Title: Straw return with diverse nitrogen fertilizer application rates modulate ecosystem services and microbial traits in a meadow soil

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Abstract:

Nitrogen (N) fertilization has received worldwide attention due to its benefits to soil fertility and productivity, but excess N application also causes an array of ecosystem dis-services, such as greenhouse gas emissions. Generally, soil microorganisms are considered to be involved in upholding a variety of ecosystem services and dis-services. However, the linkages between soil ecosystem services and microbial traits under different N fertilizer application rates remain uncertain. To address this, a 4-year in situ field experiment was conducted in a meadow soil on the Northeast China Plain after straw return with the following treatments combined with regular phosphorus (P) and potassium (K) fertilization: (i) regular N fertilizer (N+PK); (ii) 25% N fertilizer reduction (0.75N+PK); (iii) 50% N fertilizer reduction (0.5N+PK); and (IV) no N fertilizer (PK). Ecosystem services, dis-services and microbial traits responded distinctly to the different N fertilizer rates. Treatment 0.75N+PK had overall positive effects on soil fertility, productivity, straw decomposition, and microbial abundance and function and alleviated greenhouse effects due to N deficiency. Meanwhile, 0.75N+PK upregulated aboveground biomass and soil C:N and thus increased the abundance of genes encoding cellulose-degrading enzymes, which may imply the potential ability of C and N turnover. In addition, most observed changes in ecosystem services and dis-services were strongly associated with microbial modules and keystone taxa. Specifically, the Lasiosphaeriaceae-driven module 1 community promoted straw degradation and C and N release, while the Terrimonas-driven module 3 community contributed to production improvement, which was conducive to soil multifunctionality. Therefore, our results suggest that straw return with 25% chemical N fertilizer reduction is optimal for achieving ecosystem services. This study highlights the importance of abiotic and biotic factors in soil health and supports green agricultural development by optimizing N fertilizer rates in meadow soil after straw return.
Keywords: Ecosystem services; Straw return; Nitrogen fertilization; Microbial community; Crop yield

1. Introduction

Multiple soil ecosystem services are indicators of soil health (Kihara et al., 2020; Lehmann et al., 2021). Soil ecosystem services refer to the ability of soil to function as a vital living system to sustainably increase crop productivity, improve environmental quality, tackle climate change and promote plant and animal health (de Bello et al., 2010; Tang et al., 2019). In recent decades, anthropogenic activity, such as intensive agriculture, has posed a wide range of threats to agroecosystem services (Robertson et al., 2014; Allen et al., 2015). Irrational or excessive application of chemical fertilizers, especially nitrogen (N), is ubiquitous to achieve high crop yields in response to population surges globally (Shi et al., 2019). In fact, N is considered the essential macronutrient for all biota, while excessive N fertilizer inputs not only reduce soil fertility and productivity but also lead to environmental burdens (Trost et al., 2016). For example, previous studies emphasized that N fertilizer abuse may accelerate greenhouse gas emissions (Huang et al., 2006; Wu et al., 2015) and degrade groundwater quality (Rhymes et al., 2016). Therefore, how to achieve agroecosystem services by regulating N fertilizer application rates is a critical issue that needs to be fully assessed.

Straw return has also been widely applied as a major measure to moderate soil ecosystem services (Xu et al., 2021). Plant residues, as natural organic bioenergy resources, contain abundant N that further affects soil fertility and productivity (Pan et al., 2009; Liu et al., 2014). Thus, the straw-derived N released during degradation is an important source that may serve as a partial substitute for chemical N fertilizer application (Wang et al., 2017; Latifmanesh et al., 2020). However, crop fields suffering
from abundant organic materials usually have low reutilization efficiency (Hou et al., 2020). Generally, the majority of N in straw is released into the atmosphere as oxynitride, such as nitrous oxide (N₂O), resulting in lower soil organic matter (SOM) formation efficiency (Wang et al., 2019; Sun et al., 2021). Subsequent literature highlighted that straw return significantly elevates greenhouse gas emissions so that less than 15% of straw-derived N can be transformed into soil and become SOM (Yin et al., 2018; Wu et al., 2019). However, the potential for the partial substitution of straw for chemical N fertilizer application is still unclear. Revealing the mechanisms of efficient straw utilization under diverse N fertilizer input rates is essential to achieving ecosystem multifunctionality.

Compared with plants and animals, soil contains more microorganisms living in an opaque environment, making the evaluation of soil ecosystem services more complex (Handa et al., 2014; Wagg et al., 2014). Agronomic management for such “multifunctionality” has prompted research into the role that microbes play in providing desired rates of multiple ecosystem processes (Gong et al., 2020). To our knowledge, fertilization-induced changes in microbial communities and functions are fundamental to the regulation of a variety of ecosystem multifunctionalities, including SOM formation, greenhouse gas emissions, litter decomposition, and crop production (Dominati et al., 2014). To date, we still lack empirical evidence of the linkages among N fertilizers, specific microbial communities or functions and multiple ecosystem services, and the diverse cropland services driven by complex microbial traits under different N fertilizer rates are seldom clarified.

Microorganisms contribute to ecosystem services by modulating microbial function, community composition and succession, which makes understanding the consequences of the changes in microbial traits crucial for determining different N fertilizer input levels (Bradford et al., 2014). The role of microorganisms in ecosystem functioning is unequivocal, and these organisms can be recognized as the
key drivers of ecosystem services (Chen et al., 2019a). Generally, bacteria and fungi are the main
drivers of straw labile and recalcitrant component decomposition, respectively (Frey et al., 2013; Ge et
al., 2017). Therefore, the ratio of fungi to bacteria is always considered an indicator during straw
degradation periods (Hogberg et al., 2007). Specifically, the expression levels of the cbhl and GH48
genes were identified as biomarkers of cellulolytic fungi and bacteria, respectively (Zhang et al., 2017).
Previous studies revealed that the N input level dominated the associations between microbial
composition and cellulolytic gene abundance with SOM physical fractions (Duan et al., 2021). In
addition, microbial module communities and keystone taxa have been used to provide satisfactory
explanations for ecosystem services. Chen et al. (2019b) found that particular microbial modules
participated in N and phosphorus (P) accumulation and CO₂ emissions in a Cambisol. Moreover,
specific taxa are involved in agrosystem services. For example, Actinobacteria have been extensively
studied and can be considered the main degraders of straw by secreting cellulase (Bao et al., 2021).
_Mortierella_ has been proven to increase soil fertility and crop yield due in part to its strong C
sequestration capacity (Ning et al., 2020). Notably, it is also well known that microbial traits are
mediated by nutrient availability and stoichiometry (Chen et al., 2014). C, N and P stoichiometry has
profound impacts on microbial in vivo metabolism and ex vivo modification processes (Chen et al.,
2016). Multiple studies have indicated that soil C:N and N:P ratios are the key factors mediating
microbial functions and soil health (Ning et al., 2020; Duan et al., 2021). Nevertheless, the knowledge
of the microbial mechanisms that modulate ecosystem services in response to N fertilizer input levels
are still rudimentary.

As an important grain-producing region, the Northeast China Plain contributes to more than 20%
of the total grain yield in China (Li et al., 2017). However, excessive chemical N fertilizer inputs have
caused ecosystem dis-services over the past decades (Zhao et al., 2018). Therefore, a field experiment was conducted to reveal the influences of N input levels on soil ecosystem multifunctionality and associated microbial traits and to try to establish the linkages between them. In the present study, two hypotheses were tested: (i) soil ecosystem services and dis-services would show distinct responses to N fertilizer input levels, and (ii) the changes in cropland ecosystem services and dis-services would be linked to specific microbial traits.

2. Materials and methods

2.1 Site description and sampling

A field experiment under contrasting inorganic N fertilizer input levels was established in 2018 in Wenchun town (44°59′61″ N, 129°59′18″ E), Mudanjiang city, Heilongjiang Province, Northeast China Plain, which is an important grain-producing area. This region has a typical temperate continental monsoon climate with an average annual temperature of 4.3 °C and a mean annual precipitation of 579.7 mm. The soil is classified as a meadow soil according to US Soil Taxonomy (USST). The cropping system was continuous maize (*Zea mays* L.) monoculture. Four treatments received different N fertilizer input levels after straw return to the field for 4 years as follows: (1) regular chemical fertilization, N+PK (300 kg urea (N 46%) ha⁻¹ yr⁻¹, 250 kg diammonium phosphate (P₂O₅ 48%) ha⁻¹ yr⁻¹, 150 kg potassium chloride (K₂O 50%) ha⁻¹ yr⁻¹); 25% reduction of N fertilizer, 0.75N+PK (225 kg urea ha⁻¹ yr⁻¹, 250 kg diammonium phosphate ha⁻¹ yr⁻¹, 150 kg potassium chloride ha⁻¹ yr⁻¹); 50% reduction of N fertilizer, 0.50N+PK (150 kg urea ha⁻¹ yr⁻¹, 250 kg diammonium phosphate ha⁻¹ yr⁻¹, 150 kg potassium chloride ha⁻¹ yr⁻¹); and no N fertilizer, PK (250 kg diammonium phosphate ha⁻¹ yr⁻¹, 150 kg potassium chloride ha⁻¹ yr⁻¹). All straw and chemical fertilizers were applied with shallow tillage to 20 cm.
cm. Straw was cut into pieces less than 5 cm and input after the harvest in October, while the chemical fertilizers were applied during ploughing in May of the next year. All other normal management practices were consistent among treatments during the experiment. Before the experiment, the initial soil contained 18.74 g kg\(^{-1}\) SOC, 1.03 g kg\(^{-1}\) total N and 0.54 g kg\(^{-1}\) total P with a pH of 7.37 (H\(_2\)O).

The yield and some of the soil chemical properties under different bulk soil treatments during the experimental process are shown in Supplemental material Table S1.

Soils were sampled after the maize harvest in October 2021. A randomized complete block design consisting of 5 treatments with 3 replications was adopted in this study. Each field plot was 4.5 m × 15 m. We took nine soil cores (5 cm diameter) from the top 20 cm of bulk soil in each plot. Each soil sample consisted of a mixture of subsamples randomly collected at nine different positions in the same plot. In total, 12 soil samples were collected from 4 treatments with 3 replicates. Soils were sieved through a 2 mm mesh, the mineral particles and plant roots were carefully removed, and then the soils were homogenized and stored in an incubator at 4 °C in a 40% moisture environment. One part of the soil sample was air-dried to measure basal soil properties, and the other part was used for microbial molecular analysis.

### 2.2 The field straw decomposition and carbon and nitrogen release experiments

The ditch-buried straw decomposition experiment was conducted using litter nylon bags. Maize straw materials were collected after maize harvesting in 2021 and air-dried. Ten grams of maize straw was cut to 2 cm in length and put into nylon litter bags, which were then sealed via heat sealing. The nylon bags were 6 cm × 10 cm in size and were made of 200 mesh nylon fabric, which permitted the free transfer of microorganisms between the nylon bags and soil. Before maize cultivation in 2021, litter bags containing straw were buried at 10 cm depth in a spatially random design to prevent bags
associated with a given decomposition stage being placed together in space. The litter bags were collected after the harvest in October 2021.

The straw decomposition ratio was calculated based on dry weight loss as (dry initial mass - dry final mass)/dry initial mass. The straw-C concentration was measured by titrimetry after oxidation with a mixture of H$_2$SO$_4$ and K$_2$Cr$_2$O$_7$. Total N, P and K were determined using the Kjeldahl, molybdenum blue colorimetry, and flame photometry methods, respectively. All methods have been described by Lu (2000). The initial and sampled maize straw material properties are shown in Supplemental material Table S2. The amounts of total straw C and N released were calculated by the following equation:

$$\text{The amounts of total straw C and N released} = (\text{initial C (or N) content } \times \text{ dry initial mass} - \text{ final C (or N) content } \times \text{ dry final mass}) \times \text{ aboveground biomass}$$

### 2.3 Measurement of soil properties and assessment of ecosystem services

Soil pH was measured at a soil:water ratio of 1:2.5 (weight/weight). Air-dried soil and 25 ml of deionized water were shaken together for 1 min and left to settle for 30 min, and the soil pH was determined using an electrode. Soil organic carbon (SOC) was measured by titrimetry after oxidation with a mixture of H$_2$SO$_4$ and K$_2$Cr$_2$O$_7$. Total N and P were determined using the Kjeldahl and molybdenum blue colorimetric methods, respectively. All of these methods have been described by Lu (2000).

Microbial biomass C (MBC) and microbial biomass N (MBN) were analysed using the fumigation-extraction method. Ten grams of fresh soil was fumigated with chloroform in the dark for 24 h, and then the fumigated and nonfumigated soils were extracted with 0.5 M K$_2$SO$_4$ and shaken at 200 rpm for 0.5 h. Soil extracts were filtered through a 0.45-µm Millipore filter, and the C and N in the extracts were determined using a multi C/N 3100 analyser (Analytik Jena AG). The C and N contents
in extracts of the nonfumigated soil were subtracted from C and N extracted from the fumigated soil to give the C and N extracted from the soil microbial biomass. Values of 0.45 and 0.54 were used to calibrate the contents of MBC and MBN, respectively (Vance et al., 1987; Wu et al., 1990).

The activities of cellulose and N-acetyl-β-glucosaminidase (NAG) were measured using p-nitrophenyl-β-D-cellobioside and p-nitrophenyl-N-acetyl-β-D-glucosaminide as substrates, respectively. Fresh soil (1.0 g) was mixed with 2.5 mL of 0.2 M acetate buffer (pH 5.0) and 2.5 mL of 0.02 M substrates and then shaken at 200 rpm and 37 °C for 1 h. The reaction was stopped by adding 1 mL of 0.5 M CaCl₂ and 4 mL of 0.1 M Tris buffer (pH 12.0). The mixture was suspended with a vortex, the supernatant was filtered, and the concentration of p-nitrophenol (PNP) was measured by colorimetry at 400 nm. The same procedure was followed for the controls, with the exception that the substrate was added after the incubation, and CaCl₂ and Tris buffer were added (Dick, 2011; Geisseler and Horwath, 2009).

To estimate the greenhouse gas emission potential, we conducted a 60-day incubation experiment. Briefly, 20 g of fresh soil was placed in a 250-mL flask and then sealed with a gas-tight lid that had a rubber stopper in the middle. Gas samples (10 mL) were taken from the headspace of each flask at 1, 3, 7, 15, 30, and 60 days after sealing using a plastic syringe. The gas sample was immediately injected into a preevacuated 10-mL glass vial. Concentrations of methane (CH₄), N₂O and carbon dioxide (CO₂) were determined using a gas chromatograph (Agilent 7890) equipped with a flame ionization detector for CO₂ and CH₄ and a 63Ni electron capture detector for N₂O. The gas standards were provided by the National Research Center for Certified Reference Materials, Beijing, China. The precision for greenhouse gas emission concentrations was ±0.5% based on repeated measurements of gas standards. When the maize plants matured, all plants and grains were harvested from each plot, oven-dried at
60 °C for 48 h and weighed. Aboveground biomass and crop yield were converted into weight per hectare.

We selected 15 soil properties to estimate cropland ecosystem services, i.e., the soil fertility index (SOC, total N, total P, MBC and MBN), greenhouse gas emission amount (mainly CO₂, N₂O and CH₄), straw decomposition and C and N released, soil extracellular enzymes (cellulase and N-acetyl-D-glucosaminidase), and maize biomass (aboveground biomass and crop yield). Generally, SOC, total N and total P are the major soil fertility factors and indicate the present nutrient status in croplands, which can be used to explain soil fertility conditions. Microbial biomass reflects ecosystem productivity. Greenhouse gas emissions are related to climate change, which can be regulated by fertilization regimes and soil microbial activities. Soil extracellular enzymes catalyse the decomposition of a range of organic polymers, resulting in C and N turnover. Maize biomass (such as aboveground biomass and crop yield) reflects soil productivity. As a whole, all of these variables together contributed to the cropland function. To evaluate the function of the cropland ecosystem under different fertilization conditions, we calculated an integrative soil ecosystem multifunctionality index for further analysis. Due to the lack of a specific definition of multifunctionality, we first calculated the Z scores of the 15 measured variables and obtained a multifunctionality value for each plot by averaging the Z scores of the 15 variables.

2.4 DNA extraction and quantification of general fungal ITS, bacterial 16S rRNA and genes encoding cellulose-degrading enzymes

Total DNA was extracted from 0.5 g freeze-dried soil by using a Fast DNA Spin Kit for Soil (MPbio, USA) according to the manufacturer’s instructions and then dissolved in 50 μl of Tris-EDTA buffer. The success of the DNA extraction was characterized by electrophoresis on 1% (wt/vol) agarose
The quantity and quality of DNA were checked using a Nanodrop spectrophotometer (Nanodrop, PeqLab, Germany). The extracted DNA samples were stored at −80 °C before molecular analysis.

Bacterial and fungal abundances were determined to reveal the changes in microbial community compositions. The abundances of bacteria and bacteria fungi were measured according to modified procedures (Fierer and Jackson., 2005). We selected the primers 338F/518R (338F: CCTACGGGAGGCAGCAG; 518R: ATTACCGCGGCTGCTGG) and NSI1/58A2R (NSI1: GTAGTCATATGCTTGTCT; 58A2R: CATTCCCCGTTACCCGTT) for the qPCR assay. The thermal qPCR profiles for the bacteria and fungi were as follows: 95 °C 2 min, 35 cycles (95 °C 30 s, 60 °C 30 s, 72 °C 30 s, 80 °C 15 s), and data collection at 81 °C for 10 s; 95 °C 10 min, 40 cycles (95 °C 15 s, 52 °C 30 s, 72 °C 30 s, 79 °C 30 s), and data collection at 81 °C for 10 s, respectively. The initial concentrations of the two plasmids used as the standards for the bacterial and fungal abundance analyses were $1.22 \times 10^{10}$ and $9.05 \times 10^9$, respectively.

The fungal cbhI gene and bacterial GH48 gene were selected as functional biomarkers of cellulolytic fungi and bacteria, respectively. The primers GH48 F8/GH48 R5 (GH48_F8: 5' - GCCADGHTBGCG ACTACCT - 3; GH48_R5: 5' - CGCCCCABGMSWWGTACCA - 3) and cbhI F/cbhI R (cbhI F: ACCAAYTGCTAYACIRGYAA; cbhI R: GCYTCCCCAIATRTCCATC) were used for the qPCR assay. The abundance of bacterial GH48 and fungal cbhI genes was quantified according to modified procedures (Zhang et al., 2017). The thermal profiles of qPCR for the target genes of GH48 and cbhI were as follows: 95 °C for 5 min, 40 × (94 °C for 30 s, 60 °C for 45 s, and 72 °C for 90 s), and data collection at 84 °C for 10 s; and 94 °C for 4 min, 40 × (94 °C for 45 s, 50 °C for 30 s, and 72 °C for 60 s), and data collection at 81 °C for 10 s, respectively. The initial concentrations of the two plasmids as the standards for bacterial GH48 and fungal cbhI gene abundance analysis corresponded to...
1.85 × 10^{11} and 2.65 × 10^{10} copies g^{-1} dry soil, respectively. qPCR was performed in triplicate, and amplification efficiencies higher than 95% were obtained with r^2 values > 0.99.

2.5 Bacterial 16S rRNA genes and fungal ITS amplification and sequencing

High-throughput sequencing was performed with the Illumina MiSeq sequencing platform (Illumina Inc.). Both the forward and reverse primers were tagged with an adapter and linker sequence, and 8-bp barcode oligonucleotides were added to distinguish the amplicons from different soil samples. The primers 515F (5′-GTGCCAGCMGCCGCGGTAA-3′) and 907R (5′-CCGTCAATTCMTTTRAGTTT-3′) were chosen to amplify the 16S rRNA genes in the V4–V5 hypervariable region. PCR was conducted in a 50-μL reaction mixture containing 27 μL of ddH2O, 2 μL (5 μM) of each forward/reverse primer, 2.5 μL (10 ng) of template DNA, 5 μL (2.5 mM) of deoxynucleoside triphosphates, 10 μL of 5× Fastpfu buffer, 0.5 μL of bovine serum albumin, and 1 μL of TransStart Fastpfu polymerase (TransGen, Beijing, China). The PCR conditions were 94 °C for 5 min; 30 cycles of 94 °C for 30 s, 52 °C for 30 s and 72 °C for 30 s of extension; followed by 72 °C for 10 min (Caporaso et al., 2010).

The fungal ITS1 region was amplified using the primer pair ITS1F (CTTGGTCAATTAGAGGAATTA) and ITS2 (GCTGCGTCTTCATCGATGC) (Ghannoum et al., 2010). The 50-μL reaction mixture of each reaction mix consisted of 1 μl (30 ng) of DNA, 4 μl (1 μM) of each forward/reverse primer, 25 μl of PCR Master Mix, and 16 μl of ddH2O. PCR amplification was conducted at 98 °C for 3 min, followed by 30 cycles (98 °C for 45 s, 55 °C for 45 s, and 72 °C for 45 s), with a final extension at 72 °C for 7 min (Ghannoum et al., 2010). All amplicons were cleaned and pooled in equimolar concentrations in a single tube, after which they were subjected to library preparation, cluster generation, and 250-bp paired-end sequencing on an Illumina MiSeq platform.
The raw sequence data were processed using the Qualitative Insights into Microbial Ecology (QIIME) pipeline (Caporaso et al., 2010). Sequences that fully matched the barcodes were selected and distributed into separate files for the bacterial 16S rRNA and fungal ITS genes. Poor-quality sequences with lengths less than 200 bp (for fungal ITS) and 500 bp (for bacterial 16S) and quality scores less than 20 were discarded, and the chimaeras were removed using the UCHIME algorithm (Edgar et al., 2010). The remaining sequences were assigned to operational taxonomic units (OTUs) with a 97% similarity threshold using UCLUST (Edgar, 2010). Alpha diversity and Bray–Curtis distances for principal coordinate analysis of the soil microbial community were calculated after rarefying all samples to the same sequencing depth.

2.6 Statistical analysis

The soil ecosystem multifunctionality index, crop yields, microbial traits and other relevant soil variables among treatments were subjected to a chi-square test for independence of variance. Significant differences were determined by one-way analysis of variance (ANOVA) based on the post hoc Tukey test at the 5% level. Prior to ANOVA, the normality and homogeneity of variances were tested by the Kolmogorov–Smirnov test and Levene’s test, respectively. If normality was not met, log or square-root transformation was implemented. One-way ANOVA was performed using SPSS 21.0 (SPSS Inc., Chicago, IL, USA).

Nonmetric multidimensional scaling (NMDS) analysis was used to describe and evaluate the microbial community composition. Redundancy analysis (RDA) was performed to visualize the associations between the microbial community composition and selected soil properties. The NMDS and RDA were performed in the “Vegan” package of R (4.0.2). To describe the complex co-occurrence
patterns in various organisms, we constructed co-occurrence networks. We focused on the abundant microbial phylotypes (with average relative abundance > 0.01% for bacteria and fungi) for network construction. Nodes with Pearson correlations greater than 0.70 and $p < 0.05$ were retained. Network visualization between microbial taxa was conducted by Gephi software. To obtain the keystone species of each network, a Zi-Pi plot series was constructed to determine the role of each OTU. According to Deng et al. (2012), the plot includes (a) peripheral nodes ($Z \leq 0.25, P \leq 0.62$), (b) module hubs ($Z > 0.25, P \leq 0.62$), (c) connectors ($Z \leq 0.25, P > 0.62$) and (d) network hubs ($Z > 0.25, P > 0.62$). From an ecological perspective, OTUs in module hubs, connectors and network hubs may be regarded as the microbial keystone taxa of the network systems.

A heatmap was constructed to reveal the associations between microbial traits and fertilizers, soil properties, greenhouse emissions and ecosystem multifunctionality. The random forest algorithm was performed in the R package “RandomForest” to estimate the importance predictors of soil properties and microbial traits on ecosystem multifunctionality.

3. Results

3.1 Cropland ecosystem services

Data collection after a continuous 4-year in situ field experiment under different N input levels revealed changes in cropland ecosystem services (Fig. 1). In terms of soil fertility, compared with the N-limitation treatments (PK and 0.5N+PK), the SOC and total P contents were increased significantly by the N+PK and 0.75N+PK treatments (Fig. 1a, c) ($P < 0.05$), while there were no significant changes in the total N content (Fig. 1b). After straw decomposition (Fig. 1d), the amounts of straw C (Fig. 1e) and N (Fig. 1f) released showed different responses to varying N fertilizer input levels. Generally,
N-rich treatments (N+PK and 0.75N+PK) significantly increased the straw decomposition rate and achieved higher amounts of straw C and N release than the N-limitation treatments ($P < 0.05$). However, there was no significant difference between N+PK and 0.75N+PK. Microbial biomass and function were also sensitive to N fertilizer application (Fig. 1g, h, i and j). The MBC (Fig. 1g) and MBN (Fig. 1h) contents were significantly higher in the N-rich treatments than in the other treatments. However, the highest cellulase activity was observed in the 0.75N+PK treatment, which was significantly higher than that in the other treatments (Fig. 1i) ($P < 0.05$), and the N-acetyl-D-galactosaminidase activity decreased with the reduction in N application (Fig. 1j).

For the ecosystem dis-services (greenhouse gas emissions), with the increase in N fertilizer application levels, CO$_2$ and N$_2$O emissions gradually increased (Fig. 1k, m). No significant difference was observed in CH$_4$ emissions under the different fertilization treatments (Fig. 1l). In addition, the N fertilizer levels also had a strong influence on maize yields and aboveground biomass (Fig. 1n, o). Our results indicated that the N+PK and 0.75N+PK treatments resulted in higher maize yields and aboveground biomass than the other treatments ($P < 0.05$), suggesting that a 25% N fertilizer reduction could be satisfactory for maize growth. As expected, the 0.75N+PK treatment achieved the highest multifunctionality index (0.61), followed by N+PK (0.32), 0.5N+PK (-0.34) and PK (-0.59) (Fig. 1p).

However, although the 0.75N+PK treatment increased the straw N release amount and may meet the requirements for plant growth, the total N input was still dominated by inorganic N input (Fig. S1). Therefore, the N released from the straw cannot offset the deficiency of N fertilizer. Additionally, contrasting N fertilizer input levels significantly changed the stoichiometry of C, N and P (Fig. S2).

Notably, the 0.75N+PK treatment significantly increased the C:N ratio compared with the 0.5N+PK and PK treatments ($P < 0.05$). The lowest C:N ratio was shown for the 0.5N+PK treatment (Fig. S2a).
The N:P and C:P ratios showed no significant difference regardless of nutrient excess or limitation (Fig. S2b and c).

**Fig. 1** The 15 cropland variables and multifunctionality index under different N input levels after straw return. Abbreviations: N+PK, straw return plus regular inorganic N-P-K fertilizers; 0.75N+PK, straw return plus regular inorganic P-K with 25% N fertilizer reduction; 0.5N+PK, straw return plus regular inorganic P-K with 50% N fertilizer reduction; PK, straw return plus regular inorganic P-K without N fertilizer.

### 3.2 Abundances of bacteria, fungi and genes encoding cellulose-degrading enzymes

N fertilizer input levels had marked impacts on the abundances of fungi and bacteria (Table S3).

The highest fungal abundance was observed in the 0.75N+PK treatment, which was significantly
higher than that in the other treatments ($P < 0.05$). The N+PK treatment significantly increased bacterial abundance compared with the PK treatment ($P < 0.05$), while there were no obvious differences among the N+PK, 0.75N+PK and PK treatments. The ratios of fungi to bacteria also showed contrasting responses to N fertilization (Table S3). The 0.75N+PK treatment significantly increased the ratio of fungi to bacteria compared with the other treatments ($P < 0.05$), and the lowest ratio of fungi to bacteria was found in the PK treatment.

Table 1 The abundances of genes encoding cellulose-degrading enzymes across different N fertilizer level treatments after straw return

<table>
<thead>
<tr>
<th>Treatment</th>
<th>cbhI gene abundance ($\times 10^6$ copies g$^{-1}$ soil)</th>
<th>GH48 gene abundance ($\times 10^7$ copies g$^{-1}$ soil)</th>
<th>cbhI: GH48 ratio</th>
</tr>
</thead>
<tbody>
<tr>
<td>N+PK</td>
<td>4.75±0.16 a</td>
<td>1.68±0.01 a</td>
<td>0.28±0.01 a</td>
</tr>
<tr>
<td>0.75N+PK</td>
<td>4.95±0.19 a</td>
<td>1.60±0.04 a</td>
<td>0.31±0.02 a</td>
</tr>
<tr>
<td>0.5N+PK</td>
<td>4.01±0.12 b</td>
<td>1.54±0.08 a</td>
<td>0.26±0.03 b</td>
</tr>
<tr>
<td>PK</td>
<td>3.76±0.13 b</td>
<td>1.40±0.06 b</td>
<td>0.27±0.02 b</td>
</tr>
</tbody>
</table>

The results show means ± standard deviations (n = 3). Different lowercase letters after values indicate significant differences between each treatment, $P < 0.05$. N+PK, straw return plus regular inorganic N-P-K fertilizers; 0.75N+PK, straw return plus regular inorganic P-K with 25% N fertilizer reduction; 0.5N+PK, straw return plus regular inorganic P-K with 50% N fertilizer reduction; PK, straw return plus regular inorganic P-K without N fertilizer.

N fertilizer input levels led to changes in the expression levels of genes encoding cellulose-degrading enzymes (Table 1). The N-rich treatments achieved higher fungal cbhI and bacterial GH48 gene abundance than the N-limitation treatments. In contrast, the highest cbhI gene
abundance was shown in the 0.75N+PK treatment, while the highest GH48 gene abundance was shown in the N+PK treatment. Compared with the PK treatment, the ratio of the fungal cbhl gene to the bacterial GH48 gene increased significantly under the 0.75N+PK treatment ($P < 0.05$).

### 3.3 Co-occurrence network analysis of the microbial community

Regarding fungal alpha diversities, there were no significant differences in the Chao1 index across treatments. The N+PK treatment significantly increased fungal richness compared with the PK treatment ($P < 0.05$) (Table S4). In addition, the PK treatment resulted in lower bacterial richness than the other treatments ($P < 0.05$). No significant difference was observed in the bacterial Chao1 index across treatments (Table S4). NMDS plots showed that diverse N input levels significantly changed the fungal (Fig. S3a) and bacterial communities (Fig. S3b) ($P < 0.05$).

We further conducted network analysis to identify co-occurrence patterns between specific microbial taxa (Fig. 2). The cooccurrence network was aggregated into smaller coherent modules that were examined to determine important module-trait relationships. The present network comprised 1963 nodes (composed of 1520 bacterial taxa and 443 fungal taxa) and 62206 edges with 52.49% positive associations (Fig. 2a). The results showed four dominant ecological modules (1-4) that strongly co-occurred within the multitrophic network, which contributed 86.10% of the whole network. Among the four modules, bacteria accounted for the highest proportion in each module, contributing more than 70% of the total (Fig. 2b). The percentage of edges linking bacteria to bacteria (B-B) was higher than that linking fungi to fungi (F-F) and bacteria to fungi (B-F). The highest proportion of B-B (80.32%) was found in Module 3, while the highest proportion of B-F (32.66%) and F-F (6.00%) was found in Module 4 (Fig. 2c).
Fig. 2 The relationships of microbial module communities with soil ecosystem services and dis-services. Multitrophic network including multiple ecological modules. The colours of the nodes represent different ecological modules (a). OTU number proportions of bacteria and fungi (b). The proportions of the edges linking bacteria to bacteria (B-B), bacteria to fungi (B-F) and fungi to fungi (F-F) in the major ecological modules (c). Links between the specific module communities with soil ecosystem services and dis-services (d). * indicates significance at $P < 0.05$.

Abbreviations: SOC, soil organic carbon; C: N, the ratio of the SOC content to the total N content;
N: P, the ratio of the total N content to the total P content.

Individual nodes represented different roles in the microbial network based on the intramodule connectivity $Z_i$ and the intermodule connectivity $P_i$. $ZP$ plots were constructed to identify the topological roles of each node in the network (Fig. 3a). As shown in Fig. 3b, 113 microbial taxa (81 bacterial species and 32 fungal species) were regarded as connectors, and 43 microbial taxa (39 bacterial species and 4 fungal species) were regarded as module hubs. Specifically, module 2 (54) contained the most keystone taxa, followed by module 1 (38) and module 3 (32).

**Fig. 3** The topological roles of microbial taxa and their effect on the soil multifunctionality index.

The topological role of each OTU was determined according to the scatter plot of within-module connectivity ($Z$) and among-module connectivity ($P$) (a). The distribution of keystone taxa in each ecological module (b). Contribution of bacterial (c) and fungal OTUs (d) to the soil
multifunctionality index. *, ** and *** indicate significance at $P < 0.05$, 0.01 and 0.001, respectively.

3.4 Linkage between microbial traits and soil ecosystem multifunctionality

The heatmap assumed close correlations between fertilizers (N input and straw), as well as soil stoichiometry, and microbial traits (Fig. 4). Overall, the N input level, straw biomass and C:N ratio upregulated the abundance of genes encoding cellulose-degrading enzymes. In addition, N input was positively correlated with bacterial abundance, while a significant correlation was observed between straw biomass and the N input level. The random forest model was also used to identify abiotic and biotic attributes correlated with soil ecosystem multifunctionality (Fig. 5). The model explained 83.89% of the variance in ecosystem multifunctionality. The results indicated that the N input level, straw biomass and soil C:N ratio were the most prominent abiotic factors affecting the ecosystem multifunctionality index, while some biotic factors, such as the abundance of genes encoding cellulose-degrading enzymes, significantly affected the ecosystem multifunctionality index.

Moreover, to clarify the potential main specific drivers of soil ecosystem services, correlations between the microbial physiological traits and soil properties were determined to illuminate the role of the microbial community in soil ecosystem multifunctionality (Fig. 3d). The results indicated that the particular microbial module community was significantly correlated with soil ecosystem services. The communities of modules 1 and 2 and the fungal community in module 4 showed potential in soil ecosystem services (Fig. 3d). Specifically, significant correlations were observed between the SOC content, straw decomposition, straw C:N release, CO$_2$/N$_2$O mineralization and the module 1 community; the module 2 community was positively correlated with greenhouse gas emissions (except
for CH₄); and the fungal community in module 4 was positively correlated with the SOC content, straw
decomposition, straw C/N release and straw biomass. Furthermore, the bacterial and fungal
communities belonging to module 2 and the fungal community belonging to module 3 were
significantly correlated with CO₂ emission, maize yield and straw biomass.

Fig. 4 Heatmap revealing the correlation coefficients between microbial traits with fertilization
and soil stoichiometry. *, ** and *** indicate significance at $P < 0.05$, 0.01 and 0.001,
respectively. Abbreviations: C: N, the ratio of the SOC content to the total N content; N: P, the
ratio of the total N content to the total P content.

At the scale of microbial species, we selected the 20 keystone taxa (10 bacterial and 10 fungal
taxa) with the highest relative abundance for further analysis. The random forest models indicated that
the specific keystone taxa strongly influenced soil ecosystem multifunctionality (Fig. 4c and d).
Bacterial OTU21 (in module 1), OTU240 (in module 2) and OTU6346 (in module 3) were highlighted as essential predictors of soil ecosystem multifunctionality, and fungal OTU22 (module 3) was also found to be an important variable for predicting its changes. Subsequently, the relative abundances of selected keystone taxa were different across different N fertilizer level treatments after straw return (Table S5). The relative abundances of fungal OTU22 and bacterial OTU21 were higher in the N-rich treatments than in the N-limitation treatments. Moreover, compared with the N+PK treatment, the 0.75N+PK treatment increased the relative abundances of fungal OTU22 by 38.20% and bacterial OTU21 by 40.63%.

Fig. 5 Contribution of abiotic and biotic variables to the soil multifunctionality index. *, ** and *** indicate significance at $P < 0.05$, 0.01 and 0.001, respectively. Abbreviations: C: N, the ratio of the SOC content to the total N content; N: P, the ratio of the total N content to the total P content.
4. Discussion

4.1 Effect of N fertilizer reduction on cropland ecosystem services after straw return

Soil fertility, straw decomposition, C and N release amounts, and crop productivity showed an overall positive effect with the increase in the N fertilizer input level and together contributed to ecosystem services. Moreover, high N applications may also cause ecosystem dis-services due to the surge in greenhouse gas emissions (Fig. 1). Our results indicated that the soil fertility index (SOC, total N and P contents) increased under N-rich treatments as a result of high net primary production, in accordance with previous reports (Liu et al., 2010; Williams et al., 2013). Higher microbial biomass C and N, as well as relevant enzyme activities, were also observed under N-rich treatments, indicating the strong positive impact of abundant N fertilizer application (Fig. 1g, h, i, j). It was reported that straw return with N fertilizer application can stimulate microbial activity and promote biomass accumulation (Treseder, 2008). The substantially increased straw decomposition and straw C and N release under N-rich treatments may be primarily attributed to the activation of microbial activity (Fig. 1d, e, f), which is consistent with previous research (Ramirez et al., 2012). Recent studies have also proven that rational N input can stimulate microbial ex vivo production of extracellular enzymes to accelerate straw decomposition and nutrient transformation (Chen et al., 2016). Moreover, it is well known that fungi have high nutrient utilization efficiency; thus, more straw-derived C and N would be stored in soil under N-rich treatments than under N-limited treatments (Hou et al., 2020). Higher N availability is also the premise of straw decomposition and SOM formation due to the microbial “stoichiometry decomposition” theory, while the “N-mining” theory in N-limitation treatments reveals that oligotrophic species (such as K-strategists) degrade native SOM because of the lack of N fertilizer inputs (Chen et al., 2014). Finally, the increases in SOC, total N, and P contents and straw C and N
release, as well as microbial biomass and function, are commonly attributed to high aboveground biomass and maize yields (Fig. 1n, o), which are favourable from the viewpoint of ecosystem services. However, the overuse of N inputs also causes ecosystem dis-services, such as unintended environmental consequences (Tang et al., 2019). In the present study, greenhouse gas emissions were quantified to evaluate the ecosystem dis-services under different N fertilizer input levels (Fig. 1k, l, m). Straw return with N fertilizer addition might be the crucial driver of CO$_2$ and N$_2$O emissions from agroecosystems and has been widely studied in previous literature (Gregorich et al., 2005). CO$_2$ and N$_2$O emissions increased significantly compared with those under the PK treatment, likely by stimulating the activity of copiotrophs when sufficient C and N substrates were provided. For example, Dieleman et al. (2010) implied that N fertilizer addition significantly increased CO$_2$ and N$_2$O by increasing bacterial abundance through meta-analysis and field experiments, respectively. Qiu et al. (2019) indicated that the emission of CO$_2$ enhanced root and mycorrhizal N uptake and increased N$_2$O emissions, which was related to the changes in the soil denitrifier community composition in favour of N$_2$O-producing taxa (nirK- or nirS-type). In addition, there was no difference in CH$_4$ emissions among treatments, although contradictory results have been widely reported in previous literature (Tang et al., 2019). Mapanda et al. (2011) and Liu et al. (2012) indicated that the emission of CH$_4$ depended highly on the soil water content in maize crops, which is in line with our results.

In summary, compared with the N+PK treatment, the 0.75N+PK treatment supported multiple ecosystem services, including promoting soil fertility, straw nutrient release and microbial activity and alleviating greenhouse gas emissions (Fig. 1p). Therefore, a reduction of 25% in chemical N fertilizer input with straw return may be the appropriate regime to promote ecosystem services in meadow soils on the Northeast China Plain.
4.2 Responses of the microbial composition and function to straw return with N fertilizer reduction

Fungal and bacterial abundances, as well as the ratio of fungi to bacteria, were sensitive to the changes in the N fertilizer input levels (Table S3 and Fig. 2). Straw addition with N fertilizer input supplied enough C and N for microbial metabolism, thus promoting microbial proliferation (Chen et al., 2016). Generally, bacterial abundance decreased with reduced N fertilizer input. This is mainly because bacteria are more sensitive to N availability than fungi, which is in line with a previous study (Ramirez et al., 2020). Interestingly, it is worth noting that a 25% reduction of N fertilizer significantly increased fungal abundance compared with regular N inputs. This result might be attributed to the negative effect of excess N fertilizer (Wan et al., 2015). Moreover, Ning et al. (2020) demonstrated that the C:N ratio was the pivotal factor in fungal community compositions after performing 7 long-term field experiments under different fertilization conditions across China and reported a significant positive correlation between them. Therefore, the 0.75N+PK treatment with a higher C:N ratio may facilitate the proliferation of microorganisms and promote an increase in microbial abundance.

Subsequently, our results showed that N-rich treatments resulted in higher microbial cellulose-degrading gene abundances than the PK treatment (Table 1), which demonstrated the irreplaceable role of N inputs in straw degradation (Zhang et al., 2017). Additionally, compared with bacterial GH48 gene abundance, the increase in fungal cbhI gene abundance required adequate N fertilizer inputs and was regulated by the soil C:N ratio, which suggests that rational N fertilizer inputs could promote fungal function for further degradation of recalcitrant straw components (Hou et al., 2020). Therefore, the ratio of cbhI gene abundance to GH48 gene abundance was higher under 0.75N+PK than under the N-limitation treatments since the increased expression of a fungal
cellulose-degrading gene implies more straw C and N release. Our results indicated that adequate N fertilizer upregulated fungal and cbhI gene abundances, which may lead to multiple ecosystem services. It is therefore necessary to further explore the potential associations between microbial traits and ecosystem services under diverse N fertilizer input levels.

4.3 Linkages of cropland ecosystem services with microbial traits

To clarify the effect of abiotic and biotic factors on soil ecosystem services, we then quantified the contributions of abiotic and biotic attributes to the ecosystem multifunctionality index across N input treatments (Fig. 4 and 5). Biotic factors, such as cbhI and GH48 gene abundances, as well as abiotic factors, including the C:N ratio, straw biomass and N input level, are also pivotal regulators of ecosystem multifunctionality (Fig. 5). In general, promoting the rapid degradation of straw is an important way to convert straw-C into SOM, thus improving soil fertility, aboveground biomass and crop yield. In addition, fungi have a higher C utilization efficiency than bacteria; thus, a high fungal cbhI gene abundance may achieve better soil multifunctionality (Hou et al., 2020). For abiotic factors, the soil C:N ratio, straw biomass and N fertilizer input are always regarded as the main indicators of soil fertility and health, likely due to providing various nutrient accessibilities and influencing the microbial community composition (Ning et al., 2020).

Numerous studies have shown that core microbiota play a vital role in maintaining the stability of soil microbial function and the complexity of microbial networks and then promoting soil nutrient cycling ecosystem services (Ghamoum et al., 2015), and keystone species may show great explanatory power in terms of specific network (or module) structure and function (Chen et al., 2019b). Heatmaps and random forest models were used to illuminate the relationships of module communities with ecosystem services (Fig. 2d and Fig 3c, d). In the present study, Terrimonas (bacterial species in
module 1) and Lasiosphaeriaceae (fungal species in module 3) were detected as the keystone taxa in influencing soil multifunctionality of the cooccurrence network (Table S5). A previous study demonstrated that straw addition significantly increased the relative abundance of Lasiosphaeriaceae, which implied straw decomposition ability (Song et al., 2020). Afterwards, Lasiosphaeriaceae was proven to promote straw-derived C and N accumulation by secreting multiple extracellular enzymes (Guo et al., 2022). Meanwhile, Sun et al. (2023) revealed that Lasiosphaeriaceae abundance was regulated by the soil C:N ratio, especially changes in mineral N. Therefore, Lasiosphaeriaceae can effectively promote straw degradation and straw C and N release while driving the function and community of module 1, which is consistent with our results (Fig. 2d). However, relatively few studies have focused on the function of Terrimonas, so this study focused on Chitinophagaceae. As reported in the previous literature, straw return was the main method to increase Chitinophagaceae abundance (Li et al., 2021). Furthermore, Chitinophagaceae was indicated to have a strong ability to accumulate soil C and N and degrade cellulose (Zhong et al., 2022), facilitating production improvement by regulating the module 3 community and function, which is in line with our results (Fig. 2d).

Overall, straw return with sufficient N fertilizer application can increase the C:N ratio and stimulate microbial traits, which ultimately achieve soil ecosystem multifunctionality (Fig. 6). Straw return without enough N supply cannot support ecosystem services due to the decomposition of native SOM and the out-of-balance microbial community composition, according to the “N-mining” theory (Chen et al., 2014); straw return with sufficient N application (N+PK and 0.75N+PK) can promote soil fertility, straw release, microbial activity and crop productivity, which can be explained by the “stoichiometry decomposition” theory (Chen et al., 2014). Meanwhile, N+PK also caused more serious ecosystem dis-services, such as greenhouse gas emissions, than the 0.75N+PK treatment. Moreover,
compared with the N+PK treatment, the 0.75N+PK treatment increased the soil C:N ratio and stimulated microbial module 1 and 3 communities function, *cbhl* gene abundance, and keystone taxa abundances, which were significantly positively correlated with soil ecosystem multifunctionality. Our study provides evidence that a 25% reduction of chemical N fertilizer after straw return was the optimal agronomic measure for ecosystem services in meadow soil on the Northeast China Plain.

Fig. 6 A graphical sketch of the changes in ecosystem services and potential microbial mechanisms in response to different chemical N fertilizer application rates after straw return. N, nitrogen; SOM, soil organic matter

5. Conclusion

Straw return combined with different chemical N fertilizer application rates significantly changed ecosystem services and dis-services. Collectively, our work indicates that compared with the N+PK treatment, straw return with a 25% reduction in chemical N fertilizer has the potential to improve ecosystem services by maintaining soil fertility, productivity, microbial biomass and function,
promoting straw decomposition and C and N release and alleviating greenhouse gas emissions. The 0.75N+PK treatment achieved higher soil ecosystem multifunctionality than all other treatments. In addition, the N input level, straw biomass and soil C:N ratio can upregulate the abundances of the cbhl and GH48 genes, which may together achieve soil ecosystem multifunctionality.

Furthermore, the changes in multiple soil ecosystem services were strongly associated with microbial module communities and keystone taxa. The relationships between ecosystem services and microbial traits were examined here to confirm that the Lasiosphaeriaceae driving the function and structure of the module 1 community leads to the promotion of straw degradation and straw C and N release, while Terrimonas driving the function and structure of the module 3 community probably contributes to production improvement under 0.75N+PK treatment. Therefore, a 25% reduction in chemical N fertilizer with straw return might be a win–win strategy that not only produces considerable ecological benefits for the pedosphere and atmosphere but also reduces fertilizer expenditures in meadow soil on the Northeast China Plain.
Declaration of competing interests

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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References


Kandeler, E., Manning, P., Setala, H., and Jones, T.H.: Discontinuity in the responses of ecosystem
processes and multifunctionality to altered soil community composition, P Natl Acad Sci USA., 111,
Betley, J., Fraser, L., Bauer, M., Gormley, N., Gilbert, J.A., Smith, G., and Knight, R.:
Ultra-high-throughput microbial community analysis on the Illumina HiSeq and MiSeq platforms, Isme
interaction with keystone taxa induced negative priming under biochar amendments. Microbiome., 7,
Chen, L., Redmile-Gordon, M., Li, J.W., Zhang, J.B., Xin, X.L., Zhang, C.Z., Ma, D.H., and Zhou,
Y.F.: Linking cropland ecosystem services to microbiome taxonomic composition and functional
composition in a sandy loam soil with 28-year organic and inorganic fertilizer regimes, Appl Soil Ecol.,
Chen, R.R., Senbayram, M., Blagodatsky, S., Myachina, O., Dittert, K., and Lin, X.G., Blagodatskaya,
E., Kuzyakov, Y.: Soil C and N availability determine the priming effect: microbial N mining and
Chen, Y.L., Chen, L.Y., Peng, Y.F., Ding, J.Z., Li, F., Yang, G.B., Kou, D., Liu, L., Fang, K., Zhang,
B.B., Wang, J., and Yang, Y.H.: Linking microbial C: N:P stoichiometry to microbial community and
abiotic factors along a 3500-km grassland transect on the Tibetan Plateau, Global Ecol Biogeogr., 25,

de Bello, F., Lavorel, S., Diaz, S., Harrington, R., Cornelissen, J.H.C., Bardgett, R.D., Berg, M.P.,
Wardle, D.A., and Harrison, P.A.: Towards an assessment of multiple ecosystem processes and
services via functional traits, Biodivers Conserv., 19, 2873-2893, doi: http://doi.

Dieleman, W.I.J., Luyssaert, S., Rey, A., De Angelis, P., Barton, C.V.M., Broadmeadow, M.S.J.,
Broadmeadow, S.B., Chigwerewe, K.S., Crookshanks, M., Dufrene, E., Jarvis, P.G., Kasurinen, A.,
Kellomaki, S., Le Dantec, V., Liberloo, M., Marek, M., Medlyn, B., Pokorny, R., Scarascia-Mugnozza,
G., Temperton, V.M., Tingey, D., Urban, O., Ceulemans, R., and Janssens, I.A.: Soil N modulates soil
C cycling in CO2-fumigated tree stands: a meta-analysis, Plant Cell Environ., 33, 2001-2011, doi:

2011.

Dominati, E., Patterson, M., and Mackay, A.: A framework for classifying and quantifying the natural
capital and ecosystem services of soils, Ecol Econ., 69, 1858-1868, doi: http://doi.

Duan, Y., Chen, L., Li, Y.M., Wang, Q.Y., Zhang, C.Z., Ma, D.H., Li, J.Y., and Zhang, J.B.: N, P and
straw return influence the accrual of organic carbon fractions and microbial traits in a Mollisol,

Edgar, R.C.: Search and clustering orders of magnitude faster than BLAST, Bioinformatics., 26,


Rhymes, J., Jones, L., Wallace, H., Jones, T.G., Dunn, C., and Fenner, N.: Small changes in water


Figure captions

Fig. 1 The 15 cropland variables and multifunctionality index under different N input levels after straw return. Abbreviations: N+PK, straw return plus regular inorganic N-P-K fertilizers; 0.75N+PK, straw return plus regular inorganic P-K with 25% N fertilizer reduction; 0.5N+PK, straw return plus regular inorganic P-K with 50% N fertilizer reduction; PK, straw return plus regular inorganic P-K without N fertilizer.

Fig. 2 The relationships of microbial module communities with soil ecosystem services and dis-services. Multitrophic network including multiple ecological modules. The colours of the nodes represent different ecological modules (a). OTU number proportions of bacteria and fungi (b). The proportions of the edges linking bacteria to bacteria (B-B), bacteria to fungi (B-F) and fungi to fungi (F-F) in the major ecological modules (c). Links between the specific module communities with soil ecosystem services and dis-services (d). * indicates significance at $P < 0.05$. Abbreviations: SOC, soil organic carbon; C: N, the ratio of the SOC content to the total N content; N: P, the ratio of the total N content to the total P content.

Fig. 3 The topological roles of microbial taxa and their effect on the soil multifunctionality index. The topological role of each OTU was determined according to the scatter plot of within-module connectivity ($Z$) and among-module connectivity ($P$) (a). The distribution of keystone taxa in each ecological module (b). Contribution of bacterial (c) and fungal OTUs (d) to the soil multifunctionality index. *, ** and *** indicate significance at $P < 0.05$, 0.01 and 0.001, respectively.

Fig. 4 Heatmap revealing the correlation coefficients between microbial traits with fertilization and soil stoichiometry. *, ** and *** indicate significance at $P < 0.05$, 0.01 and 0.001,
respectively. Abbreviations: C: N, the ratio of the SOC content to the total N content; N: P, the ratio of the total N content to the total P content.

**Fig. 5** Contribution of abiotic and biotic variables to the soil multifunctionality index. *, ** and *** indicate significance at $P < 0.05$, 0.01 and 0.001, respectively. Abbreviations: C: N, the ratio of the SOC content to the total N content; N: P, the ratio of the total N content to the total P content.

**Fig. 6** A graphical sketch of the changes in ecosystem services and potential microbial mechanisms in response to different chemical N fertilizer application rates after straw return. N, nitrogen; SOM, soil organic matter
Table 1 The abundances of genes encoding cellulose-degrading enzymes across different N fertilizer level treatments after straw return

<table>
<thead>
<tr>
<th>Treatment</th>
<th>cbhI gene abundance (×10^6 copies g⁻¹ soil)</th>
<th>GH48 gene abundance (×10^7 copies g⁻¹ soil)</th>
<th>cbhI:GH48 ratio</th>
</tr>
</thead>
<tbody>
<tr>
<td>N+PK</td>
<td>4.75±0.16 a</td>
<td>1.68±0.01 a</td>
<td>0.28±0.01 a</td>
</tr>
<tr>
<td>0.75N+PK</td>
<td>4.95±0.19 a</td>
<td>1.60±0.04 a</td>
<td>0.31±0.02 a</td>
</tr>
<tr>
<td>0.5N+PK</td>
<td>4.01±0.12 b</td>
<td>1.54±0.08 a</td>
<td>0.26±0.03 b</td>
</tr>
<tr>
<td>PK</td>
<td>3.76±0.13 b</td>
<td>1.40±0.06 b</td>
<td>0.27±0.02 b</td>
</tr>
</tbody>
</table>

The results show means ± standard deviations (n = 3). Different lowercase letters after values indicate significant differences between each treatment, P < 0.05. N+PK, straw return plus regular inorganic N-P-K fertilizers; 0.75N+PK, straw return plus regular inorganic P-K with 25% N fertilizer reduction; 0.5N+PK, straw return plus regular inorganic P-K with 50% N fertilizer reduction; PK, straw return plus regular inorganic P-K without N fertilizer.