Dear Editor,

We would like to express our sincere thanks to the two reviewers, who did an excellent job in spotting weaknesses and minor inconsistencies in the previous version of our manuscript. We have now added soil temperature data from the field sites backing up our claim of strongest soil frost dynamics in the center of our climatic gradient and data on soil organic matter content which offers a potential explanation for the higher sensitivity to the FTC treatment by the soils from the coldest origins. Furthermore, we have clarified the experimental methods and solved minor inconsistencies. Please find below all our replies to each single point.

Best regards, Juergen Kreyling, Rhena Schumann, Robert Weigel

Anonymous Referee #1

General comment
This is a well written manuscript on a timely topic, which studies the effect of freeze thaw cycles on N and P release between snow-poorer warmer and snow-richer colder forest soils. The manuscript is very well and concisely written, clearly structured, and provides clear aims, hypotheses, and approaches. Especially the statistical analysis of the data is very strong, with a transparent use and analysis of the data. The major shortcoming of the study is lack of crucial soil data without whose, the outcome, discussion and conclusion remain speculative.

Reply: Thank you very much for this positive feedback and the thoughtful and constructive critique! Upon reading your detailed comments, we agree that additional information on soil parameters improves the interpretation of the presented results. Please compare to the replies to your specific points below for details what we have now added to the manuscript.

1. It remains unclear from which soil depth the samples had been taken and what was the criterion of the sampling. Soil temperatures in the mineral soil are known to be well buffered against air temperatures and hence it even remains unclear to which the soils studied have indeed historically experienced FTCs. Although it is described that soil temperatures had been recorded, the data are not shown except minimal winter temperatures.

Reply: To our own surprise, we indeed missed to report the sampling depth and have added this now (line 139: 0-10 cm soil depth) – thanks for spotting this! We have furthermore clarified the soil sampling design there.
We further agree that the history of FTC per site is clearly relevant for the interpretation of the site effects. Our statement of recorded soil temperatures was related to the climate chamber trials while all data provided in Table 1 of the initial submission was based on gridded climate data, which generally does not include reliable soil temperature data. However, we have now added on-site soil temperature measurements for four consecutive years across three winters (2016-2019). We acknowledge that this is a short period of observations which does not shed light on historic patterns and changes, but we assume that this recent snapshot has additional value for the interpretation of our results. As already assumed in the first submission, this additional data now suggests that FTC at the studied soil depth are rare at the western end of the gradient with the warmest winter air temperatures. More importantly, FTC are also rare at the coldest sites, very probably due to the high probability of a continuous snow cover insulating the soil against air temperature fluctuations at these sites (compare to column 'Precipitation as snow' in Table 1). The relevant on-site FTC records have been added to the reworked Table 1 and are picked up again in the discussion (lines 377ff.).

2. In addition, data about inherent soil properties are lacking. For instance, soil organic matter contents are not given. Soil organic matter is a key soil parameter driving soil microbial communities and the release of nutrients and thus of nutrients released upon lysis of microbial cells. As SOM greatly varies with climate (and which soil depth) it seems likely that it is a key co-variable which could drive the observed responses and as such it should be reported, incorporated into the statistical model and/or discussed.

Reply: SOM, pH, and C/N have been added to Table 1 and are now discussed in light of the results (lines. 380ff): "While the soil C/N-ratio appeared irresponsive to the climatic gradient in our study, soil organic matter content increased towards the coldest sites (Table 1). High organic matter content generally increases the susceptibility of soils for nutrient loss with climate change (Liu et al., 2017). Here, we cannot answer how strongly this pattern in organic matter is driven by historic winter soil temperature and occurrence of FTC, but the expectation of increased mineralization with winter soil warming (Gao et al., 2018) would fit to the observed decrease of soil organic matter content with warmer winter climate (Liu et al., 2017). Moreover, the larger pool of organically bound nutrients at the coldest sites may contribute to their observed responsiveness to FTC warming (Gao et al., 2018)."

3. Moreover, the amount of nutrients released should be normed to the amount of nutrients present in the sample. It should be made clear which drivers are/could be responsible for the observed nutrient concentrations: for example, inherent nutrient content, C/nutrient ratios that drive the net release of nutrients, or historical FTCs. The concepts and data on other drivers should be considered in the data analysis and/or ruled out in the discussion.

Reply: This is an important point and we would agree with norming to the amount of nutrients present if we would only address the impact of the experimental soil temperature manipulation on soil nutrient release rates. However, in our study and in our analysis, we aimed at analyzing the impact of experimental soil temperature manipulation scenarios under explicit consideration of the soil origin. Thus, the soil origin is included in our 3-factorial modelling approach by way of including long-term winter climate variables of the soil origin. Consequently, this accounts for the different initial nutrient contents, as they indeed are characteristic for each soil origin. Standardizing to pure nutrient release rates would, therefore, be contra-productive for our analysis. Still, we agree that this aspect should be picked up in the discussion and have added there (lines 380ff, also cited in the reply above).

4. Details from the laboratory experiment are lacking or not clear. For example, was the soil moisture kept constant during the incubation, what was the reasoning for the different incubation times or what were the equidistant temperature changes?

Reply: Soil moisture was measured directly before the experiment and the samples were kept sealed during the experiment. Based on this, we see no reason to expect a change in soil moisture during the experiment. We have added this information at line 146f.
See replies to the detailed comments below for all other aspects mentioned here.

Other comments
5. Provide information on effect sizes (increase of nutrient mobilization due to FTC treatments as compared to control). So far, the effect size is only shortly discussed for the most extreme site with the most extreme treatments. And, as far as I understood, for the models the absolute concentrations are used which, however, might be biased by inherent differences. What is the rationale behind the use of absolute concentrations?

Reply: See our reply to point 3 for the rationale of using absolute concentrations.
Concerning effect sizes, we indeed report the increase in nutrient concentrations only for the most extreme values covered as the three-factorial interactions displayed in Figures 1-3 imply that between control conditions (no FTC) and the extremes, any effect size is possible depending on the factorial combinations of the environmental parameters. We see no option to quantify this multitude of possibilities in one or a few numbers but certainly would be interested in any idea how this could be done!

6. The reading would be facilitated to have the graphs (Fig. 1-3) at the same spot. This would allow to compare the patterns for the three different nutrients. And in general, use the same perspectives for the graphs and in the same order for all three figures.

Reply: The displayed angles of the four-dimensional models (3d plus color code) have been optimized in order to ideally show the response surfaces and their patterns. Unfortunately, we did not find a single perspective and angle which would allow for optimal visibility of the surfaces across all three response parameters. Still, we agree that the displayed views should at least be ordered as comparable as possible and have re-ordered the views to achieve this. In addition, please refer to the animated gifs in the Appendix for the same views across all parameters.
While this comparison between the response parameters is of interest, we deem it more important to show each results graphic close to the text where it is presented and therefore do not combine Figures 1-3 into one Figure 1 A-C.

7. Visualization of measured analytical data is missing. The graphs from the models are great, but showing the measured data would helpful additional information for many scientists working in this field (or at least provide the information in the supplementary material).

Reply: We are happy to share our full dataset to this paper (see data availability statement). We further tried several different options but did not find a convincing way how to display the raw data of our four-dimensional datasets. Unfortunately, all displays of scatterplots in those four dimensions are hardly digestible. Displaying the data in less dimensions, however, is hardly meaningful because of the complex three-factorial interactions (as identified in the hierarchical regression analysis).

Details/Specific comments:
8. Line 43: “more than”?

Reply: compared to their own past. We have added “with climate change” in order to specify this comparison.

9. Line 45: Nutrient limited: Could you be a bit more precise: Which nutrients, is there co-limitation and is it true for all cold-temperate deciduous forests?

Reply: We specifically refer to these points in subchapter 1.3 but have specified the limitation to N and P now already in this very first paragraph (lines 45ff).

10. Line 60: “colder temperatures” than?

Reply: We have added “than 0°C” for clarity.

11. Line 61: What about cell lysis because of drying-rewetting cycles?

Reply: We acknowledge that drying-rewetting cycles can also cause microbial lysis. In fact, it appears hardly possible to distinguish between drying-rewetting and freezing-thawing as they cause similar effects at the cellular level (with freezing-thawing causing the disappearance of liquid water as stated in the text). Still, we prefer to avoid going into details if the underlying process is driven by ‘true’ temperature effects such as physical expansion of the freezing water or by the absence of liquid water here. We rather would like to focus on the ecological consequences of altered soil temperatures here in the introduction.

12. Line 62: freeze thaw cycle: abbreviation already introduced

Reply: Thanks for spotting – we are now using the abbreviation here.

13. Line 65: physiological re-adaptation to thawing conditions may lead to microbial carbon and nutrient release – not sure what this means

Reply: Reformulated to “and the physiological re-activation of microbes when soils are thawing can lead to carbon and nutrient release”

14. Line 89: would be interesting to directly give the amount of N deposition in Central Europe as comparison

Reply: added as suggested; 6 to 45 kg N ha^{-1} $year^{-1}$ for European beech forests (Rennenberg and Dannenmann, 2015).

15. Line 94: P nutrition is recently decreasing: Recently it was researched, but I think the problem is not recent. It is more likely the recent change in C:N:P stoichiometry that can push P to be a limiting factor

Reply: We agree and have reformulated this statement accordingly: “Linked to the increased growth of forest trees with N deposition, phosphorus (P) nutrition of beech appears to become another limiting factor for beech growth on nutrient poor soils.” (L 103ff)

16. There are also attempts to analyses biome patterns of FTC effects (e.g. meta-analysis document higher susceptibility of temperate ecosystem than arctic and high latitude; Gao et al., 2017 Global Change Biology)

Reply: Thanks a lot for pointing out this interesting reference which we now have cited several times in the introduction and discussion.

17. Line 98: FTC has been introduced before

Reply: Thanks for spotting – we are now using the abbreviation here.

18. Provide information on average snow heights or some measure of FTC frequency available. Mean winter temperature are not necessarily a good indicator for FTC.

Reply: We now present soil FTC data measured on site in Table 1. Together with the winter air temperatures and the Precipitation as snow (also Table 1), this data backs up our claim of few FTC at both ends of the climatic gradient due to warm air temperature in the west and due to consistent snow cover in the east.

19. Line 127: soil sub-samples? Could you be more precise? From which horizon(s) where the soils taken? What was the criterion for soil sampling? This is a very crucial information for nutrient dynamics!

Reply: We agree and are surprised ourselves that this info was not given in the first submission: Per subsample, one soil core from 0–10 cm into the A horizon was taken. The whole soil sampling is now described in more detail in the methods section (lines 132ff).

20. Line 135: Information especially about organic matter content but as well of further soil parameters like microbial biomass would be crucial to draw conclusions about the impact of FTC cycles on nutrient mobilization!

Reply: The requested information has been added to Table 1. In short, soil substrate was silty sand in all cases with pH_{CaCl2} ranging 3.1–3.5 and organic matter content ranging 3.7–8.5% in the A horizon.

21. Line 140: Table 2: Phosphate-ion is three times negatively charged, Column soil moisture (SM): change to English punctuation

Reply: We are sorry for these inconsistencies and have corrected them here and checked the whole paper for further incidents.

22. Line 146: to clarify: equidistantly between -1.2 and -12_C : deltaT= 10.8_C / 7 = 1.542_C T intervals?

Reply: Indeed, this was the plan. However, we have now added the characterization of the single levels based on the temperatures measured during the experimental treatments in the climate chambers directly at the samples, as they were less equidistant than expected. Please note that all analyses were run with the directly measured values as stated further down in the methods section (Lines 175ff).

23. Line 160: ‘Temperature directly at the soil samples’. Please provide details. Same depth, same location? How many FTCs have occurred? Why are the logged data not shown (at least in supplemental information)?

Reply: We believe that this was a misunderstanding: Here we reported the temperature measurements during the treatments in the climate chambers, not measurements at the sites of origin. The climate chamber measurements were used for all analyses. We have now added more details about the temperature measurements in the climate chamber (“Temperature was monitored for each of the 49 frequency x magnitude treatment combinations … with 7 sensors per FTC magnitude, directly at the incubated soil samples” Lines 175ff) and furthermore we provided a figure based on this data as Appendix A in the supplementary in order to unequivocally display the FTC treatments.
We agree that soil temperature from the sites of origin would also have great value for the interpretation of the data and have now added FTC measurements over four consecutive years at the sites of origin to Table 1 (compare to point 1).

24. Line 163: Just to clarify: samples with 1 FTC were extracted after one day, samples with 5 FTC after 5 days, controls after 7 days? What is the rational behind the immediate sample extraction after the treatment has finished in comparison to the extraction for all samples after 8 days – at the end of all treatments and with the same incubation time?

Reply: Yes. We decided for this standardized sampling in terms of time after the final FTC for each treatment as otherwise the period between the last FTC and the analysis could interfere with the treatment effects due to e.g. recovery or lagged responses in microbial activity and/or community composition after frost exposition.

25. Line 178: molybdenum blue

Reply: Space added, thanks for spotting this!

26. Line 180: Determination limit: Do you mean detection limit?

Reply: We specifically quantify the determination limit here, which is the same as the ‘limit of quantification’ and provides the value from which the concentration was determined with sufficient precision. The detection limit or ‘limit of detection’ would describe the value above which the analyte is considered to be detected, i.e. significantly higher than a blank value and is calculated from the standard deviation of blank values with a safety factor (= 3 for 10 blank values). In our case, the detection limit = the determination limit / 3.

27. Line 179-180: The determination limit is 0.05 umol L-1 vs. The determination limit
was slightly higher with 0.1 umol L-1? Sentences are unclear

Reply: Indeed, something went wrong here in the initial submission. We have now corrected this, it now reads: “The determination limit was 0.1 µmol l^{-1}. The combined standard uncertainty was 4.2 % for samples and the 5 µM standards.”

28. Line 235: Quality of graphs – resolution, axis labels overlap, axis numbers difficult to read

Reply: We have now increased the resolution and tried to avoid overlaps between labels and axes. However, the latter is not always possible as we rather optimized the figures to display the response surfaces and in some incidents this leads to axes labels hidden partly behind the graphs. Please also refer to the animated graphics in the appendix.

29. Line 235: Graphs: NO3 data: is this the additional release of NO3 compared to control or just the total NO3 release? Please clarify – and I would suggest to use the numbers normed to the control data

Reply: Please compare to point 3 for our rationale to rely on the absolute concentrations measured rather than standardized data.

30. Line 270: Table 4: Would be nice to have the models numbered as done in table 3

Reply: Indeed, that was the plan – added now.

31. Line 286: copy-paste error? Should probably be phosphate instead of ammonium

Reply: True and corrected, thanks a lot for spotting it!

32. Line 291: coldest site or sites?

Reply: The plural was used intentionally here, but we have now reformulated the sentence in order to improve clarity.

33. Line 316: How do we know its short-term? Like a flush? If there was only one measurement after the treatment?

Reply: True. What was meant was that we measured it directly after the FTC but the formulation was ambiguous and we have now deleted “short-term”.

34. Line 322: activating N and P? wording

Reply: Reformulated to “…processes driving the increase in N and P concentrations …”

35. Line 322: (1) minimum temperatures of -7 to -11_C were only reached for half of the treatments, but increase of nutrient concentration seems to increase linearly with increase FTC magnitude. . . which is also shown with the models with only magnitude as single factor where the linear model was the best. Would it not be expected to see a stronger increase of nutrient release when reaching the -7_C if this explanation (1) is right?

Reply: This is true, thanks for this helpful thought! We have now added this aspect here (now at lines 343ff).

36. Line 358: ‘Contrary, our coldest sites rarely experienced serious FTC in the past’. This seems likely but are there any data supporting this statement? As soil had been sampled in well-buffered subsoils, the FTC frequency and magnitude is open

Reply: Please refer to point 1 and the added data in Table 1.

Anonymous Referee #2
The manuscript addresses the effects of FTC magnitude and frequency on the shortterm release of nutrients by conducting a three-factorial gradient experiment. Although the experimental design is simple,

the hierarchical regression analysis was applied to detect the underlying response patterns in the threefold interactive gradient experiment. Therefore, the manuscript is more innovative from the perspective of analytical methods. I think that the manuscript is particularly well-written. The figures are excellent and do a great job of summarizing your results.

Reply: Thanks a lot for this positive feedback and your constructive critique below!

1. Here are some minor suggestions. Abstract Line 14: Generally speaking, we use “intensity” instead of “magnitude”

Reply: Both terms have been used in the past and a quick search through the Web of Science does not provide arguments for one or the other in terms of their frequency of occurrence. We have selected the term based on the general framework of disturbance ecology, which characterizes a disturbance by its duration, abruptness, magnitude and frequency (White & Jentsch 2001 Progress in Botany). We now stick to this wording. Not being native English speakers, though, we would be open for good arguments to change the wording.

2. Line 20: change “higher frost” to “higher FTC”

Reply: Changed as suggested.

3. Line 29-30: The unit representation is incorrect, there is no subscript and superscript.
Please check the full text.

Reply: Checked and corrected throughout the text.

4. Introduction Line 55: delete “(FTC)”

Reply: As the abbreviation was already introduced in line 43 we now use the abbreviation here and deleted the full term.

5. Line 94-96: Compared to nitrogen, there is less description of phosphorus. Could you
add more descriptions about phosphorus.

Reply: True. We have added a few lines on this topic, now at lines 100ff.

6. Materials & methods Line 104: Can you clarify what you mean by “FTC magnitude”;
delete the second “FTC”;

Reply: We have added a short definition (“the minimum temperature reached during the freezing phase of an FTC”; line 113).

7. Line 128: The collection date and depth of soil samples are not clarified. Soils sampled in different seasons have different properties, such as soil water content, soil microbial biomass, soil nutrients, and so on. Soil microorganisms also show different tolerance to changing temperature in different seasons. So, the unrealistic time of soil collection will affect the experimental results. In addition, why should the soil be stored for 16 weeks before starting the experiment? This may change the original physical and chemical properties of the soil.

Reply: To our own surprise, we indeed missed to report the sampling depth and have added this now (line 139: 0-10 cm soil depth) – thanks for spotting this! We have also specified the timing of sampling and the experimental treatments there. The timing was based on the rationale that sampling should occur before natural frost events might happen while the treatments were timed for February when typically the most intensive FTC take place in our study area. We have clarified this rationale at lines 139ff.

8. Line 130: 10 grams of soil seems to be a bit less, which leads to greater intensity and
rate of freeze-thaw cycle than under field conditions.

Reply: We specifically went for this relatively small amount in order to ensure homogeneous temperature dynamics throughout the samples. Otherwise the buffering effect that you correctly describe would interfere with our FTC treatments in a way that the exact minimum temperature per sample could not be determined, as it would differ within the sample. Such ‘controlledness’ is the basis of laboratory experiments and with our sample size we can guarantee that the sample conditions (the temperature of the sample) actually reflected the scenarios that were simulated by the climate chambers, because the samples responded almost immediately to the climate chamber temperature. As stated in the text, we aimed at exploring the discrete and causal relationship between soil temperature and nutrient release which can best be investigated with small soil samples in an ex situ approach. We have now added a short rationale for the relative small sample amounts (line 144).

9. Table 2: change “PO42–P” to “PO43–P”; The value of soil moisture is a dot instead of a comma Line 145-146, 158: Could you show the pattern of freeze-thaw cycle with a figure? (Wang, et al., 2015. Effects of freeze-thaw cycles on the soil nutrient balances, infiltration, and stability of cyanobacterial soil crusts in northern China. Plant and Soil (Figure 2))

Reply: We have corrected the inconsistencies – thanks for spotting! We have furthermore added a graphical representation of the FTC treatments to the Appendix A of the paper for clarification.

10. Line 197-198: Please explain in detail how to use the AICc to determine the best model

Reply: We have added a short explanation at lines 212ff.

11. Line 232: Please explain the abbreviation of AICc

Reply: We have now introduced this abbreviation already at line 212, but have also added it to the table captions.

12. Results Line 239: Explain abbreviations in the legend (“FTC”)

Reply: Done.

13. Line 261: Change “were” to “was”

Reply: Done.

14. Discussion Line 315-330: In the paragraph about FTC effects, you discuss potential mechanisms leading to increases in inorganic N and P. The whole discussion did not involve the discussion about phosphorus. Could you add some discussion about phosphorus.

Reply: We have added a short paragraph on implications for N:P imbalance at lines 406ff.

15. Line 336: delete the second “FTC”

Reply: Done.

Soils from cold and snowy temperate deciduous forests release more nitrogen and phosphorus after soil freeze-thaw cycles than soils from warmer, snow-poor conditions

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Abstract. Effects of global warming are most pronounced in winter. A reduction in snow cover due to warmer atmospheric temperature in formerly cold ecosystems, however, could counteract an increase in soil temperature by reduction of insulation. Thus, soil freeze-thaw cycles (FTC) might increase in frequency and magnitude with warming, potentially leading to a disturbance of the soil biota and release of nutrients.

Here, we assessed how soil freeze-thaw magnitude and frequency affect short-term release of nutrients in temperate deciduous forest soils by conducting a three factorial gradient experiment with ex-situ soil samples in climate chambers. The fully-crossed experiment included soils from forests dominated by *Fagus sylvatica* (European beech) that originate from different winter climate (mean coldest month temperature range $\Delta T > 4$ K), a range of FTC magnitudes from no ($T = 4.0$ °C) to strong ($T = -11.3$ °C) soil frost, and a range of FTC frequencies ($f = 0$–7). We hypothesized that higher FTC magnitude and frequency, respectively, will increase the release of nutrients. Furthermore, soils from cold climates with historically stable winter soil temperatures due to deep snow cover will be more responsive to FTC than soils from warmer, more fluctuating winter soil climates.

FTC magnitude and, to a lesser extent, also FTC frequency resulted in increased nitrate, ammonium, and phosphate release almost exclusively in soils from cold, snow-rich sites. The hierarchical regression analyses of our three-factorial gradient experiment revealed that the effects of climatic origin (mean minimum winter temperature) followed a sigmoidal curve for all studied nutrients and was modulated either by FTC magnitude (phosphate) or by FTC magnitude and frequency (nitrate, ammonium) in complex two- and, for all studied nutrients, in threefold interactions of the environmental drivers. Compared to initial concentrations, soluble nutrients were predicted to increase to 250 % for nitrate (up to 16 µg NO_3-N kg^{-1}DM), to 110 % for ammonium (up to 60 µg NH_4-N kg^{-1}DM), and to 400 % for phosphate (2.2 µg PO_4-P kg^{-1}DM) at the coldest site for strongest magnitude and highest frequency. Soils from warmer sites showed little nutrient release and were largely unaffected by the FTC treatments except for above-average nitrate release at the warmest sites in response to extremely cold FTC magnitude.

We suggest that currently warmer forest soils have historically already passed the point of high responsiveness to winter climate change, displaying some form of adaptation either in the soil biotic composition or in labile nutrient sources. Our data suggests that previously cold sites, which will lose their protective snow cover during climate change, are most vulnerable to increasing FTC frequency and magnitude, resulting in strong shifts in nitrogen and phosphorus release. In nutrient poor European beech forests of the studied Pleistocene lowlands, nutrients released over winter may be leached out, inducing reduced plant growth rates in the following growing season.

1 Introduction

Climate is warming over-proportionally in northern latitudes and during winter (IPCC, 2013). This has potentially important consequences for nutrient cycling and ecosystem functioning (Kreyling, 2020). Cold-temperate deciduous forests are experiencing more fluctuating soil temperatures and potentially also more frequent soil freeze-thaw cycles (FTC) with climate change because reduced or completely missing snow cover exposes them to strongly fluctuating air temperatures (Kreyling, 2020). These forests are typically nitrogen limited (Bontemps et al., 2011) with phosphorus co-limitation increasing in face of nitrogen deposition and climate change (Talkner et al., 2015; Peñuelas et al., 2013). Soil nitrogen and phosphorus release in response to FTC frequency and FTC magnitude of forests differing in their past and present climate are therefore of high ecological and economical importance.

1.1 *Winter climate change in the temperate deciduous forests of Central Europe*

Winters in temperate regions are projected to become warmer, more variable, and wetter with precipitation increasing and changing from snow to rain (Stocker, 2014; Yang and Christensen, 2012). Largest decreases in snowfall are expected for regions with winter mean air temperatures ranging from -5 to +5 °C, while colder regions (boreal, arctic) might even receive increased snowfall (Brown and Mote, 2009; Scherrer and Appenzeller, 2006). The shift from snow to rain drastically reduces soil insulation and exposes soils to the fluctuations of air temperatures (Groffman et al., 2001). While insulation by snow can prevent soil freezing even in boreal climates (Isard and Schaetzl 1998), missing snow can lead to increased soil frost in regions with sustained air frost (Groffman et al., 2001; Brown and DeGaetano, 2011; Henry, 2008), increased frequency of FTC in regions where air temperatures fluctuate around 0 °C (Henry, 2008; Campbell et al., 2010), or reduced soil frost where even minimum air temperatures rarely drop below the freezing point (Kreyling and Henry, 2011).

1.2 *Ecological consequences of altered soil temperatures*

Many relevant ecological processes are driven by winter soil temperatures such as activity and survival of organs and organisms (Kreyling, 2010; Campbell et al., 2005). Soil freezing represents an important threshold for microbial activity because of reduced availability of liquid water (Mikan et al., 2002). However, colder temperatures than 0°C are typically required to cause microbial lysis as microbial growth can continue below freezing (McMahon et al., 2009). Sub-lethal effects of freezing on soil microorganisms are not well understood, and the length of freezing, the number of FTC, and the rate of freezing can all increase cell damage for a given freezing magnitude (Elliott and Henry, 2009; Vestgarden and Austnes, 2009). In addition, soil microorganisms which survive freezing and desiccation can be lethally damaged via osmotic shock upon exposure to melt water (Jefferies et al., 2010) and the physiological re-activiation of microbes when soils are thawing can lead to carbon and nutrient release (Schimel et al., 2007). Consequently, soil freezing can disrupt soil microbial activity (Bolter et al., 2005; Yanai et al., 2004) and affect key microbial processes such as ammonification, nitrification and denitrification (Urakawa et al., 2014; Watanabe et al., 2019; Hosokawa et al., 2017). Furthermore, soil freezing can damage plant roots (Tierney et al., 2001; Reinmann and Templer, 2018; Kreyling et al., 2012a; Weih and Karlsson, 2002), induce soil nitrogen (N) leaching (Joseph and Henry, 2009; Matzner and Borken, 2008), increase soil trace gas losses (Reinmann and Templer, 2018; Matzner and Borken, 2008), reduce N uptake by trees (Campbell et al., 2014), decrease plant productivity (Göbel et al., 2019; Comerford et al., 2013; Reinmann et al., 2019) and can ultimately lead to plant mortality (Schaberg et al., 2008; Buma et al., 2017). In addition to direct frost damage, the listed consequences of soil freezing on plant performance are commonly explained by altered nutrient, mainly N and P, availabilities (Kreyling, 2020). Freezing can also affect release of these nutrients by physically breaking up soil aggregates (Oztas and Fayetorbay, 2003) or organic litter (Hobbie and Chapin, 1996) and by reducing soil water flow rates (Iwata et al., 2010).

Changes in FTC frequency can affect microbial communities, e.g. increasing saprotrophic fungal activity (Kreyling et al., 2012b). Nitrogen leaching from soil columns subjected to FTC remaining high even after 10 FTC further emphasizes the importance of FTC frequency (Joseph and Henry, 2008). A recent meta-analysis indicates that FTC increase ammonium (+19%) and nitrate (+18%) concentrations, nitrate leaching (+67%) and N_2O emissions (+145%) while soil total N (-26%) and microbial biomass N (-5%) decreased (Gao et al., 2018). Interestingly, temperate ecosystems appeared to be more responsive than arctic or alpine systems in this study. Taken together, FTC can affect soil nutrient release through damage and lysis of microbial and plant cells, through altered soil biotic activity, and/ or through physical disruption of abiotic and dead organic particles. In particular for nutrient limited ecosystems, altered occurrence of FTC with climate change could consequently affect ecosystem functioning.

1.3 *Beech forests of Pleistocene lowlands as important and potentially affected ecosystem*

Beech forests are the zonal vegetation of temperate Central Europe and face multiple anthropogenic pressures while still providing vast ecosystem services (Ammer et al., 2018). Beech (*Fagus sylvatica* L.) naturally dominates all over Central Europe under a wide range of soil conditions and occurs in regions with less than 550 to more than 2000 mm of annual rainfall on nearly all geological substrates if drainage is sufficient (Leuschner et al., 2006). Even when growing on marginal soils, beech forests have a N demand of about 100 kg N ha^{-1} $year^{-1}$ which is several times higher than current atmospheric N-deposition in European beech forests that range from 6 to 45 kg N ha^{-1} $year^{-1}$ (Rennenberg and Dannenmann, 2015). Nitrogen availability is consequently still the most limiting factor of beech growth at marginal as well as at productive sites (Bontemps et al., 2011). N availability is largely determined by internal N cycling through microbial mineralization and immobilization

(Guo et al., 2013). Any alteration in the microbial community and activity, such as in response to FTC, therefore has the potential to affect nutrient cycling and, thereby, ecosystem functioning of this ecologically and economically important ecosystem (Simon et al., 2017).

Linked to the increased growth of forest trees with N deposition, phosphorus (P) nutrition is becoming another limiting factor for beech growth in particular on nutrient poor soils (Talkner et al., 2015). As P input into unfertilized ecosystems such as forests still relies solely on bedrock weathering while N and C input strongly increases with global change, P deficiency and unparalleled imbalances in C:N:P stoichiometry occur (Peñuelas et al., 2013). Generally, increasing substrate N:P ratios are related to forest growth declines and increasing P limitation with forest age is a global phenomenon (Wardle et al., 2004). Implications of climate change on P release of beech forest soils should therefore also be investigated.

1.4 *Hypotheses*

We hypothesized that soil FTC induce nutrient release following saturation curves both with increased FTC magnitude and increased FTC frequency. We expected the combination of FTC magnitude and FTC frequency to be additive. We further hypothesized that soils from colder macroclimates which are characterized by more persistent and protective snow cover are more responsive in release of nutrients in the face of FTC than soils from warmer sites with more fluctuating winter soil temperatures.

2 Materials & methods

The effects of FTC magnitude, i.e. the minimum temperature reached during the freezing phase of an FTC, and FTC frequency, i.e. the number of consecutive FTC, on the short-term release of nutrients in temperate deciduous forest soils was assessed in a three-factorial gradient experiment with *ex-situ* soil samples in climate chambers. The fully-crossed experiment included soils from seven forests dominated by *Fagus sylvatica* (beech) that (1) originate from different winter climate (mean winter minimum temperature range $\Delta T > 4$ K) and were exposed to (2) a range of FTC magnitudes from no ($T = 4.0$ °C) to strong ($T = -11.3$ °C) soil frost, and (3) a range of FTC frequencies ($f = 0$–7).

2.1 *Forest sites and soil sample collection*

Soil samples for this study stemmed from seven sites located between Rostock (Germany) and Gdansk (Poland) which are mono-dominated by mature European beech. Along the 500-km study gradient, the sites differ markedly in winter climate with mean average winter air temperatures ($\Delta T = 4.0$ K) and mean minimum winter air temperatures ($\Delta T = 3.8$ K) decreasing towards the east, which over-proportionally drives the differences in mean annual temperature ($\Delta T = 2.8$ K; for details see Table 1). From west to east, mean annual precipitation as snow increases from 50 to 110 mm while annual precipitation is rather uniform (540 to 630 mm). With respect to winter air temperature differences, the study area is representative of a large part of the temperature range of beech as the major forest tree in Europe, while for summer precipitation, which is considered to be a major driver of beech growth (Hacket-Pain et al., 2018), differences are relatively small (Table 1).

The study sites are located in the Pleistocene lowlands with glacial deposits as bedrock. All sites share the same soil type (sandy Cambisol) and similar soil texture (sandy silt to silty sand). Sites were selected for similar forest stand structure, i.e. tree height about 30 m (ranging between 27–39 m), tree diameter about 45 cm (ranging between 37–52 cm), and canopy closure 70–80 %. In order to achieve this uniform stand structure, differences in mean tree age across sites was accepted (76-167 years). At each site, we systematically selected the sampling sites in proximity to site-representative target trees. A dendroecological pre-study (Weigel et al., 2018) identified these target tree individuals by selecting for the best correlations between individual tree-ring series and the site chronology (the mean of all individual tree-ring series of a site) during the last 30 years (three target trees out of 20 at all but coldest site, three out of 40 at coldest site). Consequently, the three selected target trees within each site showed very similar growth patterns over the past 30 years and ideally represented the growth–environment relationship of the whole stand. At each site, we randomly selected one of those three target trees and took three soil sub-samples (later on mixed) at a distance of 3 m in northeast, south, and northwest direction from each selected individual. Sampling occurred at 0-10 cm soil depth starting below the organic litter layer. The litter layer was, as it is typical for beech forests, very thin at the time of sampling in early November. Sampling was timed before natural FTC would interfere with our treatments. Samples were stored at 4 °C until the start of the treatments in early February, which is the time when, typically, the most intensive FTC happen in our study area.

The mixed samples per site were carefully homogenized and subsampled to 10 g for the subsequent FTC treatment (see below). This small amount ensured homogeneous temperature dynamics throughout the samples. Soil moisture at the start of the FTC treatment ranged between 19.4 and 36.6 % between the sites and was not significantly related to climate at site origin (correlation to mean minimum air temperature: $R^2 = 0.33$, $p = 0.103$). The samples were kept sealed during the experiment and, hence, soil moisture was assumed to stay constant. Initial values for the analyzed nutrients were also recorded at the start of the FTC treatment with the same methodology as described below and are presented in Table 2.

Table 1: Site characteristics for the seven sampled beech forest stands, ordered by decreasing winter minimum air temperature. All climatic data is display as means for the reference period 1961–1990 according to “climateEU” 4.63, (Hamann et al., 2013; Wang et al., 2012), winter refers to Dec.(prev. yr) - Feb., summer refers to Jun. – Aug. Soil parameters were measured directly on site for the years 2016-2019. ‘# FTC’ indicates the number of free-thaw cycles at the specified soil depth measured at half-hourly intervals by TMC20-HD temperature sensors connected to HOBO UX120-006M Analog Data Loggers (Onset Computer Corporation, Bourne, USA).

Site	Geography		Climate							Soil						
	Longitude (°)	Latitude (°)	Winter minimum temperature (°C)	Annual temperature (°C)	Winter temperature (°C)	Summer temperature (°C)	Annual precipitation (mm)	Precipitation as snow (mm)	Summer precipitation (mm)	Frost degree hours at soil surface (Σ h × $\text{Temp}_{<0°C}$)	Frost degree hours at -5 cm (Σ h × $\text{Temp}_{<0°C}$)	# FTC at soil surface	# FTC at -5 cm	C/N ratio	Organic matter content	pH_{CaCl2}
BH	12.32	54.12	-2.1	8.0	0.2	15.9	588	48	191	2.9	0.0	1	0	16.0	5.9	3.5
NZ	13.14	53.39	-2.9	7.8	-0.6	16.3	580	53	193	352.0	3.1	14	1	16.8	4.5	3.5
BB	13.83	53.11	-3.2	8.4	-0.8	17.1	568	51	188	363.3	96.8	19	1	17.5	3.7	3.5
GR	14.73	53.32	-3.8	8.2	-1.4	17.0	568	57	189	205.0	11.8	3	1	19.6	6.9	3.3
WE	18.08	54.72	-4.2	7.0	-2.2	15.9	623	82	204	119.2	0.0	11	0	21.8	8.5	3.1
KO	18.43	54.25	-5.5	5.6	-3.4	14.4	593	99	215	273.9	0.0	11	0	25.4	8.1	3.2
KA	18.14	54.24	-5.9	5.9	-3.8	14.8	621	107	218	68.4	0.0	1	0	17.2	8.3	3.3

Table 2: Initial nutrient concentrations (µg kg^{-1} DM; mean ± SD) and gravimetric soil moisture at the start of the FTC treatment.

Site	NO_3^--N	NH_4^+-N	PO_4^{3-}-P	SM (%)
BH	15.0±0.7	11.8±1.2	0.09±0.02	28.1
NZ	9.7±2.7	15.8±1.6	0.12±0.02	19.6
BB	15.0±0.5	14.6±0.9	0.09±0.01	19.4
GR	9.1±0.3	67.5±11.0	0.14±0.04	27.2
WE	1.2±0.3	25.5±1.1	0.48±0.52	36.6
KO	14.8±0.8	28.1±2.3	0.30±0.28	31.6
KA	6.0±1.8	55.3±8.2	0.60±0.64	36.4

2.2 *FTC treatment*

The FTC treatment was set up as a fully factorial combination of sample site, FTC magnitude, and FTC frequency in a gradient design consisting of seven sites along a gradient of winter climate (see above), seven FTC magnitudes (realized at -1.9, -2.5, -3.4, -4.6, -6.6, -7.8, -11.1°C, respectively), and seven FTC-frequencies (f = 1-7). In addition, three control samples without FTC (T = 4.0°C and f = 0) were analyzed at the end (day 8) of the experiment for each site respectively. In total, this resulted in 364 samples (7 sites x 7 FTC magnitudes x 7 FTC frequencies + 7 x 3 controls). Gradient experiments with unique (unreplicated) sampling at each factorial combination have recently been shown to outperform classical, replicated designs in terms of detecting and characterizing potentially non-linear ecological response surfaces of interacting environmental drivers (Kreyling et al., 2018). Such designs profit from expanding the range of environmental drivers and are therefore recommended to include extreme and rather unrealistic values such as the maximum FTC magnitude in our example. Soil temperatures of -12°C rarely occur in temperate forests. However, they can help elucidating response patterns and might even become possible as future warming of the Polar Ocean might increase advection of polar air masses, potentially causing unprecedented cold extremes over Europe (Petoukhov and Semenov, 2010; Yang and Christensen, 2012).

The simulated FTC followed typical FTC for temperate ecosystems with daily cycles between thawed and frozen states. The FTC treatment was realized for all samples in parallel in programmable climate chambers (Percival LT-36VLX, Percival Scientific Inc., Perry/Iowa). One FTC lasted 24 h with 2 h at the preset minimum temperature and 12 h at +1°C (sufficient for thawing but too cold for considerable microbial activity). The rates of temperature change consequently differed between FTC magnitudes but was < 3 K h^{-1} even for the coldest magnitude. Temperature was monitored for each of the 49 frequency x magnitude treatment combinations (see above) with 7 sensors per FTC magnitude, directly at the incubated soil samples (LogTag TRIX 8, LogTag Recorders Lt, Auckland, New Zealand) and the realized minimum temperatures per treatment combination rather than the preset temperature of the climate chambers were used for further analysis (see Appendix A for a visual display of the treatments). Directly after the planned FTC frequency was reached for each sample, nutrient extraction and the subsequent chemical analysis started.

2.3 *Nutrient extraction and chemical analysis*

Samples were shaken in 50 ml KCl solution (0.5 M) for 1 h and subsequently filtered through filter paper of 2-3 µm pore size. Afterwards, the filtrates were stored frozen at -20 °C upon further analysis.

Nitrate was measured after conversion to nitrite at a cadmium reductor column as an azodye (Hansen and Koroleff, 1999). Samples had to be diluted with ultrapure water (Purelab Flex, Elga) by 50 times. Nitrite was not measured, because its concentration was expected to be <10 % of nitrate. The nitrate named data are, therefore, the sum of nitrate and nitrite (NO_x). The samples were measured in a segmented flow analyser (FlowSys, Alliance Instruments) equipped with a 5 cm cuvette (Armstrong et al., 1967). Determination limit for nitrate was 0.32 µmol l^{-1} (4.5 µg nitrate N l^{-1}). The combined standard uncertainty was 4.2 % for samples and the 5 µM standards.

Ammonium was measured as an indophenol blue dye photometrically (Hansen and Koroleff, 1999). Samples had to be diluted by 50-100 times. The samples were measured in the same segmented flow analyser (Kérouel and Aminot, 1997). Determination limit for ammonium was 0.43 µmol l^{-1} (6.0 µg ammonium N l^{-1}). The combined standard uncertainty was 7.7 % for samples and the 5 µM standards.

Phosphate concentrations were measured by the molybdenum blue reaction photometrically (Murphy and Riley, 1962). This was done in the above mentioned autoanlyser (Malcolme-Lawes and Wong, 1990). The determination limit was 0.1 µmol l^{-1}. The combined standard uncertainty was 4.2 % for samples and the 5 µM standards.

Fresh weight of each sample was determined before the start of the FTC treatment. Based on the relation between dry weight and fresh weight of a further subsample, dry weight of the samples was calculated and nutrient concentrations are reported in relation to dry weight.

2.4 *Statistical analyses*

Hierarchical regression analysis was applied to detect and characterize the underlying response patterns in our threefold interactive gradient experiment according to the recommendations by Kreyling et al. (2018). In short, the hierarchical regression analysis accepts a more complex model only if it explains the data better than a simpler model, indicated by lower Akaike Information Criterion (AIC) and, for nested designs, by significant ANOVA comparing the models. Consequently, the final model of a hierarchical regression analysis contains only those parameters and interactions which help representing the underlying data, i.e. which are significant for the interpretation of the data.

We first performed linear regression for each single environmental driver (climatic origin expressed as mean minimum air temperature at the respective sampling site; FTC magnitude expressed as the minimum temperature experienced during the FTC treatment; and FTC frequency expressed as the number of FTC). Based on the hypothesized non-linear relationship of nutrient release with these environmental drivers, we then set up different non-linear candidate models for each environmental driver individually. We chose models known for their ability to describe a wide variety of ecological and biological processes, i.e. a saturating model (Michaelis-Menten function) and a sigmoidal model (Gompertz function). We used the model performance index AICc (Akaike Information Criterion corrected for small sample sizes, (Hurvich and Tsai, 1989) to determine the best model, which is indicated by the lowest AICc-value. In case of assessing model performance of linear models or comparing model performance of nested models, we also used ANOVA to test whether the more complex model explained variation significantly better than the simpler model. We continued by additive combination of the two best explaining individual models and kept this new model only if it further increased explained variation (lower AICc and significant model difference in ANOVA). Likewise, we tested whether addition of interactive terms and the third environmental driver and all other interactive terms between the three drivers to the previous best model further increased model quality. All steps and all model formulations are documented in Tables 3-5.

All analyses were performed in R 3.4.3 (R Core Team, 2017). Candidate models were fit to the data using ‘nlsLM()’ of package ‘minpack.lm’ version 1.2-1. AICc was quantified using ‘AICc()’ of package ‘AICcmodavg’ 2.2-1. The overall best model for each response parameter was visualized using ‘scatter3D()’ of package ‘plot3D’ version 1.1.1 and a correlation between measured nutrient release and predicted nutrient release by the model was used to quantify its goodness of fit.

3 Results

3.1 *Nitrate*

Variation in initial mobile nitrate concentration was large between sample sites (10.1 µg NO_3-N per kg dry matter on average ± 5.2 µg NO_3-N standard deviation across site averages). Nitrate concentrations at the end of our three-way gradient experiment followed a sigmoid increase towards colder winter minimum temperatures at the sample’s origin, which was further modulated by an interaction with FTC magnitude, an interaction between FTC magnitude with FTC frequency, and the three-way interaction between mean minimum temperature at origin, FTC magnitude, and FTC frequency (Table 3, model 15.). This model achieved a correlation between measured and predicted nitrate concentrations of 0.46. According to this model, highest nitrate concentrations and highest frost sensitivity occurred for the combination of the coldest site, the strongest FTC magnitude, and the highest FTC frequency (Figure 1) with predicted values of up to 16 µg NO_3-N per kg dry matter, i.e. a 2.5-fold increase as compared to the initial nitrate concentration before the start of the experiment at this site (site KA, Table 2). For this combination, also the maximum measured value was found with nitrate concentrations of 37.3 µg NO_3-N per kg dry matter. Single, strong FTC ($T = - 11$ and $f = 1$), however, also released above average amounts of nitrate for the warmest site. Lowest nitrate concentrations were found for all sites at the mildest FTC magnitude irrespective of FTC frequency. For mild FTC magnitudes, all sites showed below average nitrate concentrations with highest, still below-average, concentrations for the warmest site.

Individually, neither FTC magnitude nor FTC frequency were able to significantly explain nitrate concentrations, more complex saturating or sigmoid models being indistinguishable from the (non-significant) linear model for both parameters (Table 3 models 1.-6.).

Table 3. Results of the hierarchical regression analysis for nitrate concentrations of beech forest soils to changes in FTC magnitude (x_1), FTC frequency (x_2), and climatic origin (x_3; expressed as mean minimum winter temperature at origin) at the end of the FTC treatments. Tested are linear, saturating (Michaelis Menten function) and sigmoid (Gompertz function) relationships on the single environmental drivers and their interactions. Bold AICc (Akaike Information Criterion corrected for small sample sizes) values indicate best model. AICc in italics indicate best single-factor models. a_1 to a_n are the fitted parameters of the respective model. FTC = freeze-thaw cycle.

Model description	model	AICc	Notes
1. Linear, magnitude (x_1) only	$y = a_1x_1 + a_2$	*2424*	*Simplest possible start, lm: p = 0.215*
2. Saturating, magnitude only	$y = \frac{a_1 * x_1}{a_2 + x_1}$	2800	Not better than 1.
3. Sigmoid, magnitude only	$y = a_1 * e^{-a_2 * e^{-a_3x_1}}$	2426	Not better than 1.
4. Linear, frequency (x_2) only	$y = a_1x_2 + a_2$	*2425*	*Simplest possible start, lm: p = 0.537*
5. Saturating, frequency only	$y = \frac{a_1 * x_2}{a_2 + x_2}$	2485	Not better than 4.
6. Sigmoid, frequency only	$y = a_1 * e^{-a_2 * e^{-a_3x_2}}$	2427	Not better than 4.
7. Linear, climatic origin (x_3) only	$y = a_1x_3 + a_2$	2422	Simplest possible start, lm: p = 0.066
8. Saturating, climatic origin only	$y = \frac{a_1 * x_3}{a_2 + x_3}$	2383	Better than 7.
9. Sigmoid, climatic origin only	$y = a_1 * e^{-a_2 * e^{-a_3x_3}}$	*2362*	*Better than 7. and 8., best single factor model*
10. Sigmoid climatic origin and linear magnitude (additive)	$y = a_1 * e^{-a_2 * e^{-a_3x_3}} + a_4x_1$	2363	Taking the best model of the best explaining parameter so far (9.) and adding the best model of the second best explaining parameter (1.)
11. Sigmoid climatic origin and its interaction with magnitude	$y = a_1 * e^{-a_2 * e^{-a_3x_3}} + a_4x_3x_1$	2354	Interaction term instead of single factor in 10. new best model
12. Sigmoid climatic origin and its interaction with magnitude and linear frequency	$y = a_1 * e^{-a_2 * e^{-a_3x_3}} + a_4x_1x_3 + a_5x_2$	2354	Adding best model of third parameter (4.) to best model so far (11.) not better than 11.
13. Sigmoid climatic origin and its two-way interaction with magnitude and frequency, respectively	$y = a_1 * e^{-a_2 * e^{-a_3x_3}} + a_4x_1x_3 + a_5x_2x_3$	2356	Adding interaction term climatic origin x frequency to best model so far (11.) ANOVA: not different from 11. with p = 0.671 not better than 11.
14. Sigmoid climatic origin and its two-way interaction with magnitude	$y = a_1 * e^{-a_2 * e^{-a_3x_3}} + a_4x_1x_2 + a_5x_2x_3$	2352	Adding two-way interaction magnitude x frequency to best model so far (11.)

and two-way interaction magnitude x frequency			ANOVA: marginally different from 11. with p = 0.064 new best model
15. Sigmoid climatic origin and its two-way interaction with magnitude and the three-way interaction (climate origin x frequency x magnitude)	$\boldsymbol{y = a_1 * e^{-a_2 * e^{-a_3 x_3}} + a_5 x_1 x_2 + a_6 x_1 x_3 + a_7 x_2 x_3 + a_8 x_1 x_2 x_3}$	2548	**Adding three-fold interaction term to best model so far (14.)** **ANOVA: different from 14. with p = 0.007** **best model**
16. Linear magnitude and linear frequency without interaction (additive)	$y = a_1 + a_2 x_1 + a_3 x_2$	2425	Checking interaction between magnitude and frequency ANOVA not different from best single factor model (1.): p = 0.309
17. Linear magnitude and linear frequency with interaction	$y = a_1 + a_2 x_1 + a_3 x_2 + a_4 x_1 x_2$	2425	Checking interaction between magnitude and frequency ANOVA not different from best single factor model (1. p = 0.487) and additive model (16. p = 0.525)

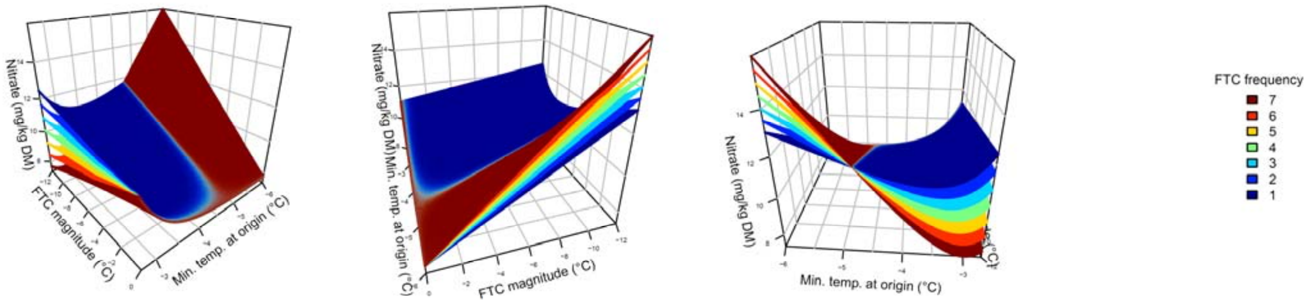

Figure 1: Nitrate concentrations were best explained by the three-fold interactive effects of winter climatic origin (expressed as longterm mean minimum winter temperature at the origin), FTC magnitude (expressed as the minimum temperature experienced during the FTC manipulation and displayed for freezing temperatures) and FTC frequency during the FTC manipulation. FTC = freeze-thaw cycle. The four dimensional representation is displayed from three different angles (see Appendix B for an animated version) and is based on the best model fit in the hierarchical regression analysis (model 15. in Table 3 with coefficients $a_1 = 7.70092$; $a_2 = -22.57795$; $a_3 = 1.52874$; $a_4 = 0.06754$; $a_5 = 0.15402$; $a_6 = 0.03231$).

3.2 *Ammonium*

Variation in initial mobile ammonium concentration was large between sample sites (31.2 µg NH_4-N per kg dry matter on average ± 21.7 µg NH_4-N standard deviation across site averages). Ammonium concentrations after the FTC treatments followed a sigmoid increase with colder winter minimum temperature at the sample's origin, an additive linear increase with FTC frequency, and were further modulated by an interaction between FTC magnitude with FTC frequency, and the three-way interaction between mean minimum temperature at origin, FTC magnitude, and FTC frequency (Table 4, model 15.). This model achieved a correlation between measured and predicted ammonium concentrations of 0.61. According to this model, highest ammonium concentrations and highest frost sensitivity occurred for the combination of the coldest site, the strongest FTC magnitude, and the highest FTC frequency (Figure 2) with predicted values of up to 60 µg NH_4-N per kg dry matter, i.e. a 10 % increase as compared to the initial ammonium concentration before the start of the experiment at this site (site KA, Table 2). For this combination, also the maximum measured value was found with ammonium concentrations of 149.7 µg NH_4-N per kg dry matter. At this site, FTC frequency had its highest and positively modulating effect while almost no effect of FTC frequency was found for mild FTC magnitude across all origins. Predicted ammonium concentrations and sensitivity to frost decreased rapidly towards the warmer sites with the inflection point of the sigmoid shape at around -3°C for high FTC magnitudes and -2°C for mild FTC magnitudes. Lowest ammonium concentrations were predicted for the warmest site almost irrespective of FTC magnitude and FTC frequency.

Individually, FTC frequency, but not FTC magnitude, was able to significantly explain ammonium concentrations, more complex saturating or sigmoid models being indistinguishable from the linear model for both parameters (Table 4 models 1.-6.). Their interaction appeared relevant and non-additive (Table 4 models 16. and 17.)

Table 4. Results of the hierarchical regression analysis for ammonium concentrations of beech forest soils to changes in FTC magnitude (x_1), FTC frequency (x_2), and climatic origin (x_3; expressed as mean minimum winter temperature at origin) at the end of the FTC treatments. Tested are linear, saturating (Michaelis Menten function) and sigmoid (Gompertz function) relationships on the single environmental drivers and their interactions. Bold AICc (Akaike Information Criterion corrected for small sample sizes) values indicate best model. AICc in italics indicate best single-factor models. a_1 to a_n are the fitted parameters of the respective model. FTC = freeze-thaw cycle.

Model description	model	AICc	Note
1. Linear, magnitude (x1) only	$y = a_1 x_1 + a_2$	3092	Simplest possible start, lm: p = 0.182
2. Saturating, magnitude only	$y = \frac{a_1 * x_1}{a_2 + x_1}$	3510	Not better than 1.
3. Sigmoid, magnitude only	$y = a_1 * e^{-a_2 * e^{-a_3 x_1}}$	3096	Not better than 1.
4. Linear, frequency (x2) only	$y = a_1 x_2 + a_2$	3088	Simplest possible start, lm: p < 0.015
5. Saturating, frequency only	$y = \frac{a_1 * x_2}{a_2 + x_2}$	3155	Not better than 3.

6.	Sigmoid, frequency only	$y = a_1 * e^{-a_2 * e^{-a_3 x_2}}$	3088	Not better than 3.
7.	Linear, climatic origin (x3) only	$y = a_1 x_3 + a_2$	2967	Simplest possible start, lm: p < 0.001
8.	Saturating, climatic origin only	$y = \frac{a_1 * x_3}{a_2 + x_3}$	2965	Better than 7.
9.	Sigmoid, climatic origin only	$y = a_1 * e^{-a_2 * e^{-a_3 x_3}}$	2954	Better than 8., best single factor model
10.	Sigmoid climatic origin and linear frequency (additive)	$y = a_1 * e^{-a_2 * e^{-a_3 x_3}} + a_4 x_2$	2946	Taking the best model of the best explaining parameter so far (9.) and adding the best model of the second best explaining parameter (4.) New best model
11.	Sigmoid climatic origin and linear frequency (with interaction term)	$y = a_1 * e^{-a_2 * e^{-a_3 x_3}} + a_4 x_2 + a_5 x_3 x_2$	2948	Adding an interaction term to 10. ANOVA: not different from 10. with p =0.570
12.	Sigmoid climatic origin and linear frequency and linear magnitude	$y = a_1 * e^{-a_2 * e^{-a_3 x_3}} + a_4 x_2 + a_5 x_1$	2947	Adding best model of third parameter (1.) to best model so far (10.) ANOVA: not different form 10. with p = 0.219
13.	Sigmoid climatic origin and linear frequency and interaction climatic origin x magnitude	$y = a_1 * e^{-a_2 * e^{-a_3 x_3}} + a_4 x_2 + a_5 x_1 x_3$	2946	Adding interaction term climatic origin x magnitude to best model so far (10.) ANOVA: not different from 10. with p = 0.219
14.	Sigmoid climatic origin and linear frequency and interaction frequency x magnitude	$y = a_1 * e^{-a_2 * e^{-a_3 x_3}} + a_4 x_2 + a_5 x_1 x_2$	2939	Adding interaction magnitude x frequency to best model so far (10.) ANOVA: different from 10. with p = 0.002 New best model
15.	Sigmoid climatic origin and linear frequency and two-way interaction frequency x magnitude and three-way interaction	$\boldsymbol{y = a_1 * e^{-a_2 * e^{-a_3 x_3}} + a_4 x_2 + a_5 x_1 x_2 + a_6 x_1 x_2 x_3}$	2937	Adding three-fold interaction term to best model so far (14.) ANOVA: different from 14. with p = 0.025 best model
16.	Linear frequency and linear magnitude without interaction (additive)	$y = a_1 + a_2 x_2 + a_3 x_1$	3090	Checking interaction between magnitude and frequency ANOVA different from best single factor model (4.): p = 0.031
17.	Linear frequency and linear magnitude with interaction	$y = a_1 + a_2 x_2 + a_3 x_1 + a_4 x_1 x_2$	3083	Checking interaction between magnitude and frequency ANOVA different from best single factor model (4. p = 0.001) and additive model (16. p = 0.003)

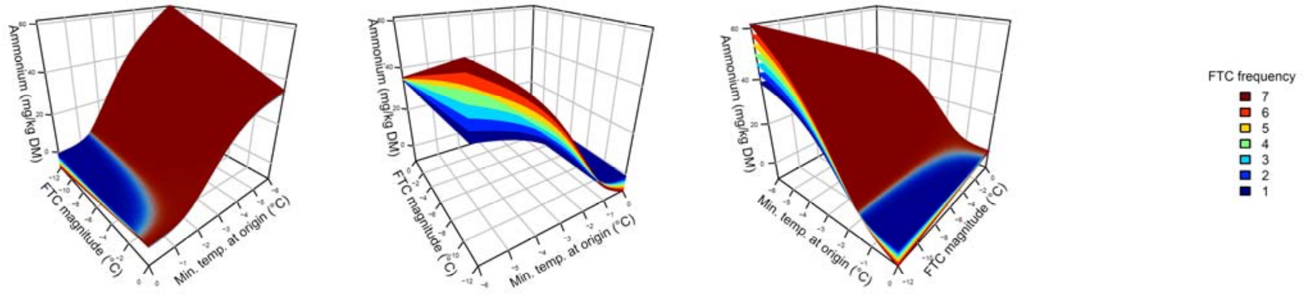

Figure 2: Ammonium concentrations were best explained by the three-fold interactive effects of winter climatic origin (expressed as longterm mean minimum winter temperature at the origin), FTC magnitude (expressed as the minimum temperature experienced during the FTC manipulation and displayed for freezing temperatures) and FTC frequency during the FTC manipulation. FTC = freeze-thaw cycle. The four dimensional representation is displayed from three different angles (see Appendix C for an animated version) and is based on the best model fit in the hierarchical regression analysis (model 15. in Table 4 with coefficients $a_1 = 35.77052$; $a_2 = 9.00972$; $a_3 = 0.94421$; $a_4 = 0.06278$; $a_5 = 0.10997$; $a_6 = 0.07065$).

3.3 *Phosphate*

Variation in initial mobile phosphate concentration was large between sample sites (0.25 µg PO_4-P per kg dry matter on average ± 0.21 µg PO_4-P standard deviation across site averages). Phosphate concentrations after the FTC treatment followed a sigmoid increase with colder winter minimum temperature at the sample's origin, modulated by an interaction with FTC magnitude, and the three-way interaction between mean minimum temperature at origin, FTC magnitude, and FTC frequency (Table 5, model 15.). This model achieved a correlation between measured and predicted phosphate concentrations of 0.49. According to this model, highest phosphate concentrations occurred for the combination of the coldest site, the strongest FTC magnitude, and the highest FTC frequency (Figure 3) with predicted values of up to 2.2 µg PO_4-P per kg dry matter, i.e. almost a four-fold increase as compared to the initial phosphate concentration before the start of the experiment at this site (site KA, Table 2). The highest measured value for the coldest site was 4.60 µg PO_4-P per kg dry matter while the absolute maximum measured occurred for the strongest FTC magnitude and the highest FTC frequency at site WE (6.70 µg PO_4-P per kg dry matter). The positively modulating effects of FTC frequency increased with decreasing winter minimum temperature at the samples' origins while almost no effect of FTC frequency was found for mild FTC magnitude across all origins. Predicted phosphate concentrations decreased rapidly towards the warmer sites with the inflection point of the sigmoid shape at around -3°C for high FTC magnitudes and -5°C for mild FTC magnitudes. Lowest phosphate concentrations were predicted for the warmest site with no visible modulation by FTC magnitude and FTC frequency.

Individually, FTC magnitude, but not FTC frequency, was able to significantly explain phosphate concentrations, more complex saturating or sigmoid models being indistinguishable from the linear model for both parameters (Table 5 models 1.-6.). Their interaction appeared relevant and non-additive (Table 5 models 16. and 17.)

Table 5. Results of the hierarchical regression analysis for phosphate concentrations of beech forest soils to changes in FTC magnitude (x_1), FTC frequency (x_2), and climatic origin (x_3; expressed as mean minimum winter temperature at origin) at the end of the FTC treatments. Tested are linear, saturating (Michaelis Menten function) and sigmoid (Gompertz function) relationships on the single environmental drivers and their interactions. Bold AICc (Akaike Information Criterion corrected for small sample sizes) values indicate best model. AICc in italics indicate best single-factor models. a_1 to a_n are the fitted parameters of the respective model. FTC = freeze-thaw cycle.

Model description	model	AICc	Note
1. Linear, magnitude (x$_1$*) only*	$y = a_1 x_1 + a_2$	*998*	*Simplest possible start, lm: p < 0.001*
2. Saturating, magnitude only	$y = \frac{a_1 * x_1}{a_2 + x_1}$	1100	Not better than 1.
3. Sigmoid, magnitude only	$y = a_1 * e^{-a_2 * e^{-a_3 x_1}}$	*991*	Best magnitude-only model
4. Linear, frequency (x$_2$*) only*	$y = a_1 x_2 + a_2$	*1025*	*Simplest possible start, lm: p = 0.369*
5. Saturating, frequency only	$y = \frac{a_1 * x_2}{a_2 + x_2}$	1028	Not better than 4.
6. Sigmoid, frequency only	$y = a_1 * e^{-a_2 * e^{-a_3 x_2}}$	1028	Not better than 4.
7. Linear, climatic origin (x_3) only	$y = a_1 x_3 + a_2$	986	Simplest possible start, lm: p < 0.001
8. Saturating, climatic origin only	$y = \frac{a_1 * x_3}{a_2 + x_3}$	993	Not better than 7..
9. Sigmoid, climatic origin only	$y = a_1 * e^{-a_2 * e^{-a_3 x_3}}$	954	Better than 7., best single factor model
10. Sigmoid climatic origin and sigmoid magnitude (additive)	$y = a_1 * e^{-a_2 * e^{-a_3 x_3}} + a_4 * e^{-a_5 * e^{-a_6 x_1}}$	955	Taking the best model of the best explaining parameter so far (9.) and adding the best model of the second best explaining parameter (3.)
11. Sigmoid climatic origin and its interaction with magnitude	$y = a_1 * e^{-a_2 * e^{-a_3 x_3}} + a_4 x_3 x_1$	940	Adding interaction instead of single effect of magnitude to 9. ANOVA: different from 11. with p < 0.001 New best model
12. Sigmoid climatic origin and its interaction with magnitude and linear frequency	$y = a_1 * e^{-a_2 * e^{-a_3 x_3}} + a_4 x_3 x_1 + a_5 x_2$	940	Adding best model of third parameter (4.) to best model so far (11.) ANOVA: not different from 11. with p = 0.128
13. Sigmoid climatic origin and its two-way interactions with magnitude and with frequency	$y == a_1 * e^{-a_2 * e^{-a_3 x_3}} + a_4 x_3 x_1 + a_5 x_2 x_3$	942	Adding interaction term climatic origin x frequency to best model so far (11.) ANOVA: not different from 11. with p = 0.802

14. Sigmoid climatic origin and its interaction with magnitude and interaction magnitude x frequency	$y == a_1 * e^{-a_2*e^{-a_3x_3}} + a_4x_3x_1 + a_5x_1x_2$	942	Adding interaction magnitude x frequency to best model so far (11.) ANOVA: not different from 11. with p = 0.701
15. Sigmoid climatic origin and interaction climate origin x magnitude and threefold interaction	$\boldsymbol{y == a_1 * e^{-a_2*e^{-a_3x_3}} + a_4x_3x_1 + a_5x_1x_2x_3}$	**937**	**Adding three-fold interaction term to best model so far (11.)** **ANOVA: not different from 11. with p = 0.044 best model**
16. Sigmoid magnitude and linear frequency without interaction (additive)	$y = a_1 * e^{-a_2*e^{-a_3x_1}} + a_4x_2$	992	Checking interaction between magnitude and frequency ANOVA: not different from best single factor model (3.): p = 0.837
17. Sigmoid magnitude and linear frequency with interaction	$y = a_1 * e^{-a_2*e^{-a_3x_1}} + a_4x_2 + a_5x_1x_2$	990	Checking interaction between magnitude and frequency ANOVA different from additive model (16. p = 0.036) but not different from best single factor model (3.): p = 0.109

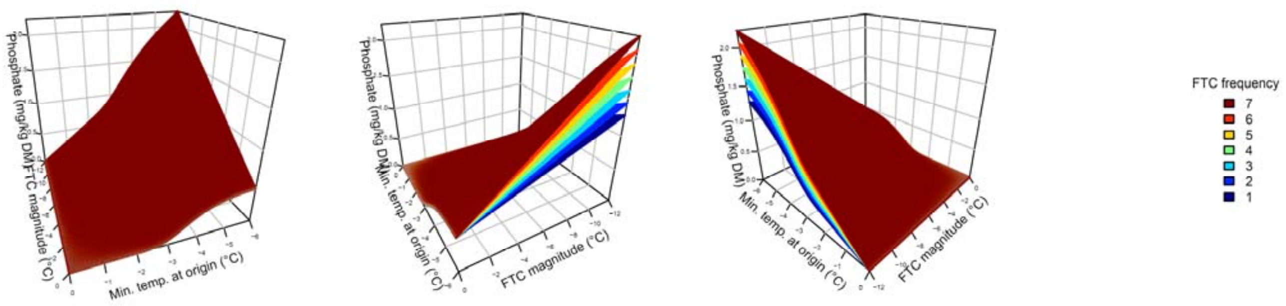

Figure 3: Phosphate concentrations depended on the three-fold interactive effects of winter climatic origin (expressed as longterm mean minimum winter temperature at the origin), FTC magnitude (expressed as the minimum temperature experienced during the FTC manipulation and displayed for freezing temperatures) and FTC frequency during the FTC manipulation. FTC = freeze-thaw cycle. The four dimensional representation is displayed from three different angles (see Appendix D for an animated version) and is based on the best model fit in the hierarchical regression analysis (model 15. in Table 5 with coefficients $a_1 = 0.49455$; $a_2 = 0.01253$; $a_3 = 1.37580$; $a_4 = 0.00890$; $a_5 = 0.00217$).

4 Discussion

4.1 *FTC induce nitrogen release but response patterns are indistinguishable from linear for increased magnitude and increased frequency*

FTC induced nutrient release at high FTC magnitude and frequency in our experiment. Increased nitrate leaching following soil freezing has been explained by decreased root uptake due to lethal or sublethal root damage (Campbell et al., 2014; Matzner and Borken, 2008) and FTC are further reported to increase ammonium production and mineralization rates (Austnes and Vestgarden, 2008; Vestgarden and Austnes, 2009; Shibata et al., 2013; Hosokawa et al., 2017). However, soil frost commonly reduces nitrification rates and nitrate production (Hosokawa et al., 2017; Hishi et al., 2014; Shibata et al., 2013) as nitrifying bacteria are sensitive to low temperatures (Cookson et al., 2002; Dalias et al., 2002). Lysis of microbial cells is reported to occur at minimum temperatures of -7°C (Skogland et al., 1988) to -11°C (Soulides and Allison, 1961) and should consequently have resulted in some form of threshold or non-linear pattern along our gradient of FTC magnitude. As no such threshold was distinguishable, our results are hardly explainable with frost-driven lysis. Based on these aspects, we assume that the processes driving the increase in N and P concentrations in our experiment are either osmotic shock upon exposure to melt water (Jefferies et al., 2010), and/ or physical destruction of organic and soil particles (Oztas and Fayetorbay, 2003; Hobbie and Chapin, 1996) rather than altered mineralization rates as those should be coupled to highest mineral N availability in the unfrozen control. However, FTC increase DON and DOC in temperate deciduous forest soils, quickly leading to enhanced growth of soil microbes and net mineralization, resulting in increased availability of ammonium (Watanabe et al., 2019). Further studies focusing on discrimination between the single processes are clearly needed in light of the strong increases in nitrate (2.5-fold increase) and phosphate (4-fold increase) concentrations over just one week of FTC treatment for the coldest site and highest FTC magnitudes and frequencies.

Here, we expected to find saturation of nutrient release both with increased FTC magnitude and frequency. However, the observed response patterns of nutrient release along these two drivers were indistinguishable from linear in our experiment. This finding has to be treated with care, though, as both drivers were involved in complex interactions with each other and site of soil origin (see below).

4.2 *The combination of magnitude and frequency of FTC on nutrient release is not additive*

We assumed FTC magnitude and frequency effects on nutrient release to be additive, but this was not supported by our data. For ammonium, we observed a significant interaction between FTC magnitude and frequency resulting in over-proportionally large release for high magnitude and frequency. However, for all three analyzed nutrients, both these drivers were further involved into significant three-way interactions with site of soil origin and should be interpreted in this sense (see below).

4.3 *Soils from colder and snowier forests are more responsive to strong and frequent FTC*

Nitrogen and phosphorus release in response to FTC was high for soils from colder and snowier sites. Warmer sites with historically low snow cover showed almost no response to FTC for ammonium and phosphate, while nitrate tended to also be released by strong frost irrespective of FTC frequency in soils from the warmest site. Overall, the strong sigmoidal increase of nutrient concentrations with soils from colder sites was modulated by FTC magnitude and frequency in all studied nutrients. Nitrate concentrations increased with FTC magnitude over the whole range of soil origins peaking for highest frequencies and the coldest sites. The effect of FTC magnitude on ammonium and phosphate concentrations over the climatic gradient was less obvious, but high FTC frequencies mattered only for the coldest sites and high FTC magnitude, then leading to maximum release. All these response shapes show that soils from warmer sites are surprisingly irresponsive to FTC while soils from colder sites are highly sensitive. All studied soils developed under comparable bedrock (sandy Pleistocene deposits) and under the same vegetation types (mono-dominant, mature beech forest with little to no understory). Still, their sensitivity to FTC differed dramatically. Over historic times, the most obvious difference with relevance for FTC sensitivity are winter soil temperature fluctuations, which are generally small at cold sites characterized by stable, insulating snow cover and which are large at the warmer sites with their soils over winter being exposed to air temperature fluctuations (Henry, 2008). Over three winters (2016-2019), our sites reflect this expectation well with strongest frost occurrence and FTC at the center of our gradient and few soil frost incidents at the warm (western) and cold (eastern) extremes (Table 1). In light of the air temperatures and the amount of precipitation as snow, the soils at the coldest sites obviously benefitted from insulation by snow (Table 1). While the soil C/N-ratio appeared irresponsive to the climatic gradient in our study, soil organic matter content increased towards the coldest sites (Table 1). High organic matter content generally increases the susceptibility of soils for nutrient loss with climate change (Liu et al., 2017). Here, we cannot answer how strongly this pattern in organic matter is driven by historic winter soil temperature and occurrence of FTC, but the expectation of increased mineralization with winter soil warming (Gao et al., 2018) would fit to the observed decrease of soil organic matter content with warmer winter climate (Liu et al., 2017). Moreover, the larger pool of organically bound nutrients at the coldest sites may contribute to their observed responsiveness to FTC warming (Gao et al., 2018).

Higher magnitude of FTC changes microbial community composition and functioning, leading to increased tolerance to FTC in temperate forest soils (Urakawa et al., 2014). In light of these results, we suggest that our warmer sites already experienced high winter soil temperature fluctuations with past warming and their microbial community and soil organic matter content adapted to these conditions, making them comparably irresponsive to our FTC treatments. Contrary, our coldest sites rarely experienced serious FTC in the past, exposing a non-adapted microbial community and large pools of organic matter to FTC stress and leading to high rates of mortality and release in consequence. These spatial differences in adaptation or legacy of past conditions might also help explaining why microbial responses to mild FTC appear highly divergent with either little to no effects on microbial biomass and nutrient dynamics (Lipson & Monson 1998; Grogan et al. 2004) or temperature fluctuations in FTC down to only -4°C affecting microbial biomass and nutrient leaching (Larsen et al. 2002; Joseph & Henry

2008). In consequence, largest effects of winter climate change on microbial communities and nutrient dynamics are to be expected for sites where snow cover is currently disappearing (Kreyling, 2020).

The fate of the nutrients released in response to FTC in those regions where snow cover is disappearing is of crucial importance for ecosystem functioning, e.g. tree growth and nitrogen leaching. An increase in available nutrients could increase plant growth. But if the fluctuations in soil temperature lead to lethal or sublethal damage of plant roots (Tierney et al., 2001; Reinmann and Templer, 2018; Kreyling et al., 2012a; Weih and Karlsson, 2002) in parallel to lysis of microbes, the excess nutrients might be leached out of the ecosystem due to reduced root uptake (Matzner and Borken, 2008; Campbell et al., 2014). The projected increase in winter rain for temperate ecosystems (Stocker, 2014) could then further exacerbate nutrient leaching with the downward flow of the additional water (Bowles et al., 2018).

Phosphate is much less mobile in the soil than nitrate and, consequently, leaching of phosphate is not to be expected. Stoichiometric imbalance between N and P nutrition is a global phenomenon, mainly because of the atmospheric deposition of active N and no comparable analogue for P in unfertilized ecosystems (Peñuelas et al., 2013). In light of the surprisingly high mobilization of P in our study and the potential leaching losses of nitrate, an aggravation of the imbalance between N and P of temperate deciduous forests in response to altered winter soil temperature regimes is therefore not to be expected.

The applied gradient design analyzed by hierarchical regression analysis (Kreyling et al., 2018) proofed instrumental for the detection and characterization of non-linear response shapes modulated by complex interactions of the environmental drivers. A traditional, replicated design at few treatment levels along the environmental drivers would not have provided these insights about the complexity of the relationships of the studied drivers.

4.4 *Conclusions*

FTC magnitude and, to a lesser extent, also FTC frequency resulted in increased nitrate, ammonium, and phosphate release almost exclusively in soils from cold, snow-rich sites with high organic matter content while soils from warmer sites characterized by a history of infrequent snow cover and largely fluctuating soil temperatures were comparably irresponsive to FTC. We propose that currently warmer forest soils have historically already passed the point of high responsiveness to winter climate change and might have lost organic matter, displaying some form of adaptation either in the soil biotic composition or in labile nutrient sources. This suggests that previously cold sites losing their protective snow cover with climate change are most vulnerable to strong shifts in nitrogen and phosphorus release. In nutrient poor European beech forests of the studied Pleistocene lowlands, nutrients released over winter may get lost when microbes and plant roots are damaged by soil frost and induce reduced plant growth and increased nutrient leaching rates.

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Code and data availability

The data and the R code to reproduce the analyses are available and can be processed at Dryad: https://doi.org/10.5061/dryad.rxwdbrv5n.

Author contributions

JK and RW designed the study, RW conducted the field work and the experiment, RS performed the chemical analyses. JK analyzed the data and wrote the manuscript with contributions from all co-authors.

Competing interests

The authors declare that they have no conflict of interest.

References

Ammer, C., Fichtner, A., Fischer, A., Gossner, M. M., Meyer, P., Seidl, R., Thomas, F. M., Annighöfer, P., Kreyling, J., Ohse, B., Berger, U., Feldmann, E., Häberle, K.-H., Heer, K., Heinrichs, S., Huth, F., Krämer-Klement, K., Mölder, A., Müller, J., Mund, M., Opgenoorth, L., Schall, P., Scherer-Lorenzen, M., Seidel, D., Vogt, J., and Wagner, S.: Key ecological research questions for Central European forests, Basic and Applied Ecology, 32, 3–25, doi:10.1016/j.baae.2018.07.006, 2018.

Armstrong, F.A.J., Stearns, C.R., and Strickland, J.D.H.: The measurement of upwelling and subsequent biological process by means of the Technicon Autoanalyzer® and associated equipment, Deep Sea Research and Oceanographic Abstracts, 14, 381–389, doi:10.1016/0011-7471(67)90082-4, 1967.

Austnes, K. and Vestgarden, L. S.: Prolonged frost increases release of C and N from a montane heathland soil in southern Norway, Soil Biol Biochem, 40, 2540–2546, 2008.

Bolter, M., Soethe, N., Horn, R., and Uhlig, C.: Seasonal development of microbial activity in soils of northern Norway, Pedosphere, 15, 716–727, 2005.

Bontemps, J.-D., Hervé, J.-C., Leban, J.-M., and Dhôte, J.-F.: Nitrogen footprint in a long-term observation of forest growth over the twentieth century, Trees, 25, 237–251, doi:10.1007/s00468-010-0501-2, 2011.

Bowles, T. M., Atallah, S. S., Campbell, E. E., Gaudin, A. C. M., Wieder, W. R., and Grandy, A. S.: Addressing agricultural nitrogen losses in a changing climate, Nature Sustainability, 1, 399–408, doi:10.1038/s41893-018-0106-0, 2018.

Brown, P. J. and DeGaetano, A. T.: A paradox of cooling winter soil surface temperatures in a warming northeastern United States, Agricultural and Forest Meteorology, 151, 947–956, doi:10.1016/j.agrformet.2011.02.014, 2011.

Brown, R. D. and Mote, P. W.: The response of northern hemisphere snow cover to a changing climate, J Climate, 22, 2124–2145, doi:10.1175/2008JCLI2665.1, 2009.

Buma, B., Hennon, P. E., Harrington, C. A., Popkin, J. R., Krapek, J., Lamb, M. S., Oakes, L. E., Saunders, S., and Zeglen, S.: Emerging climate-driven disturbance processes: Widespread mortality associated with snow-to-rain transitions across 10° of latitude and half the range of a climate-threatened conifer, Glob Change Biol, 23, 2903–2914, doi:10.1111/gcb.13555, 2017.

Campbell, J. L., Mitchell, M. J., Groffman, P. M., Christenson, L. M., and Hardy, J. P.: Winter in northeastern North America: a critical period for ecological processes, Front Ecol Environ, 3, 314–322, 2005.

Campbell, J. L., Ollinger, S. V., Flerchinger, G. N., Wicklein, H., Hayhoe, K., and Bailey, A. S.: Past and projected future changes in snowpack and soil frost at the Hubbard Brook Experimental Forest, New Hampshire, USA, Hydrol. Process., 19, n/a-n/a, doi:10.1002/hyp.7666, 2010.

Campbell, J. L., Socci, A. M., and Templer, P. H.: Increased nitrogen leaching following soil freezing is due to decreased root uptake in a northern hardwood forest, Glob Change Biol, 20, 2663–2673, 2014.

Comerford, D. P., Schaberg, P. G., Templer, P. H., Socci, A. M., Campbell, J. L., and Wallin, K. F.: Influence of experimental snow removal on root and canopy physiology of sugar maple trees in a northern hardwood forest, Oecologia, 171, 261–269, doi:10.1007/s00442-012-2393-x, 2013.

Cookson, W.R., Cornforth, I.S., and Rowarth, J.S.: Winter soil temperature (2–15°C) effects on nitrogen transformations in clover green manure amended or unamended soils; a laboratory and field study, Soil Biology and Biochemistry, 34, 1401–1415, doi:10.1016/S0038-0717(02)00083-4, 2002.

Dalias, P., Anderson, J. M., Bottner, P., and Coûteaux, M.-M.: Temperature responses of net nitrogen mineralization and nitrification in conifer forest soils incubated under standard laboratory conditions, Soil Biology and Biochemistry, 34, 691–701, doi:10.1016/S0038-0717(01)00234-6, 2002.

Elliott, A. C. and Henry, H. A. L.: Freeze-thaw cycle amplitude and freezing rate effects on extractable nitrogen in a temperate old field soil, Biol Fert Soils, 45, 469–476, 2009.

Gao, D., Zhang, L., Liu, J., Peng, B., Fan, Z., Dai, W., Jiang, P., and Bai, E.: Responses of terrestrial nitrogen pools and dynamics to different patterns of freeze-thaw cycle: A meta-analysis, Glob Change Biol, 24, 2377–2389, doi:10.1111/gcb.14010, 2018.

Göbel, L., Coners, H., Hertel, D., Willinghöfer, S., and Leuschner, C.: The role of low soil temperature for photosynthesis and stomatal conductance of three graminoids from different elevations, Front Plant Sci, 10, 330, doi:10.3389/fpls.2019.00330, 2019.

Groffman, P. M., Driscoll, C. T., Fahey, T. J., Hardy, J. P., Fitzhugh, R. D., and Tierney, G. L.: Colder soils in a warmer world: A snow manipulation study in a northern hardwood forest ecosystem, Biogeochemistry, 56, 135–150, 2001.

Guo, C., Simon, J., Gasche, R., Naumann, P. S., Bimüller, C., Pena, R., Polle, A., Kögel-Knabner, I., Zeller, B., Rennenberg, H., and Dannenmann, M.: Minor contribution of leaf litter to N nutrition of beech (Fagus sylvatica) seedlings in a mountainous beech forest of Southern Germany, Plant Soil, 369, 657–668, doi:10.1007/s11104-013-1603-6, 2013.

Hacket-Pain, A. J., Ascoli, D., Vacchiano, G., Biondi, F., Cavin, L., Conedera, M., Drobyshev, I., Liñán, I. D., Friend, A. D., Grabner, M., Hartl, C., Kreyling, J., Lebourgeois, F., Levanič, T., Menzel, A., van der Maaten, E., van der Maaten-Theunissen, M., Muffler, L., Motta, R., Roibu, C.-C., Popa, I., Scharnweber, T., Weigel, R., Wilmking, M., and Zang, C. S.: Climatically controlled reproduction drives interannual growth variability in a temperate tree species, Ecol Lett, 21, 1833–1844, doi:10.1111/ele.13158, 2018.

Hamann, A., Wang, T., Spittlehouse, D. L., and Murdock, T. Q.: A comprehensive, high-resolution database of historical and projected climate surfaces for Western North America, B Am Meteorol Soc, 94, 1307–1309, doi:10.1175/BAMS-D-12-00145.1, 2013.

Hansen, H. P. and Koroleff, F.: Determination of nutrients, in: Methods of seawater analysis, 3rd, completely rev. and extended ed., Grasshoff, K., Anderson, L. G. (Eds.), Wiley VCH, Weinheim, 159–228, 1999.

Henry, H. A. L.: Climate change and soil freezing dynamics: historical trends and projected changes, Clim Change, 87, 421–434, 2008.

Hishi, T., Urakawa, R., Tashiro, N., Maeda, Y., and Shibata, H.: Seasonality of factors controlling N mineralization rates among slope positions and aspects in cool-temperate deciduous natural forests and larch plantations, Biology and Fertility of Soils, 50, 343–356, doi:10.1007/s00374-013-0863-x, 2014.

Hobbie, S. E. and Chapin, F. S.: Winter regulation of tundra litter carbon and nitrogen dynamics, Biogeochemistry, 35, 327–338, doi:10.1007/BF02179958, 1996.

Hosokawa, N., Isobe, K., Urakawa, R., Tateno, R., Fukuzawa, K., Watanabe, T., and Shibata, H.: Soil freeze–thaw with root litter alters N transformations during the dormant season in soils under two temperate forests in northern Japan, Soil Biol Biochem, 114, 270–278, doi:10.1016/j.soilbio.2017.07.025, 2017.

Hurvich, C. M. and Tsai, C.-L.: Regression and time series model selection in small samples, Biometrika, 76, 297–307, doi:10.1093/biomet/76.2.297, 1989.

IPCC (Ed.): Climate Change 2013: The Physical Science Basis. Contribution of Working Group I to the Fifth Assessment Report of the Intergovernmental Panel on Climate Change, Cambridge University Press, Cambridge, United Kingdom and New York, NY, USA, 2013.

Iwata, Y., Hayashi, M., Suzuki, S., Hirota, T., and Hasegawa, S.: Effects of snow cover on soil freezing, water movement, and snowmelt infiltration: A paired plot experiment, Water Resour Res, 46, 2010.

Jefferies, R. L., Walker, N. A., Edwards, K. A., and Dainty, J.: Is the decline of soil microbial biomass in late winter coupled to changes in the physical state of cold soils?, Soil Biol Biochem, 42, 129–135, 2010.

Joseph, G. and Henry, H. A. L.: Soil nitrogen leaching losses in response to freeze-thaw cycles and pulsed warming in a temperate old field, Soil Biol Biochem, 40, 1947–1953, 2008.

Joseph, G. and Henry, H. A. L.: Retention of surface nitrate additions in a temperate old field: implications for atmospheric nitrogen deposition over winter and plant nitrogen availability, Plant Soil, 319, 209–218, 2009.

Kérouel, R. and Aminot, A.: Fluorometric determination of ammonia in sea and estuarine waters by direct segmented flow analysis, Marine Chemistry, 57, 265–275, doi:10.1016/S0304-4203(97)00040-6, 1997.

Kreyling, J.: Winter climate change: A critical factor for temperate vegetation performance, Ecology, 91, 1939–1948, doi:10.1890/09-1160.1, 2010.

Kreyling, J.: The ecological importance of winter in temperate, boreal, and arctic ecosystems in times of climate change, in: Progress in Botany Vol. 81, Cánovas, F. M., Lüttge, U., Leuschner, C., Risueño, M.-C. (Eds.), Progress in Botany, Springer International Publishing, Cham, 377–399, 2020.

Kreyling, J., Haei, M., and Laudon, H.: Absence of snow cover reduces understory plant cover and alters plant community composition in boreal forests, Oecologia, 168, 577–587, doi:10.1007/s00442-011-2092-z, 2012a.

Kreyling, J. and Henry, H. A. L.: Vanishing winters in Germany: soil frost dynamics and snow cover trends, and ecological implications, Clim Res, 46, 269–276, doi:10.3354/cr00996, 2011.

Kreyling, J., Persoh, D., Werner, S., Benzenberg, M., and Wöllecke, J.: Short-term impacts of soil freeze-thaw cycles on roots and root-associated fungi of Holcus lanatus and Calluna vulgaris, Plant Soil, 353, 19–31, 2012b.

Kreyling, J., Schweiger, A. H., Bahn, M., Ineson, P., Migliavacca, M., Morel-Journel, T., Christiansen, J. R., Schtickzelle, N., and Larsen, K. S.: To replicate, or not to replicate - that is the question: How to tackle nonlinear responses in ecological experiments, Ecol Lett, 21, 1629–1638, doi:10.1111/ele.13134, 2018.

Leuschner, C., Meier, I. C., and Hertel, D.: On the niche breadth of Fagus sylvatica: soil nutrient status in 50 Central European beech stands on a broad range of bedrock types, Ann Forest Sci, 63, 355–368, doi:10.1051/forest:2006016, 2006.

Liu, Y., Wang, C., He, N., Wen, X., Gao, Y., Li, S., Niu, S., Butterbach-Bahl, K., Luo, Y., and Yu, G.: A global synthesis of the rate and temperature sensitivity of soil nitrogen mineralization: latitudinal patterns and mechanisms, Glob Change Biol, 23, 455–464, doi:10.1111/gcb.13372, 2017.

Malcolme-Lawes, D. J. and Wong, K. H.: Determination of orthophosphate in waters and soils using a flow analyser, Analyst, 115, 65–67, doi:10.1039/AN9901500065, 1990.

Matzner, E. and Borken, W.: Do freeze-thaw events enhance C and N losses from soils of different ecosystems?: A review, Eur J Soil Sci, 59, 274–284, 2008.

McMahon, S. K., Wallenstein, M. D., and Schimel, J. P.: Microbial growth in Arctic tundra soil at-2 degrees C, Environ Microbiol Rep, 1, 162–166, 2009.

Mikan, C. J., Schimel, J. P., and Doyle, A. P.: Temperature controls of microbial respiration in arctic tundra soils above and below freezing, Soil Biol Biochem, 34, 1785–1795, 2002.

Murphy, J. and Riley, J.P.: A modified single solution method for the determination of phosphate in natural waters, Analytica Chimica Acta, 27, 31–36, doi:10.1016/S0003-2670(00)88444-5, 1962.

Oztas, T. and Fayetorbay, F.: Effect of freezing and thawing processes on soil aggregate stability, Catena, 52, 1–8, 2003.

Peñuelas, J., Poulter, B., Sardans, J., Ciais, P., van der Velde, M., Bopp, L., Boucher, O., Godderis, Y., Hinsinger, P., Llusia, J., Nardin, E., Vicca, S., Obersteiner, M., and Janssens, I. A.: Human-induced nitrogen-phosphorus imbalances alter natural and managed ecosystems across the globe, Nature Communications, 4, 2934, doi:10.1038/ncomms3934, 2013.

Petoukhov, V. and Semenov, V. A.: A link between reduced Barents-Kara sea ice and cold winter extremes over northern continents, J Geophys Res-Atmos, 115, 10, doi:10.1029/2009JD013568, 2010.

R Core Team: R: A Language and Environment for Statistical Computing.: R version 3.4.3, R Foundation for Statistical Computing. URL http://www.R-project.org, Vienna, Austria, 2017.

Reinmann, A. B., Susser, J. R., Demaria, E. M. C., and Templer, P. H.: Declines in northern forest tree growth following snowpack decline and soil freezing, Glob Change Biol, 25, 420–430, doi:10.1111/gcb.14420, 2019.

Reinmann, A. B. and Templer, P. H.: Increased soil respiration in response to experimentally reduced snow cover and increased soil freezing in a temperate deciduous forest, Biogeochemistry, 140, 359–371, doi:10.1007/s10533-018-0497-z, 2018.

Rennenberg, H. and Dannenmann, M.: Nitrogen nutrition of trees in temperate forests - the significance of nitrogen availability in the pedosphere and atmosphere, Forests, 6, 2820–2835, doi:10.3390/f6082820, 2015.

Schaberg, P. G., Hennon, P. E., D'Amore, D. V., and Hawley, G. J.: Influence of simulated snow cover on the cold tolerance and freezing injury of yellow-cedar seedlings, Glob Change Biol, 14, 1282–1293, 2008.

Scherrer, S. C. and Appenzeller, C.: Swiss Alpine snow pack variability: Major patterns and links to local climate and large-scale flow, Clim Res, 32, 187–199, doi:10.3354/cr032187, 2006.

Schimel, J., Balser, T. C., and Wallenstein, M.: Microbial stress-response physiology and its implications for ecosystem function, Ecology, 88, 1386–1394, doi:10.1890/06-0219, 2007.

Shibata, H., Hasegawa, Y., Watanabe, T., and Fukuzawa, K.: Impact of snowpack decrease on net nitrogen mineralization and nitrification in forest soil of northern Japan, Biogeochemistry, 116, 69–82, doi:10.1007/s10533-013-9882-9, 2013.

Simon, J., Dannenmann, M., Pena, R., Gessler, A., and Rennenberg, H.: Nitrogen nutrition of beech forests in a changing climate: importance of plant-soil-microbe water, carbon, and nitrogen interactions, Plant Soil, 418, 89–114, doi:10.1007/s11104-017-3293-y, 2017.

Skogland, T., Lomeland, S., and Goksoyr, J.: Respiratory burst after freezing and thawing of soil - experiments with soil bacteria, Soil Biol Biochem, 20, 851–856, 1988.

Soulides, D. A. and Allison, F. E.: Effect of drying and freezing soils on carbon dioxide production, available mineral nutrients, aggregation, and bacterial population, Soil Science, 91, 291–298, 1961.

Stocker, T. (Ed.): Climate change 2013: The physical science basis Working Group I contribution to the Fifth assessment report of the Intergovernmental Panel on Climate Change, Cambridge University Press, Cambridge, 1535 pp., 2014.

Talkner, U., Meiwes, K. J., Potočić, N., Seletković, I., Cools, N., Vos, B. de, and Rautio, P.: Phosphorus nutrition of beech (Fagus sylvatica L.) is decreasing in Europe, Ann Forest Sci, 72, 919–928, doi:10.1007/s13595-015-0459-8, 2015.

Tierney, G. L., Fahey, T. J., Groffman, P. M., Hardy, J. P., Fitzhugh, R. D., and Driscoll, C. T.: Soil freezing alters fine root dynamics in a northern hardwood forest, Biogeochemistry, 56, 175–190, 2001.

Urakawa, R., Shibata, H., Kuroiwa, M., Inagaki, Y., Tateno, R., Hishi, T., Fukuzawa, K., Hirai, K., Toda, H., Oyanagi, N., Nakata, M., Nakanishi, A., Fukushima, K., Enoki, T., and Suwa, Y.: Effects of freeze–thaw cycles resulting from winter climate change on soil nitrogen cycling in ten temperate forest ecosystems throughout the Japanese archipelago, Soil Biol Biochem, 74, 82–94, doi:10.1016/j.soilbio.2014.02.022, 2014.

Vestgarden, L. S. and Austnes, K.: Effects of freeze-thaw on C and N release from soils below different vegetation in a montane system: a laboratory experiment, Glob Change Biol, 15, 876–887, 2009.

Wang, T., Hamann, A., Spittlehouse, D. L., and Murdock, T. Q.: ClimateWNA—High-resolution spatial climate data for Western North America, J. Appl. Meteor. Climatol., 51, 16–29, doi:10.1175/JAMC-D-11-043.1, 2012.

Wardle, D. A., Walker, L. R., and Bardgett, R. D.: Ecosystem properties and forest decline in contrasting long-term chronosequences, Science (New York, N.Y.), 305, 509–513, doi:10.1126/science.1098778, 2004.

Watanabe, T., Tateno, R., Imada, S., Fukuzawa, K., Isobe, K., Urakawa, R., Oda, T., Hosokawa, N., Sasai, T., Inagaki, Y., Hishi, T., Toda, H., and Shibata, H.: The effect of a freeze-thaw cycle on dissolved nitrogen dynamics and its relation to dissolved organic matter and soil microbial biomass in the soil of a northern hardwood forest, Biogeochemistry, 142, 319–338, doi:10.1007/s10533-019-00537-w, 2019.

Weigel, R., Muffler, L., Klisz, M., Kreyling, J., van der Maaten-Theunissen, M., Wilmking, M., and van der Maaten, E.: Winter matters: Sensitivity to winter climate and cold events increases towards the cold distribution margin of European beech (Fagus sylvatica L.), J Biogeogr, 45, 2779–2790, doi:10.1111/jbi.13444, 2018.

Weih, M. and Karlsson, P. S.: Low winter soil temperature affects summertime nutrient uptake capacity and growth rate of mountain birch seedlings in the subarctic, Swedish lapland, Arct Antarct Alp Res, 34, 434–439, 2002.

Yanai, Y., Toyota, K., and Okazaki, M.: Effects of successive soil freeze-thaw cycles on soil microbial biomass and organic matter decomposition potential of soils, Soil Sci Plant Nutr, 50, 821–829, 2004.

Yang, S. and Christensen, J. H.: Arctic sea ice reduction and European cold winters in CMIP5 climate change experiments, Geophys Res Lett, 39, 890, doi:10.1029/2012GL053338, 2012.