

# 1           Interactive effects of soil amendments (biochar and gypsum) and 2           salinity on ammonia volatilization in coastal saline soil

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30     **Abstract:** Ammonia ( $\text{NH}_3$ ) volatilization is a major route of nitrogen (N) loss from soil, especially in saline soil.  
31     Biochar and gypsum are two important soil amendments that are widely used in coastal saline farmland.  
32     However, little is known about the interactive effects of soil amendments and salinity on  $\text{NH}_3$  volatilization. In  
33     this study, five soil salinity levels, three N sources (urea, monoammonium phosphate (MAP), and manure) and  
34     two soil amendments (biochar and gypsum, both applied at two rates) were selected to conduct incubation  
35     experiments. Nitrogen transformation experiments were conducted simultaneously with  $\text{NH}_3$  volatilization  
36     experiments. The results showed that cumulative  $\text{NH}_3$  volatilization increased with salinity due to the  
37     accumulation of ammonium, which resulted from the inhibition of nitrification by salinity, and also the effect  
38     of salinity on soil properties related to  $\text{NH}_3$  volatilization. Among the tested N sources, the highest  $\text{NH}_3$   
39     volatilization was observed for urea, followed by MAP and manure, in soils with three different salinity levels.  
40     Biochar application increased  $\text{NH}_3$  volatilization in saline soil, as salt ions constrained the  $\text{NH}_3/\text{NH}_4^+$  adsorption  
41     capacity of biochar, and the inhibition of nitrification by biochar was aggravated in saline soil. Overall,  $\text{NH}_3$   
42     volatilization increased with the biochar application rate in saline soil, whereas the effect of a low rate of biochar  
43     application on  $\text{NH}_3$  volatilization was not significant. Gypsum decreased  $\text{NH}_3$  volatilization in saline soil,  
44     whereas the contribution of different gypsum rates to  $\text{NH}_3$  volatilization showed no difference. The conclusion  
45     could be drawn from the above results that low rates of biochar and gypsum may prevent or reduce  $\text{NH}_3$   
46     volatilization and nitrogen losses in coastal saline soil.

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48     **Keywords:** Ammonia volatilization; Soil salinity; Biochar; Gypsum; Nitrogen sources; Interactive effects

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52     **1. Introduction**

53       Nitrogen (N) is a vital dominant nutrient for crop growth that undergoes a series of transformations in soil,  
54       including ammonia ( $\text{NH}_3$ ) volatilization, nitrification, denitrification and immobilization. During the various  
55       transformation processes, some nitrogen enters the atmosphere and water through runoff, leaching,  $\text{NH}_3$   
56       volatilization and nitrous oxide emissions, resulting in a reduction in the available nitrogen in farmland. Many  
57       investigations conducted in China have indicated that the recovery efficiency of N fertilizer is only 18%~30%  
58       (Wang et al., 2001; Peng et al., 2006, 2010) and is lower than the ideal N recovery efficiency (30%~50%)  
59       indicated by Dobermann (2005). In addition to the adverse impacts on crop productivity, the loss of nitrogen to  
60       the atmosphere and water will lead to atmospheric pollution and water eutrophication.

61        $\text{NH}_3$  volatilization is one of the major pathways of nitrogen loss in arable land (Rochette et al., 2013), and  
62       the proportion of  $\text{NH}_3$  volatilization loss from applied nitrogen fertilizers ranges from 1~50% (Sommer et al.,  
63       2004). In some rice-wheat rotation systems in South China,  $\text{NH}_3$  volatilization can even become the main route  
64       of nitrogen fertilizer loss (Liu et al., 2015). Beusen et al. (2008) investigated the  $\text{NH}_3$  emissions from agricultural  
65       systems on a worldwide scale and found that 27-38 million tons of  $\text{NH}_3$  are emitted into the atmosphere every  
66       year, which results in the loss of nitrogen in agro-ecological systems. On the other hand,  $\text{NH}_3$  is a vital  
67       atmospheric contaminant involved in atmospheric processes such as the formation of fine particulate matter  
68       ( $\text{PM}_{2.5}$ ), which is the key facilitating factor in haze formation (Ho et al., 2016; Yang et al., 2011). Furthermore,  
69       the redeposition of  $\text{NH}_3$  to the land surface is an important cause of soil acidification (Van Breemen et al., 1982)  
70       and the eutrophication of water ecosystems (Beusen et al., 2008).

71        $\text{NH}_3$  volatilization is affected by a series of factors, and the key influencing factor varies under different  
72       environmental conditions. In saline soil, soil salinity may become the main influencing factor. Soil salinization  
73       is a worldwide problem affecting approximately 10 million  $\text{km}^2$  of land (Setia and Marschner, 2013). Some

74 studies have indicated that NH<sub>3</sub> volatilization shows a positive correlation with salinity (Duan and Xiao, 2000).  
75 However, NH<sub>3</sub> volatilization is affected by related nitrogen transformation mechanisms such as urea hydrolysis  
76 and nitrification. Kumar and Wagenet (2007) reported that the amount of urea hydrolyzed was lower in saline  
77 soil than in non-saline soil. On the other hand, some studies have investigated the effect of salinity on  
78 nitrification. For example, Akhtar et al. (2012) indicated that the first and second steps of nitrification were both  
79 inhibited by salinity, and Reddy and Crohn (2014) reported that salinity had a negative effect on nitrification.  
80 Some studies have also indicated that microorganisms related to nitrogen transformation are inhibited by salinity  
81 (Yuan et al., 2007; Rietz and Haynes, 2003). These studies have indicated that urea hydrolysis and nitrification  
82 are both inhibited by salinity. However, urea hydrolysis and nitrification have adverse effects on the ammonium  
83 content in soil. The inhibition of nitrification will lead to the accumulation of ammonium and, consequently, the  
84 aggravation of NH<sub>3</sub> volatilization. In contrast, the inhibition of hydrolysis may reduce the NH<sub>4</sub><sup>+</sup>-N concentration  
85 and thus reduce NH<sub>3</sub> volatilization relative to other nitrogen fertilizers which releases nitrogen directly without  
86 hydrolysis. Therefore, the effect of salinity on the NH<sub>3</sub> volatilization of urea may be different from that in other  
87 nitrogen fertilizers and needs to be further studied.

88 Biochar, an amendment pyrolyzed from organic material, has attracted much scientific and commercial  
89 attention in recent years regarding its function in sequestering carbon and the improvement of nutrient use  
90 efficiency (Mandal et al., 2016; Liu et al., 2013) due to its properties of C-richness, porosity, alkalinity and high  
91 adsorbability (Cao et al., 2011). Numerous studies involving field, pot and incubation experiments have been  
92 conducted to investigate the effects of biochar application on soil physical-chemical properties (Suliman et al.,  
93 2017), soil nutrient retention (Knowles et al., 2011; Bai et al., 2015), crop productivity and nutrient uptake  
94 (Zhang et al., 2012; Gruber et al., 2010), greenhouse gas emission (Case et al., 2012; Nelissen et al., 2014), NH<sub>3</sub>  
95 volatilization (Mandal et al., 2016; Feng et al., 2017), and soil microbial activities (Lin et al., 2017; Bi et al.,

96 2017). The results have indicated that biochar is beneficial to most of the aforementioned processes. However,  
97 the conclusions about the effect of biochar addition on NH<sub>3</sub> volatilization have been inconsistent. For example,  
98 Mandal et al. (2016) reported that 5% biochar addition significantly reduced NH<sub>3</sub> volatilization in non-saline  
99 soil. In contrast, Feng et al. (2017) found that low biochar treatment (0.5 wt%) did not significantly increase  
100 NH<sub>3</sub> volatilization, while higher biochar treatment (3 wt%) significantly increased NH<sub>3</sub> volatilization in non-  
101 saline paddy soils. Furthermore, Sun et al. (2017) found that biochar applied at higher rates (>2%) aggravated  
102 NH<sub>3</sub> volatilization in coastal saline soil. The effect of biochar addition on NH<sub>3</sub> volatilization depends on the  
103 interactions of several properties including adsorbability, pH and the impact on nitrification (Feng et al., 2017).  
104 However, the dominant property varies under different conditions. In saline soil, the NH<sub>3</sub> adsorption capacity  
105 of biochar may be affected by the higher salt ion content. Additionally, its effect on nitrification may be  
106 influenced by the inhibition of nitrification by salinity. Therefore, the effect of biochar on NH<sub>3</sub> volatilization in  
107 saline soil may be different from that in non-saline soil. However, few studies have compared the effect of  
108 biochar addition on NH<sub>3</sub> volatilization in saline and non-saline soils. Thus, the interactive effects of biochar  
109 addition and salinity on NH<sub>3</sub> volatilization warrant further study.

110 Gypsum, a byproduct of flue-gas desulfurization, is widely used as a saline-alkali soil amendment due to  
111 its low cost and high production (Stamford et al., 2015). Various studies have been conducted to explore the  
112 effect of gypsum addition on soil physical-chemical properties (Yu et al., 2003), crop yields (Hamza and  
113 Anderson, 2003; Caires et al., 2010), soil respiration (Wong et al., 2009) and so on. However, its effect on NH<sub>3</sub>  
114 volatilization in soils is poorly documented. Only Zia et al. (1995) has compared NH<sub>3</sub> volatilization in soil with  
115 or without gypsum amendment, and they found that gypsum application significantly reduced NH<sub>3</sub> volatilization  
116 loss. In addition, some researchers have investigated the effect of gypsum addition on NH<sub>3</sub> volatilization during  
117 the composting of livestock manure and biosolids (Mishra et al., 2013; Burt et al., 2017; Tubail et al., 2008).

118 Most researchers believe that the effect of gypsum addition on reducing NH<sub>3</sub> volatilization might be attributed  
119 to its lower pH (Zia et al., 1999). However, a recent study indicated that gypsum addition inhibited the  
120 abundance of nitrification functional genes (Li et al., 2016), thus inhibiting nitrification, which might lead to  
121 the accumulation of ammonium. In saline soil, the inhibitory effect of gypsum on nitrification may be  
122 aggravated by the inhibition of nitrification by salinity. The interactive effects of gypsum and salinity on NH<sub>3</sub>  
123 volatilization are still ambiguous and require further study.

124 The coastal saline soil region is one of the major saline soil regions in China and is distributed across 11  
125 provinces in eastern China. In salinized farmland, soil salinity influences nitrogen transformation and NH<sub>3</sub>  
126 volatilization. Biochar and gypsum are two widely used amendments in saline soil. Their interactive effects with  
127 salinity on NH<sub>3</sub> volatilization are poorly studied in saline soil. Therefore, we selected five soil salinities, three  
128 nitrogen sources and two amendments (biochar and gypsum) (1) to investigate the influence of soil salinity on  
129 NH<sub>3</sub> volatilization, (2) to determine the NH<sub>3</sub> volatilization of different nitrogen sources, and (3) to assess the  
130 interactive effects of soil amendments (biochar and gypsum) and salinity on NH<sub>3</sub> volatilization in saline soil.

## 131 **2. Materials and methods**

### 132 **2.1. Soil, nitrogen sources, biochar and gypsum**

133 Soil samples were collected from the 0~10 cm soil layer in coastal saline farmland in eastern China  
134 (32°38'42.01"N, 120°54'8.04"E) in June 2018 after the wheat harvest. This farmland is located approximately  
135 1 km from the Yellow Sea in the east. The soils exhibited five different salt contents and were collected from  
136 five sites in the same farmland. All the soils were typical saline alluvial soils (Fluvisols, FAO). We designated  
137 the soils NS (non-saline soil), LS (low-salinity soil), MS (moderately saline soil), HS (high-salinity soil) and  
138 SS (severely saline soil). Then, the soil samples were air dried and ground to pass through a 2 mm sieve prepared  
139 for the incubation experiment. Nitrogen sources including urea, monoammonium phosphate (MAP) and manure

140 were collected from a local poultry farm. Biochar was derived from wheat straw under oxygen-limited  
141 conditions. The temperature was increased to 550°C at a rate of 5 °C min<sup>-1</sup> and then maintained for 4 h. Gypsum  
142 was purchased from a local gypsum factory (Zhenhua gypsum Co., Ltd, Jiangsu, China).

143 The pH of the soils and amendments (1:5 w/v) was measured using a combined pH electrode (Mettler-  
144 Toledo Ltd., Shanghai, China); the electrical conductivity (EC) was measured using a conductivity electrode  
145 (Mettler-Toledo Ltd., Shanghai, China); the total organic carbon (TOC) content of the soils was measured using  
146 the potassium dichromate external heating method; the total N content was measured using the Kjeldahl method;  
147 the soil texture was measured using the hydrometer method; and the cation exchange capacity (CEC) was  
148 measured using the ammonium acetate extraction-flame photometry method (Lu, 2000). Proximate analysis of  
149 the biochar was performed by thermogravimetry using a muffle furnace (Liu et al., 2018). The properties of the  
150 soils and amendments are listed in Table 1.

151 **2.2. Experimental treatments and management**

152 Three incubation experiments were conducted in this study. For all experiments, 100 g of soil (on an oven-  
153 dried basis) was weighed and transferred to a 500 ml Mason jar. All jars were incubated in the dark at 25±1°C.  
154 During the incubation period, soil moisture was maintained at 60% of water-filled pore space (WFPS), which  
155 was calculated from the bulk density and the gravimetric moisture content (Yu et al., 2016). Before incubation,  
156 all jars were preincubated at 25°C in a constant-temperature incubator for three days to activate the soil  
157 microorganisms and ensure water diffusion uniformity. The first experiment was conducted to measure the effect  
158 of soil salinity on NH<sub>3</sub> volatilization. All five soils with urea as the nitrogen source were used to conduct a 65-  
159 day experiment. The second experiment was performed to measure the effect of nitrogen sources on NH<sub>3</sub>  
160 volatilization. Three soils (NS, LS and MS) were selected from the five soils included in experiment 1, and urea,  
161 MAP and manure were added to each soil for a 29-day experiment. The third experiment was conducted to

162 measure the effect of biochar and gypsum on NH<sub>3</sub> volatilization. Two soils (NS and MS) were selected to  
163 conduct a 29-day experiment with urea as the nitrogen source. For all experiments, nitrogen sources were added  
164 at a rate of 200 mg N kg<sup>-1</sup> soil. The urea and MAP were dissolved in deionized water and added to the soils, and  
165 manure was added as dry material and mixed with the soil thoroughly, as were biochar and gypsum. Biochar  
166 was added at rates of 1% and 2% (equivalent to 15 t ha<sup>-1</sup> and 30 t ha<sup>-1</sup>) because 15–30 t ha<sup>-1</sup> of biochar is the  
167 range that is widely used (Taghizadeh-Toosi et al., 2012a, b). Gypsum was added at rates of 0.36 g kg<sup>-1</sup> and 0.72  
168 g kg<sup>-1</sup> (equivalent to 5 t ha<sup>-1</sup> and 10 t ha<sup>-1</sup>, respectively). Each treatment included three replications.

169 **2.3. NH<sub>3</sub> volatilization measurement**

170 For the measurement of NH<sub>3</sub> volatilization, a 25 ml glass beaker containing 10 ml of 0.1 M H<sub>2</sub>SO<sub>4</sub> was  
171 placed in a Mason jar to absorb NH<sub>3</sub>. Then, the NH<sub>3</sub> absorbed by H<sub>2</sub>SO<sub>4</sub> was measured using the indigo blue  
172 colorimetry method (Lu, 2000). NH<sub>3</sub> volatilization was measured every day in the first week, then every two  
173 days in the second week, every three days in the third week, and every four days or once per week in the next  
174 few weeks. Cumulative NH<sub>3</sub> volatilization was calculated from the sum of the daily NH<sub>3</sub> volatilization  
175 measurements.

176 **2.4. Soil N transformation**

177 Nitrogen transformation experiments were conducted synchronously with the NH<sub>3</sub> volatilization  
178 experiments. We set up a control for each of the soils and then determined NH<sub>4</sub><sup>+</sup>-N and nitrate NO<sub>3</sub><sup>-</sup>-N  
179 concentrations by calculating the difference between the soils with and without nitrogen addition.

180 NH<sub>4</sub><sup>+</sup>-N and NO<sub>3</sub><sup>-</sup>-N were extracted using a 1 M potassium chloride solution at a ratio of 1:10 w/v in a 250  
181 ml conical flask, and the flask was then subjected to shock using a 200 rpm end-over-end shaker for 1 h at 25°C.  
182 The extracted solution was filtered for the measurement of NH<sub>4</sub><sup>+</sup>-N and NO<sub>3</sub><sup>-</sup>-N concentrations. NH<sub>4</sub><sup>+</sup>-N was  
183 measured using the indigo blue colorimetry method, and NO<sub>3</sub><sup>-</sup>-N was measured using the ultraviolet dual

184 wavelength method (Lu, 2000).

185 **2.5. Statistical analysis**

186 The means and standard deviations were calculated using Excel 2016. The statistical tests were conducted  
187 using SPSS 16.0. The figures were drawn using Excel 2016 and Origin 8.0.

188 **3. Results**

189 **3.1. Effect of soil salinity on NH<sub>3</sub> volatilization**

190 Different NH<sub>3</sub> volatilization patterns were observed in the soils with different salinities during the 65-day  
191 incubation experiment (Fig. 1a). The NH<sub>3</sub> volatilization rate, the length of the NH<sub>3</sub> volatilization period and  
192 cumulative NH<sub>3</sub> volatilization were all different between the treatments. Regarding the NH<sub>3</sub> volatilization rate,  
193 at the early stage, the NH<sub>3</sub> volatilization rate in the higher-salinity soils (HS and SS) was lower than that in the  
194 lower-salinity soils (NS, LS and MS), while in the later stage, the NH<sub>3</sub> volatilization rate in the higher-salinity  
195 soils (HS and SS) was greater than that in the lower-salinity soils (NS, LS and MS). NH<sub>3</sub> volatilization in NS,  
196 LS, MS, HS and SS reached the peak rate on the third day, fourth day, seventh day, thirteenth day and twenty-  
197 fifth day, respectively, and the length of the NH<sub>3</sub> volatilization period in NS, LS, MS, HS and SS was 6 days,  
198 11 days, 13 days, 44 days and 65 days, respectively. The cumulative NH<sub>3</sub> volatilization increased with salinity,  
199 with cumulative amounts of 1.50 mg kg<sup>-1</sup>, 4.16 mg kg<sup>-1</sup>, 5.03 mg kg<sup>-1</sup>, 27.71 mg kg<sup>-1</sup> and 37.73 mg kg<sup>-1</sup> being  
200 recorded in NS, LS, MS, HS and SS, respectively, and a significant difference existed between the treatments  
201 ( $P<0.05$ ) (Fig. 1b). A correlation analysis was performed to evaluate the relationship between soil salinity and  
202 cumulative NH<sub>3</sub> volatilization, and the results showed that cumulative NH<sub>3</sub> volatilization was positively linearly  
203 correlated with soil salinity ( $R^2 = 0.977$  at  $P<0.05$ ) (Fig. 1c).

204 **3.2. Effect of N sources on NH<sub>3</sub> volatilization**

205 Different NH<sub>3</sub> volatilization patterns were observed in the soils treated with different N sources (Fig. 2).

206 The length of the NH<sub>3</sub> volatilization period and cumulative NH<sub>3</sub> volatilization differed under the different N  
207 sources in the three soils. The NH<sub>3</sub> volatilization period of all nitrogen sources increased with salinity, which  
208 confirmed the finding described in section 3.1 that salinity aggravated NH<sub>3</sub> volatilization. In all three soils (NS,  
209 LS and MS), the NH<sub>3</sub> volatilization rate of manure was always the lowest throughout the NH<sub>3</sub> volatilization  
210 process. The mean NH<sub>3</sub> volatilization rate of urea was higher than that of MAP. However, MAP exhibited a  
211 longer NH<sub>3</sub> volatilization period than urea, and the NH<sub>3</sub> volatilization rate of urea was higher than that of MAP  
212 in the early stage, while an opposite pattern appeared in the later stage. The cumulative NH<sub>3</sub> volatilization of  
213 different nitrogen sources displayed the order of urea>MAP>manure in the three saline soils, and the difference  
214 in cumulative NH<sub>3</sub> volatilization increased with salinity.

### 215 **3.3. Effect of biochar and gypsum on NH<sub>3</sub> volatilization**

216 Two biochar and gypsum application rates were selected to measure their effects on NH<sub>3</sub> volatilization in  
217 two soils (NS and MS), and low cumulative NH<sub>3</sub> volatilization was observed under all treatments in non-saline  
218 soil compared with moderately saline soil (Fig. 3). In non-saline soil, neither high (BCH) nor low biochar  
219 addition (BCL) significantly affected cumulative NH<sub>3</sub> volatilization compared with the control, and the  
220 cumulative NH<sub>3</sub> volatilization under BCL was slightly higher than that under BCH. Both high (SGH) and low  
221 gypsum addition (SGL) significantly decreased cumulative NH<sub>3</sub> volatilization compared with the control, and  
222 the cumulative NH<sub>3</sub> volatilization under SGL was lower than that under SGH. The difference between the high  
223 and low dosages of the amendments was not significant. In moderately saline soil, biochar addition increased  
224 the cumulative NH<sub>3</sub> volatilization compared with the control, with BCH significantly increasing the cumulative  
225 NH<sub>3</sub> volatilization. In contrast, gypsum addition decreased the cumulative NH<sub>3</sub> volatilization compared to the  
226 control; however, the difference was not significant under either high or low gypsum addition.

### 227 **3.4. Effect of salinity on N transformation**

228       The  $\text{NO}_3^-$ -N and  $\text{NH}_4^+$ -N concentrations in soils with different salinities were measured in parallel with the  
229        $\text{NH}_3$  volatilization experiment (Fig. 4). Similar change patterns for both  $\text{NO}_3^-$ -N and  $\text{NH}_4^+$ -N were observed in  
230       the lower-salinity soils (NS, LS and MS). In the first week, the  $\text{NH}_4^+$ -N concentration reached its peak, after  
231       which it decreased sharply in the second week due to nitrification, while the  $\text{NO}_3^-$ -N concentration reached its  
232       peak in the second week. The  $\text{NH}_4^+$ -N concentrations among the three soils displayed the order of MS>LS>NS  
233       in the first week, which was consistent with the cumulative  $\text{NH}_3$  volatilization (Fig. 1b) observed in NS, LS and  
234       MS.

235       The dynamics of the  $\text{NO}_3^-$ -N and  $\text{NH}_4^+$ -N concentrations in the higher-salinity soils (HS and SS) were  
236       different from those in the above three soils. The higher-salinity soils exhibited longer nitrogen transformation  
237       periods. The  $\text{NH}_4^+$ -N concentration continuously increased in the first 3 weeks for HS and the first 4 or 5 weeks  
238       for SS. The  $\text{NO}_3^-$ -N concentration continued to increase throughout the period. At the end of the incubation  
239       experiment, the  $\text{NO}_3^-$ -N concentrations in HS and SS were significantly lower than those in the three lower-  
240       salinity soils.

241       **3.5. Effect of N sources on N transformation**

242       The  $\text{NO}_3^-$ -N and  $\text{NH}_4^+$ -N concentrations of the soils treated with different N sources are shown in Fig. 5.  
243       Among the tested N sources, the  $\text{NH}_4^+$ -N and  $\text{NO}_3^-$ -N concentrations of the soils with manure addition were  
244       always the lowest and were significantly lower than those of the soils with urea and MAP addition throughout  
245       the incubation experiment. The same pattern was found in all three soils. For urea and MAP, similar change  
246       patterns of  $\text{NO}_3^-$ -N and  $\text{NH}_4^+$ -N concentrations were also found in all three soils. The  $\text{NH}_4^+$ -N concentration in  
247       the soils with MAP addition was always higher than that in the soils with urea addition during the incubation  
248       period. Additionally, the  $\text{NH}_4^+$ -N concentration in the soils with urea addition decreased more rapidly than that  
249       in the soils with MAP addition, and the  $\text{NO}_3^-$ -N concentration in the soils with urea addition increased more

250 rapidly than that in the soils with MAP addition.

251 **3.6. Effect of biochar and gypsum on N transformation**

252 The  $\text{NO}_3^-$ -N and  $\text{NH}_4^+$ -N concentrations of the soils amended with biochar and gypsum are shown in Table  
253 2 (non-saline soil) and Table 3 (moderately saline soil). Regarding biochar addition, in the non-saline soil, even  
254 when the  $\text{NH}_4^+$ -N concentration sharply decreased to a low level in the first week due to nitrification, the  $\text{NO}_3^-$   
255 -N concentration was lower than that in the control throughout the incubation experiment. In the moderately  
256 saline soil, nitrification was slower than in the non-saline soil. The  $\text{NH}_4^+$ -N concentration in the soils with  
257 biochar addition was significantly higher than that in the control in the first two weeks. The  $\text{NO}_3^-$ -N  
258 concentration was lower than that in the control throughout the incubation experiment, and the  $\text{NO}_3^-$ -N  
259 concentration in the soil with high biochar addition was lower than that in the soil with low biochar addition.

260 Under gypsum addition, in the non-saline soil, the  $\text{NH}_4^+$ -N concentration was low due to rapid nitrification,  
261 and the  $\text{NO}_3^-$ -N concentration in the soil with gypsum addition was lower than that in the control in the first  
262 week. In the moderately saline soil, the  $\text{NH}_4^+$ -N concentration under gypsum addition was also significantly  
263 higher than that in the control in the first two weeks, and the  $\text{NO}_3^-$ -N concentration was significantly lower than  
264 that in the control in the first two weeks.

265 **3.7. Effect of biochar and gypsum on soil pH**

266 The pH levels of the soils with biochar and gypsum addition after the incubation experiment are shown in  
267 Fig. 6. In the non-saline soil, high biochar addition (BCH) significantly reduced the soil pH. However, the  
268 difference in pH between the low biochar addition (BCL) and the control treatments was not significant, and  
269 the difference in pH between BCH and BCL was also not significant. In the moderately saline soil, the  
270 differences were similar to those in the non-saline soil, except that the pH under BCH was significantly lower  
271 than that under BCL. Regarding the gypsum treatments, both high (SGH) and low gypsum addition (SGL)

272 significantly reduced the soil pH, and the pH decreased with the gypsum addition rate in the non-saline soil, as  
273 also in the moderately saline soil.

274 **4. Discussion**

275 **4.1. Effect of soil salinity on NH<sub>3</sub> volatilization**

276 NH<sub>3</sub> volatilization is one of the most important routes of nitrogen loss in farmland and causes large nitrogen  
277 losses around the world every year (Sommer et al., 2004). In recent years, NH<sub>3</sub> volatilization under different  
278 conditions has been widely studied. These studies have covered large scales and have examined factors such as  
279 soil reactions (Mandal et al., 2016), soil amendments (Martines et al., 2010; Chen et al., 2013), water and  
280 nitrogen management (Zhou et al., 2016), tillage management (Rochette et al., 2009), and biochemical inhibitors  
281 (Soares et al., 2012). However, few studies have focused on soil salinity, even though salinization is a common  
282 problem in agricultural land and 50% of all arable land is predicted to be affected by salinization by 2050 (Wang  
283 et al., 2003). Additionally, in some related nitrogen transformation studies, a salinity gradient has been achieved  
284 by adding salt, which does not reflect actual field conditions well. Therefore, we selected in situ soils with  
285 different salinities to conduct our experiments.

286 In our study, NH<sub>3</sub> volatilization was positively correlated with soil EC. However, other soil properties,  
287 including the soil pH, texture, organic matter content and CEC, might also have affected NH<sub>3</sub> volatilization  
288 because in situ soils were selected. For instance, Mandal et al. (2016) indicated that an increase in pH increased  
289 NH<sub>3</sub> volatilization. Fenn and Kissel (1976) found that NH<sub>3</sub> volatilization was negatively correlated with CEC.  
290 Reynolds and Wolf (1987) reported that NH<sub>3</sub> volatilization was negatively correlated with clay content and  
291 organic carbon. Therefore, we analyzed the correlations between cumulative NH<sub>3</sub> volatilization and these soil  
292 properties. In our study, the soil pH and clay content were not correlated with NH<sub>3</sub> volatilization because these  
293 properties were similar in all soils. The TOC content was negatively correlated with NH<sub>3</sub> volatilization ( $R^2=0.50$ ,

294 P<0.01). However, the difference in TOC might have resulted from the difference in soil salinity. Na<sup>+</sup>  
295 accumulation degrades the soil aggregate structure, which leads to a decrease in organic carbon (Guo et al.,  
296 2019). On the other hand, higher soil salinity decreases plant root biomass, which is an important source of soil  
297 carbon. CEC was also negatively correlated with NH<sub>3</sub> volatilization ( $R^2=0.57$ , P<0.01) in the present study.  
298 Previous studies have indicated that soil CEC is determined by the soil clay and organic matter contents and pH  
299 (Lu, 2000). However, in our study, the soil pH and clay content showed no correlation with CEC, while soil  
300 organic carbon showed a significant correlation with CEC ( $R^2=0.73$ , P<0.01), which indicated that soil CEC  
301 was also affected by soil salinity. These results confirmed that soil salinity was the dominant factor under our  
302 study conditions.

303 As shown in Fig. 1, NH<sub>3</sub> volatilization presented different change patterns between lower-salinity soils  
304 (NS, LS and MS) and higher-salinity soils (HS and SS). In the lower-salinity soils, NH<sub>3</sub> volatilization increased  
305 rapidly and reached a peak, then decreased dramatically over a short time (two weeks), which was consistent  
306 with other studies, such as that of Feng et al. (2017), who found that NH<sub>3</sub> volatilization occurred for a short time  
307 and was nearly complete within one week, as also indicated by Sun et al. (2013). This is because urea hydrolysis  
308 is a rapid process if there is no influence of inhibitory factors (Vlek and Carter, 1983). In our study, cumulative  
309 NH<sub>3</sub> volatilization in the lower-salinity soils accounted for 0.75 to 2.53% of the nitrogen applied in the  
310 incubation experiment. This result is similar to those reported by Zhou et al. (2016), who found that 3.2 to 3.8%  
311 of nitrogen was lost via NH<sub>3</sub> volatilization. Similar to the urea application method with drip fertigation employed  
312 by these researchers, we added urea in solution form. When the urea solution was added to the soils, it infiltrated  
313 into the soils quickly and reached the deeper layer of the soils, consequently minimizing NH<sub>3</sub> volatilization  
314 (Haynes, 1985).

315 In the higher-salinity soils (HS and SS), the NH<sub>3</sub> volatilization pattern was different from the conventional

316 short-duration NH<sub>3</sub> volatilization pattern found in most previous studies. The NH<sub>3</sub> volatilization period was  
317 significantly longer than that in lower-salinity soils. The peak of NH<sub>3</sub> volatilization was delayed due to the  
318 inhibition of urea hydrolysis by soil salinity (Kumar et al., 1988). When NH<sub>3</sub> volatilization reached its peak, it  
319 was maintained for a longer time than in the lower-salinity soils and then decreased slowly until NH<sub>3</sub>  
320 volatilization was complete (Fig. 1a). NH<sub>3</sub> volatilization is directly determined by the ammonium concentration  
321 in soil (Bosch-Serra et al. 2014). Ammonium is the intermediate product of N transformation, which connects  
322 urea hydrolysis and nitrification. We determined the NH<sub>4</sub><sup>+</sup>-N and NO<sub>3</sub><sup>-</sup>-N concentrations in the soils (Fig. 4) and  
323 found that the NH<sub>4</sub><sup>+</sup>-N concentration was still high from the third to eighth week relative to that in the lower-  
324 salinity soils, which contained no NH<sub>4</sub><sup>+</sup>-N at the end of the second week. Additionally, the NO<sub>3</sub><sup>-</sup>-N concentration  
325 was always low in the first 5 weeks (Fig. 4). The results proved that nitrification was significantly inhibited by  
326 soil salinity, as indicated by Ye et al. (2009). Similar results were found in the lower-salinity soils; the measured  
327 NH<sub>4</sub><sup>+</sup>-N and NO<sub>3</sub><sup>-</sup>-N concentrations demonstrated that nitrification was inhibited by soil salinity, as the NH<sub>4</sub><sup>+</sup>-N  
328 concentration increased with the increase in soil salinity, while the NO<sub>3</sub><sup>-</sup>-N concentration was inversely related  
329 to soil salinity in the first week (Fig. 4). Overall, we may conclude that soil salinity aggravates NH<sub>3</sub> volatilization  
330 by inhibiting nitrification and affecting soil properties related to NH<sub>3</sub> volatilization.

331 **4.2. Effect of N sources on NH<sub>3</sub> volatilization**

332 In coastal saline farmland, manure is widely used as a soil amendment; MAP is usually selected as the  
333 basic fertilizer; and urea is usually used as an additional fertilizer. In our study, the cumulative NH<sub>3</sub> volatilization  
334 was highest for urea, which was consistent with the findings of Mandal et al. (2016), who inferred that urea  
335 hydrolysis increased the soil pH and thereby aggravated NH<sub>3</sub> volatilization. The NH<sub>4</sub><sup>+</sup>-N concentration in urea-  
336 treated soil was always lower than that in MAP-treated soil (Fig. 5), although urea-treated soil still showed the  
337 highest cumulative NH<sub>3</sub> volatilization, which may confirm the existence of inference. Some differences also

338 existed. The difference in cumulative  $\text{NH}_3$  volatilization between MAP-treated soil and manure-treated soil was  
339 inconsistent in the study by Mandal et al. (2016). We found that the cumulative  $\text{NH}_3$  volatilization in manure-  
340 treated soil was always lower than that in MAP-treated soil. The nitrogen in manure is mainly organic nitrogen  
341 and is released slowly by mineralization. The released  $\text{NH}_4^+$ -N will be nitrated to  $\text{NO}_3^-$ -N quickly and thus result  
342 in a low  $\text{NH}_4^+$ -N concentration, which can be indicated by the  $\text{NO}_3^-$ -N and  $\text{NH}_4^+$ -N concentrations in Fig. 5.

343 **4.3. Interactive effect of salinity and biochar on  $\text{NH}_3$  volatilization**

344 Biochar is one of the most popular soil amendments used for soil improvement and nutrient utilization.  
345 Recently, biochar has also been widely used in saline soil to alleviate the negative effect of salinity on soil  
346 properties and plant growth (Thomas et al., 2013). However, its effect shows high dependence on the  
347 experimental conditions and its own agronomic properties (Liu et al., 2013; Cely et al., 2015). With respect to  
348 the effect of biochar on  $\text{NH}_3$  volatilization, the results are also inconsistent under different experimental  
349 conditions. In our study, two soils (NS and MS) and two biochar application rates were selected to measure their  
350 interactive effect on  $\text{NH}_3$  volatilization.

351 In the non-saline soil, the difference in cumulative  $\text{NH}_3$  volatilization between biochar-amended soils and  
352 the control was not significant, regardless of whether the biochar was applied at a high level (2%) or a low level  
353 (1%) (Fig. 3). These findings were partially consistent with those of Feng et al. (2017), who found that the  
354 difference in  $\text{NH}_3$  volatilization between the control and treatment at a 0.5% biochar application rate was not  
355 significant; however,  $\text{NH}_3$  volatilization was significantly greater than in the control under a biochar application  
356 rate of 3% in paddy soil. Chen et al. (2013) indicated that biochar affected  $\text{NH}_3$  volatilization via two properties.  
357 The first is the adsorption capacity of biochar, which is determined by the biochar BET surface and CEC,  
358 between which the BET surface is the dominant factor related to the adsorption of  $\text{NH}_3$  (Taghizadeh-Toosi et  
359 al., 2012b). The second property is the pH of biochar, which might affect the soil pH. Therefore, it is likely that

360 the effects of the two properties were nearly balanced when biochar was applied at a 0.5% rate and that pH  
361 increases played a major role when biochar was applied at a 3% rate (Subedi et al., 2015; Sun et al., 2016).  
362 Mandal et al. (2016) found that NH<sub>3</sub> volatilization was alleviated under biochar application, which was in  
363 agreement with the findings of Malinska et al. (2014), who indicated that high-pH biochar reduced NH<sub>3</sub>  
364 volatilization, which could be attributed to its high sorption capacity. Interestingly, we obtained a finding that  
365 differed from those of these previous studies: soil pH did not increase with biochar addition and was even lower  
366 than in the control (Fig. 6), which was similar to the results of Cely et al. (2014), who found that the addition of  
367 biochar with different pH levels did not change the soil pH significantly, regardless of whether the biochar pH  
368 was higher than or close to the soil pH. Yang (2015) also reported that alkaline biochar could reduce the pH of  
369 soda-saline soil to some extent. Feng et al. (2017) indicated that the effect of biochar addition on nitrification  
370 also affected NH<sub>3</sub> volatilization and that the inhibition of nitrification by biochar resulted in the accumulation  
371 of ammonium. Biochar addition also increased soil EC (Fig. S1), as also reported by Cely et al. (2014), which  
372 would exacerbate the inhibition of nitrification. In our study, pH was not the driving factor, and the effects of  
373 the adsorption capacity and nitrification inhibition were nearly balanced. Therefore, the effect of biochar on  
374 NH<sub>3</sub> volatilization was not significant in non-saline soil.

375 In moderately saline soil, the cumulative NH<sub>3</sub> volatilization in high-biochar-treated soil was significantly  
376 greater than that in the control, while the cumulative NH<sub>3</sub> volatilization in low-biochar-treated soil was slightly  
377 higher than that in the control, although the difference was not significant. This result was in agreement with  
378 those reported by Sun et al. (2017), who conducted an experiment in coastal saline soil and found that the  
379 difference in NH<sub>3</sub> volatilization between the control and treatment at a 0.5% or 1% biochar application rate was  
380 not significant; however, NH<sub>3</sub> volatilization was significantly greater than in the control when the biochar  
381 application rate was greater than 2%. Similar to non-saline soil, biochar addition did not increase the soil pH

382 (Fig. 6), while biochar addition significantly increased soil EC in moderately saline soil (Fig. S1). Nitrification  
383 was also measured in the moderately saline soil. In contrast to the rapid nitrification process observed in non-  
384 saline soil, the  $\text{NH}_4^+$ -N concentration in biochar-amended soils was higher than that in the control in the first  
385 two weeks, and the  $\text{NO}_3^-$ -N concentration in biochar-amended soils was lower than that in the control throughout  
386 the incubation period (Table 3). The results proved that nitrification was inhibited in soils subjected to biochar  
387 addition, which is consistent with Feng et al. (2017). Ameloot et al. (2014) showed that biochar addition  
388 restricted soil microbial activity, and Wang et al. (2015) found that biochar addition reduced the abundance of  
389 ammonia-oxidizing bacteria and thereby inhibited nitrification. The inhibitory effect of biochar addition on  
390 nitrification may have become stronger due to the interactive effect of biochar and salinity in moderately saline  
391 soil. With respect to the adsorption capacity, the salt ion concentration in moderately saline soil was significantly  
392 higher than that in non-saline soil, and the adsorption of salt ions decreases the  $\text{NH}_3/\text{NH}_4^+$  adsorption capacity.  
393 Therefore, nitrification inhibition played an important role between the effects of the adsorption capacity and  
394 nitrification inhibition in moderately saline soil, and biochar addition therefore aggravated  $\text{NH}_3$  volatilization  
395 in moderately saline soil. Regarding the difference in  $\text{NH}_3$  volatilization between BCH and BCL, it was likely  
396 that nitrification inhibition was significantly increased with higher biochar addition; however, the rate of the  
397 increase in the adsorption capacity was lower than that for nitrification inhibition. Overall,  $\text{NH}_3$  volatilization  
398 increased with the biochar application rate in moderately saline soil, whereas a low rate could prevent the  
399 increase in  $\text{NH}_3$  volatilization, so biochar application at a low rate is suggested.

400 **4.4. Interactive effect of salinity and gypsum on  $\text{NH}_3$  volatilization**

401 Gypsum is a traditional amendment used for the amelioration of saline soil. In our study, we selected two  
402 application rates for the two soils. The cumulative  $\text{NH}_3$  volatilization in soils treated with gypsum was  
403 significantly lower than that in the control in non-saline soil. A similar result was found in moderately saline

404 soil, although the difference was not significant (Fig. 3). This result was consistent with those of Zia et al. (1999),  
405 who demonstrated that gypsum addition could reduce NH<sub>3</sub> volatilization due to the effect of decreasing pH. A  
406 similar result was indicated by Burt et al. (2017) in composting. The pH mainly exerts its effect by influencing  
407 the equilibrium of NH<sub>4</sub><sup>+</sup> and NH<sub>3</sub>. A decreased pH reduces the transformation of NH<sub>4</sub><sup>+</sup> to NH<sub>3</sub> and thus reduces  
408 NH<sub>3</sub> volatilization. In our study, the pH in soils amended with gypsum was significantly lower than that in the  
409 control (Fig. 6), which was in agreement with this previous study. We also found that the pH decreased more  
410 significantly with more gypsum addition.

411 Nitrification is another important process influencing NH<sub>3</sub> volatilization that is affected by gypsum. In our  
412 study, gypsum addition significantly increased soil EC (Fig. S1), which was confirmed by Burt et al. (2017) and  
413 Tubail et al. (2008), who attributed this effect to the increase in sulfate. An increase in the soil salt content will  
414 aggravate the inhibition of nitrification. Ponette et al. (1996) found that nitrification was depressed by gypsum  
415 application, and Li et al. (2016) reported that the copy number of nitrification functional genes was significantly  
416 decreased by gypsum addition. The lower NO<sub>3</sub><sup>-</sup>-N and higher NH<sub>4</sub><sup>+</sup>-N concentrations found in soils with gypsum  
417 addition (Table 2 and Table 3) also indicated that nitrification was inhibited in our study. Furthermore, the  
418 inhibition was more significant under higher gypsum addition.

419 Overall, gypsum influenced both pH and nitrification, and its effect on NH<sub>3</sub> volatilization depended on the  
420 interaction between the decreasing pH and the inhibition of nitrification. Which of these factors is dominant in  
421 such situations will sometimes lead to different results. In the non-saline soil, the cumulative NH<sub>3</sub> volatilization  
422 in the soil with added gypsum was significantly lower than that in the control, and we might conclude that the  
423 effect of decreasing pH was dominant because nitrification occurred for a short duration (one week). In the  
424 moderately saline soil, the cumulative NH<sub>3</sub> volatilization in the soil with added gypsum was also lower than  
425 that in the control, although the difference was not significant ( $P=0.54$  for both SGH and SGL relative to the

control). This might have occurred because the NH<sub>3</sub> volatilization period grew longer due to the inhibition of nitrification by salinity; the inhibitory effect of gypsum addition on nitrification may have become stronger due to their interactive effect; and the effect of decreasing pH on NH<sub>3</sub> volatilization was alleviated compared with that in non-saline soil. Therefore, the difference between the soil with gypsum addition and the control was not significant in moderately saline soil. We also found that the difference in NH<sub>3</sub> volatilization between SGH and SGL was not significant; it was likely that because gypsum affected both the decrease in pH and the inhibition of nitrification, even though more gypsum addition could decrease the soil pH more significantly, its inhibitory effect on nitrification was also significantly increased. Thus, the two effects were still nearly balanced regardless of whether gypsum was added at a high or low rate. Overall, gypsum decreased NH<sub>3</sub> volatilization, although the contribution of different gypsum rates to NH<sub>3</sub> volatilization showed no difference, so gypsum application at a low rate is suggested.

## 5. Conclusions

The results clearly indicated that soil salinity significantly exacerbated cumulative NH<sub>3</sub> volatilization and even changed the NH<sub>3</sub> volatilization pattern when soil salinity exceeded a certain threshold. These effects were mainly ascribed to the accumulation of ammonium, which resulted from the inhibition of nitrification by salinity, and also the effect of salinity on soil properties related to NH<sub>3</sub> volatilization. Among the tested N sources, urea showed the highest cumulative NH<sub>3</sub> volatilization, followed by MAP and then manure in soils with three different salinity levels. The reason for these results lies in the fact that urea hydrolysis increases soil pH. Biochar addition had no significant influence on NH<sub>3</sub> volatilization in non-saline soil because the adsorption capacity and nitrification inhibition were in balance. Biochar application increased NH<sub>3</sub> volatilization in moderately saline soil because salt ions decreased the NH<sub>3</sub>/NH<sub>4</sub><sup>+</sup> adsorption capacity of biochar, and the interactive effect of biochar and salinity exacerbated the inhibition of nitrification. Overall, NH<sub>3</sub> volatilization

448 increased with the biochar application rate in saline soil, and a low biochar application rate could prevent the  
449 increase of NH<sub>3</sub> volatilization, so biochar addition at a low rate is suggested. Gypsum application significantly  
450 decreased NH<sub>3</sub> volatilization in non-saline soil by decreasing the soil pH. However, the differences were not  
451 significant in moderately saline soil due to the interactive effect of gypsum and salinity aggravating the  
452 inhibition of nitrification. Overall, gypsum decreased NH<sub>3</sub> volatilization in saline soil, whereas the contribution  
453 of different gypsum rates to NH<sub>3</sub> volatilization showed no difference. A conclusion could be drawn from the  
454 above results that low rates of biochar and gypsum application should be suggested to prevent or reduce NH<sub>3</sub>  
455 volatilization and nitrogen losses in coastal saline soil.

## 456 **Conflict of interest**

457 The authors declare that there is no conflict of interests regarding the publication of this paper.

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## Figure caption list

Fig.1. The rate of NH<sub>3</sub> volatilization (a), cumulative NH<sub>3</sub> volatilization (b) in different saline soils and the correlation relationship between soil salinity and cumulative NH<sub>3</sub> volatilization (c). NS = non-saline soil, LS = low-salinity soil, MS = moderately saline soil, HS = high-salinity soil and SS = severely saline soil. Error bars denote the standard deviation for three replications. Different letters denote significant difference ( $P < 0.05$ ) between treatments

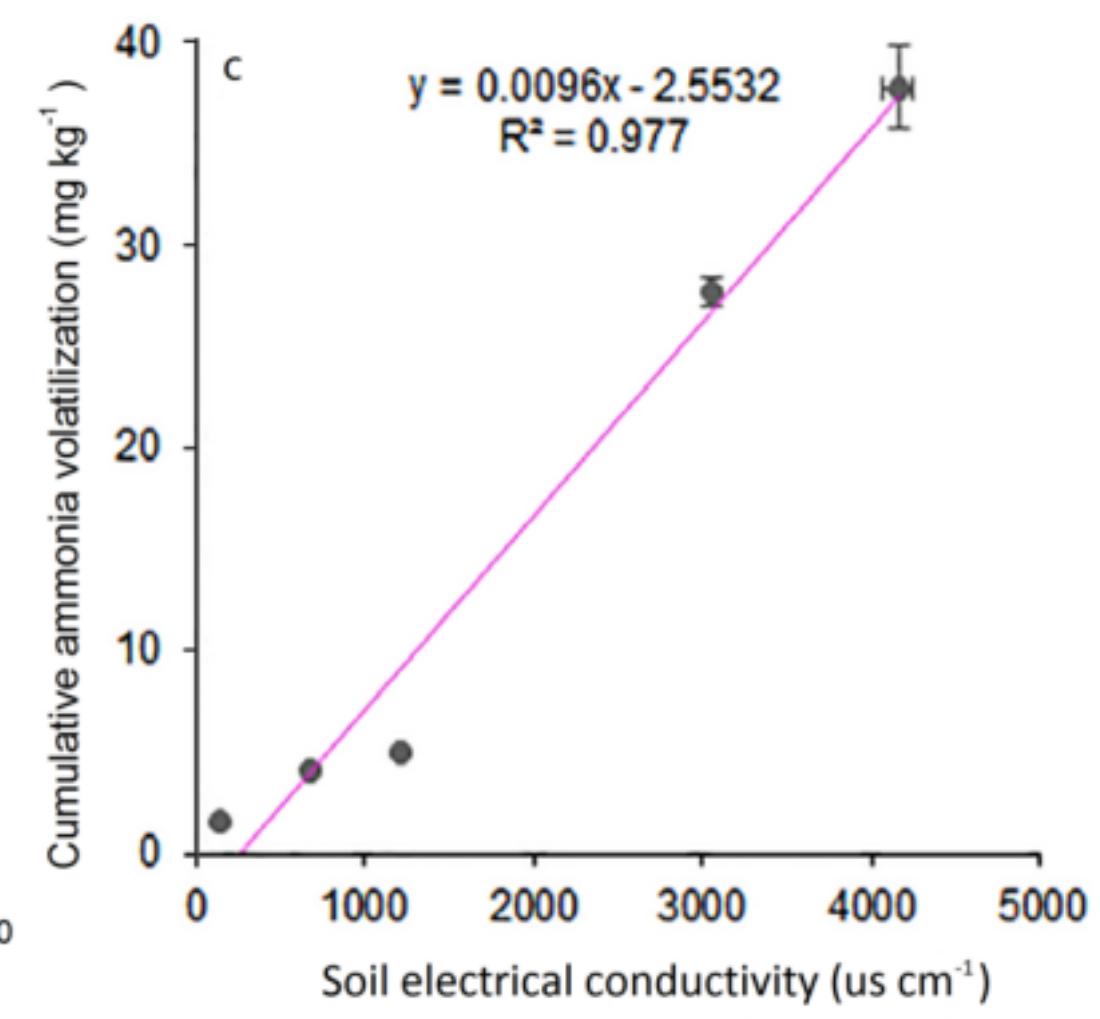
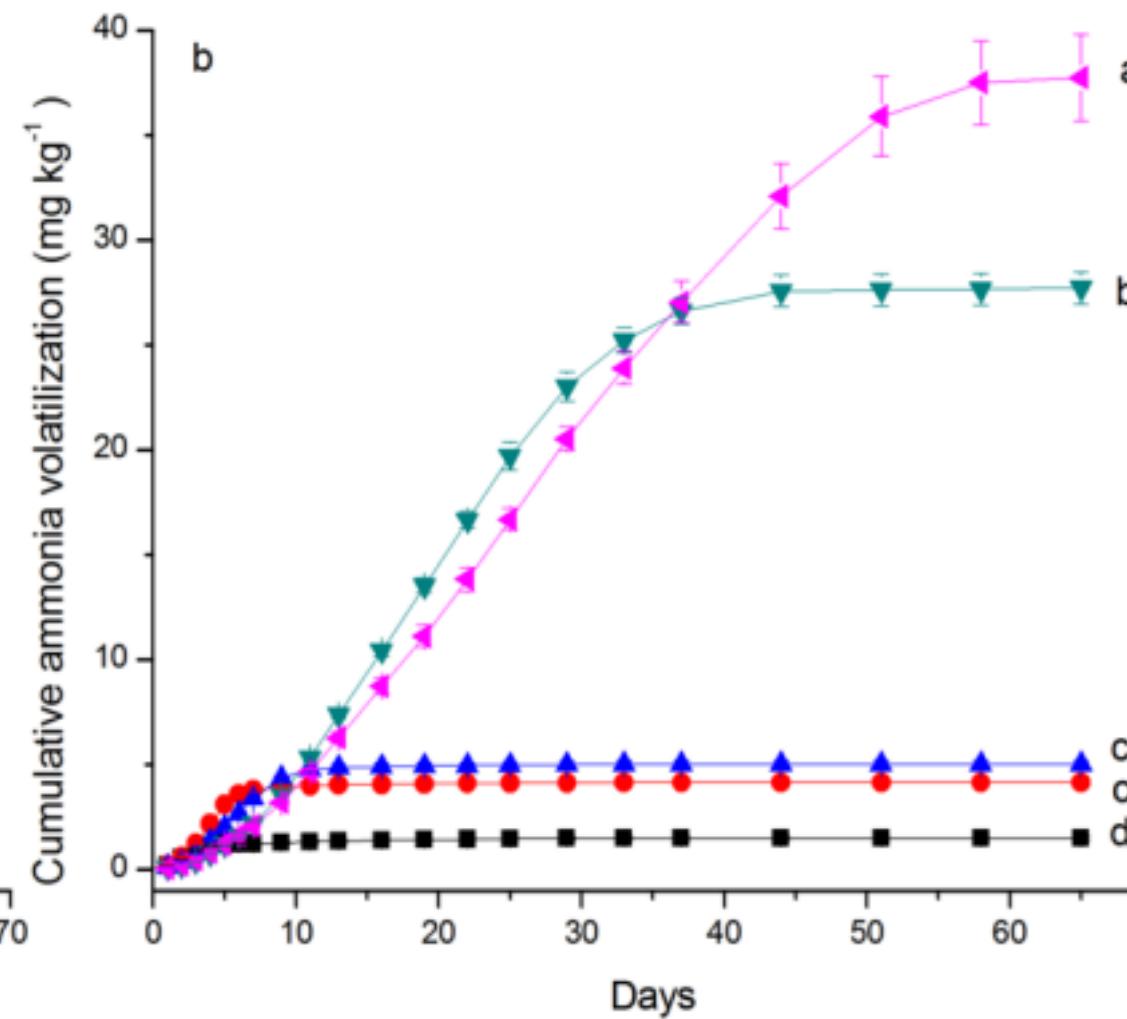
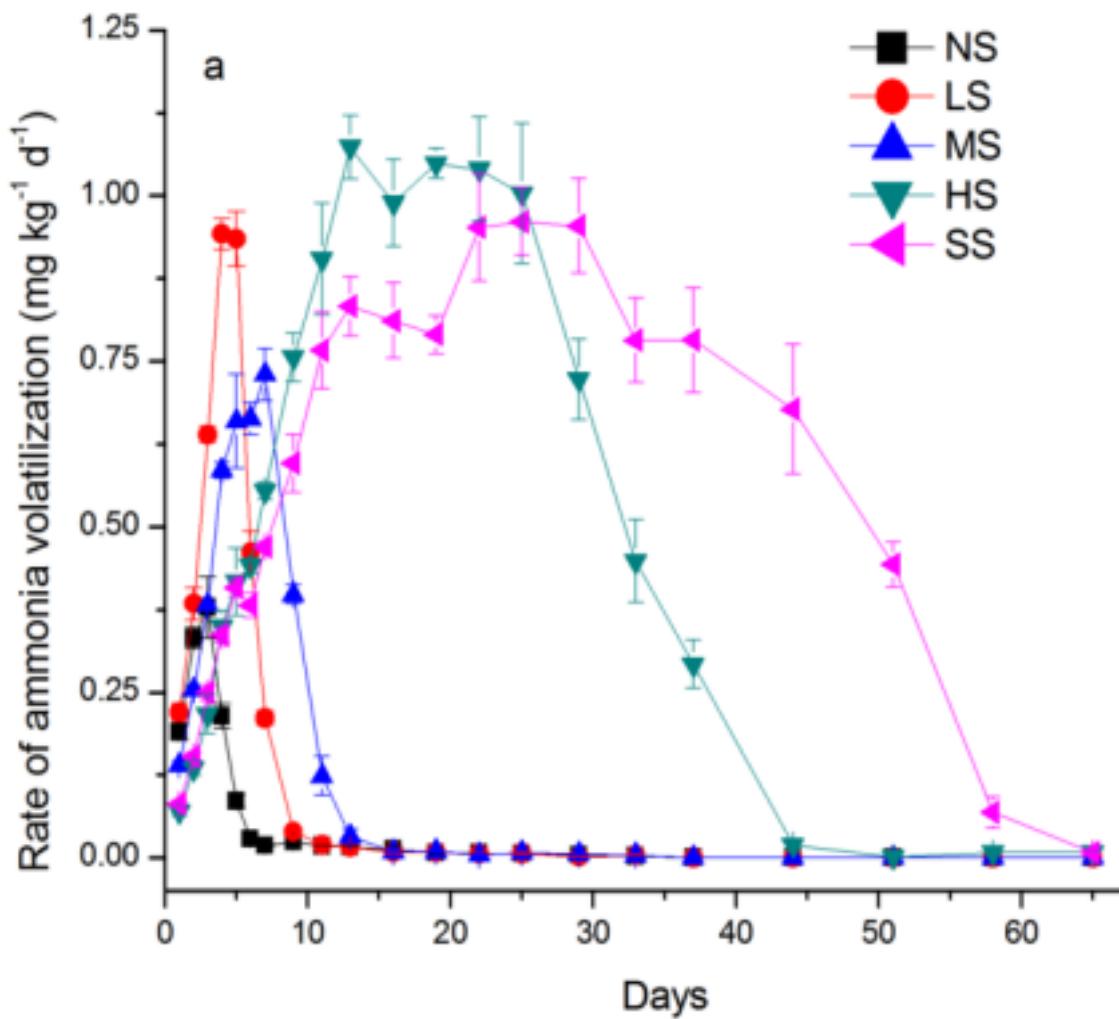
Fig.2. The NH<sub>3</sub> volatilization rate and cumulative NH<sub>3</sub> volatilization of different N sources in non-saline soil (a), low-salinity soil (b) and moderately saline soil (c). Error bars denote the standard deviation for three replications. Different letters denote significant difference ( $P < 0.05$ ) between treatments

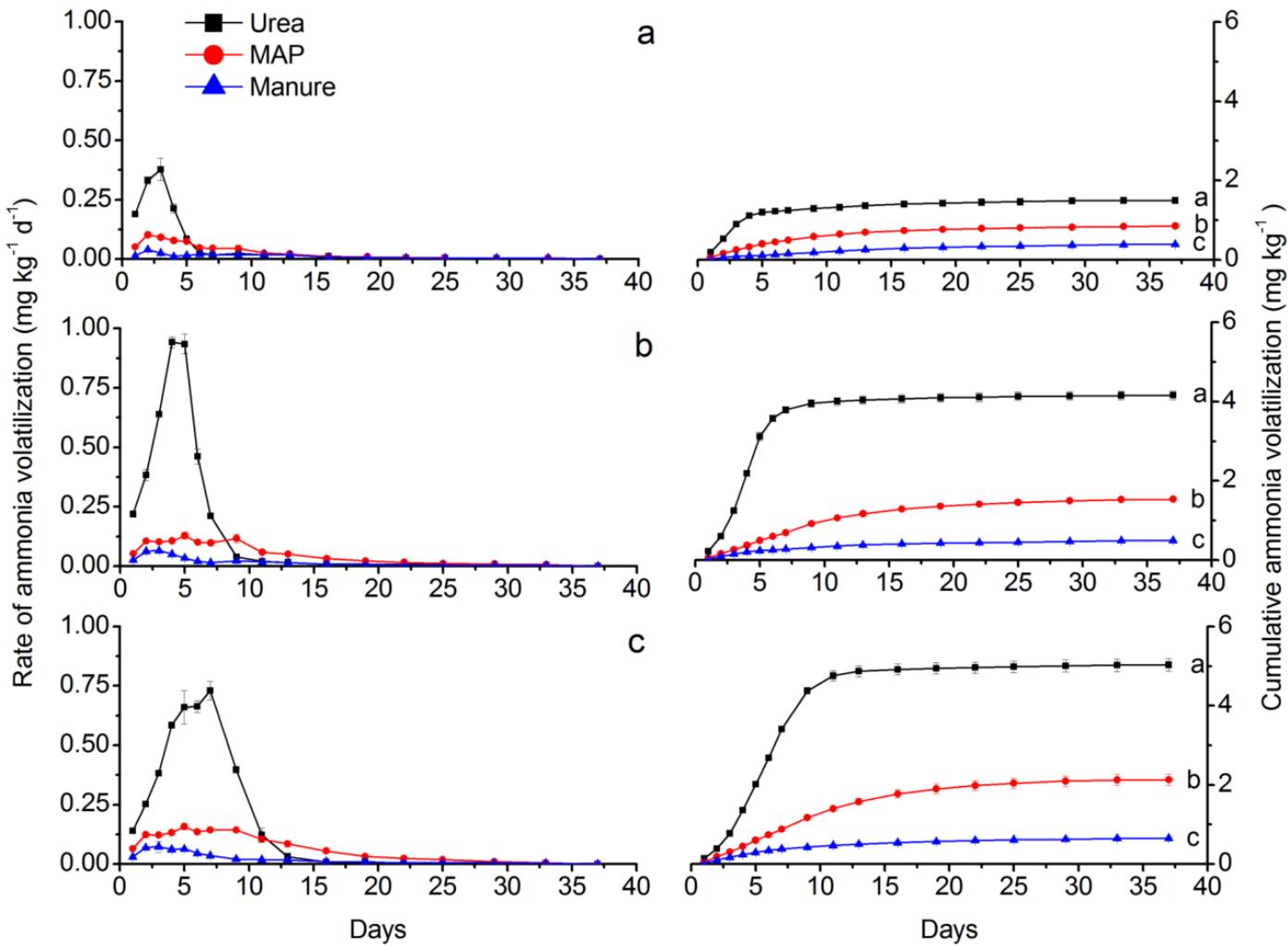
Fig.3. The cumulative NH<sub>3</sub> volatilization of non-saline soil (a) and moderately saline soil (b) added with amendments. BCH = high biochar addition, BCL = low biochar addition, SGH = high gypsum addition, SGL = low gypsum addition, CK = soil without amendments addition. Error bars denote the standard deviation for three replications. Different letters denote significant difference ( $P < 0.05$ ) between treatments.

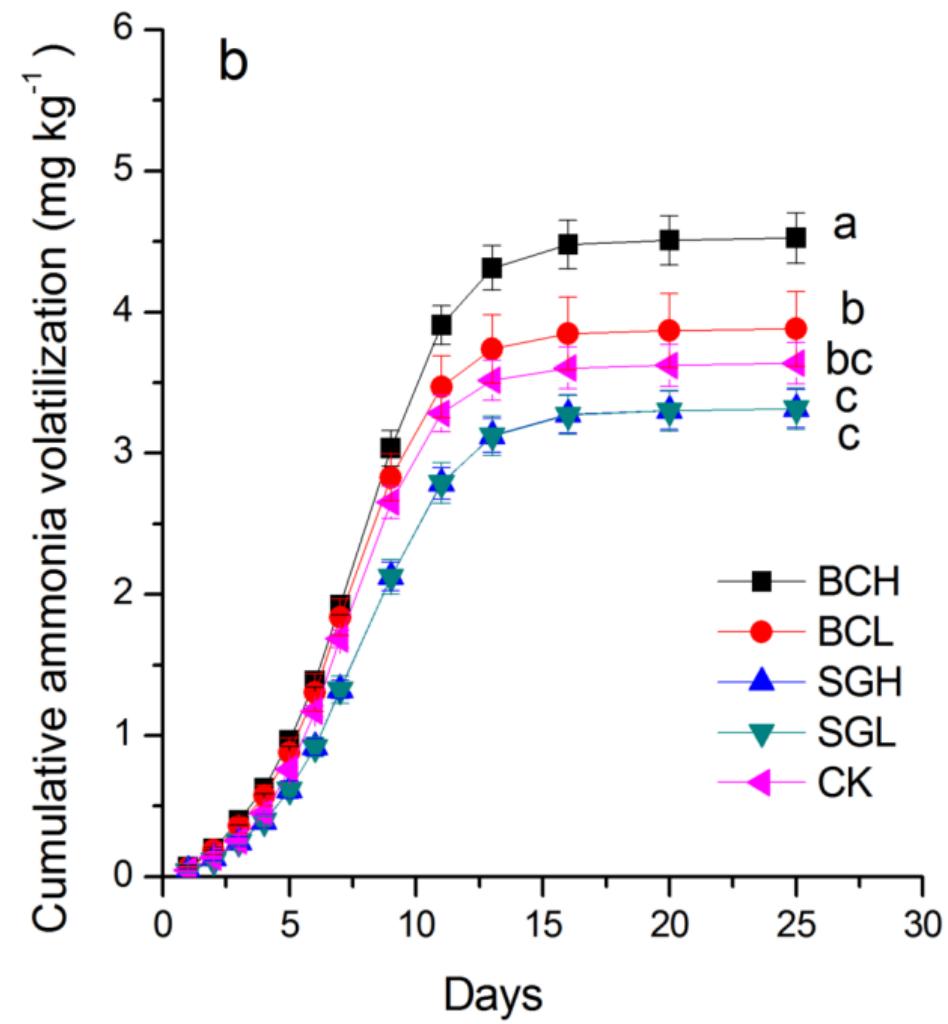
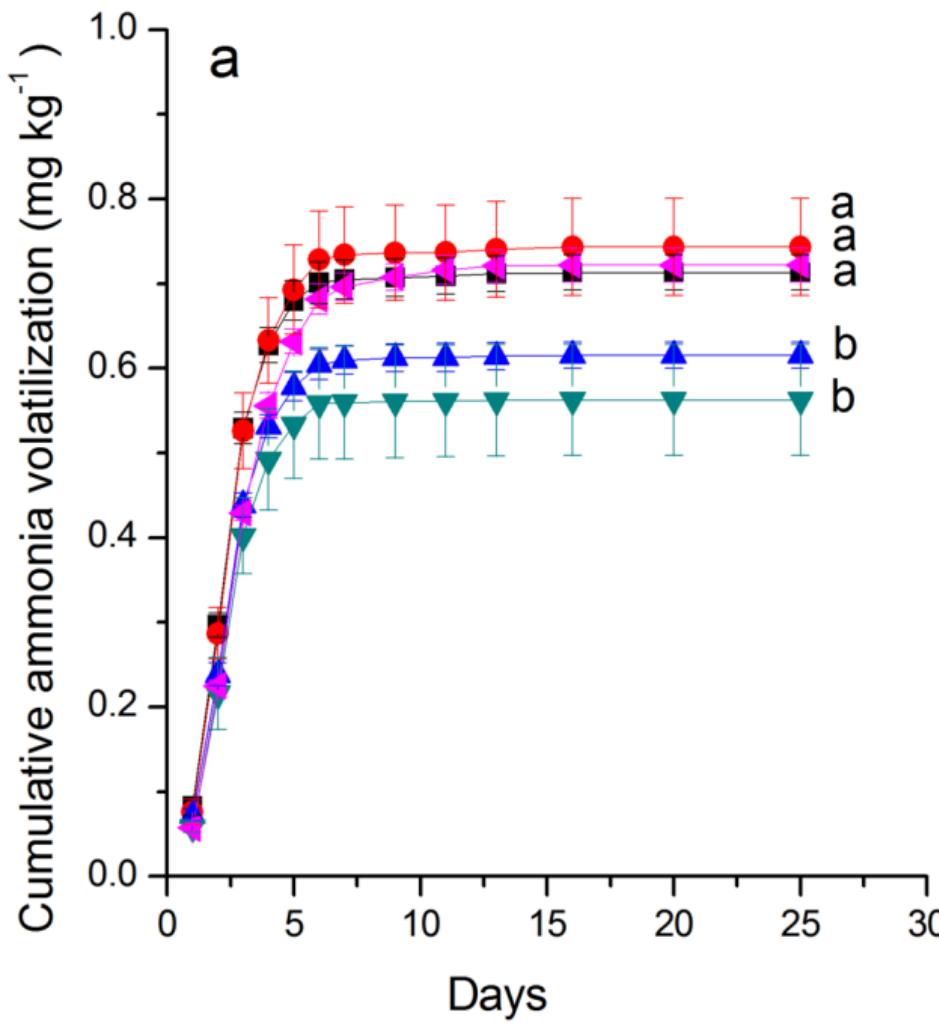
Fig.4. The NO<sub>3</sub><sup>-</sup>-N (a) and NH<sub>4</sub><sup>+</sup>-N (b) concentration in different saline soils. NS = non-saline soil, LS = low-salinity soil, MS = moderately saline soil, HS = high-salinity soil and SS = severely saline soil. Error bars denote the standard deviation for three replications.

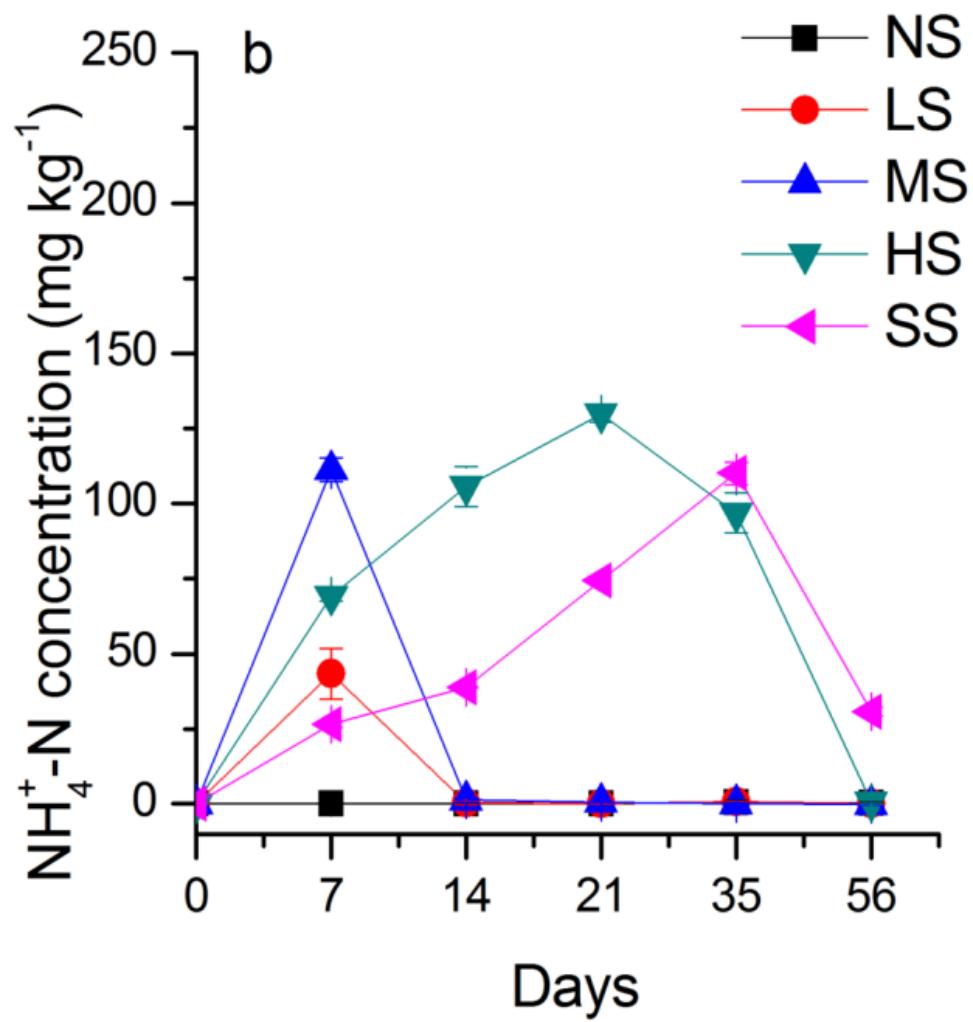
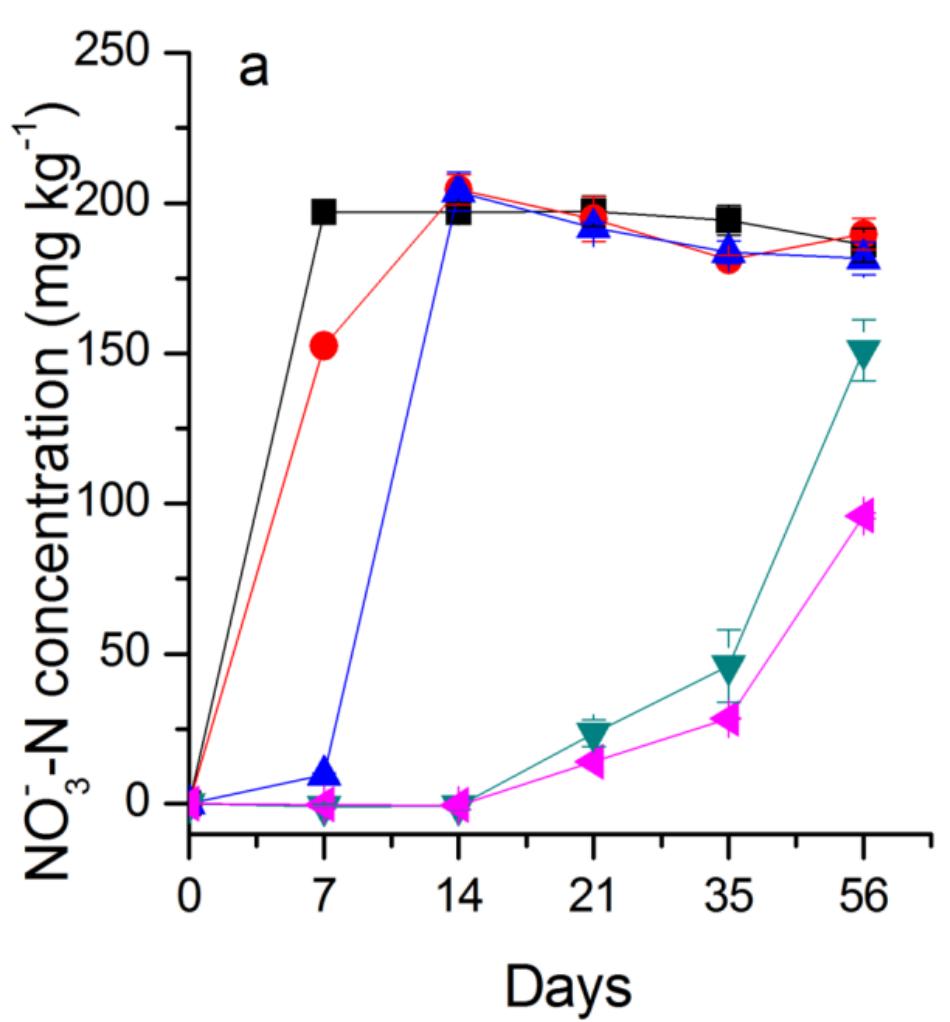
Fig.5. The NO<sub>3</sub><sup>-</sup>-N and NH<sub>4</sub><sup>+</sup>-N concentration of different N sources in non-saline soil (a), low-salinity soil (b) and moderately saline soil (c). Error bars denote the standard deviation for three replications.

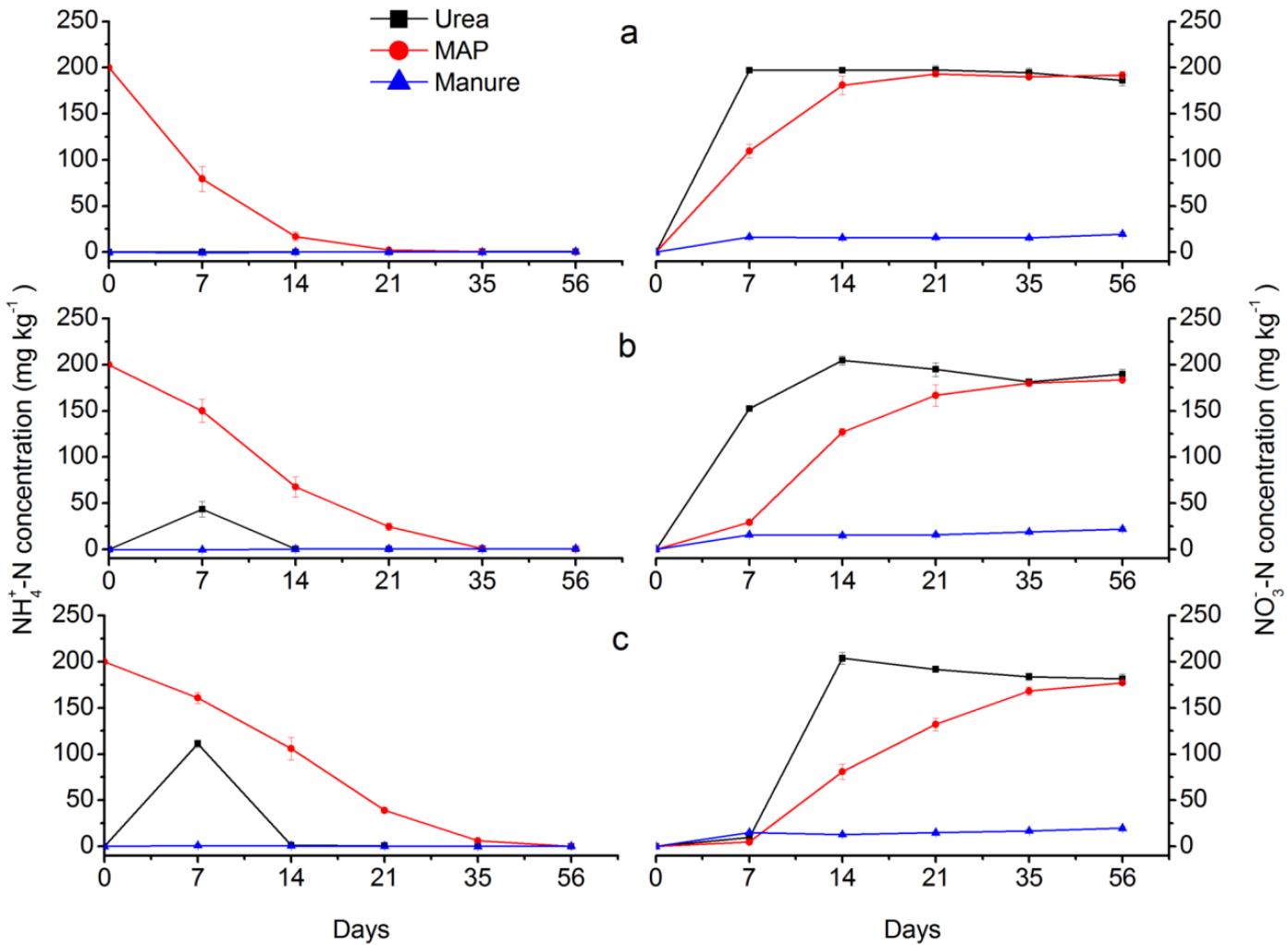
Fig.6. The pH of non-saline soil (a) and moderately saline soil (b) added with amendments after the incubation experiment. BCH = high biochar addition, BCL = low biochar addition, SGH = high gypsum addition, SGL = low gypsum addition, CK = soil without amendments addition. Error bars denote the standard deviation for three replications. Different letters denote significant difference ( $P < 0.05$ ) between treatments.

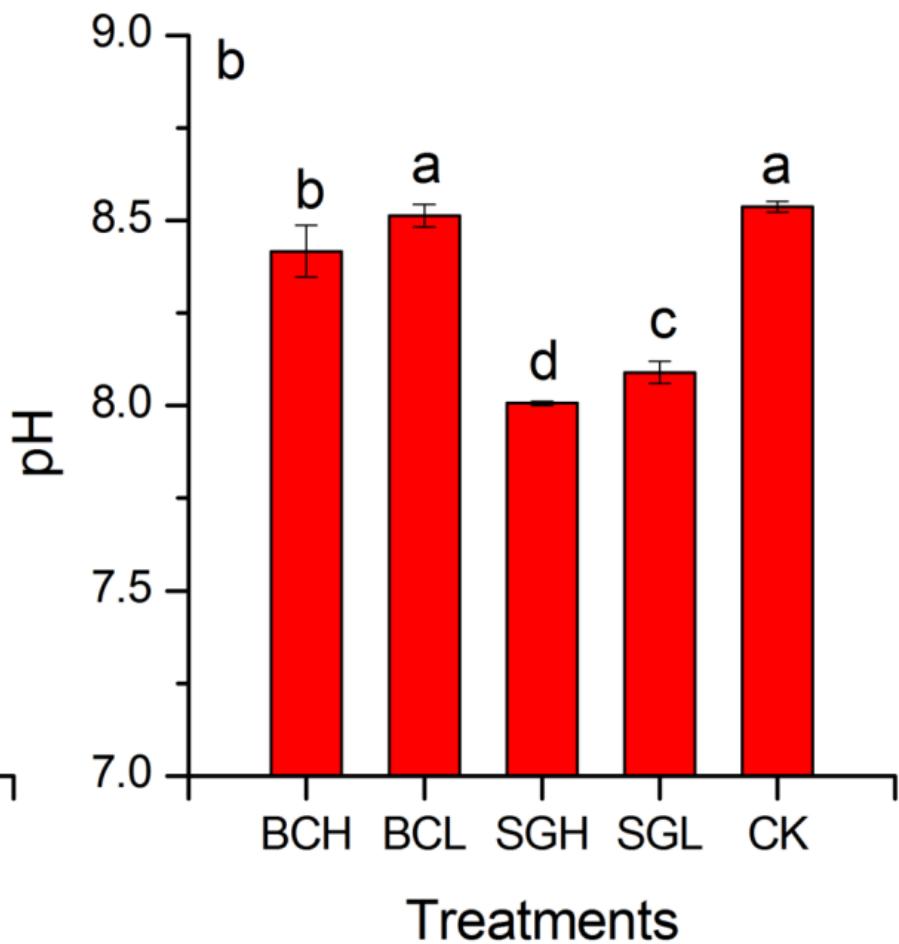
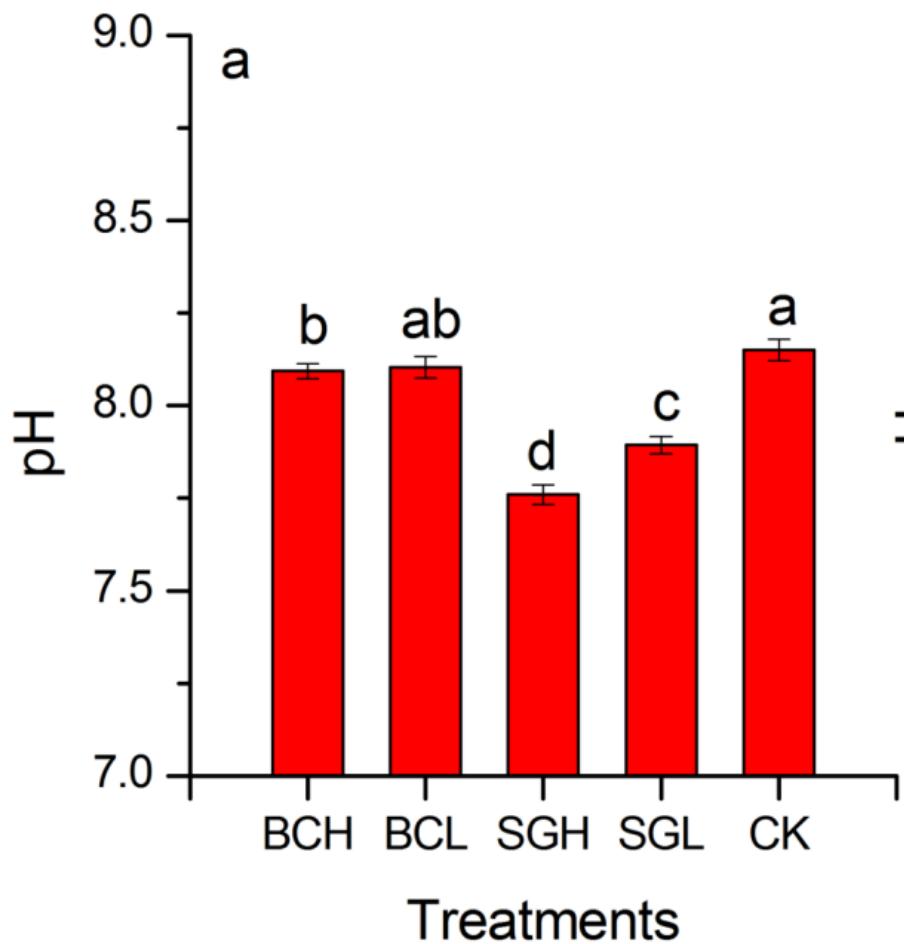












**Table 1** Physicochemical characteristics of soils, biochar and gypsum.

|    | Sand (%)   | Silt (%)   | Clay (%)  | EC (μs cm <sup>-1</sup> ) | pH        | Total N (g kg <sup>-1</sup> ) | TOC (g kg <sup>-1</sup> ) | NH <sub>4</sub> <sup>+</sup> -N (mg kg <sup>-1</sup> ) | NO <sub>3</sub> <sup>-</sup> -N (mg kg <sup>-1</sup> ) | CEC (cmol kg <sup>-1</sup> ) | ES (cmol kg <sup>-1</sup> ) | BET surface (m <sup>2</sup> g <sup>-1</sup> ) | PV (cm <sup>3</sup> g <sup>-1</sup> ) | VM (%) | Ash (%) | FC (%) |
|----|------------|------------|-----------|---------------------------|-----------|-------------------------------|---------------------------|--|--|------------------------------|-----------------------------|---|---------------------------------------|--------|---------|--------|
| NS | 30.73±1.59 | 62.24±1.68 | 7.03±0.55 | 147.90±7.78               | 8.57±0.08 | 0.31±0.01                     | 3.48±0.03                 | 17.79±0.44   | 5.39±0.19  | 6.23±0.31                    | 0.06±0.03                   |   |                                       |        |         |        |
| LS | 31.95±1.08 | 61.60±1.72 | 6.45±0.83 | 686.70±16.12              | 8.88±0.01 | 0.24±0.02                     | 2.69±0.11                 | 7.09±0.30  | 4.63±0.13  | 5.69±0.15                    | 0.25±0.03                   |   |                                       |        |         |        |
| MS | 31.10±2.67 | 61.18±1.35 | 7.72±1.33 | 1215.50±17.68             | 8.97±0.02 | 0.22±0.01                     | 2.43±0.03                 | 16.75±0.16   | 4.85±0.06  | 5.79±0.15                    | 0.51±0.03                   |   |                                       |        |         |        |
| HS | 30.06±1.78 | 61.81±0.97 | 8.13±0.86 | 3048.50±14.85             | 8.45±0.04 | 0.22±0.01                     | 2.65±0.08                 | 25.90±0.19   | 5.17±0.38  | 5.76±0.20                    | 0.49±0.07                   |   |                                       |        |         |        |
| SS | 33.30±1.43 | 59.48±0.97 | 7.21±0.89 | 4155.50±91.22             | 8.42±0.01 | 0.19±0.01                     | 1.90±0.10                 | 50.89±0.47   | 4.94±0.32  | 4.78±0.12                    | 0.43±0.04                   |   |                                       |        |         |        |
| SG | —          | —          | —         | 27160±1.41                | 6.36±0.17 | 0.06±0.00                     | 0.56±0.02                 | 1.34±0.02  | 7.84±0.24  | —                            | —                           |   |                                       |        |         |        |
| BC | —          | —          | —         | 7581.50±81.32             | 9.13±0.05 | 6.64±0.12                     | —                         | 35.01±1.65   | 2.40±0.09  | 46.60±3.72                   | 0.67±0.03                   | 20.968  | 0.025                                 | 13.71  | 19.52   | 66.77  |

Note: Data were mean ± standard deviation (n = 3), NS = non-saline soil, LS = low-salinity soil, MS = moderately saline soil, HS = high-salinity soil and SS = severely saline soil, SG=gypsum, BC=biochar, PV = pore volume, ES=.exchangeable sodium,

VM=volatile matter, FC=fixed carbon.

**Table 2** The  $\text{NO}_3^-$ -N and  $\text{NH}_4^+$ -N concentration of non-saline soil added with biochar and gypsum.

| Days | NO <sub>3</sub> <sup>-</sup> -N concentration (mg kg <sup>-1</sup> ) |             |             |             |             | NH <sub>4</sub> <sup>+</sup> -N concentration (mg kg <sup>-1</sup> ) |            |            |            |            |
|------|--|-------------|-------------|-------------|-------------|--|------------|------------|------------|------------|
|      | BCH  | BCL         | SGH         | SGL         | CK          | BCH  | BCL        | SGH        | SGL        | CK         |
| 3    | 64.79±1.28   | 61.51±1.67  | 53.90±4.95  | 66.86±0.99  | 78.73±1.85  | 17.49±1.35   | 17.99±0.50 | 19.25±0.54 | 18.28±0.23 | 18.32±0.58 |
| 7    | 221.79±14.53   | 220.40±9.98 | 228.50±5.63 | 235.36±9.05 | 235.06±1.01 | 2.54±0.11  | 3.21±0.58  | 2.35±0.23  | 2.43±0.20  | 2.33±0.24  |
| 14   | 207.54±5.23  | 213.56±8.13 | 220.10±1.12 | 222.90±3.68 | 218.35±5.79 | 1.53±0.44  | 1.72±0.00  | 1.92±0.34  | 1.62±0.22  | 1.83±0.11  |
| 21   | 206.73±3.39  | 212.50±1.93 | 228.12±4.79 | 222.97±2.97 | 212.67±4.82 | 2.02±0.40  | 2.02±0.40  | 1.80±0.22  | 1.58±0.16  | 1.65±0.41  |
| 35   | 208.16±10.02   | 208.94±4.16 | 226.20±6.87 | 226.13±8.13 | 224.77±6.09 | 2.04±0.55  | 2.07±0.11  | 2.04±0.19  | 1.70±0.13  | 2.18±0.22  |

Note: BCH = high biochar addition, BCL = low biochar addition, SGH = high gypsum addition, SGL = low gypsum addition, CK = soil without amendment

addition. Data in the table are average values and standard deviations (n=3).

**Table 3** The  $\text{NO}_3^-$ -N and  $\text{NH}_4^+$ -N concentration of moderately saline soil added with biochar and gypsum.

| Days | NO <sub>3</sub> <sup>-</sup> -N concentration (mg kg <sup>-1</sup> ) |             |             |             |             | NH <sub>4</sub> <sup>+</sup> -N concentration (mg kg <sup>-1</sup> ) |             |             |             |              |
|------|--|-------------|-------------|-------------|-------------|--|-------------|-------------|-------------|--------------|
|      | BCH  | BCL         | SGH         | SGL         | CKU         | BCH  | BCL         | SGH         | SGL         | CKU          |
| 3    | 15.36±0.22   | 16.06±0.15  | 16.6±0.18   | 16.61±0.18  | 17.43±0.08  | 12.87±0.08   | 12.07±0.17  | 11.95±0.18  | 11.47±0.12  | 12.99±0.15   |
| 7    | 28.94±0.99   | 30.52±0.53  | 27.05±1.08  | 28.28±1.37  | 33.60±0.42  | 117.57±4.38  | 117.21±5.57 | 109.66±0.54 | 105.26±3.55 | 102.39±13.59 |
| 14   | 194.65±1.22  | 204.13±8.09 | 173.52±7.43 | 181.09±5.71 | 222.71±3.14 | 17.27±4.33   | 13.51±2.30  | 41.87±5.86  | 31.67±6.74  | 10.57±3.23   |
| 21   | 206.27±3.51  | 214.09±3.72 | 217.23±4.62 | 225.13±3.65 | 223.33±4.81 | 1.94±0.29  | 1.81±0.24   | 2.62±0.31   | 2.43±0.23   | 2.16±0.21    |
| 35   | 186.27±3.11  | 203.80±2.99 | 200.15±3.58 | 203.99±9.00 | 208.17±5.08 | 2.11±0.08  | 2.20±0.23   | 1.98±0.14   | 2.46±0.21   | 2.73±0.18    |

Note: BCH = high biochar addition, BCL = low biochar addition, SGH = high gypsum addition, SGL = low gypsum addition, CK = soil without amendment

addition. Data in the table are average values and standard deviations (n=3).



### A soil incubation study

