Response to comments by anonymous referee #1

We thank the reviewer for her/his insightful comments and questions.

This work studied the effects of drying-wetting, soil aggregate size, litter addition and plant on N_2O flux from floodplain soils. The authors used model soils and mesocosm experiments to conduct the research. As far as I can say, there are still much more space can be improved for this manuscript.

In general, it is interesting to know how soil N_2O flux are controlled by different environmental factors. However, there are already many studies conducted in no matter drying-wetting and soil aggregation, or litter addition and vegetation effects. What the knowledge gaps do you want to fill? It should be clarified in the introduction part.

Reply: We concur with the reviewer that the specific objectives of this study were not sufficiently well stated. While the effects of microhabitats related to soil aggregates, the detritusphere and plant-soil interactions in the rhizosphere on N_2O emissions from soils have been studied individually, little is known about their relative effects and interactions. In a mesocosm study, we investigated this aspect for the hot moments of N_2O emissions from floodplain soils during the drying phase after flooding. In particular, aggregate size effects have not been investigated in this context (as stated on lines 80ff). A particular novel aspect of the study is the minimization of the potentially confounding factor “soil structure” by mixing a given aggregate size fraction with inert material replacing the removed smaller or larger fraction. As stated on line 71ff, previous studies employing isolated aggregate size fractions have provided partially inconsistent results.
The innovative aspects of objectives,will be further clarified in the introduction of the revised manuscript, with emphasis on the relevance of the research, and addressing also the potential with regards to filling knowledge gaps.

Here are technical questions:

1. Line 14-15, it is not accurate to write the buried organic matter and rhizosphere processes. Actually, the experiments were about litter addition and plant vegetation. It still takes several steps from litter to organic matter. And also, you didn’t took the rhizosphere samples.

R1: We agree that the term “buried organic matter” is too unspecific. Although, sensu stricto, litter is “organic matter” as well, it indeed might be confused with further decomposed and transformed “soil organic matter”. We therefore have replaced “buried organic matter” with “buried litter”. We checked the entire manuscript and this is the only place where we used this rather unspecific term.
We also agree that at this point “rhizosphere processes” should be replaced by “plant-soil interactions”, even though in the later discussion mainly rhizosphere processes per se are invoked to explain the observed plant effects.

2. L148, for soil pH measurement, normally it is 10 g soil was mixed with 25 mL solution. The authors used 20 mL of solution,any references? The solution can be water or CaCl2, as far as I know, for alkaline soil, it it better to use water. In this study, the soil pH were ~ 8, any reasons to choose CaCl2?

R2: There are several soil-to-solution ratios recommended in the literature, among them also 1:2.5 (Blume et al. 2010. Scheffer/Schachtschabel – Lehrbuch der Bodenkunde, 16th ed., p. 151), or 1:1 (Thomas G.W. 1996. «Soil pH and Soil acidity» In: Sparks et al. (eds.) Methods of soil analysis – 3. Chemical methods. SSSA Book Series 5, pp. 475ff.). A soil-to-solution ratio of 1:2 for mineral soil samples – as has been used in our laboratories since more than 30 years – is also recommended by one of the newest method handbooks: Hendershot et al.

(2008) “Soil reaction and exchangeable acidity” In: Carter, M.R. (ed.) Soil sampling and methods of analysis. 2nd ed., Can. Soc. Soil Sci., chapter 16. Furthermore, this handbook, citing several individual studies, recommends $CaCl_2$ as suspending solution with several advantages over water, in particular also for agricultural soils whose pH is often comparatively high. There is no mentioning in this, or any of the other cited references, of a disadvantage in using $CaCl_2$ for carbonate containing soils.

More generally, soils are heavily buffered systems and the measured pH should be virtually independent of such small variation in ionic strength.

3. Have the authors ever considered the emission/uptake of N2O by the aboveground of plant? There are already many studies in this field, such as: Smart D R, Bloom A J. Wheat leaves emit nitrous oxide during nitrate assimilation[J]. Proceedings of the National Academy of Sciences, 2001, 98(14): 7875-7878. In this study, the authors measured N2O flux from the mesocosm have both soil and plant. This flux cannot be called soil flux, but may be soil/plant flux?

R3: In the introduction (line 96ff) we considered potential bypassing of the soil matrix by N_2O fluxes via plant-internal aeration channels (aerenchyma). This phenomenon is well documented for *Poaceae* such as the Genus *Oryza* or *Phalaris arundinacea*. However, for willows (*Salix sp.)* such a process has, to our knowledge, not been documented yet. Although, considering that also adventitious roots of *Salix* species contain aerenchyma, we cannot exclude this process to occur in our case, our results do not indicate an enhanced N_2O emission via the plant, since we observed the lowest flux rates as well as lowest total integrated emissions in the mesocosms with plants. Therefore we conclude that in our experiment, such a process, if present, was of minor importance in terms of modulating net N_2O fluxes to the atmosphere. However, we agree that the possibility that part of the N_2O fluxes from the planted soils occurred via plant-internal channels should be mentioned in discussion section 4.3. We also agree that emission fluxes should be termed “soil/plant flux” or “ecosystem flux” instead of “soil flux”. Although nowhere in the manuscript we have used the term “soil flux”, we agree that we need to clarify at respective prominent places in the manuscript that in the case of the treatments with willow emissions/fluxes relate to the whole soil/plant system and not to the soil alone.

4. L274, the author can show the data in support information.

R4: we will upload a file containing the supplementary information and adjust the text accordingly.

5. L313-315, the authors didn’t check the statistics difference of soil chemical/physical properties between different treatments. Therefore, the hypothesis is not really correct before statistics analysis were done.

R5: The comparison of the initial physicochemical properties by t-tests with Welch’s correction showed statistically significant differences for the C:N ratio and pH. However, C:N ratios of 12 and 16 can be considered ecologically similar in terms of soil organic matter degradability, in particular since both Corg and total N do not differ that much. The higher pH in the macroaggregated model soil is probably due to a higher carbonate content, which also is not expected to strongly affect biogeochemical processes of the N cycle. These remarks will be added in the revised manuscript, and a new column will be added in table 1 with the results of the statistical analyses.

6. L346-347, Actually WFPS-SA value were not decreased to pre-flood even until the end of experiments (Fig. 2 a and b). The explanation might be low diffusion rate of N2O in SA treatments caused reduction of N2O to N2?

R6: Considering the high WFPS in the SAU treatment, the referee's remark represents a valid explanation for the observed low fluxes under the given circumstances. However, the relatively high redox potentials, which we invoke here, argue against sufficient anoxia for complete reduction of N_2O to N_2. Nevertheless, we will include this aspect in the discussion in section 4.1. of the revised manuscript.

7. L409, delete one dot

8. L457, delete DOI

9. Table 2, it would be better to explain the meanings of LAU, SAU....in the table caption.

10. L638-639, no dotted line in Fig. 3?

11. Fig. 2, it would be better to put WFPS in the right Y axis. And put WFPS-LA, WFPS-SA....in the figure legend.

12. Fig. 3e, the data are not completely shown.

13. Fig. 4, would be better to have the same unit (μM) for nitrate and nitrite/ammonium

R7-13: the authors consent with all these remarks and will make changes to the revised manuscript accordingly.

Response to comments by Y. A. Teh (Referee)

We thank the reviewer for his supportive evaluation, insightful comments and questions. Addressing them will strongly improve the manuscript.

GENERAL COMMENTS

This is a creative and interesting process-based experiment that uses different aggregate treatments (i.e. micro- versus macro-aggregate dominated) and plant-soil treatments (i.e. a gradient of "plant influence," from rhizosphere to detritus-affected soil to plant-free soil) to determine how differences in soil structure and various levels of plant influence potentially influence N2O dynamics in soil. The factorial experimental design is powerful because it enables the investigators to assess not only main effects, but also evaluate the potential importance of synergistic effects among different treatments. Overall, it is my view that this paper was clearly written, with a well-justified experimental design, and a logical analysis of the data. The introduction to the paper clearly explains the basis and wider significance of this research, while the methods section explains the overall approach taken with clarity. The results section documents the main findings of the work succinctly, while the discussion takes a reasonable (and not overly speculative) approach to data interpretation, informed by the authors' grasp of the current literature. The investigators' comprehensive measurement of a range of environmental parameters is to be commended and enables them to make logical inferences about the role of different treatments and environmental factors in regulating N2O dynamics during different parts of the simulated water cycle. In particular, the investigators make good use of redox potential measurements to evaluate how changes in redox/O2 availability could be driving N dynamics along the "plant influence" gradient that they have created in the laboratory.

However, while I am generally supportive of this research and believe it will make a valuable contribution to the wider body of knowledge on this topic, I do have a few general remarks that I believe need to be addressed before this paper can go forward to publication. First, I think the authors need to be open and transparent about the potential limitations of their research. For example, the soil structure treatments represent two extremes (large versus small aggregates), whereas in reality micro- and macro-aggregates would be mixed together. The authors need to explain how their experimental treatment could relate or correspond to real-world conditions, drawing if possible on pre-existing field or laboratory data (see points 1 and 5 below).

R I: We agree that including a discussion of the implicit limitations of our experimental approach with respect to natural conditions will contribute to a better evaluation of the results of our study, and we thus will include this in a revised version of the manuscript.

By investigating two pedogenetically well-defined aggregate size fractions (4000 – 250 µm and 250 – 0 µm; Tisdall and Oades, 1982) separately – but with soil structure

kept similar by replacing the removed fraction by inert material of the same size - , we aimed at evaluating the individual potential of these fractions to offer conditions for the soil microbial community to form N_2O. Following the reviewer's suggestion, we propose to include a discussion of how these conditions relate to real-world conditions as follows. As detailed in our response R1 below, these two size fractions represent significant "components" both of our investigated original soil and of most other soils. However, we intentionally excluded interactions between the two soil aggregate size factions to assess the individual potential of each faction separately. Therefore we can neither assess any interactions between large and small aggregates, nor such with soil structures larger than 4mm, which all may also be important for N_2O emissions under natural conditions. Since we have no data related to this, we prefer not to speculate about such effects in our paper.

Likewise, the authors need to be clearer about the limitations underlying their rhizosphere (Salix) treatment. It is difficult to generalise more widely about the effects of plant rhizospheres on N dynamics without examining a range of different plants (including single and multi-species mixtures), in order to tease-apart individual species effects from generic rhizosphere effects (see point 6 below); I think it is important, in the revised version of this text, that the authors acknowledge this limitation and spend a bit more time exploring what they believe could be more widely generalisable from their study, rather than what is species-specific.

R II: For a reply the reader is kindly referred to R6

Second, I do not believe that the authors have fully exploited their experimental design in the analysis of their data, and sincerely believe that more could be done to examine these data in greater depth. For example, as mentioned above, one of the strengths of a factorial experimental design is that the investigators can establish if there are synergistic interactions among different experimental treatments (e.g. aggregate X rhizosphere effects). However, the investigators do not appear to have examined if interactions among treatments occurred, or at least these findings are not reported if these tests were conducted. Moreover, I would suggest that the authors try more complex multivariate models to analyse their data; for instance, using approaches such as analysis of co-variance (ANCOVA), generalized linear models, or mixed effects models. The benefit of these more comprehensive multivariate models is that they enable the investigator to establish the relative importance of different treatments and continuous environmental variables in regulating flux.

R III: We fully agree that an experiment has to be analyzed according to its experimental design. In our case, this includes the interaction of aggregate size and soil treatment (unamended, litter addition, plant presence). We in fact have included this term in all ANOVA models, but failed to report the results when the term was not statistically significant or only weakly significant. We will fix this in the revised version. The structure of our experimental treatments is not hierarchical so that no mixed model is required. Such a model would only be necessary if one would analyse the

time series data, i.e. if one had several values per microcosm. We have considered this but decided not to do so, for the following reasons:
(1) our focus was on the *average response* during distinct phases that we have identified in our time series, in particular during "hot moments" after wetting; working with average time-series data provides an answer to hypotheses about whether total emissions during this period, for example, differ between treatments; in other words, our hypotheses were about cumulated fluxes during a period, and we therefore carried out these analyses at this level.
(2) the processes we observed are extremely dynamic; fitting a full time series model would almost certainly have resulted in significant time x treatment interactions – such effects would primarily be driven by the peak values of e.g. N_2O emissions after wetting; whether treatment differences for these single measurements reflect true differences in time and extent of peak fluxes is uncertain… it in fact is very likely that the true peak occurred a short time before or after these measurements, and this may be treatment specific. Again, we were not interested in whether the maximum flux occurred a bit earlier or later in time (this may not be reproducible anyways), but whether total emissions during the hot moment changed. Working with such aggregated data solves the problem of subtle shifts in emission timing, and gives extreme values much less weight.
(3) the proper modelling of the time series is very complicated: this involved heterogeneous variances (because large values scatter more) and the modelling of serial correlations (because subsequent values are not independent). On the time-aggregated scale, these problems do not occur. We also could log-transform the data to compare the treatments, which was not possible on the raw data because (a) negative values occurred due to measurement error, and (b) we were asking questions about total fluxes (e.g. grams of N_2O emitted) and not relative effects.
In summary, we agree that more complex analyses can potentially be done. However, we have deliberately focused on (1) the aggregation level that matched the questions we were asking, and (2) the aggregation level at which statistical procedures were robust. We agree that we did not document this very well and propose to address this in the revision.

Third, I agree with the first referee that the authors need to spend a bit more time clearly highlighting what knowledge gaps this paper fills. As the first referee indicates, there are already existing studies that have examined the individual effects of all the variables discussed here. In order to make this paper more impactful, the authors need to articulate how this specific study is unique or advances our current state-of-knowledge (e.g. does the factorial design add knowledge or insight?). Specific comments are provided in the section below.

R IV: We concur with both reviewers that the specific objectives of this study were not sufficiently well stated. As mentioned in our response to Reviewer 1, this aspect will be addressed. We will clarify that, while the effects of microhabitats related to soil aggregates, the detritusphere and plant-soil interactions in the rhizosphere on N_2O emissions from soils have been studied individually, little is known about their relative

effects and interactions. In our mesocosm study, we investigated this aspect for the hot moments of N_2O emissions from floodplain soils during the drying phase after flooding. In particular, aggregate size effects have not been investigated in this context (as stated on lines 79f). A particular novel aspect of the study is the minimization of the potentially confounding factor "soil structure" by mixing a given aggregate size fraction with inert material replacing the removed smaller or larger fraction. As stated on line 71ff, previous studies employing isolated aggregate size fractions have provided partially inconsistent results possibly linked to some extent to the changes in soil structure by aggregate separation.
The better specified objectives and novel aspects will be included in the introduction of the revised manuscript.

SPECIFIC COMMENTS
1. Lines 136-137: For experimental purposes, the investigators have created quasi-artificial system conditions, with treatments either containing macro- or microaggregates. While I fully understand why this was done, it would be useful to understand (even qualitatively) how close or far from reality these treatments are. For example, what was the proportion of macro- and micro-aggregates under natural conditions?

R1: The original floodplain soil consisted of 18.5 ± 4.6 % aggregates smaller than 250 µm and 81.5 ± 4.6 % macroaggregates (mean ± sd; n = 10). We composed our model soils of a 1:1 mixture of isolated aggregates and inert matrix material. This is different from the original soil composition, but well within the range of published top soil aggregate size distributions (e.g. Cantón et al., 2009; Gajić et al., 2010; Six et al., 2000). 50% microaggregates may be more than what is found in most natural or agricultural soils. Nevertheless, we chose to use equal amounts of small and large aggregates to be able so separate effects of aggregate size from effects of aggregate amount (soil mass). To reflect these reasonings, we propose to discuss the distribution of small and large aggregates in the original soil (material and method section of the revised manuscript). The discussion of relevance would be added to the discussion in section 4.1 and in the conclusions. For additional considerations on the effect of flood disturbance on small-scale heterogeneity and dynamics of aggregate size distribution see R5 below.

2. Line 173: Clarity of expression; consider revising this section to read "The mesocosm experiment had a factorial experimental design consisting of two factors (model soil and plant-soil treatment), with the first factor containing two levels (macroaggregates, microaggregates) and the second factor containing three levels (unamended, litter added, plant present). This experimental design resulted in six treatments, each replicated six times."

R2: The authors concur with this remark and will adjust this part accordingly

3. Line 179-180: What was the rationale for autoclaving the leaves? Under natural conditions, these leaves would contain their own microbial community which could contribute to N_2O dynamics, and autoclaving means that the results will be biased towards the activity of the soil community (or, spore-forming phyllosphere microbes able to resist the effects of autoclaving).

R3: Since we specifically wanted to test the effect of additional labile C available to the N_2O producing or consuming soil microbial community, we decided to eliminate, or at least reduce the effect of and interaction with the phyllosphere of the collected leaves by sterilization. We are aware that this introduces a certain bias. However, so far there are no direct effects of the phyllosphere community on N_2O production described in the literature. The only role of these organisms in plant-atmosphere interactions reported in the literature is in capturing/consuming methane and/or volatile organic carbon compounds (Bringel and Couée, 2015). On the other hand, we cannot say anything about potential effects of interactions between the phyllosphere and soil communities on N_2O production/consumption. These remarks will be added to the discussion section of the litter effects, 4.2., in the revised version of the manuscript.

4. Lines 221-232: Further detail on the statistical analyses are required here. For example, what were the independent variables used in the ANOVA? Did the model include interaction terms? Given that sampling was conducted over different periods of time, did the authors use a repeated measures ANOVA, to account for the effects of time?

R4: The independent variables for the two way ANOVA were SOIL TREATMENT (unamended, litter addition, plant presence) and AGGREGATE SIZE. The ANOVA model also included interactions, which were indeed significant for some of the parameter. However, we did not report the cases where the interaction was not or only weakly statistically significant. We will address this in the revision.
Our hypotheses were related to total fluxes during hot moments, which is why we did not analyze the time series but aggregated data. The rationale for this was already explained in detail above (R III).

5. Lines 300-353: This is an interesting and well-written part of the discussion. However, I do think that this part of the discussion could be improved by trying to link back the findings from the experiment to natural conditions (see point 1). For example, under natural conditions, what is the relative distribution of macro- or micro-aggregates? Based on your understanding/knowledge of the natural aggregate distributions, what patterns or processes do you think will dominate in a natural setting? While I realise this might be somewhat speculative (unless other data, such as field measurements, are available), I think it's an important talking point, as it will enable the reader to relate these findings (derived under somewhat artificial conditions) to the real world.

R5: For our assessment and evaluation of the relative distribution of macro- and micro-aggregates in our experimental soil and other soils reported in the literature see R1.
Furthermore, the frequent hydrological disturbance in floodplains creates a highly dynamic and small-scaled spatial mosaic of different aggregate size distributions. Therefore, the results on the individual potentials of differently sized aggregates to emit N_2O and their respective interactions with plant roots and litter accumulation could help to better understand the seemingly erratic spatial and temporal distribution of enhanced N_2O emissions from floodplain areas. Considering our results, one could speculate that zones with a relatively high percentage of macroaggregates would be prone to particularly high emissions during hot moments. In a revised manuscript, these considerations would be added also to the discussion in section 4.1.

6. Lines 380-406: The discussion of potential direct and indirect effects facilitated by the presence of an active root system is interesting and well-reasoned. However, I was left wondering as to how generalizable these findings are, given the wide range of traits displayed by different plants? I.e. to what extent are the trends identified here unique to Salix, and to what extent are these patterns more widely generalizable? I think it is important that the authors develop this section a bit further, in particular acknowledging this limitation more frankly.

R6: Different plant species may indeed exert different rhizosphere effects (for an overview of potential rhizosphere effects see the current manuscript lines 81 to 101). Thus, strictly speaking, this study is directly relevant only for salix sp.. However, this is an important plant genus adapted to temporary flooding and thus often found in river floodplains. While oxygen depletion by root exudation stimulated microbial respiration, discussed as one process potentially reducing N_2O emissions in our study, likely occurs in the rhizosphere of any plant, rhizosphere aeration as alternative process is restricted to plants possessing aerenchyma. However, the latter is a trait of many plants adapted to temporary flooding. It has been described also for the grass family of poaceae, or for ash, and It would not be surprising to find this trait in other Salicaceae like poplar sp. and other species of softwood floodplain forests.

References:

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List of all relevant changes

Adjustments according to our responses to comments by anonymous referee #1

Line 14: Terminology was specified in the abstract (Ref. 1, R1)
Line 120 – 127: An improved objectives section has been added to the introduction section (Ref. 1, general reply; Ref. 2, RIV)
Line 157: An appropriate reference was added (Ref. 1, R2)
Line 295: Added reference to supplementary information (Ref. 1, R4)
Line 335 – 337: remarks concerning statistical analysis of the initial physicochemical soil properties and adjustments to Table 1 added in to the manuscript (Ref. 1, R5)
Line 372 – 374: In response to the referees remark concerning the effect of diffusion limitation on N_2O reduction, according text was added to the discussion in section 4.1. (Ref. 1, R6)
Line 431 – 437: Additional paragraph concerning potential bypassing of the soil matrix by N_2O fluxes via plant-internal aeration channels added in discussion section 4.3. (Ref. 1, R3)

References/Figures/Tables: several minor adjustments on graphs and figure captions according the referee's remarks (Ref. 1, R7-13)

Adjustments according to our responses to comments by anonymous referee #2

Line 138 – 139: Additional information on pre-experimental conditions provided (Ref. 2, R1)
Line 148 – 153: Additional information on experimental conditions and limitations provided (Ref. 2, RI, R1)
331 – 335
Line 182 – 185: Adjustment of phrasing according the referee's remarks (Ref. 2, R2)
Line 192 – 195: Rationale for sterilization of leaf litter added (Ref. 2, R3)
Line 234 – 237: Statistics section specified (Ref. 2, RIII, R4)
241 – 245
Line 380 – 384: Discussion paragraph added about extrapolating our findings to natural conditions (Ref. 2, R5)
Line 442 – 447: Discussion paragraph added about the generalizability of our findings in the plant treatments (Ref. 2; R6)

Alteration of nitrous oxide emissions from floodplain soils by aggregate size, litter accumulation and plant–soil interactions

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Abstract. Semi–terrestrial soils such as floodplain soils are considered potential hotspots of nitrous oxide (N_2O) emissions. Microhabitats in the soil, such as within and outside of aggregates, in the detritusphere, and/or in the rhizosphere, are considered to promote and preserve specific redox conditions. Yet, our understanding of the relative effects of such microhabitats and their interactions on N_2O production and consumption in soils is still incomplete. Therefore, we assessed the effect of aggregate size, buried leaf litter, and plant–soil interactions on the occurrence of enhanced N_2O emissions under simulated flooding/drying conditions in a mesocosm experiment. We used two model soils with equivalent structure and texture, comprising macroaggregates (4000–250 µm) or microaggregates (< 250 µm) from a N-rich floodplain soil. These model soils were planted either with basket willow (*Salix viminalis* L.), mixed with leaf litter, or left unamended. After 48 hours of flooding, a period of enhanced N_2O emissions occurred in all treatments. The unamended model soils with macroaggregates emitted significantly more N_2O during this period than those with microaggregates. Litter addition modulated the temporal pattern of the N_2O emission, leading to short-term peaks of high N_2O fluxes at the beginning of the period of enhanced N_2O emissions. The presence of *S. viminalis* strongly suppressed the N_2O emission from the macroaggregated model soil, masking any aggregate-size effect. Integration of the flux data with data on soil bulk density, moisture, redox potential and soil solution composition suggest that macroaggregates provided more favorable conditions for spatially coupled nitrification–denitrification, which are particularly conducive to net N_2O production, than microaggregates. The local increase in organic carbon in the detritusphere appears to first stimulate N_2O emissions, but ultimately, respiration of the surplus organic matter shifts the system towards redox conditions where N_2O reduction to N_2 dominates. Similarly, the low emission rates in the planted soils can be best explained by root exudation of low-molecular weight organic substances supporting complete denitrification in the anoxic zones, but also by the inhibition of denitrification in the zone, where rhizosphere aeration takes place. Together, our experiments highlight the importance of microhabitat formation in regulating oxygen (O_2) content and the completeness of denitrification in soils during drying after saturation. Moreover, they will help to better predict the conditions under which hotspots, and moments, of enhanced N_2O emissions are most likely to occur in hydrologically dynamic soil systems like floodplain soils.

Kommentar [MaL1]: Ref. 1, R1

1. Introduction

Nitrous oxide (N_2O) is a potent greenhouse gas with a global warming potential over a 100 year time horizon 298 times higher than the one of carbon dioxide (Forster et al., 2007). Given its role as climate-relevant gas and in the depletion of stratospheric ozone (Ravishankara et al., 2009), the steady increase of its average atmospheric concentration of 0.75ppb yr^{-1} (Hartmann et al., 2013) asks for a quantitative understanding of its sources and the factors that control its production. On a global scale, vegetated soils are the main natural terrestrial sources of N_2O. Agriculture is the main anthropogenic source and the main driver of increasing atmosphere N_2O concentrations (Ciais et al., 2013).

In soils, several biological nitrogen (N) transformation processes produce N_2O either as a mandatory intermediate or as a by-product (Spott et al., 2011). Under oxic conditions, the most important process is obligate aerobic nitrification that yields N_2O as by-product when hydroxylamine decomposes (Zhu et al., 2013). Under low oxygen (O_2) availability, nitrifier denitrification and heterotrophic denitrification with N_2O as intermediate become more relevant (Philippot et al., 2009). At stably anoxic conditions and low concentrations of nitrate (NO_3^-), complete denitrification consumes substantial amounts of previously produced N_2O by further reduction to N_2 (Baggs, 2008; Vieten et al., 2009). In environments that do not sustain stable anoxia, but undergo sporadic transitions between oxic and anoxic conditions, the activity of certain N_2O reductases can be suppressed by transiently elevated O_2 concentration and thus can lead to the accumulation of N_2O (Morley et al., 2008).

Nitrous oxide emissions from soils depend on the availability of carbon (C) and N substrates that fuel the involved microbial processes. On the other hand, given its dependency on O_2, N_2O production is also governed by the diffusive supply of O_2 through soils. Similarly, soil N_2O emissions are modulated by diffusive N_2O transport from the site of production to the soil surface (e.g. Böttcher et al., 2011; Heincke and Kaupenjohann, 1999). Substrate availability, gas diffusivity, and the distribution of soil organisms are highly heterogeneous in soils at a small scale, with micro-niches in particular within soil aggregates, within the detritusphere, and within the rhizosphere. These can result in “hot spots” with high denitrification activity (Kuzyakov and Blagodatskaya, 2015).

Soil aggregate formation is a key process in building soil structure and pore space. Soil aggregates undergo different stages in their development, depending on the degradability of the main binding agent (Tisdall and Oades, 1982). Initially, highly persistent primary organo–mineral clusters (20–250 µm) are held together by root hairs and hyphae, thus forming macroaggregates (> 250 µm). Upon decomposition of these temporary binding agents and the subsequent disruption of the macroaggregates, microaggregates (< 250 µm) are released (Elliott and Coleman, 1988; Oades, 1984; Six et al., 2004). These consist of clay-encrusted fragments of organic debris coated with polysaccharides and proteins. This multi-stage development leads to a complex relationship between aggregate size, intra-aggregate structure and soil structure (Ball, 2013; Totsche et al., 2017), which influences soil aeration, substrate distribution and pore water dynamics (Six et al., 2004). Often, micro-site heterogeneity increases with aggregate size, thus fostering the simultaneous activity of different N_2O producing microbial communities with distinct functional traits (Bateman and Baggs, 2005). Aggregate size effects on N_2O production and consumption have generally been studied in static batch incubation experiments with a comparatively small number of isolated aggregates of uniform size, at constant levels of water saturation (Diba et al., 2011; Drury et al., 2004; Jahangir et al., 2011; Khalil et al., 2005; Sey et al., 2008), and through modelling approaches (Renault and Stengel, 1994; Stolk et al., 2011). Previous work provided partially inconsistent results, which led to an ongoing discourse about the interplay of physicochemical properties and different aggregate

sizes in controlling N_2O emission. Such inconsistencies may in parts be attributed to the use of different aggregate size classes, changes in soil structure by aggregate separation, other methodological constraints (water saturation, redox potential), and differences in microbial communities. The effects of specific aggregate sizes within a simulated soil structure, in combination with fluctuating water saturation, on soil N_2O emissions have, to our knowledge, not been addressed specifically.

Similar to soil aggregates, the detritusphere and the rhizosphere (the zone of the soil that is affected by root activity) (Baggs, 2011; Luster et al., 2009), can be considered biogeochemical hot spots (Kuzyakov and Blagodatskaya, 2015; Myrold et al., 2011). Here, carbon availability is much higher than in the bulk soil and thus rarely limiting microbial process rates. The detritusphere consists of dead organic material, which spans a wide range of recalcitrance to microbial decomposition. Spatially confined accumulations of variably labile soil litter form microhabitats that are often colonized by highly active microbial communities (Parkin, 1987). Aggregation of litter particles has been shown to affect N_2O emissions (Loecke and Robertson, 2009). Hill (2011) identified buried organic-rich litter horizons in a stream riparian zone as hot spots of N cycling. Similarly, in the rhizosphere, root exudates and exfoliated root cells provide ample degradable organic substrate for soil microbes (Robertson and Groffman, 2015). Yet, plant growth may also affect soil microbial communities through competition for water and nutrients (e.g., fixed N) (Bender et al., 2014; Myrold et al., 2011). The combined effects of these plant–soil interactions on N_2O production have been reviewed by Philippot et al. (2009). Root-derived bioavailable organic compounds can stimulate heterotrophic microbial activity, specifically N mineralization and denitrification. Nitrification in turn can be enhanced by the elevated N turnover and mineralization rates, but may also be negatively affected by specific inhibitors released from the root or through plant-driven ammonium depletion. The ability of some plants adapted to water-saturated conditions to „pump“ air into the rhizosphere via aerenchyma (gas conductive channels in the root) leads to an improved oxygenation of the rhizosphere and a stimulation of nitrification (Philippot et al., 2009). Surrounded by otherwise anoxic sediments, such aerated micro-environments may create optimal conditions for coupled nitrification–denitrification (Baldwin and Mitchell, 2000; Koschorreck and Darwich, 1998). On the other hand, transport of N_2O produced in the soil to the atmosphere is may be facilitated via these plant-internal channels, bypassing diffusive transport barriers and enhancing soil–atmosphere gas fluxes (Jørgensen et al., 2012).

The dynamics of N_2O emissions are strongly coupled to the dynamics of pore water. Re-wetting of previously dried soil can lead to strong N_2O emissions (Goldberg et al., 2010; Ruser et al., 2006), likely fostered by a wetting-induced flush in N mineralization (Baldwin and Mitchell, 2000). On the other hand, the drying-phase after water saturation of sediments and soils can lead to a period of enhanced N_2O emissions (e.g. Baldwin and Mitchell, 2000; Groffman and Tiedje, 1988; Rabot et al., 2014; Shrestha et al., 2012) when water-filled pore space (WFPS) exceeds 60% (Beare et al., 2009; Rabot et al., 2014). The increased N_2O production has been attributed to enhanced coupled nitrification–denitrification (Baldwin and Mitchell, 2000). Depending on the spatial distribution of water films around soil particles and tortuosity (which is a function of aggregate size and soil structure), the uneven drying of the soil after full saturation may generate conditions that are conducive to the formation of anaerobic zones in otherwise oxic environments (Young and Ritz, 2000). Pore water thereby acts as a diffusion barrier for gas exchange, limiting the O_2 availability in the soil pore space (Butterbach-Bahl et al., 2013). Moreover, pore water serves as a medium for the diffusive dispersal of dissolved C and N substrates, e.g. from the site of litter decomposition to spatially separated N_2O producing microbial communities (Hu et al., 2015). Therefore, fluctuations in water saturation efficiently promote the development of hot spots and hot

moments of N_2O emissions in floodplain soils and other semi-terrestrial soils (Hefting et al., 2004; Shrestha et al., 2012).

The main objective of the present experimental study was to assess both the relative and combined effects of soil microhabitats associated with soil aggregates, the detritusphere and plant–soil interactions on N_2O emissions from floodplain soils under changing pore-space saturation. We simulated a flooding event in mesocosm experiments with main focus on the dynamics of N_2O emissions during hot moments in the drying phase after flooding. To isolate the effect of aggregate-size and to minimize confounding effects of differences in soil structure, we prepared model soils by mixing aggregate size fractions of a floodplain soil with suitable inert material. The combined effects of soil aggregate size and plant detritus or plant-soil interactions were addressed by mixing the model soils with leaf litter or by planting them with willow cuttings (*Salix viminalis* L.).

Kommentar [MaL2]: Ref 1, Reply; Ref 2, R IV

We demonstrate that the level of soil aggregation significantly affects N_2O emission rates from floodplain soils through its modulating control on the model soil's physicochemical properties. We further show that these effects can be modified by the presence of a detritusphere and by root–soil interactions, changing carbon and N substrate availability and redox conditions.

2. Material and methods

2.1 Model soils

In February 2014, material from the uppermost 20 cm of a N-rich gleyic Fluvisol (calcaric, humic siltic) with 20% sand and 18% clay (Samaritani et al., 2011) was collected in the restored Thur River floodplain near Niederneunforn (NE Switzerland 47°35' N, 8°46' E, 453 m.a.s.l.; MAT 9.1 °C; MAP 1015 mm). After removing plant residues such as roots, twigs and leaves, the soil was mixed and air-dried to a gravimetric water content of 24.7 ± 0.4 %. In the next step, the original floodplain soil material, consisting of 18.5 ± 4.6 % aggregates smaller than 250 µm and 81.5 ± 4.6 % macroaggregates (mean ± SD; n = 10), was separated into a macroaggregate fraction (250–4000 µm) and a microaggregate fraction (< 250 µm) by dry sieving. The threshold of 250 µm between macroaggregates and microaggregates was chosen based on Tisdall and Oades (1982). Soil aggregate fractions were then used to re-compose model soils. In order to preserve soil structure, the remaining aggregate size fractions were complemented with an inert matrix replacing the removed aggregate size fraction of the original soil. Model Soil 1 (LA) was composed of soil macroaggregates mixed in a 1:1 (w/w) ratio with glass beads of 150–250 µm size serving as inert matrix material replacing the microaggregates of the original soil. Similarly, Model Soil 2 (SA) was composed of soil microaggregates mixed at the same ratio with fine quartz gravel of 2000–3200 µm size. To generate an even mixture of original soil aggregates and the respective inert matrix a Turbula mixer (Willy A. Bachofen AG, Muttenz, Switzerland) was used. The proportions of the aggregate size fractions in the model soils were different from the original soil, and 50% microaggregates may be more than what is found in most natural or agricultural soils (often less than 10 %). Nevertheless, we chose to use equal amounts of micro- and macroaggregates, in order to be able to separate the effects of aggregate size from effects of aggregate amount (soil mass). These proportions were still well in the range of common top soils (e.g. Cantón et al., 2009; Gajić et al., 2010; Six et al., 2000). The physicochemical properties of the two soils were determined by analysing three random samples of each model soil. Texture of the complete model soils was determined using the pipette method (Gee and Bauder, 1986) and pH was measured potentiometrically in a stirred slurry of 10 g soil in 20 ml of 0.01 M $CaCl_2$, as recommended in Hendershot et al. (2007). Additionaly

Kommentar [MaL3]: Ref 2, R1

Kommentar [MaL4]: Ref 2; RI, R1

Kommentar [MaL5]: Ref 1, R2

organic carbon (C_{org}) and total nitrogen (TN) were analysed in both aggregate size fractions without the inert material, using the method described by Walthert et al. (2010). The two model soils displayed very similar physicochemical properties (Table 1), except for the C:N ratio that was lower in macroaggregates than in microaggregates. The latter was due to the slightly lower organic carbon content in concert with slightly higher TN values in the macroaggregates. The high calcium carbonate ($CaCO_3$) content of the source material of our model soils (390 ± 3 g $CaCO_3$ kg^{-1}; Samaritani et al., 2011) buffered the systems at an alkaline pH of 8.00 ± 0.02 for LA and 7.56 ± 0.01 for SA respectively (Table 1), ensuring that the activity of key N-transforming enzymes was not hampered by too low pH, and that the potential for simultaneous production and consumption of N_2O in our experiment was fully intact (Blum et al., 2018; Frame et al., 2017).

2.2 Mesocosms

For the mesocosm experiments, transparent polyvinyl chloride (PVC) cylinders with polymethyl methacrylate (PMMA) couplings were used. A mesocosm comprised a bottom column section, containing the soil material and a drainage layer as described below, and the upper headspace section with a detachable headspace chamber (Fig. 1). Each column section was equipped with two suction cups (Rhizon MOM Soil Moisture Samplers, Rhizosphere Research Products, Netherlands; pore size 0.15 µm) for soil solution sampling. The suction cups were horizontally inserted at 5 cm and 20 cm below soil surface. For redox potential measurements, two custom-made Pt electrodes (tip with diameter of 1 mm and contact length of 5 mm) were placed horizontally at a 90° angle to the suction cups at the same depths, with the sensor tip being located 5 cm from the column wall. A Ag/AgCl reference electrode (B 2820, SI Analytics, Germany) was installed as shown in Fig. 1. A volumetric water content (VWC) sensor (EC-5, Decagon, USA) was installed 15 cm below the soil surface. To avoid undesired waterlogging, each column section contained a 5 cm thick drainage layer composed of quartz sand with the grain size decreasing with depth from 1 mm to 5.6 mm (Fig. 1). The upper cylinder section was equipped with three way valves for gas sampling, and an additional vent for pressure compensation.

2.3. Experimental setup

The mesocosm experiment had a factorial experimental design consisting of two factors (MODEL SOIL and TREATMENT), with the first factor containing two levels (macroaggregates, microaggregates) and the second factor containing three levels (unamended, litter added, plant presence). This experimental design resulted in six treatments, each replicated six times (Table 2). As basic material, each mesocosm contained 8.5 kg of either of the two model soils. Unamended model soils were used to investigate exclusively the effect of aggregate size, abbreviated as LAU (large aggregates, unamended) and SAU (small aggregates, unamended), respectively. In order to specifically assess the effect of enhanced availability of labile C in the detritusphere for the N_2O producing or consuming soil microbial community, two sets of mesocosms were amended with freshly collected leaves of Basket Willow (*Salix viminalis* L.). Those leaves were cut into small pieces, autoclaved, and then added to the model soil components (8 g kg^{-1} model soil) during the mixing procedure to create treatments LAL (large aggregates, litter) and SAL (small aggregates, litter), respectively. The sterilization step was included to create equal starting conditions in all litter treatments by reducing any potential effect of, and interaction with, the phyllosphere microbial community even though a direct involvement of the phyllosphere community in N_2O production was unlikely according to the literature (Bringel and Couée, 2015). A third set of mesocosms was planted with cuttings collected from the same *Salix viminalis* creating treatments LAP (large aggregates, plant)

Kommentar [MaL6]: Ref. 2, R2

Kommentar [MaL7]: Ref 2, R3

and SAP (small aggregates, plant), respectively to evaluate the effects of root–soil interactions in the respective model soils. For each mesocosm one cutting was inserted 10 cm into the soil, protruding from the surface about 3 cm.

The addition of leaf litter to the model soils led to an increase of C_{org} and TN in LAL relative to LAU by 41 % and 35 %, respectively, and in SAL relative to SAU by 58 % and 44 % respectively. The bulk density of the unamended model soil SAU (1.27 ± 0.01 g cm^{-3}) was slightly higher than the one of LAU (1.22 ± 0.01 g cm^{-3}; adj. *P*: < 0.0001). Regarding the litter addition treatments, the bulk density of LAL (1.13 ± 0.01 g cm^{-3}) was significantly smaller than the one of LAU (adj. *P*: < 0.0001), whereas the bulk density of SAL (1.27 ± 0.02 g cm^{-3}) did not differ significantly from the one of SAU. The soils in the treatments with plants exhibited a similar bulk density (LAP: 1.23 ± 0.02 g cm^{-3}; SAP: 1.24 ± 0.01 g cm^{-3}) as in the respective unamended treatments.

The experiments were conducted inside a climate chamber set to constant temperature (20 ± 1 °C) and relative air humidity (60 ± 10%), with a light/dark cycle of 14/10 h (PAR 116.2 ± 13.7 µmol m^{-2} s^{-1}). The experimental period was divided into four consecutive phases: The conditioning phase (Phase 1) lasted for 15 weeks and allowed the model soils to equilibrate and the plants to develop a root system. This was followed by the first experimental phase of nine days (Phase 2), serving as a reference period under steady-state conditions. During Phases 1 and 2, the soils were continuously irrigated with artificial river water (Na^+: 0.43 µM; K^+: 0.06 µM; Ca^{2+}: 1.72 µM; Mg^{2+}: 0.49 µM; Cl^-: 4.04 µM; NO_3^-: 0.16 µM; HCO_3^-: 0.5 µM; SO_4^{2-}: 0.11 µM; pH: 7.92) via suction cups, to maintain a volumetric water content of 35 ± 5 %. In Phase 3, the mesocosms were flooded by pumping artificial river water through the drainage vent at the bottom into the cylinder (10 mL min^{-1}, using a peristaltic pump; IPC-N-24, Ismatec, Germany) until the water level was 1 cm above the soil surface. After 48 h of flooding, the water was allowed to drain and the soil to dry for 18 days without further irrigation (Phase 4).

2.4 Sampling and analyses

During the entire experiment, water content and redox potential were automatically logged every 5 minutes (EM5b, Decagon, USA and CR1000, Campbell scientific, USA, respectively).

At selected time points during the experiment, soil-emitted gas and soil solution were sampled. For N_2O flux measurements, 20, 40 and 60 minutes after closing the mesocosms, headspace gas samples (20 mL) were collected using a syringe and transferred to pre-evacuated exetainers. The samples were analyzed for their N_2O concentration using a gas chromatograph (Agilent 6890, Santa Clara, USA; Porapak Q column, Ar/CH_4 carrier gas, micro-ECD detector). Measured headspace N_2O concentrations were converted to moles using the ideal gas law and headspace volume. The N_2O efflux rates were calculated as the slope of the linear regression of the N_2O amounts at the three sampling times, relative to the exposed soil surface area (Fig. 1, Shrestha et al., 2012).

For soil water sampling, 20 mL of soil solution were collected using the suction cups. Water samples were analyzed for dissolved organic carbon (DOC) and TN concentrations with an elemental analyzer (Formacs$^{HT/TN}$, Skalar, The Netherlands). Nitrate and ammonium concentrations were measured by ion chromatography (IC 940, Metrohm, Switzerland), and nitrite (NO_2^-) concentrations were determined photometrically (DR 3900, Hach Lange, Germany).

2.5 Data analyses

We were interested in effects on cumulated N_2O emissions during hot moments following flooding. We therefore analyzed data aggregated over this period rather than the raw full time series data. This procedure also

avoided potential issues with small shifts in the timing of emissions that might have been significant but which were irrelevant for the total fluxes we focused on. The total amount of N_2O emitted during the period of enhanced N_2O fluxes in Phase 4, Q_{tot}, was calculated by integrating the N_2O fluxes between day 11 and 25 of the experiment as follows:

$$Q_{tot} = \frac{1}{2} \sum_{n=1}^{n_{max}} [\Delta_n \times (q_n + q_{n+1})] \quad (1)$$

where Δ_n is the time period between the n^{th} and the $n+1^{th}$ measurement, and q_n and q_{n+1} the mean flux on the n^{th} and $n+1^{th}$ measurement day, respectively. "n=1" refers to day 11, and n_{max} to day 25 of Phase 4. The integrated N_2O fluxes, as well as the average DOC and N-species concentrations in the soil solution during this period were analyzed by performing two-way ANOVAs with the fixed terms TREATMENT and MODEL SOIL including their interaction. In case of significant MODEL SOIL, TREATMENT or MODEL SOIL × TREATMENT effects, their causes were inspected with the Tukey's honestly significant difference (HSD) post hoc test. For all data, the residuals of the ANOVA models were inspected, and the Shapiro–Wilk normality test was applied to ensure that the values follow a Gaussian distribution. In case that this requirement for ANOVA was not met, the respective data set was log-transformed. Significance and confidence levels were set at $\alpha < 0.05$. The results of the performed ANOVAs are summarized in Table 3. For the statistical analyses we used GraphPad Prism (GraphPad Software Inc., 2017) and R (R Core Team, 2018).

Kommentar [MaL8]: Ref 2, R III

Kommentar [MaL9]: Ref 2, R III, R4

3. Results

3.1 Soil moisture and redox potential

During Phase 1 and 2, saturation levels stabilized at 53.0 ± 2.1% WFPS (water filled pore space) in the treatments with LA soils, and were slightly higher in SA treatments (57.8 ± 2.0%) (Fig. 2). The flooding of the mesocosms for 48 h with artificial river water raised the WFPS for all LA soils to 87.8 ± 0.1%, significantly exceeding the increase of WFPS in SA soils (80.6 ± 0.1%). The water release from the system after the simulated flood resulted in an immediate drop of the WFPS, except for the LAU treatment (Fig. 2). This was followed by slow drying for 1 week, and a more marked decrease in WFPS during the second week after the flood. During the latter period, the plant treatments dried faster than the other treatments. As a result, at the end of the experiment, WFPS was still above pre-flood values in unamended and litter treatments, while WFPS levels in the treatments with plants were lower than before the flooding.

The time course of the redox potential measured in 5 cm and 20 cm depth exhibited distinct patterns depending on the respective model soil (Fig. 3). In all treatments, flooding induced a rapid decrease of the redox potential to values below 250 mV within 36 hours. Upon water release, the redox potential returned rapidly to pre-flood values at both measurement depths only in SA soils. In the LA treatments (most pronounced in LAL), soils at 20 cm depth underwent a prolonged phase of continued reduced redox condition, returning to the initial redox levels only towards the end of the experiment.

3.2 Hydrochemistry of soil solutions

Considering individual treatments, DOC concentrations varied only little with time. Yet, the DOC concentrations were generally much higher in treatments with LA than with SA soils. This main effect of MODEL SOIL was

highly significant, as was the interaction with TREATMENTS due to a smaller difference in the litter addition treatments than in the unamended and plant treatments (Table 3). Nitrate was the most abundant dissolved reactive N species in the soil solution, with pre-flood concentrations of 1 to 5 mM (Fig. 4d–f). In the unamended and plant treatments, NO_3^- concentrations were markedly higher in SA than in LA soils, whereas they were similar in both litter addition treatments. Two distinct temporal patterns in the evolution of NO_3^- concentration could be discerned. In the unamended and litter-addition treatments, NO_3^- concentrations decreased after the flooding, consistently reaching a minimum on day 19, in the case of the litter treatments below the detection limit of 0.2 µM, before increasing again during the latter drying phase (Fig. 4d,e). In contrast, in the treatments with plants, NO_3^- concentrations steadily declined from concentrations of 1–2 mM to around 0.5 mM at the end of the experiment (Fig. 4f). Nitrite was found at significant concentrations only in LA soils, with highest concentrations in the LAU treatment right after the flooding (33.6 µM) and decreasing concentrations throughout the remainder of the experiment (Fig. 4g–i). In SA soils NO_2^- concentration was always < 5 µM, without much variation. Similarly, in most treatments except SAL, ammonium (NH_4^+) concentrations were < 10 µM, and particularly towards the end of the experiment very close to the detection limit (Fig. 4j, 4l). In the SAL treatment, NH_4^+ concentrations peaked 5 days after the flood with concentrations of around 70 µM (Fig. 4k). This deviation from the other temporal patterns prompted a significant interaction effect between MODEL SOIL and TREATMENTS.

3.3 Nitrous oxide emissions

During Phase 2 (i.e., before the flooding), N_2O fluxes were generally low (< 1 µmol m^{-2} h^{-1}; Fig. 2), however, fluxes in the LAL treatment were significantly higher than in the other treatments (adj. P = 0.002–0.039; Fig. 2). The flooding triggered the onset of a "hot moment", defined here as period with strongly increased N_2O emissions, which lasted for about one week independent of the treatment (Fig. 2). The maximum efflux was observed immediately after the flood. The subsequent decline in N_2O emission rates followed different patterns among the various treatments. Normalizing the N_2O flux to the maximum measured efflux for each replicated treatment revealed a slower decrease with time for the unamended soils than for the litter and plant treatments (Fig. S1). The strongest peak emissions were observed in the LAL treatment (91.6 ± 14.0 µmol m^{-2} h^{-1}; mean ± SD). Throughout most of the drying phase, the LAU and LAL treatments exhibited higher N_2O emissions than the corresponding SAU and SAL experiments. In contrast, there was no such difference in the treatments with plant cuttings, and peak N_2O emissions were overall lower than in the other treatments. The integrated N_2O fluxes during the hot moment (days 11 to 25 of the experiment) were significantly higher for the LAU and LAL than for all other treatments (Fig. 5), and the aggregate size effect was also significant within the unamended (adj. P = 0.045) and litter-addition treatments (adj. P = 0.008). The integrated N_2O emissions in the two plant treatments did not differ significantly from each other, but were significantly smaller than in the LAU (adj. P = 0.001), and the LAL (adj. P = 0.005) treatments. Overall, the effects of MODEL SOIL and TREATMENTS were significant, as was the interaction between the two factors due to the different aggregate size effect in the plant compared to the unamended and litter addition treatments (Table 3).

Kommentar [MaL10]: Ref 1; R4

4. Discussion

In our experiment, we could confirm the occurrence of periods of enhanced N_2O emissions in the drying phase shortly after flooding, as expected based on previous research (Baldwin and Mitchell, 2000; Groffman and

Tiedje, 1988; Rabot et al., 2014; Shrestha et al., 2012). We observed that the six treatments had a substantial effect on the magnitude and temporal pattern of N_2O emissions that could only be captured by observations at relatively high temporal resolution. The fast occurrence of strong N_2O fluxes over a comparatively short period in the litter-amended treatment on the one side, and the relatively weak response to the flooding in the plant treatment on the other, suggests complex interactive mechanisms related to distinct microhabitat effects leading to characteristic periods of enhanced N_2O emission. Rabot et al. (2014) explained N_2O emission peaks during the desaturation phase with the release of previously produced and entrapped N_2O. Such a mechanism may partly contribute to high N_2O emissions in our experiment initially, but the continuing depletion of NO_3^- and NO_2^- during the phase of high N_2O emissions indicates that the flooding and drying has strong effects on N transformations mediated by microorganisms in the soil (e.g., the balance and overall rates of nitrification, nitrifier–denitrification, and denitrification). Hence, physical controls alone clearly do not explain the observed timing and extent of hot moments with regard to N_2O emission. In the following sections we will discuss how the effect of flooding on microbial N_2O production is modulated by differential microhabitat formation (and hence redox conditions) in the various treatments.

4.1 Effect of aggregate size on N_2O emissions

LA model soils exhibited both higher peak and total N_2O emissions during the hot moment in the drying phase than SA model soils (Figs. 2 and 5). By contrast, in the presence of a growing willow, there was no detectable effect of aggregate size on the overall N_2O emission (further discussion below). The aggregate size effects observed in the unamended and litter treatments can be explained by factors controlling (i) gas diffusion (e.g. water film distribution, tortuosity of the intra-aggregate pore space) and (ii) decomposition of encapsulated soil organic matter (SOM) regulating the extent of N_2O formation (Neira et al., 2015). In order to isolate the effect of aggregate size (i.e., to minimize the effect of other factors that are likely to influence gas diffusion), we created model soils of similar soil structure and texture (see Materials and Methods). We thereby implicitly accepted that potential interactions of the two size fractions with each other, or with soil structures larger than 4 mm could not be assessed in this experiment. Although this approach thus represents only an approximation of real-world conditions it was still an improvement compared to experiments where no attempts were made to conserve soil structure. Similarly, the bulk soil chemical properties of the two aggregate size fractions, such as C_{org} content and TN, are essentially the same. Differences in the initial C:N ratio and pH, although statistically significant, can be considered equivalent in the ecological context, e.g., in terms of organic matter degradability. Therefore, we assume in the following that the differences in N_2O emissions among the treatments can mainly be attributed to size-related aggregate properties and their interactions with litter addition or rhizosphere effects.

Kommentar [MaL11]: Ref 2; RI

Kommentar [MaL12]: Ref 1, R5

During Phase 3 with near-saturated conditions, no aggregate size effect was observed. High WFPS seem to have limited the gas diffusion (O_2 and N_2O) independent of the aggregate size, limiting soil–atmosphere gas exchange in both model soils equally (Neira et al., 2015; Thorbjørn et al., 2008). As a consequence of inhibited gas exchange/soil aeration, a sharp drop in the redox potential was observed in all treatments, indicating a rapid decline in O_2 availability to suboxic/anoxic conditions. Together with an incipient decrease in soil solution NO_3^-, this indicates that N_2O production is primarily driven by denitrification in this phase.

The aggregate size effects on the formation of moments of enhanced N_2O emission became evident during the subsequent drying period. During the initial drying phase, when a heterogeneous distribution of water films around soil particles/aggregates develops (Young and Ritz, 2000), the macroaggregates in the LA model soils

appear to foster micro-environmental conditions that are more beneficial to N_2O production. This could be related to the longer diffusive distances for re-entering O_2 caused by the higher tortuosity of the intra-aggregate pore space of macroaggregates, as reported by Ebrahimi and Or (2016). This may have helped to maintain, or even extend, reducing conditions due to microbial activity inside the core of macroaggregates during drying. Thus, on the one hand, large aggregates favor the emergence of anoxic microhabitats expanding the zones where denitrification occurs. On the other hand, the overall higher porosity of the LA soils supports a better aeration in drained parts of the soil (Sey et al., 2008), and aerobic processes (e.g., nitrification) are supported. As a result, ideal conditions for spatially coupled nitrification–denitrification are created (Baldwin and Mitchell, 2000; Koschorreck and Darwich, 1998). Indeed, the emergence of heterogeneously distributed, spatially confined oxygen minimum zones during soil drying may be reflected by the high variability of the redox conditions observed in replicate mesocosms and, on average, the tendency towards lower redox potentials for a prolonged period of time in the subsoils of the LA model soils (Fig. 3 d–f). In this context, the relevance of water films for the emergence of periods of enhanced N_2O emissions is further highlighted by the fact that elevated flux rates were only observed as long as the WFPS was above 65%. This is consistent with work by Rabot et al. (2014) and Balaine et al. (2013), who found similar soil water saturation thresholds for elevated N_2O emissions from soils, attributing this phenomenon to suboptimal environmental conditions for both nitrification and denitrification at lower saturation levels.

Given the arguments above, we assume that N_2O emissions during the drying phase originate to a large degree from heterotrophic denitrification, and that they are governed mainly by the aggregate-size dependent redox conditions within the semi-saturated soils. This conclusion stands in good agreement with findings from Drury et al. (2004), who found higher production of N_2O due to enhanced denitrification with increasing size of intact arable soil aggregates in a laboratory incubation study. In contrast, the much lower emissions from the SA treatments can best be explained by a rapid return to pre-flood, i.e. oxic redox conditions in most of the pore space, under which N_2O production driven by denitrification is inhibited. Enhanced reduction of N_2O to N_2 in the SA versus LA treatments seems less likely as an explanation for lowered net N_2O emission rates, since the relatively high redox potential represents an impediment to complete denitrification to N_2. Furthermore, according to Manucharova et al. (2001) and Renault and Stengel (1994), aggregates smaller than 200 µm are simply not large (and reactive) enough (i.e., molecular diffusive distances for oxygen are too short) to develop suboxic or anoxic conditions in the center, let alone denitrifying zones. Hence, only a relatively small fraction of the total number of microaggregates in the SA soils would have been large enough (between 200 and 250 µm) to host denitrification and act as site of anaerobic N_2O production.

Under natural conditions, frequent hydrological disturbance in floodplains creates a highly dynamic and small-scaled mosaic of different aggregate size distributions. In this regard, our results, demonstrating the effect aggregate size has on N_2O emissions, may help to understand the seemingly erratic spatial and temporal distribution of enhanced N_2O emissions from floodplain areas. Moreover they imply that zones with a relatively high percentage of macroaggregates would be particularly prone to high emissions of N_2O after a flood event.

Kommentar [MaL13]: Ref 1, R6

Kommentar [MaL14]: Ref 2, R 5

4.2 Litter effect on N_2O emissions

We expected that litter addition would increase N_2O emissions from model soils with both small and large aggregates, as was found earlier (e.g. Loecke and Robertson, 2009; Parkin, 1987). The addition of litter to the model soils changed the temporal dynamics of the N_2O emission substantially, but its effect on the net integrated

N_2O emission was rather minor (Fig. 5). More precisely, highest peak emission rates of all treatments were observed in the LAL treatment, but peak emission rates were followed by a faster return to low pre-flood emission rates in the LAL and the SAL treatments relative to the unamended treatments (Fig 2). This confirms that surplus organic carbon can, on short-term, boost N_2O emissions, particularly in the large-aggregate treatment. The fast mid-term return to low N_2O emission suggests that N_2O production by heterotrophic denitrification either becomes limited by substrates other than carbon, and/or that the carbon added to the soils affects the redox-biogeochemistry in a way that shifts the balance between N_2O production and consumption in favor of consumption. Loecke and Robertson (2009) reported similar temporal N_2O emission patterns in field experiments with litter-amended soil, and attributed the observed dynamic of a rapid decline after peak emission to an increased demand for terminal electron acceptors during denitrification shortly after the carbon addition. Nitrate/nitrite limitation leads, under stable anoxic conditions, ultimately to the complete reduction of produced N_2O to N_2 decreasing net N_2O emission. Indeed, the rapid decrease in N_2O emissions after the emission rate peak in the litter addition treatments was accompanied by the complete depletion of NO_3^- in the soil solution at low redox potential, suggesting nitrate limitation. The increased demand for electron acceptors can be attributed to the increased availability of labile C compounds and nutrients provided by the mineralization of litter, and the concomitant stimulation of aggregate-associated microbial communities during the flooding (Li et al., 2016). At the same time, the litter-stimulated soil respiration increases the soil's oxygen demand, maintaining stable low redox conditions for a longer period of time during the drying phase. Since high activity of N_2O reductase requires very low O_2 concentrations (Morley et al., 2008), such conditions may be particularly favorable for complete denitrification to N_2, an additional, or alternative, explanation for the low N_2O emission rates shortly after the N_2O emission peak.

4.3 Effects of *Salix viminalis*

Planted willow cuttings resulted in relatively low maximum N_2O emission rates (LAP: 19.75 ± 9.31 µmol m^{-2} h^{-1}; SAP: 15.07 ± 12.07 µmol m^{-2} h^{-1}; mean ± SD), independent of aggregate size. The high values for WFPS throughout the hot moment, and a low redox potential in the subsoil, imply optimal conditions for denitrification or nitrifier denitrification, but compared to unamended and litter-addition treatments, only little N_2O was emitted (both during peak N_2O emission rates and with regards to the integrated N_2O flux). *S. viminalis* suppressed peak N_2O emissions, overriding the positive effect of large aggregates on N_2O emissions observed otherwise. The specific mechanisms involved are uncertain. Fender et al. (2013) found in laboratory experiments with soil from a temperate broad-leaved forest planted with ash saplings (*Fraxinus excelsior* L.) N_2O fluxes and plant effects very similar to the ones observed in our study. They attributed reduced N_2O emissions in presence of ash partly to plant uptake of nutrients that reduced NO_3^- availability to denitrifiers. Fast-growing plant species like *Salix* are particularly effective in removing soil inorganic N (Kowalik and Randerson, 1994). Such a causal link between reduced N_2O emissions and plant growth is, however, not supported by our data. More precisely, the NO_3^- concentrations during the hot moment of N_2O emissions were always relatively high (> 0.5 mM) and above the levels observed in the litter treatments.

An alternative explanation for the reduced N_2O emissions in the plant treatments could be rhizosphere aeration by aerenchyma, a physiological trait of *Salix viminalis* roots, which prevents the formation of anoxia in their close vicinity (Blom et al., 1990; Randerson et al., 2011), and thus inhibits anaerobic N_2O production. Indeed, redox potentials in the topsoil were higher in SAP and LAP compared to the other treatments. By contrast, the

redox potential in the saturated subsoil below was even lower than observed for the unamended soils. This indicates that the aeration effect by aerenchyma is constrained to the upper soil, or is, in the deeper soil portions, compensated by respiratory rhizosphere processes. On the other hand, aerenchyma can also aid in the gas exchange between the soil and the atmosphere, leading to an accelerated transport of N_2O by bypassing the soil matrix. This phenomenon is well documented for various grasses such as *Oryza* (Baruah et al., 2010), *Triticum* (Smart and Bloom, 2001) or *Phalaris arundinacea* (Jørgensen et al., 2012). However, we are not aware of any reports on enhanced N_2O emissions via aerenchyma by willows (*Salix sp.*), and indeed, our results do not indicate any increased N_2O emission via plants. In fact, we observed the lowest ecosystem flux rates and lowest total integrated N_2O emissions in the mesocosms with *S. viminalis*.

Kommentar [MaL15]: Ref 1; R3

According to Fender et al. (2013), in vegetated soils, microbial respiration is stimulated by deposition of root exudates, which in concert with root respiration in a highly saturated pore space, leads to severe and ongoing oxygen depletion. Under such stable anoxic conditions complete denitrification would take place generating N_2 and not N_2O as the dominant final product and therefore N_2O emissions would be low.

While oxygen depletion by root-exudation-stimulated microbial respiration likely occurs in the rhizosphere of any plant, rhizosphere aeration is restricted to plants possessing aerenchyma. However, the latter is a characteristic of many plants adapted to temporary flooding, and has been described also for *Poaceae*, or for ash. Furthermore, it is reasonable to expect this trait to be found in other *Salicaceae* like *Populus sp.* and other species of softwood floodplain forests. In areas with monospecific stands of, for example *Salix sp.*, which are often found on restored river banks, this N_2O-emission reducing trait can be a welcome side effect.

Kommentar [MaL16]: Ref 2; R6

5. Conclusions

In this study, we investigated the distinct effects of aggregate size, surplus organic carbon from litter and vegetation on N_2O emission from model soils after flooding. Flooding and drying were always associated with hot moments of N_2O production, most likely due to heterotrophic denitrification as result of suboxic O_2 levels at high WFPS. Our results demonstrate that aggregate size is a very important factor in modulating N_2O emission from soils under changing pore space water saturation. Aggregates of a diameter > 250 µm appear to foster suboxic microhabitats that favor denitrification and associated N_2O emission. This soil aggregate size effect may be amplified in the presence of excess carbon substrate, as long as heterotrophic denitrification, as the main N_2O producing process, is not electron-acceptor limited, and extremely reducing conditions in organic rich soils do not promote complete denitrification leading to further reduction of N_2O to N_2. On the other hand, the higher porosity of the soils with macroaggregates may aid in the formation of microsites at the surface of aggregates where nitrification is re-initialized during drying, supporting favorable conditions for spatially coupled nitrification–denitrification. The mechanisms by which processes in the rhizosphere of *Salix viminalis* effectively suppress N_2O emissions, and thus mask any aggregate size effect, remain ambiguous. Distinct physiological features of *Salix viminalis*, its root metabolism, in combination with microbial respiration can lead to the simultaneous aeration of some parts of the rhizosphere, and the formation of strongly reducing zones in others. In both cases, redox conditions seem to be impedimental for extensive net N_2O production.

Our results demonstrate the importance and complexity of the interplay between soil aggregate size, labile organic C availability, respiratory processes in the rhizosphere, and plant-induced aeration of soils under changing soil water content. Those interactions emerged as modulators of N_2O emissions by controlling the O_2

distribution in the soil matrix. Indeed, O_2 appears as the unifying master variable that ultimately sets the boundary conditions for N_2O production and/or consumption.

The main scope of this work was to expand our knowledge on the controls on net N_2O emissions from floodplain soils. The systematic relationships observed in this study are likely to help anticipating where and when hotspots and hot moments of N_2O emissions are most likely to occur in hydrologically dynamic soil systems like floodplain soils. Further understanding of the complex interaction between plants and soil microorganisms, the detritusphere, and soil aggregation, as well as their influence on N turnover and N_2O accumulation in soils, should focus on how the parameters tested affect the actual activity of the nitrifying and denitrifying communities, with an in-depth investigation into the biogeochemical pathways involved.

Data availability. Data will be openly available at https://datadryad.org/

Competing interests. The authors declare that they have no conflict of interest.

Authors contributions. The initial concept of the experiment was developed by JL, MFL and PAN. ML planned the experiment in detail, set it up and performed it. PAN supervised the measurement of N_2O gas concentrations, whereas ML conducted all other measurements and data analyses. ML wrote the manuscript with major contributions by JL, MFL and PAN.

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Table 1: Physicochemical properties of the two aggregate size fractions (macroaggregates and microaggregates) and added leaf litter. C_{org} and TN of the aggregates were measured in triplicates. The leaf litter was analyzed in quadruplicates. Final pH and texture of model soil 1 and 2 were measured in duplicates (means ± SD). Significant differences in the t-tests ($P < 0.05$) are highlighted in bold.

		Macroaggregates	Microaggregates	Macroaggregates vs. Microaggregates	Litter (*Salix v.* L.)
C_{org}	g kg^{-1}	19.22 ± 0.55	21.56 ± 2.39	P = 0.229	459.9 ± 2.55
Total N	g kg^{-1}	1.58 ± 0.02	1.35 ± 0.14	P = 0.106	27.39 ± 0.15
C:N ratio		12.16 ± 0.22	15.99 ± 0.71	**P = 0.007**	16.79 ± 0.06
		Model soil 1	Model soil 2	Model soil 1 vs. Model soil 2	
pH ($CaCl_2$)		8 ± 0.02	7.56 ± 0.01	**P = 0.009**	
sand	%	71.25 ± 0.05	70.7 ± 0.50	P = 0.469	
silt	%	20 ± 0.30	21.1 ± 0.60	P = 0.285	
clay	%	8.75 ± 0.25	8.2 ± 0.10	P = 0.240	

Table 2: Overview of treatments in the flooding–drying experiment. Model Soil 1, containing soil macroaggregates is abbreviated LA, whereas Model Soil 2 contains soil microaggregates and is abbreviated SA. The last character of each abbreviation stands for unamended (U), litter addition (L) and plant presence (P). Each treatment was replicated six times.

	LAU	SAU	LAL	SAL	LAP	SAP
Model Soil 1 (LA)	+	-	+	-	+	-
Model Soil 2 (SA)	-	+	-	+	-	+
Leaf litter (*Salix v.*)	-	-	+	+	-	-
Salix v.	-	-	-	-	+	+

Table 3: Results of the two-way analysis of variance (ANOVA) of the integrated fluxes (Q_{tot}) and the mean concentrations of chemical properties in soil solution (n=6) during the period of enhanced N_2O emissions (from day 11 to day 25). Shown are *P* values with significant differences ($P < 0.05$) highlighted in bold characters.

	Q_{tot}	DOC	NO_3^-	NO_2^-	NH_4^+
TREATMENT	**0.0003**	**0.0133**	0.0988	**< 0.0001**	**0.0007**
MODEL SOIL	**0.0002**	**< 0.0001**	0.2181	**< 0.0001**	**0.0004**
TREATMENT × MODEL SOIL	**0.0145**	**< 0.0001**	0.0668	0.1174	**< 0.0001**

Figure Captions

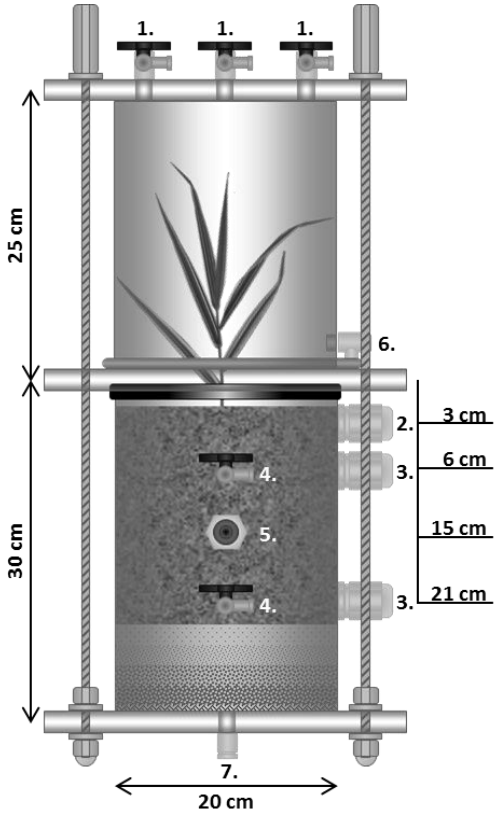
Figure 1: Schematic of a mesocosm with gas sampling valves (1), Ag/AgCl reference electrode (2), Pt redox electrodes (3), suction cups (4), volumetric water content sensors (5), vent (6), and water inlet/outlet (7). The top part is only attached during gas sampling.

Figure 2: Mean N_2O emission during the flooding–drying experiment from large-aggregate model soil (LA; filled circles) and small-aggregate model soil (SA, open circles). The corresponding water-filled pore space (WFPS) in LA (filled triangles) and SA (open triangles) are depicted on the right Y-axis. Unamended soils (A), litter addition (B) and plant treatment (C). Flooding phase indicated by the grey area. Symbols indicate means; error bars are SE; n= 6.

Figure 3: Redox potential relative to standard hydrogen electrode during the flooding–drying experiment in 5 cm and 20 cm depth (mean ± SE; n=6). Unamended soils (a and d, respectively), litter addition (b and e, respectively), plant treatment (c and f, respectively). LA (filled circles) and SA (open circles); the dotted line at 250 mV marks the threshold, below which denitrification is expected to occur.

Figure 4: DOC (circles), nitrate (squares), nitrite (diamonds) and ammonium (triangles) concentrations in pore water during the flooding–drying experiment. LA (filled symbols) and SA (empty symbols). Unamended soils (a, d, g and j, respectively), litter addition (b, e, h and k, respectively) and plant treatment (c, f, j and l, respectively).; (mean ± SE; n=6).

Figure 5: Integrated N_2O fluxes over the 14 days period of elevated N_2O emissions in the drying phase of the flooding–drying experiment (mean ± SE; n= 6). Black bars represent Model Soil 1 (macroaggregates 250-4000µm) whereas Model Soil 2 (microaggregates < 250µm) is depicted as white bars. Significant differences among the six treatments are denoted by different lower case letters at adj. $P < 0.05$.



715

716 Figure 1

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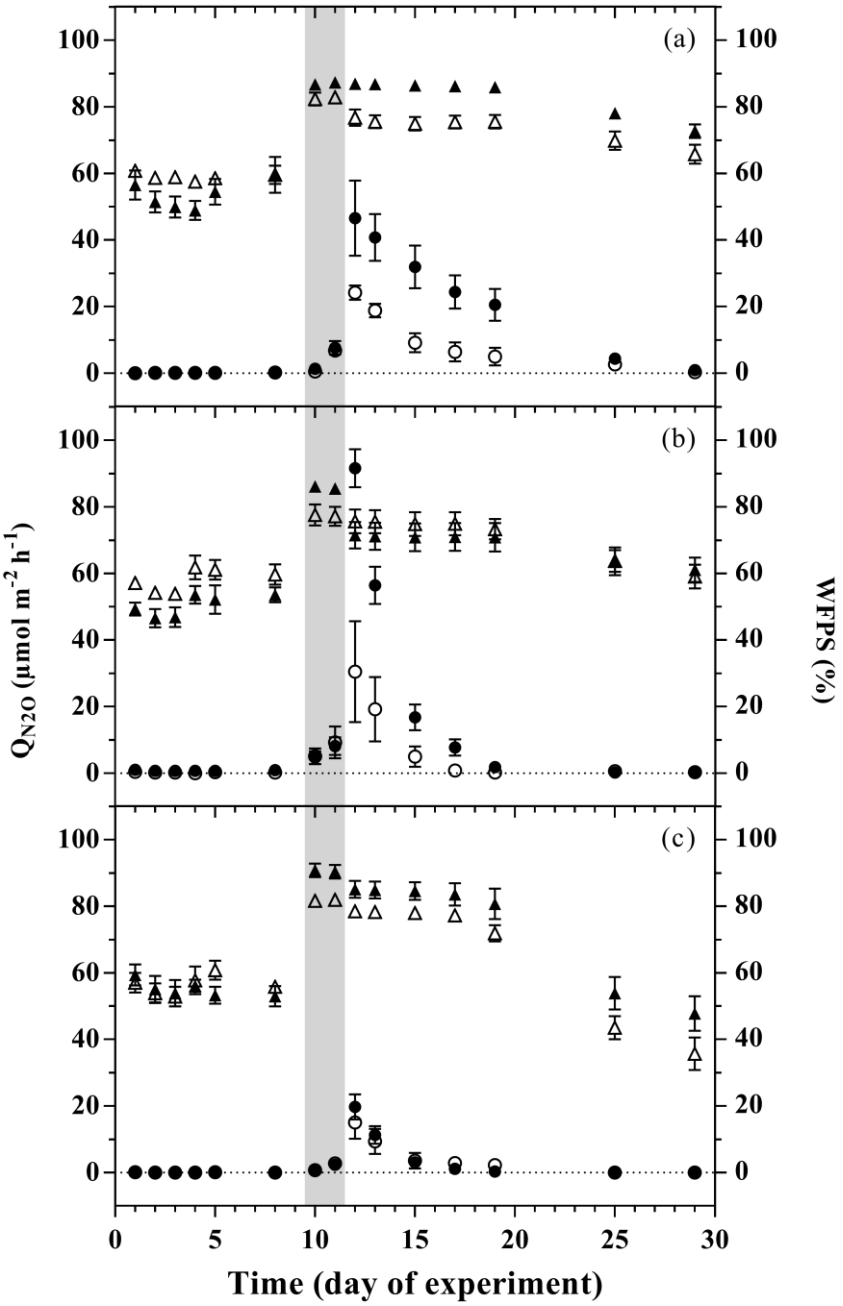

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719 Figure 2

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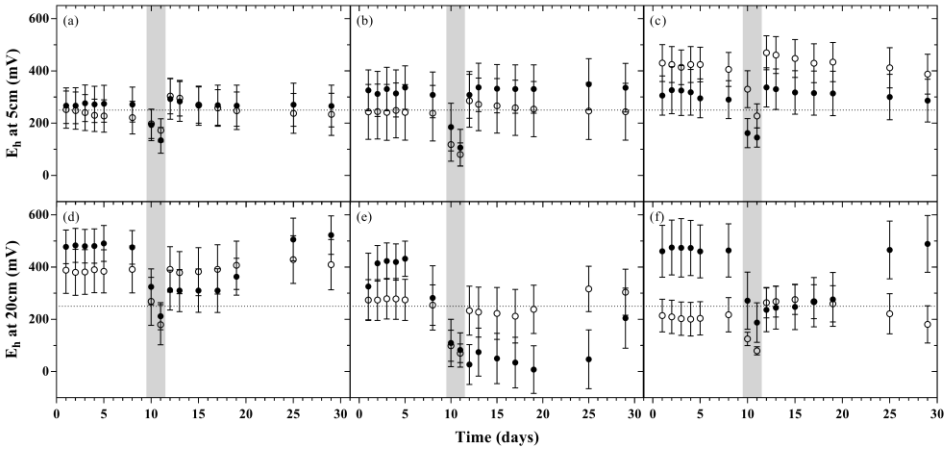

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722 Figure 3

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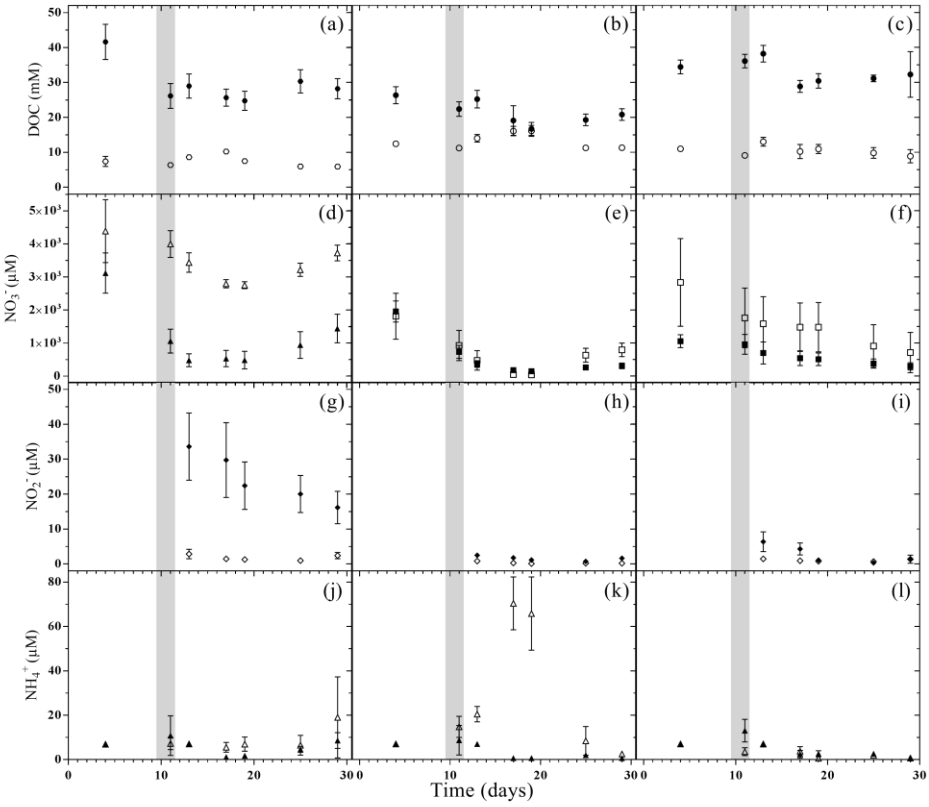

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725 Figure 4

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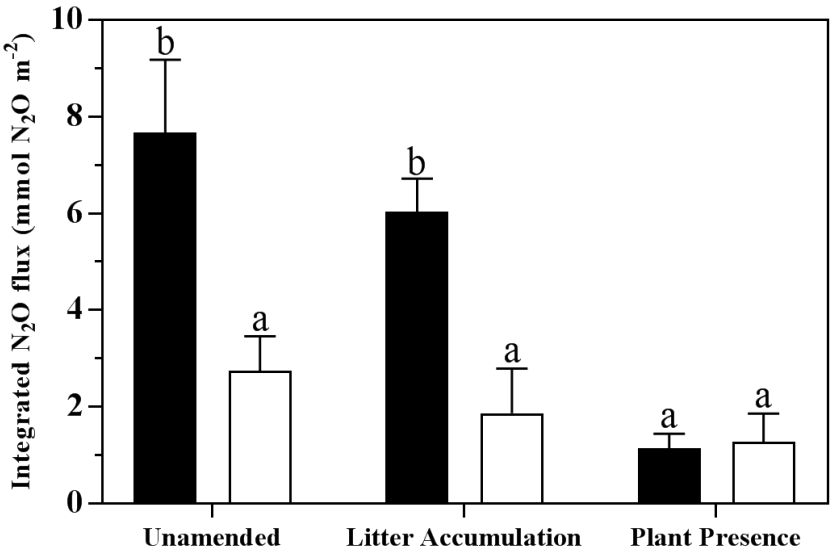

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728 Figure 5