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Nitrogen Cycling and Soil Amelioration in *Camellia oleifera* Plantations

Bangliang Deng and Ling Zhang

Abstract

Camellia oleifera Abel. is one of the four woody edible oil trees around the world, which is also an important economic species in subtropical China. It is mainly cultivated in subtropical region, where the soil constrains the yield of *C. oleifera* oil due to its low fertility and pH. Thereby, intensive management including fertilization practice, especially intensive nitrogen (N) input, has been developed as a vital way to enhance oil yield in *C. oleifera* plantations. However, excessive nitrogen input increases soil nitrous oxide (N₂O) emissions and soil acidification, limiting sustainable development of economic forests. As one of the important greenhouse gases, N₂O is 265 times greater than carbon dioxide in global warming potential on 100-year scale. To mitigate soil N₂O emissions and soil acidification, soil amelioration, including applications of biochar, nitrification inhibitors, and urease inhibitors, played an important role in sustainable management of *C. oleifera* plantations. This chapter reviewed soil nitrogen cycling, N₂O emissions, and soil amelioration in *C. oleifera* plantations, which will benefit the sustainable management of *C. oleifera* plantations and hence the development of *C. oleifera* industries.

Keywords: *Camellia oleifera*, biochar, nitrification inhibitor, soil amelioration, sustainable forest management, urease inhibitor

1. *Camellia oleifera*

Camellia oleifera Abel. as a native edible oil tree has a long cultivation history in subtropical China [1]. It is a perennial and evergreen species with synchronous flowers and fruits. The cultivation area and total product value of *C. oleifera* have reached 4.47 million ha and 102.4 billion Chinese yuan, respectively [2]. With rapid development, the *C. oleifera* oil accounted for 80% domestic high-end vegetable edible oils in 2018 from China. High habitat suitability area for *C. oleifera* cultivation in China has been up to 4.94% [3].

Specially, *C. oleifera* oil and oils derived from palm, olive, and coconut are the four major woody edible oils in the world [4]. The *C. oleifera* oil is characterized by remarkable antioxidant activity [5] and high content of unsaturated fatty acids (about 83%) [6].

Camellia oleifera can survive and adapt to low-fertility acid soil. Generally, it usually is used in the conservation of soil and water as well as afforestation in barren hill. Therefore, *C. oleifera* is an excellent species with both ecological and

economic advantages. Development of *C. oleifera* industry would be beneficial for the safety of edible oil and the conservation of soil and water in China.

As a typical economic tree, intensification such as water management, fertilization, and trimming takes an important part in the management of *C. oleifera* plantations. Notably, organic matter, available phosphorus, and pH value was low in *C. oleifera* plantation soils [7], constraining the yield of *C. oleifera* oil. Therefore, intensive management with fertilization is often performed in *C. oleifera* plantations [1].

2. Challenges

Fertilization is the major way of intensive management, efficiently improving the yield of oil in *C. oleifera* plantations. However, a large amount of nitrogen (N) input increased the risk of soil N leaching and gaseous N (e.g., nitrous oxide (N_2O), nitric oxide (NO), ammonia (NH_3)) losing [8]. In addition, excessive N input induced soil acidification [9].

2.1 Nitrous oxide emissions

Nitrous oxide, as the major ozone-depleting substance [10], has been recognized to be an important greenhouse gas. Especially, the potential of N_2O for global warming is 265 times than that of carbon dioxide [11]. The concentration of N_2O is ranging from 270 ppb in pre-industrial period to 329.9 ppb in 2017 [12].

Soil systems contributed the largest source of N_2O emissions ($13 \text{ Tg N}_2\text{O-N yr}^{-1}$), of which human activities accounted for 54% [13]. Nitrogen input such as N deposition and N fertilization often increased N_2O emissions and altered the process of N transformation [14–17]. Generally, soil N_2O emissions had a positive and linear relationship with N input [18]. About 120 Tg N was contributed by human activities per year [13]. Therefore, intensive N input often leads to high emissions of soil N_2O [19].

2.1.1 Nitrification and denitrification

Nitrification and denitrification are the two main pathways of N_2O emissions (Figure 1) [20–22], which produced global 70% soil N_2O emissions [13].

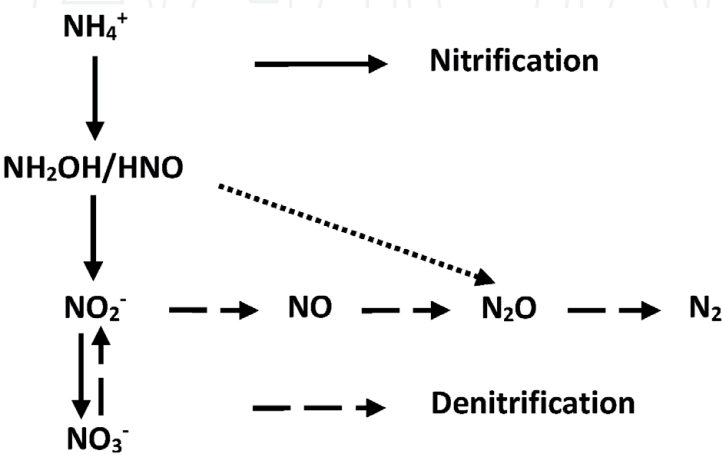


Figure 1.
Nitrification- and denitrification-related pathways [20–22].

2.1.2 Influence factors

Soil N₂O emissions can be influenced by soil environmental factors such as soil moisture, temperature, oxygen (O₂), and pH condition [23, 24].

2.1.2.1 Soil moisture

Soil moisture is a vital factor that affects soil N₂O emissions. Generally, soil N₂O emission rates reached the peak when soil water-filled pore space (WFPS) was 60–70% [25]. For example, soil N₂O emissions were significantly higher under 60% WFPS conditions than that under flooded conditions [26].

2.1.2.2 Soil temperature

Effects of soil temperature on N₂O emissions were more complex than that of soil moisture. For example, warming increased soil N₂O emissions from boreal peatland [27] and alpine meadow [28]. Soil N₂O emissions had an exponential increased relationship with incubation temperatures [29]. A significant positive correlation was presented in N₂O emissions and soil temperature from different soil types (paddy, orchard, forest, and mountain) [30]. Although warming did not affect soil N₂O emissions from northern peatlands, it suppressed N₂O emissions under N addition conditions [31]. By contrast, the effects of warming on soil N₂O emissions from alpine meadow soil were not observed [32]. Consistently, no significant increase of soil N₂O emissions was found with increasing incubation temperatures [33]. Previous study reported that soil moisture and temperature can explain 86% of soil N₂O emissions [34].

2.1.2.3 Soil O₂ concentration

Soil O₂ concentration was closely related with soil moisture and soil mechanical composition. Generally, soil with higher water content and larger clay fraction had lower soil O₂ concentrations. Lower soil O₂ concentrations mainly promoted soil N₂O emissions via denitrification [20, 35]. The production of N₂O and NO was increased when O₂ concentration decreased from 21% to 0.5% in a laboratory study [36]. Similarly, field study reported that soil N₂O emissions increased with increasing soil O₂ concentrations in wetland [37].

2.1.2.4 Soil pH

pH played an important role in the activity of microbes [38]. Indeed, soil acidification [39] and soil pH amelioration [40] significantly influenced soil N₂O emissions. However, other researchers reported that there was no significant correlation between N₂O emissions and pH [41].

pH influenced the activity of nitrification- and denitrification-related enzymes [42]. Generally, soil acidification increased N₂O emissions [42]. Compared with ammonia-oxidizing bacteria (AOB), ammonia-oxidizing archaea (AOA) were higher in activity and resistance from acid soil [43]. However, the domination of AOB was increased by increasing soil pH [44]. Additionally, archaeal *amoA* genes had a wide pH range of about 3.7–8.65, which had high activity in extreme environments such as high temperature and extreme acid [45].

2.1.3 Nitrous oxide emissions from *Camellia oleifera* plantation soils

Our previous field study (1 year) found that soil N_2O emissions were $92.14 \pm 47.01 \text{ mg m}^{-2}$ in control treatment and were $375.10 \pm 60.30 \text{ mg m}^{-2}$ in fertilization treatment ($400 \text{ kg NH}_4\text{NO}_3\text{-N ha}^{-1}$) from *C. oleifera* plantations [1].

2.2 Soil acidification

Acid soil ($\text{pH} < 5.5$) as a main soil type covers about 30% free ice land [46]. However, soil acidification has been becoming more and more serious [47]. Soil acidification should be taken into consideration due to its constraint in the sustainable development of agricultural sector [48]. In China, soil pH (except alkaline soils at $\text{pH} 7.10\text{--}8.80$) from crop fields reduced by $0.13\text{--}0.76$ during the year 1980–2000 [49]. For example, soil pH (surface layer) decreased by 0.30 units from 1981 to 2012 in Sichuan Province, China [47].

With a long cultivation history, *C. oleifera* was widely cultivated in acid or strongly acid soil in subtropical China [7]. The optimum pH for the growth of *C. oleifera* is $5.5\text{--}6.5$ [50]. However, acid deposition [51] and intensive N input [49] may stimulate soil acidification from *C. oleifera* plantations. Additionally, long-term N input may also increase the toxicity of aluminum (Al) [52], limiting the sustainable development of *C. oleifera*.

Soil acidification from *C. oleifera* plantations is mainly related to the following factors.

2.2.1 Precipitation

Long-term precipitation increased the loss of base cations such as Ca^{2+} , Mg^{2+} , K^+ , and Na^+ , reducing the soil pH buffering capacity. In addition, long-term precipitation promoted the accumulation of Al^{3+} and Fe^{3+} in soil, which could further hydrolyze to $\text{Fe}(\text{OH})_3$ or $\text{Al}(\text{OH})_3$ and release 3H^+ .

2.2.2 Plant physiology

When plant roots absorb a NH_4^+ from soil, an H^+ will release into soil; in turn, absorbing a NO_3^- from soil will release an OH^- into soil [53].

Organic acid (R-COOH) from root exudates can release an H^+ after hydrolysis. In addition, anions of organic acids (e.g., citric acid and malic acid) can chelate with Al^{3+} in the soil and inhibit the root system that absorbs Al^{3+} , alleviating Al^{3+} toxicity to plant growth [48, 54, 55].

Plants such as *C. oleifera* [56] can uptake Al^{3+} by roots, promoting the accumulation of Al^{3+} in surface soil via litter decomposition [57]. The accumulation of Al^{3+} can replace the base cations such as Ca^{2+} , Mg^{2+} , K^+ , and Na^+ and accelerate leaching, hence reducing the pH buffering capacity of top soil.

2.2.3 Microbial-mediated nitrification

For example, NH_4^+ transfers to NO_3^- along with the 2H^+ release ($\text{NH}_4^+ + 2\text{O}_2 \rightarrow \text{NO}_3^- + \text{H}_2\text{O} + 2\text{H}^+$) [53]. AOB, AOA, and fungi can participate in the process of nitrification [20]. Nitrification includes the pathway of ammonia oxidation to hydroxylamine, the pathway of hydroxylamine oxidation to nitrite, and the pathway of nitrite oxidation to nitrate (**Figure 1**) [58]. Ammonia can be oxidized by AOA or AOB to hydroxylamine via ammonia monooxygenase (*amo*).

Hydroxylamine can be oxidized to nitrite by hydroxylamine oxidoreductase. Nitrite can be oxidized to nitrate by nitrite oxidoreductase.

2.2.4 Oxidation of sulfur-containing organics

Oxidation of sulfur mineral, for example, oxidation of FeS_2 , will produce 2H^+ ($2\text{FeS}_2 + 7\text{O}_2 + 2\text{H}_2\text{O} \rightarrow 2\text{Fe}^{2+} + 4\text{SO}_4^{2-} + 4\text{H}^+$).

Oxidation of sulfur-containing organics will release 4H^+ ($2\text{Organic-S} + 3\text{O}_2 + 2\text{H}_2\text{O} \rightarrow 2\text{SO}_4^{2-} + 4\text{H}^+$).

2.2.5 Intensive nitrogen fertilization

Intensive NH_4^+ input can replace the base cations such as Ca^{2+} , Mg^{2+} , K^+ , and Na^+ and accelerate leaching, reducing the pH buffering capacity of top soil [59]. Hydrolysis of soil NH_4^+ will generate NH_3 (gas) and consume an OH^- ($\text{NH}_4^+ + \text{OH}^- = \text{NH}_3\uparrow + \text{H}_2\text{O}$) [60].

Acidic fertilizers such as $\text{Ca}(\text{H}_2\text{PO}_4)_2$ will gradually release H^+ , hence increasing soil acidification ($\text{Ca}(\text{H}_2\text{PO}_4)_2 \rightarrow \text{CaHPO}_4 + \text{H}_3\text{PO}_4$, $\text{H}_3\text{PO}_4 \rightarrow \text{H}^+ + \text{H}_2\text{PO}_4^- \rightarrow 2\text{H}^+ + \text{HPO}_4^{2-} \rightarrow 3\text{H}^+ + \text{PO}_4^{3-}$).

2.2.6 Acid deposition

Acid deposition (water-soluble acid gases such as CO_2 and sulfur dioxide) and N deposition (especially $\text{NH}_4^+\text{-N}$) increased soil acidification [51]. Precipitation with H^+ can replace the soil base cations such as Ca^{2+} , Mg^{2+} , K^+ , and Na^+ , which directly reduce the soil pH buffering capacity [51].

2.2.7 Other factors

For example, deforestation and other land uses can reduce litter accumulation in surface soil, hence declining the accumulation of base cations such as Ca^{2+} , Mg^{2+} , K^+ , and Na^+ that generate from litter decomposition [61].

2.3 Effects of soil acidification on nitrous oxide emissions

Acid soils have been facing an increased risk of acidification due to human activities, especially intensive N fertilization [47, 49, 62]. For example, after 6 years of application of $600 \text{ kg Urea-N ha}^{-1} \text{ yr}^{-1}$, soil pH was significantly decreased (soil pH in control and fertilization treatment was 5.1 and 4.9, respectively) from a tea plantation in Yixing City, Jiangsu Province, China [63]. A meta-analysis of 1104 field data showed that a negative correlation between soil N_2O emissions and pH (3.34–8.7) ($\text{N}_2\text{O-N} = -0.67x + 6.55$, $R = 0.22$) is negatively related with N fertilization [9]. Moreover, deposition of sulfur dioxide increased soil acidification, stimulating soil N_2O emissions [64].

The mechanism of soil acidification on the stimulation of soil N_2O emissions is complex, which may include (but not limited to) the following points.

2.3.1 Chemical decomposition of nitrous acid

Under acidic conditions, $\text{pH} < 5.5$, NO_2^- (HNO_2 , $\text{pK}_a = 3.3$) will naturally decompose into NO and/or NO_2 ($3\text{HNO}_2 \rightleftharpoons 2\text{NO} + \text{HNO}_3 + \text{H}_2\text{O}$ or $2\text{HNO}_2 \rightleftharpoons \text{NO} + \text{NO}_2 + \text{H}_2\text{O}$) [65]. Soil NO can be further transformed to N_2O with Fe^{2+} when it was not escaping soil [65].

2.3.2 Shifts in microbial communities and abundance

Generally, the abundance of AOB was lower in soil pH < 5.5 than that in neutral soil pH. Here, nitrification was weak and almost disappears at soil pH < 4 [66]. However, AOA could mediate the process of ammonia oxidation in extremely strong acidity soil (pH: 4.2–4.47) [43]. Another study reported that the abundance of AOB was positively correlated with pH ($R^2 = 0.2807$), while the abundance of AOA was negatively correlated with pH ($R^2 = 0.2141$) [67]. For example, AOA dominated in acid paddy soil (pH 5.6), while AOB dominated in alkaline soil (pH 8.2) [68]. Previous research indicated that fungi were the main microbial community that mediated N_2O emissions in acid soil [69, 70]. Additionally, fungi-mediated denitrification accounted for 70% soil N_2O emissions from a 100-year-old tea plantation (soil pH 3.8) [71].

In acid soils, the activity of N_2O reductase was inhibited, leading to higher N_2O emissions in lower soil pH [72]. Indeed, there was a positive correlation between the abundance of *nirS*, *nirK*, or *nosZ* and soil pH (4.0–8.0) and a negative correlation between $N_2O/(N_2O + N_2)$ and soil pH [73]. In agreement, $N_2O/(N_2O + N_2)$ was negatively correlated with soil pH (3.7–8.0) ($R^2 = 0.759$, $P < 0.001$), and lime addition decreased $N_2O/(N_2O + N_2)$ [74]. The ratio of $N_2O/(N_2O + N_2)$ increased with decreasing pH (5.57–7.06) ($R^2 = 0.82$) [75]. Consistently, soil pH was negatively correlated with N_2O/N_2 [76]. Intensive management consistently decreased soil pH and increased the ratio of $N_2O/(N_2O + N_2)$ [77]. Increasing dolomite dosage increased soil pH and hence increased the transcription of *nosZ* genes and reduced the potential of N_2O production in acid soils [26].

2.3.3 Microbes increased resistance to soil acidification

Laboratory study showed that the potential of soil N_2O emissions was increased with decreasing pH (soil pH ranging from 2.96 to 6.26) from tea plantations in Japanese [78]. In addition, higher soil N_2O emissions and lower abundance of *nosZ* genes were observed in soil pH at 3.71 (control) than in pH at 5.11, 6.19, and 7.41 (lime amelioration) under NO_3^- -N fertilization (50, 200, and 1000 mg kg^{-1}) from a 100-year-old tea plantation [79]. Field study found a negative correlation between soil N_2O emissions and pH (pH 3.6–5.9) ($N_2O-N = 636.6 * e^{-0.8028 * pH}$, $R = -0.93$) from *Betula pendula* Roth forest [80]. Thus, denitrifying microorganisms may have been adapted extremely to acid soil environments, resulting in high N_2O emissions when soil acidification happened.

3. Sustainable forest management

Soil amelioration (e.g., application of lime, biochar nitrification inhibitors, and urease inhibitors) plays an important role in mitigation of soil acidification and N_2O emissions.

3.1 Lime

Lime as an ameliorant was often used to amend acid soils in southern China due to increasing soil pH. It can relieve the toxic effect of soil Al^{3+} on plant growth by reducing soil exchangeable H^+ [81]. Lime addition increased soil pH and salt saturation [82]. In addition, application of lime can reduce soil N_2O emissions [40]. For example, under 60% WFPS or flooded conditions, dolomite addition at medium- or high-dose levels (1 or 2 g kg^{-1} soil) can reduce N_2O emissions and increase the transcription of *nosZ* genes ($N_2O \rightarrow N_2$) by increasing acid soil pH from a rice-rapeseed

rotation system [26]. However, lime addition reduced the content of soluble organic carbon in the soil layer 10–30 cm [83]. Consistently, long-term lime addition increased the soil pH but stimulated the decomposition of soil organic carbon [84].

3.2 Biochar

Biochar was stable in the soil from Amazon basin of Brazil, and biochar input improved soil fertility [85]. This discovery accelerated the development of technologies for biochar application in soil amelioration.

Biochar is a carbon (C)-rich solid material by pyrolyzing of organic biomass such as crop straw, forestry by-products, urban waste, industrial by-products, animal manure, and urban sludge at low oxygen and high temperature (250–700°C) condition [86]. Biochar has been characterized by a high pH, specific surface area, degree of aromatization, and porosity. In addition, biochar is rich in C-containing functional groups (e.g., C–H, C–O, C=C and C=O) and relatively stable organic C. The physicochemical properties of biochar were mainly determined by pyrolysis temperature [87].

Presently, biochar was widely used as a soil ameliorant in agriculture and forestry field. For example, our previous studies reported that *C. oleifera* fruit shells are ideal feedstock for producing biochar as they are rich in C and N [1, 88]. Biochar includes the following advantages:

1. Carbon recalcitrance of biochar can increase soil C pool. The potential of biochar in mitigation of greenhouse gas emissions was 1.0–1.8 Pg CO₂-C_{eq} yr⁻¹ [89].
2. Biochar had excellent physicochemical characteristics in soil nutrient retention and utilization [90, 91] and water conservation [92]. Additionally, biochar can increase the plant resistance to Al³⁺ toxicity [81], the clone of arbuscular mycorrhizal fungi, and crop yield [93, 94]. It can decrease continuous cropping obstacles such as root-knot nematode [95] and *Ralstonia solanacearum* [96].
3. Biochar is rich in macro- and microelements [97], which can reduce the dosage of fertilizer.

3.2.1 Effects of biochar on soil nitrous oxide emissions

The physicochemical properties of biochar and soil can interactively influence soil N₂O emissions [98]. However, the effects of biochar on soil N₂O emissions varied, including positive effects [99], negative effects [100], and no effects [101].

Biochar addition increased soil N₂O emissions with the release of N from biochar [102]. By contrast, biochar reduced soil N₂O emissions with (1) increased NO₃⁻-N immobilization [103]; (2) increased copy numbers of *nosZ* gene [104, 105]; and (3) increased toxic effects of polycyclic aromatic hydrocarbons and other toxic substances (pyrolysis by-products) on N-cycle microorganisms [106].

3.2.2 Effects of biochar on soil pH buffer capacity

Biochar that increased soil pH buffer capacity may predominantly correlate with biochar rich in oxygen-containing functional groups in surface. The anions of weakly acidic functional groups can associate with H⁺, hence increasing soil pH. Meanwhile, exchangeable base cations can release into the solution, thus increasing soil pH buffer capacity [107, 108]. In addition, soluble silicon (Si) such as H₃SiO₄⁻ (present at a high pH) can combine with H⁺ and generate H₂SiO₃ precipitation [107, 108].

3.3 Nitrification inhibitor

Nitrification inhibitors are a class of organic compounds that can inhibit the activity of nitrifying bacteria.

Nitrification inhibitors, especially synthetic nitrification inhibitors (e.g., dicyandiamide (DCD) and 3,4-dimethylpyrazole phosphate (DMPP)), were widely used in agriculture for improving N use efficiency. Ammonia-oxidizing bacteria and AOA are the major microbial communities in nitrification and denitrification, and both contain *amo* enzyme that can catalyze ammonia oxidation ($\text{NH}_4^+ - \text{N} \rightarrow \text{NH}_2\text{OH}$). Synthetic nitrification inhibitors such as DCD and DMPP mainly inhibit nitrification by suppressing the activity of *amo* enzyme (a Cu-copper cofactor enzyme). In addition, biological nitrification inhibitors also can inhibit soil nitrification [109, 110]. In the mid-1980s, researchers found that *Brachiaria humidicola* cv. Tully (CIAT 679), a single community forage, had lower nitrification rates than a single legume community or bare land [111]. This phenomenon stimulated further studies on biological nitrification inhibitors. The first biological nitrification inhibitor (methyl 3-(4-hydroxyphenyl) propionate: MHPP) was identified from the root exudate of *Sorghum bicolor* in 2008, which mainly inhibited the activity of *amo* enzyme [112]. Subsequently, biological nitrification inhibitor (brachialactone) from the root exudate of *Brachiaria humidicola* was found to inhibit the activity of *amo* enzyme [113]. The Nanjing Soil Research Institute of China firstly found and identified a biological nitrification, 1,9-decanediol, from the root exudate of rice, which can inhibit the activity of *amo* enzyme [114].

Ammonium N can be adsorbed by soil colloids, while soil $\text{NO}_3^- - \text{N}$ (the end product of nitrification) easily can be leached to groundwater by precipitation. In addition, microbial-mediated nitrification is closely related with soil N_2O emissions [20–22]. Nitrification inhibitors can effectively inhibit soil nitrification, slowing the transformation of $\text{NH}_4^+ - \text{N}$ to $\text{NO}_3^- - \text{N}$ and hence reducing the $\text{NO}_3^- - \text{N}$ leaching and N_2O emissions.

An evaluation from 62 field studies showed that although nitrification inhibitors increased 20% NH_3 emissions, they reduced 48% inorganic N leaching, 44% N_2O emissions, and 24% NO emissions and increased 58% plant N utilization, 9% grain yield, 5% straw yield, and 5% vegetable yield [115]. Consistently, other studies evaluated that nitrification inhibitors decreased by 38% [116], 50% [117], or 73% [118] N_2O emissions and decreased by $0.3 \text{ t CO}_2\text{e ha}^{-1} \text{ yr}^{-1}$ [119]. Similarly, DCD did not increase crop yields but reduced 35% N_2O emissions [120]. A meta-analysis showed that DCD rather than DMPP significantly increased 6.5% crop yield as well as DCD and DMPP decreased N_2O emissions by 44.7% and 47.6%, respectively [121].

Therefore, application of nitrification inhibitors could reduce N_2O emissions and mitigate environmental pollution after intensive N inputs.

3.4 Urease inhibitors

Urease inhibitors are a class of compounds that can slow soil urease activity (Figure 2). Addition of urease inhibitors after urea input can inhibit the hydrolysis

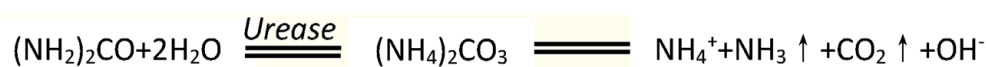


Figure 2.
The chemical equation of urea hydrolysis with urease catalysis.

of urea via inhibiting the activity of urease, hence reducing NH_3 volatilizations and N_2O emissions. Additionally, the application of urease inhibitors also contributes to increase N utilization efficiency and reduce NO_3^- -N leaching. *N*-(*n*-butyl) thiophosphoric triamide (NBPT) is one of the most wide and effective urease inhibitors.

Urease, a Ni-copper enzyme, has two Ni–O bidentate ligands, specifically catalyzing urea into NH_3 and CO_2 . Urea only can bind with one specific Ni–O ligand of urease, but NBPT can bind with two Ni–O bidentate ligands of urease and generate a tridentate ligand [122], hence inhibiting the activity of urease.

Presently, a meta-analysis reported that a nonlinear response was presented in soil NH_3 volatilizations and N input [123]. Application of NBPT can effectively inhibit NH_3 volatilizations. For example, 530 mg NBPT kg^{-1} urea treatment delayed NH_3 volatilizations and decreased accumulation of NH_3 volatilizations compared with the control treatment. NH_3 volatilizations were linearly related with the NBPT dosage in the range of 0–1000 mg NBPT kg^{-1} Urea (0, 530, 850, 1500, and 2000 mg NBPT kg^{-1} Urea) [124]. Other study reported that NBPT increased 27% oat yield and 33% crop N uptake [120].

The effects of NBPT on N_2O emissions were controversial. For example, NBPT can reduce 80% N_2O emissions [117]. No effects of NBPT (0.07%, NBPT/Urea-N, w/w) on N_2O emissions were observed [125]. Similarly, there was no change of N_2O emissions with NBPT (250 mg NBPT kg^{-1} Urea) addition from urea-fertilized (50 kg Urea-N ha^{-1}) soil [126].

Additionally, NBPT can reduce N_2O emissions from alkaline soils but has no effects on acidic soils [127], which indicated that pH plays a key role in the regulation of NBPT effects on N_2O emissions. Further laboratory study showed that NBPT inhibited nitrification, stimulating N_2O emissions from alkaline soils (pH 8.05) but not affecting N_2O emissions from acid soils (pH 4.85). This finding suggested that the effect of NBPT on soil N_2O emissions is not only influenced by pH but also by other unknown factors [127].

Generally, urease inhibitors correlated with nitrification inhibitor could mitigate N_2O emissions. A meta-analysis showed that urease inhibitors and nitrification inhibitors interactively reduced 30% N_2O emissions [116]. For example, a field study reported that the combination of NBPT (0.3%, NBPT/Urea-N, w/w) and DCD (0.3%, DCD/Urea-N, w/w) reduced 32.1% soil N_2O emissions with the addition of 519 kg Urea-N ha^{-1} from banana plantation, but did not affect the yield of banana [128].

4. Sustainable management in *Camellia oleifera* plantations

Our previous incubation study found that although biochar application increased N_2O emissions, DCD addition decreased soil N_2O emissions under urea fertilization from *C. oleifera* field [88]. Our field study showed that N_2O emission rates were inhibited by biochar or DCD application and the effects of biochar application on mitigation of cumulative N_2O were comparable to DCD addition in *C. oleifera* plantations [1]. Compared with control treatment, available N (NH_4^+ -N and NO_3^- -N) was not affected by NH_4NO_3 , NH_4NO_3 + DCD, or NH_4NO_3 + biochar treatment [1]. In addition, the seed yield of *C. oleifera* was higher in NH_4NO_3 or NH_4NO_3 + biochar treatment than that in control or NH_4NO_3 + DCD treatment (Figure 3). Soil amelioration is necessary and improves N use efficiency and pH, mitigating N_2O emissions. Soil amelioration plays an important role in the sustainable management of oil safety in *C. oleifera* plantations.

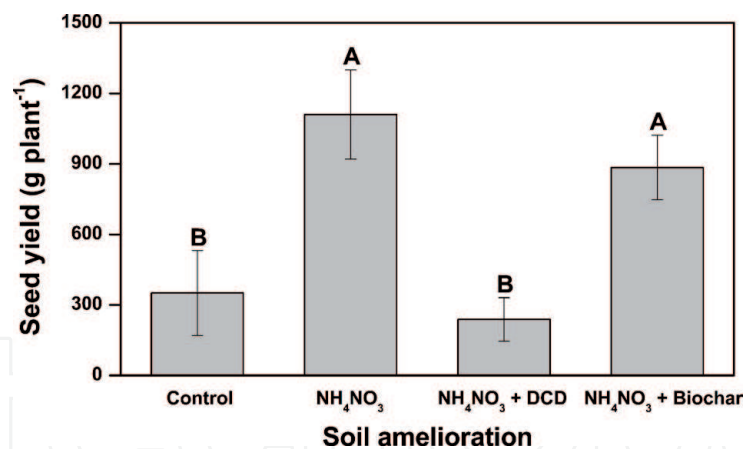


Figure 3. The seed yield of *Camellia oleifera* with nitrogen fertilization, in combination with nitrification inhibitor (DCD) or biochar. Bars connected by different letters indicate significant difference in post-hoc tests at $\alpha = 0.05$ (means \pm se).

5. Conclusions

Soil acidification, especially induced by N fertilization, will inhibit the activity of N₂O reductase and increase the abundance of N₂O-producing fungi as well as the acid resistance of N₂O-producing microorganisms, hence the ratio of N₂O/(N₂O + N₂). In addition, NO₂⁻ will generate NO under soil pH < 5.5 condition, which will further transform into N₂O. Under the background of global acidification, the soil from *C. oleifera* forest also suffers the potential risks of soil acidification and N₂O emissions. Mitigation of soil acidification and N₂O emissions by soil amelioration is necessary and improves N use efficiency and soil pH from *C. oleifera* plantations. Soil amelioration such as biochar and nitrification inhibitor plays an important role in sustainable forest management in *C. oleifera* plantations.

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Conflict of interest

The authors declare no conflict of interest.

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