating DOC losses from N-rich ecosystems, in contrast to that of N-limited

426 ecosystems. We further found that N addition decreased greatly annual DOC  
427 effluxes below the primary rooting zones, and increased water-extractable DOC in  
428 soils. It is suggested that DOC constitutes an important carbon efflux to forested  
429 mineral soils (Schwesig et al., 2003), and DOC adsorbed by soils may contribute to  
430 the stock of organic C accumulating during soil development (Qualls and Bridgham,  
431 2005). Therefore, our results indicate that long-term N deposition could increase soil  
432 C sequestration in the upper soils by decreasing DOC efflux in N-rich forests, which  
433 may support a novel mechanism responsible for continuing to accumulate C in  
434 old-growth forests (Zhou et al., 2006; Luyssaert et al., 2008). Thus, this study may  
435 give us a new understanding on forests ecosystem C cycling and possible C  
436 sequestration, and also support data bases for model predictions in N-rich ecosystems,  
437 with the globalization of N deposition. Although our findings would be typical for  
438 other N-rich sites, however, our results and corresponding control mechanism should  
439 be further validated in various tropical ecosystems in the future with elevated N  
440 deposition.

441

## 442 **Acknowledgements**

443 This study was founded by the National Basic Research Program of China  
444 (2010CB833502), National Natural Science Foundation of China (No. 30900202,  
445 30970521), and the Knowledge Innovation Program of the Chinese Academy of  
446 Sciences (Grant No.KSCX2-EW-J-28). We wish to thank Dinghushan Forest  
447 Ecosystem Research Station for the strong support in the field work, and Drs. Guoyi  
448 Zhou, Deqiang Zhang, Sandra Brown, Zhi'an Li, Weixing Zhu, Wei Zhang and Juan  
449 Huang for invaluable suggestions in this paper, and Ms. Shaowei Chen and Ms.  
450 Hongying Li for their skilful assistance in laboratory work.

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704 **Tables**

705

706 **Table 1** Effects of N addition on average concentrations of DOC,  $\text{NO}_3^-$ -N,  $\text{NH}_4^+$ -N,  
 707 and pH, and annual DOC efflux in soil solutions below the primary rooting zones  
 708 (0-20 cm soils) during the periods from July 2009 to June 2010. The different  
 709 lowercase letters indicate significant differences at  $P<0.05$  level, and no letters  
 710 indicate no significant differences among N treatment levels, respectively (Tukey's  
 711 HSD test); *Contrast Test* is conducted between N treatments and the Controls using  
 712 planned contrast analysis. Values are mean with S.E. in parentheses.

N treatments	DOC (mg L <sup>-1</sup> )	DOC efflux kg C ha <sup>-1</sup> yr <sup>-1</sup>	$\text{NO}_3^-$ -N (mg L <sup>-1</sup> )	$\text{NH}_4^+$ -N (mg L <sup>-1</sup> )	pH
Control	23.96(2.18)a	99.61(2.63)a	9.74(0.92)	0.24(0.01)	3.81(0.01)a
Low-N	20.19(1.16)ab	63.62(6.25)b	11.26(0.40)	0.23(0.01)	3.78(0.01)a
Medium-N	17.10(0.92)b	60.99(8.87)b	11.74(0.82)	0.24(0.03)	3.78(0.00)a
High-N	14.98(0.46)b	79.06(3.73)ab	14.04(2.05)	0.45(0.20)	3.70(0.01)b
<i>Contrast Test</i>	<i>P</i> =0.003	<i>P</i> =0.002	<i>P</i> =0.099	<i>P</i> =0.58	<i>P</i> =0.001

713 **Table 2** Responses of soil chemistry in the primary rooting zones (0-20 cm soils) to N  
 714 addition in the lowland tropical forest of southern China in August, 2009. The  
 715 different lowercase letters indicate significant differences at  $P<0.05$  level, and no  
 716 letters indicate no significant differences among N treatment levels, respectively  
 717 (Tukey's HSD test). *Contrast Test* is conducted between N treatments and the  
 718 Controls using planned contrast analysis.

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Parameters	N treatments				<i>Contrast Test</i>
	Control	Low-N	Medium-N	High-N	
Total N (mg g <sup>-1</sup> )	1.90(0.11)	2.05(0.08)	2.13(0.09)	2.22(0.06)	$P=0.045$
Total C (mg g <sup>-1</sup> )	21.88(0.40)	24.17(0.97)	24.82(0.50)	25.64(1.44)	$P=0.023$
C/N ratio	11.27(0.45)	11.53(0.83)	11.45(0.59)	11.29(0.44)	$P=0.83$
WDOC(mg Kg <sup>-1</sup> )	107.43(8.24)	160.92(25.55)	140.10(10.97)	179.20(20.30)	$P=0.032$
Al <sup>3+</sup> (m mol kg <sup>-1</sup> )	30.50(1.31)	30.54(2.78)	31.49(1.66)	31.54(1.60)	$P=0.67$
Fe <sup>3+</sup> (m mol kg <sup>-1</sup> )	0.12(0.013)a	0.17(0.018)ab	0.17(0.020)ab	0.20(0.010)b	$P=0.012$
pH (H <sub>2</sub> O)	3.87(0.02)	3.84(0.00)	3.75(0.05)	3.75(0.04)	$P=0.045$

720 Notes: Total C means total soil organic carbon; WDOC means water-extracted  
 721 dissolved organic carbon; Values are means with SE in parentheses.

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731 **Figure Legends**  
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733 **Figure 1** Responses of DOC concentration (a) and its relative concentration (b), and  
734 DOC efflux (c) and its relative efflux (d) to long-term N addition below the  
735 primary rooting zone in the lowland tropical forests of Southern China. **Soil**  
736 **leachate data were available from July 2009 to June 2010.** Notes: Asterisk (\*)  
737 indicates that there are significant differences at  $P<0.05$  level between N  
738 treatments and the Controls using planned contrast analysis.

739 **Figure 2** Responses of  $\text{NO}_3^-$ -N (a),  $\text{NH}_4^+$ -N (b) and pH (c) dynamics to long-term N  
740 addition in soil solution below the dominant rooting zone in the lowland tropical  
741 forests of Southern China. Asterisk (\*) indicates that there are significant  
742 differences at  $P<0.05$  level between N treatments and the Controls using planned  
743 contrast analysis.

744 **Figure 3** Monthly dynamics of litterfall with elevated N addition in the lowland  
745 tropical forests of Southern China during the study period.

746 **Figure 4** Relationships between DOC concentrations and pH in soil solutions across  
747 all plots during the study period. Notes: Triangles ( $\Delta$ ) indicate DOC concentration  
748 at control plots, and solid circles ( $\bullet$ ) indicate DOC concentration at N-treatments  
749 plots.

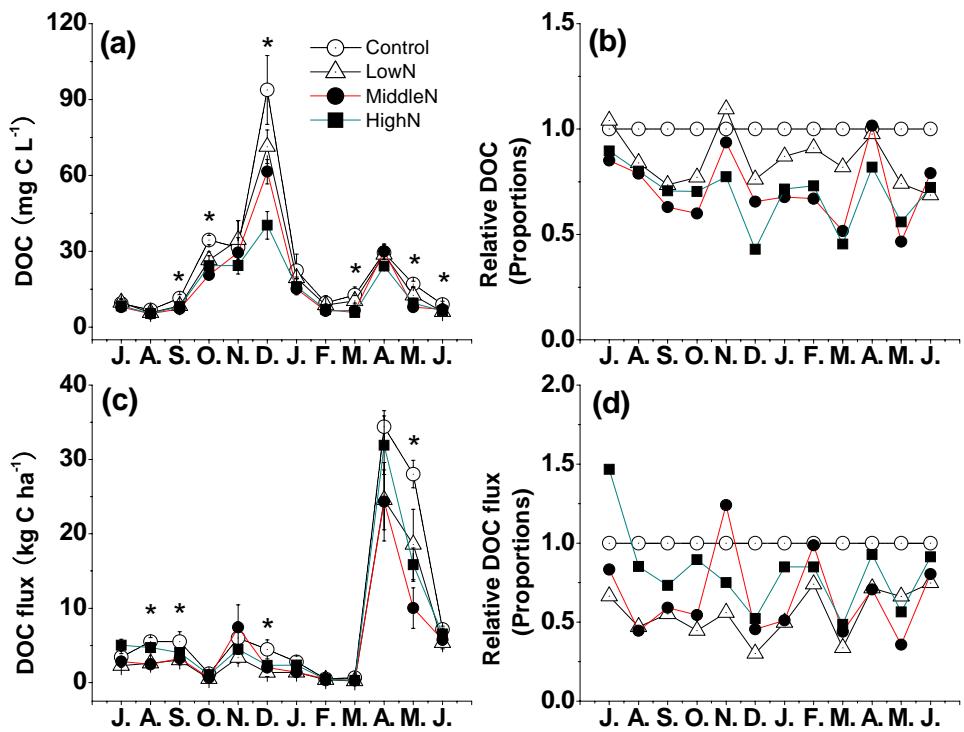
750 **Figure 5** Relationships between mean DOC concentrations in soil solution during the  
751 study period and soil extractable Al and Fe pools in upper 0-20cm mineral soils.

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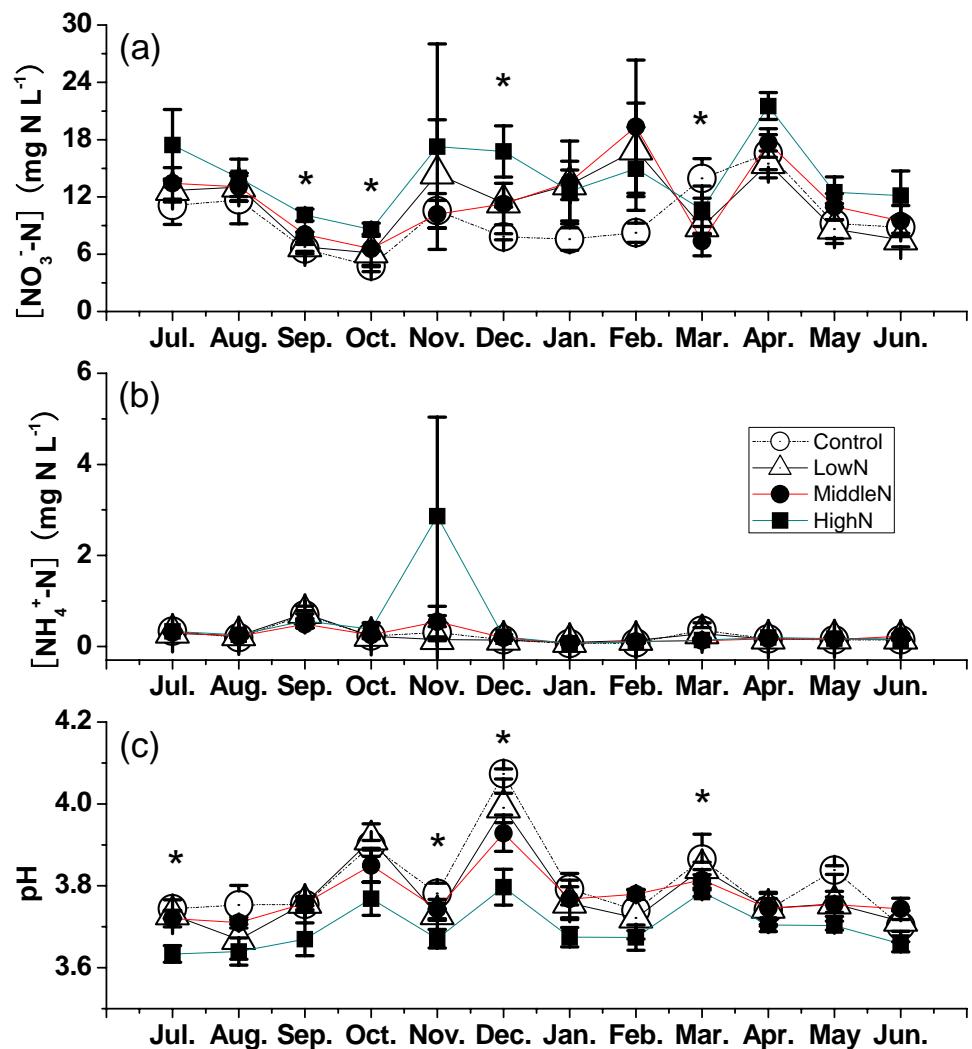
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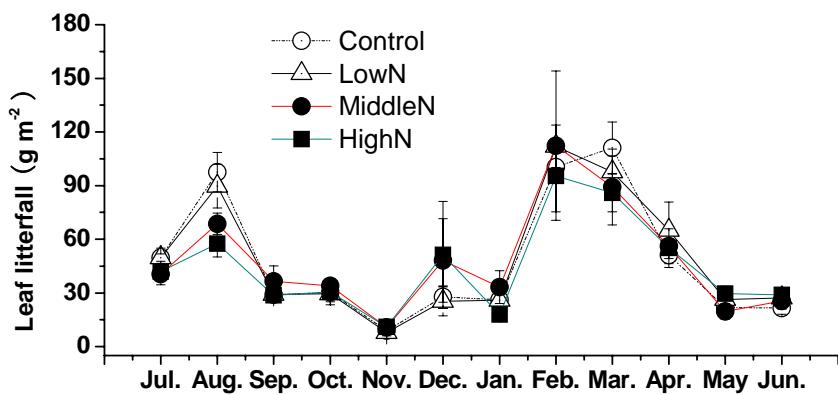
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757 **Figure 1**



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Figure 2



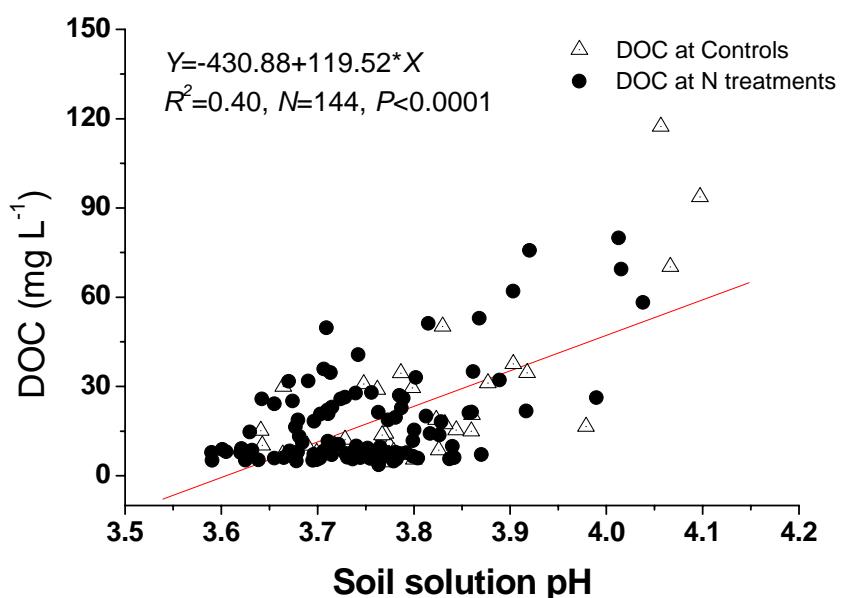
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761 **Figure 3**

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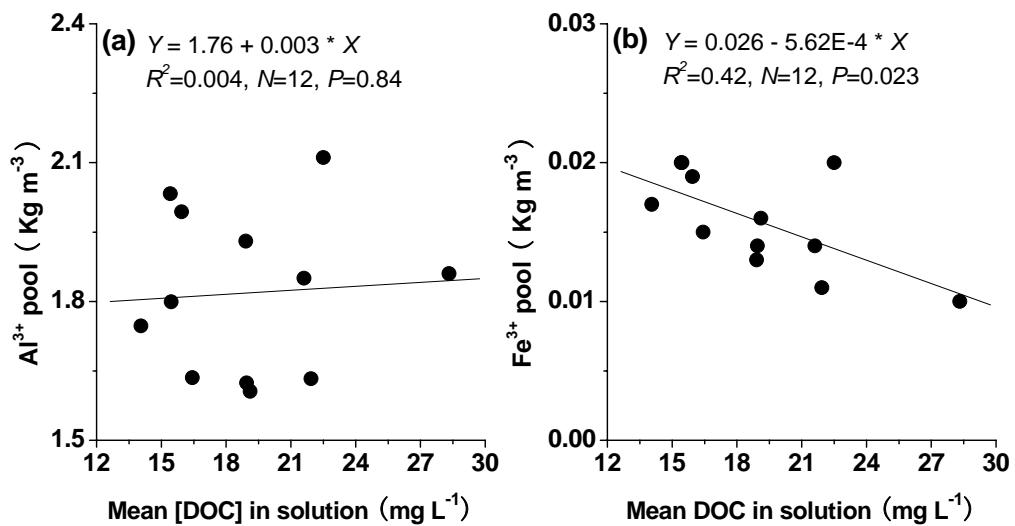


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767 **Figure 4**

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**Figure 5**

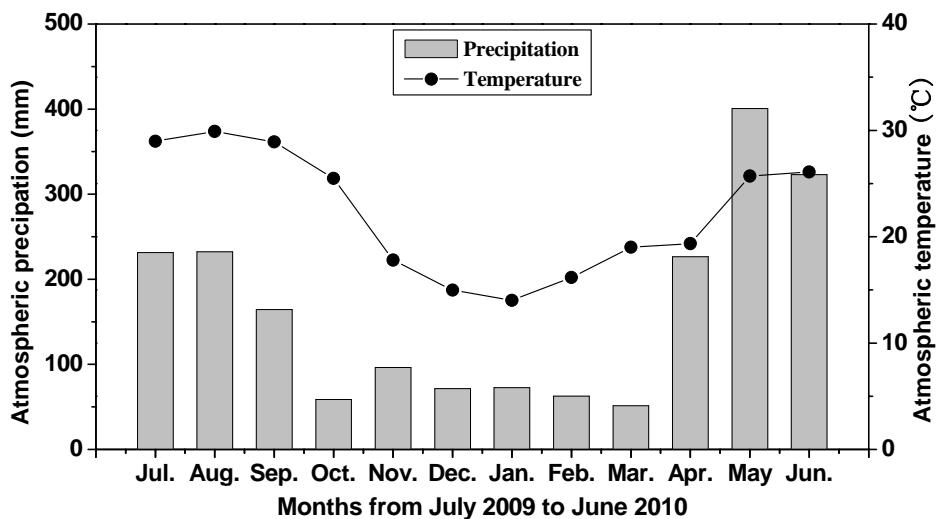
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**Appendix 1** Monthly precipitation and monthly mean air temperature at Dinghushan

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Biosphere Reserve, southern China, during this study period.