Structure of the author's response

Please note that the line numbers indicated in the replies correspond to the line numbers in the track_changed version of the manuscript

I would like to thank you the authors for the meticulous review and response to the comments on the first manuscript version. In this new version authors have addressed most of the issues raised during the interactive revision process including additional information that helps to understand methods and results, as is the case for the temporal analysis of the canopy N field measurements and the improved statistical processes. However I still see some weak points in the paper:

Thank you for your kind words

1. In my opinion authors have not jet provided a proper justification on the usefulness of the statistical analysis using resampled MTCI images to lower spatial resolution. In fact, there is a kind of contradiction in the manuscript between this analysis and the information provided by the authors in the introduction and discussion about the future potential of canopy N estimation form RS using new generation of sensor with improved spatial resolution. I would find the analysis useful if the authors wanted to demonstrate that sensors with lower spatial resolution can be potentially used to obtain global estimations of canopy N, but, as this is not the case, I would find more convenient to undertake an analysis that allows to demonstrate the sensitivity of the statistical relationships found to the field data (sample size and distribution). This is an important issue raised by the authors in the discussion (section4-5). Ground canopy N observations are necessary to calibrate and validate models at regional-global scales. In this context, an interesting (and I would say feasible) output of this work could be a sensitivity analysis on the model performance according to field data availability.

All analyses in the manuscript use MTCI data at the original 1 km resolution. In Sections 2.3.3 and 3.2, however, we included an additional analysis using resampled MTCI data. We would like to explain here why we did this additional analysis where both data sets (MTCI and forest plots) were resampled to the same lower spatial resolutions. The objective of our study is not to prove the usefulness of the resampled MTCI images, but rather to investigate the relationship between the MTCI product (1 km) and canopy N from forest plots at regional scale. We realize that our analysis is based on data at different spatial scale. We therefore resampled (in the additional analysis mentioned above) the datasets to the same support size to be able to study the relationship between MTCI and canopy N independently of the initial difference in support size. When the relationship between our datasets is still present. Given the similarity in the relationship between canopy N values and MTCI values at the original and lower spatial resolutions, the confidence in the relationship is increased.

This was not clearly explained in the manuscript. In the revised manuscript, a paragraph was added in the Material and Methods part, section 2.3.2 (Line 242 – 250):

This was done because of the initial difference in support size between MTCI spatial resolution and the forest plots size (i.e. 1 km and 6 m, respectively). This enabled us to investigate the relationships between MTCI and canopy N data when the spatial discrepancy was accounted for independently of differences in initial support size. The statistical basis of this approach is that we bring both datasets (forest plots and MTCI values) to the same support size or representative area (Bierkens, 2000). By

averaging out forest plot values within this support size, we calculate the mean of the canopy N value at that support size. By resampling the MTCI values to that same support size, the obtained result consist of a mean of the MTCI value at that support size. We then regressed the expected canopy N values (at the new support size) against the expected MTCI values (at the new support size).

We do not see a contradiction between resampling both our datasets to the same support size and our statement that the advent of sensors with high spatial resolution is expected to improve the observed relationship. Given that the sample size of the forest plots (6 m) will unlikely be larger in the future, using higher spatial resolution data is the only way to decrease the initial scale discrepancy between the forests plots size and the original pixel size.

We think that a sensitivity analysis of the global vegetation model performance according to field data density and distribution is important to assess the validity of the model output, however, it is outside of the scope of our study and study objectives, mainly because we do not have the enormous amount of plot data that would be required to do a proper sensitivity analysis. To still take your comment into account, we have now mentioned this aspect regarding model validation in the revised version of the manuscript in the future perspective section 4.5 (Line 459 – 461):

Obtaining reliable ground based canopy N data over larger areas and for diverse and globally distributed vegetation types would also be necessary to calibrate and validate global vegetation models, as the model performance will depend on the ground data availability and distribution.

2. I still miss in the discussion a more "quantitative" consideration on the potential of the results obtained to feed global vegetation models. Authors argue in their response that their study contributes to the ongoing discussion on canopy N estimations on larger areas using RS but this is, in my opinion, a quite diffuse argument. I would expect a more detailed discussion on how much the estimations should be improved to provide useful input to those models (what is the uncertainty in canopy N that can be considered acceptable for the models? And specifically for Mediterranean environments?)

We have included a consideration about how foliar nitrogen prediction in the model LPJ-Guess could benefit from canopy N estimates from remote sensing in section 4.5 (Line 461 – 465): *Remotely sensed canopy N estimates would support calibration of such models. In a recent study the global vegetation model LPJ-Guess was able to simulate the differences in foliar nitrogen between different PFTs but not within one PFT (Fleischer et al., 2015). In this context, improving remotely sensed canopy N estimates for homogeneous vegetation types would be a beneficial development for such models.*

3. In the discussion authors compare their results (in terms of r2) with other works were similar relationships have been found between canopy N and vegetation indices but they do not mention that other studies do not include the temporal dimension. Temporal variability of vegetation due to phenology should not be ignored when estimates are based in secondary relationships as is the case with N vs vegetation indices and, therefore, studies that including or not this temporal dimension are not fully comparable.

We agree that phenology is important and should not be ignored. To address the temporal dimension, we 1) analyzed the influence of the temporal discrepancy between the plot sampling campaign and the period of MTCI acquisition data in the inter-annual variation of canopy N (Fig. A1). The graph shows that this inter-annual variation is not strong. 2) Forest plots are linked to a 10 year average of MTCI values measured during the same month, i.e. plots measured in July are linked to a 10 years average of MTCI values measured at the same location in July. In this case, the influence of phenology is thus present in both the plot data and the remote sensing data. We think that the main influence of using data from a forest inventory that was carried out during the whole growing season over several years is that the range of canopy N values included is larger.

Regarding the studies we reference and compare our results to, the canopy at the forest sites are indeed sampled once or during a short period (Cho et al., 2013;Ramoelo et al., 2012;Wang et al., 2016), the studies carried out in crops (Tian et al., 2011;Li et al., 2014) were sampled during the whole growing season over several years.

A mention was added to the manuscript, section 4.2.1 (Line 358- 360):

In these comparisons, it should be taken into account that most previous studies were based on a short sampling campaign while our study incorporates canopy N data from a forest inventory that was carried out during the entire growing season and therefore includes differences in phenology.

I have also some comments addressing technical/formal issues referred to manuscript version 3:

Abstract line 12. Remote sensing and vegetation indices are not excluding terms, I would recommend rephrasing.

This was replaced by: "Remotely sensed vegetation indices"

Abstract line 19. I would say "original" instead of "initial higher" This was replaced at the mentioned occurrence and later in text

Section 2.2.1 Authors mention that "all foliar cohorts in the canopy were included in the leaf sample" but, was the % of new-old leaves in the crown taken into account during the sampling or the data processing? The N content can greatly differ depending on the leaf age so, in certain phonological periods this need to be considered to obtain an accurate estimation of canopy N.

The % of new-old leaves in the crown was not taken into account. During the sampling campaign, all the foliar cohorts were pooled together in the same sample and this percentage was not recorded. As all the foliar cohorts present on the selected sampled branches are included, we expect that the measurements still represent the plot canopy N value with acceptable accuracy.

Section 2.2.2. I think authors should mention here Sentinel-3 OLCI sensor as the most direct inheritor of MERIS ENVISAT.

This was edited in the text, section 2.2.2. (Line 200 – 205):

While the ESA ENVISAT satellite mission producing MERIS data came to an end in 2012, MERIS products and MTCI in particular are still relevant because the new ESA Sentinel-2 and Sentinel-3 satellite missions haves improved band settings compared to those of MERIS. and increased the spatial resolution to 20 m MTCI can be calculated from Sentinel-2 reflectance data with increased spatial resolution to 20 m (Drusch et al., 2012). The Sentinel-3 mission also releases a level 2 chlorophyll product, the OLCI Terrestrial Chlorophyll Index (OTCI), which calculation is directly based on MTCI. OTCI continues the time series already available for MTCI (Dash and Vuolo, 2010;Vuolo et al., 2012).

Section 2.3.1 line 207. It is not clear why you need to resample the landcover map to the MTCI images resolution. If I properly understand you just want to identify and mask the field plots that changed from forest to other non forest covers. If so, you would just mask those field plots located in a landcover map pixel classified as those covers excluded from the analysis.

We decided to resample the Globcover landcover map (with an original resolution of 300 m) to MTCI original spatial resolution of 1 km before using it as a selection criterion for our forest plots to be on the safe side regarding plot selection and leave out plots located on heterogeneous MTCI pixels. For example, if one plot was located in the only 300 m natural vegetation area of the 1 km MTCI pixel, e.g. a small forest patch surrounded by agricultural crops, the MTCI pixel value would also be influenced by the non-forested area surrounding the plot. Resampling the Globcover landcover map was carried out using the majority option. That way, by using the resampled landcover map (1 km) as the selection criterion, we make sure that the plot located in the isolated vegetation patch is excluded. Also, the number of plots selected for both the analysis with and without resampling of MTCI pixels, i.e. the analysis at the original spatial resolution of 1 km and the analysis at the resampled spatial resolution, is equal. The difference in the results cannot be attributed to the difference in the plots selected.

However, when the analysis is conducted using the original landcover map instead of the resampled landcover map, the number of plot selected is almost the same (n = 866 instead of n = 846) and the relationships observed between the variables are almost not affected and we thus propose to leave out these results as they provide negligible additional information that could be relevant for the interpretation of our results.

Line 222. In the title of this section and all through the manuscript I recommend to replace "initial higher" by "original 1Km" spatial resolution. This has been done.

Line 281. Authors mean here statistically significant? Yes, this has been replaced

Line 288. P-value of this relationship?

The p-value was added: *Quercus ilex (r2 = 0.10, p-value < 0.000, n = 160).*

Lines 407-408. Consider rephrasing to avoid repetition (addition...adding..additional)

The sentence has been changed to (Line 456 – 458): In addition to more detailed remote sensing data, supplementary ground based canopy N observations could better constrain the regression models as well.

Figure 1. I would recommend to add a couple of zoom windows showing the MERIS MTCI 1 km grid on areas with high and low density of field sampling points.

This has been added on the map of Catalonia (Figure 1):



Figure 1. Map showing the forest plots (n = 846) location in the region of Catalonia, north eastern Spain. Two zoom windows are included showing the density of the plots, one with high density and one with low density, relatively to the MTCI 1 km pixel grid. DBF = Deciduous Broadleaf Forest, EBF = Evergreen Broadleaf Forest, ENF = Evergreen Needleleaf Forest, mixed = mixed forest.

References

Cho, M. A., Ramoelo, A., Debba, P., Mutanga, O., Mathieu, R., van Deventer, H., and Ndlovu, N.: Assessing the effects of subtropical forest fragmentation on leaf nitrogen distribution using remote sensing data, Landscape Ecology, 28, 1479-1491, doi:10.1007/s10980-013-9908-7, 2013.

Li, F., Miao, Y., Feng, G., Yuan, F., Yue, S., Gao, X., Liu, Y., Liu, B., Ustin, S. L., and Chen, X.: Improving estimation of summer maize nitrogen status with red edge-based spectral vegetation indices, Field Crops Research, 157, 111-123, doi:10.1016/j.fcr.2013.12.018, 2014.

Ramoelo, A., Skidmore, A. K., Cho, M. A., Schlerf, M., Mathieu, R., and Heitkönig, I. M. A.: Regional estimation of savanna grass nitrogen using the red-edge band of the spaceborne rapideye sensor, International Journal of Applied Earth Observation and Geoinformation, 19, 151-162, doi:10.1016/j.jag.2012.05.009, 2012.

Tian, Y. C., Yao, X., Yang, J., Cao, W. X., Hannaway, D. B., and Zhu, Y.: Assessing newly developed and published vegetation indices for estimating rice leaf nitrogen concentration with ground- and space-based hyperspectral reflectance, Field Crops Research, 120, 299-310, doi:10.1016/j.fcr.2010.11.002, 2011.

Wang, Z., Wang, T., Darvishzadeh, R., Skidmore, A. K., Jones, S., Suarez, L., Woodgate, W., Heiden, U., Heurich, M., and Hearne, J.: Vegetation indices for mapping canopy foliar nitrogen in a mixed temperate forest, Remote Sensing, 8, doi:10.3390/rs8060491, 2016.

Associate Editor Decision: Publish subject to minor revisions (review by editor) (01 Mar 2018) by Sönke Zaehle

Comments to the Author:

Dear authors,

my apologies for the delay in coming to a decision. I failed to find a second reviewer, therefore I reviewed the manuscript myself. I believe that the manuscript can be publishable in Biogeosciences, if you decided to further revise the manuscript according to the suggestions and comments by reviewer #1 and my comments below. Best wishes,

Sönke

While I find that the manuscript has improved, I still see need for further improvements Major comments:

My main worry with this manuscript is still that the authors state that they show their relationships to be robust against spatial upscaling, while they leave out the critical scale jump from the plot level to the 1km resolution level. I don't think that this invalidates the results of the study per se, but I do believe that the design of this study is unsuitable to make claims about the scalability of the results, because for this one would need to address the scaling from the proximity of the forest plots to the 1km as well.

All analyses in the manuscript use MTCI data at the original 1 km resolution. In Sections 2.3.3 and 3.2, however, we included an additional analysis using resampled MTCI data. We would like to explain here why did this additional analysis where both data sets (MTCI and plot samples) were resampled to the same lower spatial resolutions. The objective of our analysis was not to make claims about the scalability of the results. Our main objective was to study the relationship between the MTCI time series (1 km) and canopy N data from forest plots (6 m). As our two datasets present a difference in scale, in the additional analysis mentioned above, we resampled the two datasets to the same support size to be able to study the relationship between MTCI and canopy N independently of the initial difference in support size. The results show that the correlation between MTCI and canopy N after resampling is not strongly influenced by the resampling. Given the similarity in the relationship between canopy N values and MTCI values at the original and lower spatial resolutions, the confidence in the relationship is increased.

This was not clearly explained in the manuscript. In the revised manuscript, a paragraph was added in the Material and Methods part to make it more clear, section 2.3.2 (Line 242 – 250):

This was done because of the initial difference in support size between MTCI spatial resolution and the forest plots size (i.e. 1 km and 6 m, respectively). This enabled us to investigate the relationships between MTCI and canopy N data when the spatial discrepancy was accounted for. The statistical basis of this approach is that we bring both datasets (forest plots and MTCI values) to the same support size or representative area (Bierkens et al., 2000). By averaging out forest plot values within this support size, we calculate the mean of the canopy N value at that support size. By resampling the MTCI values to that same support size, the obtained result consist of a mean of the MTCI value at that support size. We then regressed the expected canopy N values (at the new support size) against the expected MTCI values (at the new support size).

We have also downscaled the claims about the scalability of the results. In the result part, section 3.2, we have replaced (Line 298 – 299):

This was done to investigate the relationship between MTCI and canopy N data independently of difference in support size

By:

This was done to investigate the relationship between MTCI and canopy N data when the initial spatial discrepancy between the two datasets was accounted for.

In the discussion part section 4.1, we have removed part of the text where the claims were too strong (Line 342 – 344):

This showed that, when the spatial discrepancy between the original datasets, i.e. 6 m and 1 km, was taken into account, MTCI and canopy N data were linked and that the MTCI-canopy N relationship was not strongly affected by the resampled spatial resolution.

The use of the word detection is inappropriate. The paper demonstrates a sometimes significant log-linear correlation between these two variables, but does not attempt to disentangle the possible signal from canopy nitrogen from confounding factors, it does therefore not allow for a detection of canopy N trends.

The word "detection" was replaced when it appeared in text when describing the results we obtained (but not when describing results obtained by others, e.g. in the Introduction part).

The results of the manuscript are presented in a misleading fashion, because it states that (L221) that MTCI and canopy values are related via linear regression, whereas infact the authors use a log-linear regression. This needs to be made clear at every instance (for ease of writing possible by introducing two new symbols referring to the log-transformed canopy values), since this affects the interpretation of the regression (i.e. the connection is not linear as written in the text) as well of the r2 value. I would also expect a reasoning as to why the authors believe that the use of a log-linear relationship between MTCI and canopy N is to be expected.

The mentions to linear regression have been converted to "log-linear regression" when it appeared in text.

We use a log normal relationship because the canopy N concentration show outliers at higher values, as shown by its distribution (Figure 3 in the manuscript). Several phenomena in nature show outliers towards higher values, compared to lower values, in particular when bounded to values > 0 (Limpert et al., 2001).



Figure 2. The upper right part of this figure shows the Pearson correlation matrix between canopy $N_{[2^{\prime}]}$ (%N), canopy $N_{[area]}$ (g m⁻²) and foliar biomass (g m⁻²) variables for the whole dataset, n = 841. The diagonal presents the histograms of the variables on the x-axis, while the y-axis represents the number of counts. The lower left part of this figure represents the scatterplots between the variables. PFT = Plant Functional Type, DBF = Deciduous Broadleaf Forest, EBF = Evergreen Broadleaf Forest, ENF = Evergreen Needleleaf Forest, mixed = mixed forest.

Moreover, the log transformation of the canopy N variables was carried out to fulfill linear regression assumptions. This is mentioned in the section 2.3.4 "Statistical analysis" (Line 267-269):

Preliminary analysis showed that using a natural logarithm transformation (log) of the canopy N variables was necessary to fulfil linear regression model assumptions, namely normality and homogeneity of variance of the residuals.

This is also mentioned in the section 3.1 "Descriptive statistics" (Line 293-294):

As canopy $N_{[\%]}$ and canopy $N_{[area]}$ distributions are positively skewed, a logarithmic transformation was applied to these variables to fulfil linear model assumptions.

The correlation between the variables was stronger when the canopy N variables were log-transformed and the scatterplots did not show a linear pattern without log-transforming the variables.

The reference list is incomplete (e.g. Bontemps 2011 and ESA 2010 are missing).

Our apologies, we updated and rechecked the reference list.

Minor comments:

please carefully check the usage of hyphen and punctuation.

L 10: I find this sentence confusing, because canopy concentration and content are nothing else then leaf concentration and content accumulated to canopy scale. Why then is it necessary to point to processes at "leaf and canopy scales"

This mention has been removed.

L 17 specify whether the unit is per unit leaf area or per unit ground area

N g m-2 of ground area was added.

L20 here and thereafter (as noted above): note that this is a significant, log-linear relationship. This has been changed.

L25: starting with ", ultimately..." should be removed, because the paper, while showing that a log-linear correlation exists, does not provide the ground for such a mapping. The sentence part has been removed

L31: "such as" before Zaehle & Friend. This has been added

L64:68: irrelevant here, can be removed. This was removed

L72ff: It seems to me that this paragraph would be better suited to follow L48, with the redundant sentences at the beginning of the paragraph currently beginning in L49 removed. The paragraph has been replaced. I do not see which sentence are redundant.

L96-100 can be removed This has been removed

L154 how was fbiom determined?

This question is a bit confusing because the "fbiom" mention appears in the section where we explain how the foliar biomass was measured (Line 169 - 177). Fbiom was replaced by "foliar biomass" in the equation 1 to make it clearer.

L202: Rather: forest plots for which the dominant vegetation type of the gridcell in the MTCI data-set did not correspond to forest were excluded?

This was changed by: "The Globcover 2009 land cover map was used to exclude forest plots for which the dominant vegetation type of the MTCI pixel did not correspond to natural vegetation." (Line 231 – 232).

L203: Sentence beginning with "It was downloaded". Remove sentence as add reference to previous sentence. Both references are missing in the reference list. This was done

L211-215: This does not explain, why this aggregation study was performed. Of course you average across more forest plots when aggregating, but you also average across more MTCI grid-cells. I do not see why this would increase the confidence in the regression analysis. Please be more explicit as to why this should be of interest.

We have now explained in more detail why we performed the resampling for both our datasets in the section 2.3.2 (Line 242 - 250):

This was done because of the initial difference in support size between MTCI spatial resolution and the forest plots size (i.e. 1 km and 6 m, respectively). This enabled us to investigate the relationships between MTCI and canopy N data when the spatial discrepancy was accounted for. The statistical basis of this approach is that we bring both datasets (forest plots and MTCI values) to the same support size or representative area (Bierkens et al., 2000). By averaging out forest plot values within this support size, we calculate the mean of the canopy N value at that support size. By resampling the MTCI values to that same support size, the obtained result consist of a mean of the MTCI value at that support size. We then regressed the expected canopy N values (at the new support size) against the expected MTCI values (at the new support size).

We resampled the datasets to the same lower spatial resolution to be able to study the relationship between MTCI and canopy N independently of the initial difference in support size.

L290-294. Not necessary. Please remove,

This was removed.

L296-303. This partially repeats the results section. Isn't the more relevant question to discuss uncertainties in the upscaling of 6m plots to 1km resolution?

The discussion section was changed to mention the existence of such uncertainties. We have also downscaled the claims about the scalability of the results (Line 335 – 342):

This pre-analysis was undertaken to study the MTCI-canopy N relationships when taking the discrepancy between MTCI original spatial resolution (1 km) and the size of the forest plots (diameter of 6 m) into account. By resampling both datasets to a lower spatial resolution, i.e. 5 km, 10 km, 15 km and 20 km, the obtained values were less impacted by small-scale variations because they were obtained by averaging several values over a larger area. The results showed that the relationship between MTCI and canopy N data was significant and consistent across the resampled spatial resolutions investigated: 5 km, 10 km, 15 km and 20 km. This, however, does not give any indication about the uncertainties resulting from the initial spatial discrepancy between both datasets and about the influence of such uncertainties on the MTCI-canopy N relationship.

L321 and all other places: is the R2 the adjusted r2 to account for the large variance in sample size?

The r2 values were replaced by the adjusted r2 in the tables, graphs and when it appeared in the text. For most of these values, the difference between the r2 and the adjusted r2 was small.

This was also mentioned in the revised manuscript (Line 270 - 271): All the coefficients of determination (r2) presented are the adjusted r2 to account for the differences in sample sizes.

L323: the use of may is not appropriate here. Either the correlation is mainly driven by the deciduous and mixed plots, or it is not. No need for speculation.

The sentence was changed:

This indicates that, the relationship observed for all the forest plots was mainly driven by DBF and mixed plots. (Line 366)

L362: Does this not suggest that the nitrogen signal is confounded by other factors, therefore limiting the ability to "detect" nitrogen with MTCI?

We have modified the sentence to stress that the biomass was a confounder of the MTCI-canopy N relationship in (Line 407 - 408):

This suggests that biomass had an influence on and was a confounder of the MTCI-canopy N log-linear relationship.

And in Line 417 – 418:

In this context, our analysis showed that the PFTs of the plots and the biomass had an influence on the MTCI canopy N relationship in a specific type of ecosystem, namely Mediterranean forests.

L380 I don't understand what you want to say here that is new compared to the preceding sentences. I was trying to summarize the preceding sentences. The sentence was removed.

L387: "...might reveal laborious". Please be more explicit, do you mean to say that this is infeasible? The sentence was changed (Line 434 - 437):

However, due to the different treatments required as well as the limited swath width associated with the high spatial resolution (from 3 m to 30 m for Hyspex airborne and Hyperion spaceborne sensors, respectively, Wang et al., 2016;Smith et al., 2003), applying imaging spectrometry at a broader scale, although feasible, might reveal time-consuming.

Reference

Limpert, E., Stahel, W. A., and Abbt, M.: Log-normal Distributions across the Sciences: Keys and CluesOn the charms of statistics, and how mechanical models resembling gambling machines offer a link to a handy way to characterize log-normal distributions, which can provide deeper insight into variability and probability—normal or log-normal: That is the question, BioScience, 51, 341-352, 10.1641/0006-3568(2001)051[0341:LNDATS]2.0.CO;2, 2001.

Regional detection<u>Remote sensing</u> of canopy nitrogen<u>at regional scale</u> in Mediterranean forests using the spaceborne MERIS Terrestrial Chlorophyll Index

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Abstract. Canopy nitrogen (N) concentration and content are linked to several vegetation processes. at leaf and canopy levels. Therefore, canopy N concentration is a state variable in global vegetation models with coupled carbon (C) and N cycles. While there is ample C data available to constrain the models, widespread N data are lacking. Remote sensing and Remotely sensed vegetation indices have been used to detect canopy N concentration and canopy N content at the local scale in grasslands and forests. Vegetation indices could be a valuable tool to detect canopy N concentration and canopy N content at larger scale. In this paper we conducted a regional case-study analysis to investigate the relationship between the Medium Resolution Imaging Spectrometer (MERIS) Terrestrial Chlorophyll Index (MTCI) time series from ESA ENVISAT at 1 km spatial resolution and both canopy N concentration (%N) and canopy N content (N g m⁻² of ground areag m⁻²) from a Mediterranean forests inventory in the region of Catalonia, NE of Spain. The relationships between the datasets were studied after resampling both datasets to lower spatial resolutions (20 km, 15 km, 10 km and 5 km) and at the initial higher original spatial resolution of 1 km. The results at the higher spatial resolution (1 km) yielded significant log-linear relationships between MTCI and both canopy N concentration and content, $r^2 = 0.32$ and $r^2 = 0.17$, respectively. We also investigated these relationships per plant functional type. While the relationship between MTCI and canopy N concentration was strongest for deciduous broadleaf and mixed plots $(r^2 = 0.254)$ and $r^2 = 0.474$, respectively), the relationship between MTCI and canopy N content was strongest for every needleleaf trees ($r^2 = 0.1920$). At the species level, canopy N concentration was strongly related to MTCI for European Beech plots ($r^2 = 0.6974$). These results present a new perspective on the application of MTCI time series for canopy N detection. ultimately leading towards the generation of canopy N maps that can be used to constrain global vegetation models.

Keywords: vegetation index, MERIS, foliar nitrogen concentration, foliar nitrogen content, plant functional types, 30 Mediterranean forest, remote sensing

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1 Introduction

Canopy nitrogen (N) concentration is an essential state variable in regional (Ollinger and Smith, 2005) and global vegetation models including both the carbon (C) and the N cycles (such as Zaehle and Friend, 2010;Smith et al., 2014). This variable has been linked to several vegetation traits and processes at the leaf and canopy levels. At the leaf level, leaf N concentration, which represents the leaf N status expressed as a percentage of leaf dry matter (%N, N g 100g⁻¹ DM), has been related to photosynthetic capacity (Evans, 1989;Reich et al., 1995;Reich et al., 1997;Reich et al., 1999;Wright et al., 2004) and light use efficiency (Kergoat et al., 2008). Leaf N concentration expressed on a leaf area basis, also called leaf N content (N g m⁻²), has also been linked with chlorophyll content, Rubisco content (Evans, 1989) and photosynthetic capacity (Evans, 1989;Reich et al., 1995). At stand scale, canopy nitrogen concentration, which

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represents the leaf N concentration averaged over the stand canopy, has also been found to correlate with above ground Net Primary Productivity (NPP) (Reich, 2012), while canopy N content has been linked with the canopy light use efficiency (Green et al., 2003).

Given their links to many vegetation processes, leaf and canopy N variables could be used to constrain N cycle modules in global vegetation models. At the global scale, ample data is available to constrain models for the C cycle; however, data to

- 45 constrain the N cycle are limited. Currently, canopy N data are not widely available and canopy N sampling campaigns are time-consuming and thus expensive tasks. Moreover, upscaling from local sampling campaign measurements represents an additional limitation. In this perspective, local, regional or even global remotely sensed canopy N estimates will be a valuable addition, enabling us to collect information in a less time intensive and expensive manner than traditional on-field sampling campaigns. Such near global canopy N estimates will be beneficial as input in global vegetation models or to calibrate and the provide these models.
- 50 validate these models.

Remote detection of foliage N status has been extensively studied at the leaf scale (Hansen and Schjoerring, 2003;Ferwerda et al., 2005;Li et al., 2014) and few studies have investigated the processes underlying the relationships between vegetation indices and foliar N (Pacheco-Labrador et al., 2014). Detection of foliage N status with vegetation indices is attributed to the strong link between foliar nitrogen and chlorophyll content (Schlemmer et al., 2013) and is often based on the NIR and red-edge region of the spectrum, hence similar to the ones used for chlorophyll detection (Filella and Penuelas, 1994;Dash and Curran, 2004;Clevers and Gitelson, 2013). At canopy level, however, spectral reflectance is a complex function of vegetation cover, plant activity, water content, illumination angle, viewing angle and atmospheric composition (Kumar et al., 2006) and it is not straightforward to disentangle the influence of nitrogen from other contributions in the spectra. It is thus not clear how the relationships observed at the leaf level translate at the canopy level. The mechanisms possibly modifying the remote sensing of foliage N status at the canopy scale are still not clearly understood (Ollinger, 2011). High correlation between canopy N and both NIR reflectance and albedo has been reported in boreal forests (Ollinger et al., 2008). However, the mechanism behind these findings is still controversial. Knyazikhin et al. (2013) argued that the observed correlation solely resulted from canopy structural differences between broad and needleleaf forests and was thus spurious. Other authors, although agreeing that canopy

structure was a confounding factor to account for, stated that the NIR - canopy N relationship was not necessarily spurious and

65 stemmed from an association between canopy N and structural traits (Ollinger et al., 2013;Townsend et al., 2013). Canopy traits are interrelated (Wright et al., 2004) and have been known to covary due to evolutionary convergence, as stated by Ollinger (2011).

Different remote sensing techniques have been applied to detect canopy N in terrestrial vegetation. Imaging spectrometry has proven efficient in improving N sensing capabilities at the local scale. Imaging spectrometry images are acquired from either airborne or spaceborne sensors and are analysed with different methods, including partial least squares regression (PLS), continuum removal, spectral unmixing or vegetation indices (Smith et al., 2003;Ollinger et al., 2008;Huber et al., 2008;Martin et al., 2008;Schlerf et al., 2010;Wang et al., 2016). Among other techniques, ratios or normalized differences of reflectance bands in the Red and Near Infrared (NIR) regions of the spectrum, the so called vegetation indices (VI) (Glenn et al., 2008), are one of the most straightforward methods for canopy N detection. Combined with in situ hyperspectral devices, vegetation

75 indices have been extensively used for leaf or canopy N detection in agricultural systems (Peñuelas et al., 1994;Filella et al., 1995;Hansen and Schjoerring, 2003;Tian et al., 2011;Schlemmer et al., 2013;Li et al., 2014). Vegetation indices have also been applied to airborne or spaceborne acquired imagery in natural environments (Ramoelo et al., 2012;Wang et al., 2016).

A particular vegetation index, the MERIS Terrestrial Chlorophyll Index (MTCI) has been proposed for detecting canopy N (Clevers and Gitelson, 2013). MTCI was originally computed from three reflectance bands from the Medium Resolution
 Imaging Spectrometer (MERIS) aboard the European Space Agency (ESA) ENVISAT satellite at a spatial resolution of 1 km. However, it can also be obtained from other sensors' reflectance data and a similar product will be available from the ESA Sentinel-2 satellite mission (Drusch et al., 2012). It was first developed to estimate chlorophyll content (Dash and Curran, 2004, 2007). Since then, other applications of this index have been described, among which the possibility to estimate Gross Primary Productivity (GPP) from natural (Harris and Dash, 2010, 2011;Boyd et al., 2012) and cultivated lands (Peng and Gitelson, 2011). Furthermore, MTCI has been used to discriminate between C3 and C4 grasses (Foody and Dash, 2007) and to monitor vegetation phenology at the sub regional (Boyd et al., 2011) and continental scales (Rodriguez Galiano et al., 2015;Crabbe et al., 2016). Regarding canopy N detection, most studies were carried out in agricultural crops using MTCI values computed from

- in situ hyperspectral reflectance data (Tian et al., 2011;Clevers and Gitelson, 2013;Li et al., 2014). A few were directed towards sensing N concentration in natural environments using airborne data, e.g. in temperate forests (Wang et al., 2016), or spaceborne
 data, for example in grasslands (Ramoelo et al., 2012;Ullah et al., 2012) or sub-tropical forests (Cho et al., 2013).
- Remote detection of foliage N status has been extensively studied at the leaf scale (Hansen and Schjoerring, 2003;Ferwerda et al., 2005;Li et al., 2014) and few studies have investigated the processes underlying the relationships between vegetation indices and foliar N (Pacheco Labrador et al., 2014). Detection of foliage N status with vegetation indices is attributed to the strong link between foliar nitrogen and chlorophyll content (Schlemmer et al., 2013) and is often based on the NIR and red edge region of the spectrum, hence similar to the ones used for chlorophyll detection (Filella and Penuelas, 1994;Dash and Curran, 2004;Clevers and Gitelson, 2013). At canopy level, however, spectral reflectance is a complex function of vegetation cover,
 - plant activity, water content, illumination angle, viewing angle and atmospheric composition (Kumar et al., 2006) and it is not

straightforward to disentangle the influence of nitrogen from other contributions in the spectra. It is thus not clear how the relationships observed at the leaf level translate at the canopy level. The mechanisms possibly modifying the remote detection of foliage N status at the canopy scale are still not clearly understood (Ollinger, 2011). High correlation between canopy N and both NIR reflectance and albedo has been reported in boreal forests (Ollinger et al., 2008). However, the mechanism behind these findings is still controversial. Knyazikhin et al. (2013) argued that the observed correlation solely resulted from canopy structure was a confounding factor to account for, stated that the NIR – canopy N relationship was not necessarily spurious and stemmed from an association between canopy N and structural traits (Ollinger et al., 2013;Townsend et al., 2013). Canopy traits are interrelated (Wright et al., 2004) and have been known to covary due to evolutionary convergence, as stated by Ollinger (2011).

In this context, there are several knowledge gaps that we would like to address in this paper. First, although 1 km spatial resolution spaceborne MTCI time series are available from the ESA, MTCI has mainly been employed to detect canopy N in 110 agricultural applications with in situ devices and rarely in a broader range of natural ecosystems and scales using spaceborne data. Due to its almost global coverage, MTCI time series could be applied to estimate canopy N over a larger spatial extent Moreover, Mediterranean forests have specific functional characteristic due to their great forest ecosystems diversity, influenced by contrasting climatic and topographic conditions, and their high tree species richness (Vilà-Cabrera et al., 2018). However, to our knowledge, limited research has been conducted to sense canopy N in Mediterranean ecosystems (Serrano et al., 2002) 115 and even more so in Mediterranean forests. In addition, although in a temperate forest the reflectance spectrum of individual plant functional types (PFT) has been shown to be different (Wang et al., 2016), the relationship between MTCI and canopy N has seldom been studied and compared between PFTs. Moreover, investigating the influence of PFTs on this relationship might give further insight into the influence of structural effects in canopy N detection. Finally, the difference between sensing canopy N concentration ($N_{I_{22}I_{2}}$, %N) and canopy N content ($N_{I_{appeal_{7}}}$ g m⁻²) has rarely been investigated. The relationship between MTCI and both N concentration ($N_{1\%1}$, %N) and canopy N content (N_{lareal} , g m⁻²) of these variables has been studied separately (Clevers 120 and Gitelson, 2013; Wang et al., 2016), but very few analyses (Mirik et al., 2005; Ullah et al., 2012) have compared the ability

to detect canopy N concentration and canopy N content simultaneously, especially in forest ecosystems.
 The objective of our study is thus to investigate the relationship between the spaceborne MTCI remote sensing product and canopy N in Mediterranean forests at the regional scale. More specifically, the relationships between MTCI and both canopy N
 concentration and canopy N content are investigated and compared. We then also examine these relationships per PET and at

125 concentration and canopy N content are investigated and compared. We then also examine these relationships per PFT and at the species level.

<u>Remote sensingDetection</u> of canopy N is often limited to local scale studies due to the spatial restrictions associated with N data acquisition in the field and treatment of high spatial resolution remote sensing imagery with limited spatial coverage (Lepine et al., 2016). Our case-study exploits the broadly and readily available MTCI time series at 1 km spatial resolution from the ESA
 ENVISAT mission and combines it with canopy N data, both concentration and content, from 846 forest plots measured between 1988 and 2001 by the Catalonian National Forest Inventory (Gracia et al., 2004). First, we develop a methodology to overcome

the time discrepancy between our two sets of data. Next, both data sets are resampled to the same, lower, spatial resolutions, i.e. 5 km, 10 k, 15 km and 20 km, in order to overcome the initial spatial discrepancy between MTCI spatial resolution (1 km) and the size of the forest plots (6 m). Subsequently, we analyse the relationship between MTCI and both canopy N concentration

135 and canopy N content variables, both at the resampled and initial spatial resolutions. The relationships at the initial spatial resolution are then stratified according to the PFT of the plots. The results are presented and discussed. Finally, we address the implications for future research and draw a conclusion.

2 Material and methods

2.1 Study area

- Our study area corresponds to the region of Catalonia (Fig. 1) which is located in north eastern Spain and has a spatial extent of 32,114 km² (Sardans et al., 2011). While the region is characterised by a Mediterranean climate, the presence of the Pyrenees to the northwest and the Mediterranean Sea to the east creates contrasting climate conditions with an altitudinal gradient from north to south and a continental gradient from west to east. Following this pattern, the mean annual temperature varies from 1 °C in the north to 17 °C in the south (Sardans et al., 2011). While mean annual precipitation (MAP) is 1400 mm in the Pyrenees, in the south, the MAP is lower than 350 mm (Sardans et al., 2011), leading to seasonal drought (Lana and Burgueño,
- 145 Pyrenees, in the south, the MAP is lower than 550 min (Sardans et al., 2011), leading to seasonal drought (Lana and Burgueno 1998) and fires (González and Pukkala, 2007), impacting the vegetation (Liu et al., 2015).

2.2 Data collection

2.2.1 Canopy N data

The canopy N data used in this research was collected by the Ecological and Forestry Applications Research Centre (CREAF),
Universitat Autònoma de Barcelona. The data included 2300 closed canopy forest plots sampled between 1988 and 2001 by the Catalonian National Forest Inventory (Gracia et al., 2004).

The forest plots (Fig. 1) had a minimum diameter of 6 m, which varied depending on the tree density in order to include between 15 and 25 trees with a diameter at breast height (DBH) of at least 5 cm. The DBH was recorded for all the trees present on the plot with a DBH of minimum 5 cm. The plots were investigated for canopy N concentration ($N_{[\%]}$, %N) defined as g of N per

- 155 100 g of leaf dry matter. The leaf samples were collected from the upper central part of the crown using extensible loppers. All foliar cohorts present in the canopy were included in the leaf sample. Each leaf sample was constituted by the leaves of at least three different trees of the dominant tree species in the canopy. The species dominance was determined by the tallest individual. A proportion of 96% of the plots included in this analysis were monospecific (Sardans et al., 2011). 4% of the plots (n = 30) had two codominant species. For these plots, two leaf samples were collected, one for each of the codominant species found on the plots.
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The leaf samples were dried and then ground using a Braun Mikrodismembrator-U (B. Braun Biotech International, Melsungen, Germany). They were analysed for foliar N concentration using the combustion technique coupled to gas chromatography using a Thermo Electron Gas Chromatograph (model NA 2100, CE Instruments-Thermo Electron, Milan, Italy) (Gracia et al., 2004). To scale from leaf to canopy level, we used the leaf nitrogen concentration averaged over three individuals as the plot level value (Schlerf et al., 2010). We did not weight the average by species abundance (Smith and Martin, 2001) as only 4% of the

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plots had two different species.

Along with the canopy $N_{[\%]}$ data, we used foliar biomass data (dry matter g per square meter of ground area, g m⁻²) acquired during the same forest inventory (n = 2286). The foliar biomass data were obtained for each plot from allometric equations relating the diameter at breast height to the leaves dry weight. The allometric equations were species specific (Sardans et al. (2015), Table A1). The foliar biomass data were used to calculate canopy N content (N_[area], g of N per m⁻² of ground) for each plot following Eq. (1):

$$canopy N_{[area]} = \frac{canopy N_{[\%]} * foliar biomassfbiom}{100},$$
(1)

where *canopy* $N_{[area]}$ is the canopy N content (N g per square meter of ground area, g m⁻²), *canopy* $N_{[\%]}$ is the canopy N concentration (%N) and *fbiom*<u>the *foliar biomass*</u> is the foliar biomass<u>expressed in</u> (dry matter g per square meter of ground area, g m⁻²).

For the plots with two codominant species, the concentration measurements were done separately. The obtained foliar N concentration and biomass values were then averaged to obtain a single canopy $N_{[\%]}$ and canopy_[area] value for each plot with two codominant species. Among these 30 plots with codominant species, 16 plots had codominant species from different PFT. Their PFT is thus labelled as mixed while the plots with several codominant species from the same PFT are labelled according to their PFTs.

Catalonian forests include both deciduous and evergreen broadleaf as well as evergreen needleleaf tree species. These three PFTs are referred to as Deciduous Broadleaf Forest (DBF), Evergreen Broadleaf Forest (EBF) and Evergreen Needleleaf Forest (ENF), respectively. The main tree species are *Pinus halepensis* Mill., *Pinus sylvestris* L., *Quercus ilex* L., *Pinus uncinata* Ramond ex DC., *Pinus nigra* J.F. Arnold, *Quercus suber* L., *Quercus cerrioides* Willk. & Costa., *Quercus petraea* Liebl. and

185 Fagus sylvatica L. These species accounted for 92% of the sampled forest plots. The 15 tree species included in this analysis are listed in Table 1. Plots with a rare dominant tree species, i.e. species that were detected in only one single plot, were excluded from the analysis. This applied to plots with these dominant species: Abies alba Mill., Fraxinus augustifolia Vahl, Fraxinus excelsior L., Pinus radiata D. Don, Populus nigra L., Populus tremula L., Quercus robur L.

2.2.2 MTCI product

190 The MERIS Terrestrial Chlorophyll Index (MTCI) was first developed to estimate chlorophyll content in canopies. MTCI is sensitive to high chlorophyll content while presenting low sensitivity to soil brightness (Curran and Dash, 2005). Its calculation, presented in Eq. (2), is based on three reflectance bands, located around the red edge point (REP) (Dash and Curran, 2004):

$$MTCI = \frac{R_{band10} - R_{band9}}{R_{band9} - R_{band8}} = \frac{R_{753.75} - R_{708.75}}{R_{708.75} - R_{681.25}}$$
(2)

- where *R_{band8}*, *R_{band9}* and *R_{band10}* represent the 8th, 9th and 10th bands of MERIS, respectively. Following MERIS standard
 bands settings, the centres of the bands were located at 681.25 nm, 708.75 nm and 753.75 nm on the electromagnetic spectrum. While the ESA ENVISAT satellite mission producing MERIS data came to an end in 2012, MERIS products and MTCI in particular are still relevant because the new ESA Sentinel-2 and Sentinel-3 satellite missions haves improved band settings compared to those of MERIS, and increased the spatial resolution to 20 m MTCI can be calculated from Sentinel-2 reflectance data with increased spatial resolution to 20 m -(Drusch et al., 2012). The Sentinel-3 mission also releases a level 2 chlorophyll product, the OLCI Terrestrial Chlorophyll Index (OTCI), which calculation is directly based on MTCI. OTCI continues the time series already available for MTCI. In this study, we put emphasis on ENVISAT-MERIS as our field data are closer to the MERIS acquisition period.
- MTCI level 3 imagery was obtained from the NERC Earth Observation Data Centre (NEODC, 2015) for the region of Catalonia
 between 2002 and 2012. The original data were provided by the European Space Agency and then processed by Airbus Defence and Space. The original MERIS reflectance images, following ENVISAT specifications, have a revisit time of three days and a spatial resolution of 300 m. Compared to the original reflectance images, the MTCI processed imagery has been corrected for atmospheric influences and cloud cover (Curran and Dash, 2005) and is available as an either weekly or monthly averaged product <u>almost globally (Curran et al., 2007)</u>. The spatial resolution of the processed data is approximately 1 km. As there is no temporally averaged product available at full resolution, we chose to carry out this analysis with the MTCI monthly averaged processed imagery. This was done to decrease the uncertainty resulting from the use of single daily reflectance values. An MTCI time series of 10 years is available almost globally. One MTCI monthly averaged imagery product covering the entire study area was obtained for every month between June 2002 and March 2012, except for October 2003, when no valid product was available.

215 2.3 Data handling

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2.3.1 Methodology to link canopy N data to MTCI values

There is a discrepancy between the timing of the ground truth sampling and the satellite image acquisition period. While the plot sampling campaigns were carried out between 1988 and 2001, the ENVISAT satellite mission was launched in 2002 and ended in 2012. To overcome the discrepancy, MTCI images were averaged by month over the 10 years of the satellite mission period. This process yielded twelve MTCI averaged images, one for each month. The averaged MTCI images were then linked to the forest plots based on the forest plot coordinates and sampling month, as the exact sampling date was known for each plot. The period between the 1st of June and the 31st of October was determined to be the growing season after a pre-analysis, where we studied yearly temporal variation of MTCI in several locations and forest types in Catalonia. This extended period was chosen to encompass the different vegetation phenology types corresponding to the contrasted climate conditions in this region.

225 The forest plots sampled outside of the growing season were excluded from the analysis. The inter-annual variation of canopy N_{1%1} data was analysed for each month included in the analysis to ensure that the ground data could be related with MTCI data (Figure A1). The Globcover 2009 land cover map was used to exclude forest plots for which the dominant vegetation type of the MTCI pixel did not correspond to natural vegetation. to exclude forest plots located on unsuitable land surface. The Globcover map was created by ESA using MERIS reflectance data from 2009 (Bontemps et al., 2011). The Globcover mapHt 230 was downloaded from the ESA data user elements website (ESA, 2010). This map comprises 22 land cover classes and has a spatial resolution of 300 m. Using this map, we excluded forest plots that had undergone a land cover change since the sampling period and did not have a natural vegetation cover any more at the time of remote sensing image acquisition. To do so, the landcover map was first resampled to a spatial resolution of 1 km to be in accordance with MTCI spatial resolution. Then, the plots located on land area classified as either rainfed cropland, mosaic between croplands and natural vegetation, sparse 235 vegetation or artificial surfaces were excluded from the analysis.

2.3.2 Relationship between MTCI and canopy N data at lower spatial resolution

In a first step, the relationships between MTCI and canopy N data values were investigated after resampling both datasets to the same, lower, spatial resolution. The resampled spatial resolutions were 5 km, 10 km, 15 km, and 20 km. This was done because of the initial difference in support size between MTCI spatial resolution and the forest plots size (i.e. 1 km and 6 m, 240 respectively). This enabled us to investigate the relationships between MTCI and canopy N data when the spatial discrepancy was accounted for. independently of differences in initial support size. The statistical basis of this approach is that we bring both datasets (forest plots and MTCI values) to the same support size or representative area (Bierkens et al., 2000). By averaging out forest plot values within this support size, we calculate the mean of the canopy N value at that support size. By resampling the MTCI values to that same support size, the obtained result consist of a mean of the MTCI value at that support size. We 245 then regressed the expected canopy N values (at the new support size) against the expected MTCI values (at the new support size).

The monthly averaged MTCI images obtained previously (section 2.3.1) were resampled successively to 5 km, 10 km, 15 km, and 20 km. Beforehand, the Globcover 2009 land cover map was used to exclude from the resampling computation the MTCI pixels located on land surface without natural vegetation cover. As for the forest plots, MTCI pixels whose land cover class corresponded to rainfed cropland, mosaic between croplands and natural vegetation, sparse vegetation or artificial surfaces were excluded from the upscaling analysis. Forest plots data were then averaged per month over the newly obtained pixel. The relationship between the resampled MTCI values and canopy N data was analysed using log-linear regression.

2.3.3 Relationship between MTCI and canopy N data at initial higheroriginal spatial resolution (1 km)

In a second step, the relationships between MTCI and canopy N data, both canopy N_[%] and canopy N_[area], were examined at the original spatial resolution of 1 km. This allowed us to investigate the influence of PFT and species on the relationships as

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this information was lost in the resampling process. The relationships between MTCI and canopy N at 1 km-spatial resolution were analysed with log-linear regression for the whole dataset, for each PFT separately as well as for individual species.

2.3.4 **Statistical analysis**

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After applying the selection criteria as explained in the section 2.3.1, i.e. plots measured between June 1st and October 31st, exclusion of plots with infrequent species and selection based on Globcover 2009, 846 forest plots were available for analysis, including 841 plots with foliar biomass and canopy N content information. Descriptive statistics of canopy N_[%], foliar biomass and canopy $N_{\text{[area]}}$ were produced for each of the tree species and PFT included in the analysis. The <u>log-linear</u> regressions between MTCI and canopy N were performed for both resampled and non-resampled datasets. Preliminary analysis showed that using a natural logarithm transformation (log) of the canopy N variables was necessary to fulfil linear regression model 265 assumptions, namely normality and homogeneity of variance of the residuals. The minimum number of data points needed to carry out the linear regression analysis was fixed at 10. All the coefficients of determination (r^2) presented are the adjusted r^2 to account for the differences in sample sizes. We calculated the Relative Root Mean Square Error of cross-validation (RRMSEcv, %) using the leave-one-out cross validation method (Clevers and Gitelson, 2013). Its calculation is presented in Eq. (3) following (Yao et al., 2010):

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$$RRMSEcv = \sqrt{\frac{1}{n} \times \sum_{i=1}^{n} (P_i - O_i)^2} \times \frac{100}{\overline{O_i}}$$
(3)

where P_i represents the predicted value, O_i , the observed value, \overline{O}_i the mean of all observed value and n the total number of measurement. Resampling both datasets as well as linking the plots to the MTCI pixels was done with the PCRaster software (Karssenberg et al., 2010). The statistical analyses were performed in the R environment (R Development Core Team, 2014) and the ggplot2 package was used for the graphics (Wickham, 2009).

3 Results 275

3.1 **Descriptive statistics**

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in the dataset (Table 1). The four most abundant species (Pinus halepensis, Pinus sylvestris, Quercus ilex and Pinus uncinata) dominated 667 plots i.e. almost 80% of the plots. The cumulated abundance percentages of ENF, EBF and DBF species were equal to 66 %, 22 % and 9 %, respectively. From this data, it is clear that the forests plots were mainly dominated by ENF species. On average, *Pinus uncinata* plots had the highest biomass values while *Quercus suber* plots showed the lowest mean value for this variable. Descriptive statistics were also analysed by PFT. The mean canopy $N_{[\%]}$ was lowest for ENF species, 0.97 %N, and highest for DBF trees, 2.17 %N (Fig. 2a). Canopy N_[%] value ranges were equal to 1.91 %N, 2.06 %N, 1.68 %N and 1.42 %N for DBF, EBF, ENF and mixed plots, respectively. The canopy N_[area] statistics were analysed by PFT as well (Fig.

Descriptive statistical analysis of canopy $N_{[m]}$, canopy $N_{[area]}$ and foliar biomass were performed for each tree species included

- 2b) and the averaged canopy N_[area] values ranged from 1.82 g m⁻² to 4.61 g m⁻². A Pearson correlation matrix (Fig. 3) was 285 computed between the variables for the whole dataset. The correlation between each pair of variables was significant and the correlation between canopy $N_{\text{[area]}}$ and foliar biomass was strongest (r = 0.88). This result was expected as the foliar biomass was included in the $N_{\text{[area]}}$ calculation. This matrix also shows distribution histograms of the three variables. As canopy $N_{\text{[\%]}}$ and canopy N_{fareal} distributions are positively skewed, a logarithmic transformation was applied to these variables to fulfil linear 290 model assumptions. Correlation matrices for each DBF, EBF and ENF plots are presented in the Appendix (Fig. A 2-4).

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3.2 Relationship between MTCI and canopy N data at lower spatial resolution

The relationships between MTCI and both canopy $N_{1\%1}$ and canopy N_{lareal} were studied after resampling both datasets to the same, lower, spatial resolution. This was done to investigate the relationship between MTCI and canopy N data when the initial spatial discrepancy between the two datasets was accounted for. <u>independently of differences in support size</u>. The results showed that the <u>log-linear</u> relationships between MTCI and either canopy $N_{[3e]}$ or canopy $N_{[area]}$ were all highly significant (p<0.000). Moreover, the relationship between MTCI and canopy $N_{[\%]}$ was always stronger than the relationship for MTCI and canopy $N_{\text{[area]}}$ for each resampling factor. The r² values of the relationship between MTCI and canopy $N_{\text{[\%]}}$ were equal to 0.33, 0.37, 0.34 and 0.42 for 5 km, 10 km, 15 km and 20 km resampled spatial resolution, respectively. The r² values of the relationship between MTCI and canopy $N_{\text{[area]}}$ were equal to 0.20, 0.20, 0.1920 and 0.187 at 5 km, 10 km, 15 km and 20 km spatial resolution. The relationship between MTCI and canopy $N_{[\%]}$ at 20 km spatial resolution is shown in Figure 6. Table 2 shows the number of plots per pixel for different pixel sizes (km). As expected, the number of plots per pixel increased with the pixel size, with a mean of 4.1 plots at 20 km spatial resolution. The descriptive statistics of the number of different PFT, species and sampling years per pixel spatial resolution are provided in the Appendix (Table A 2 - A 4).

3.3 Relationship between MTCI and canopy N data at original higher spatial resolution (1 km)

305 3.3.1 **Relationship between MTCI and canopy N concentration**

The relationships between MTCI and canopy N data were studied at the initial higheroriginal spatial resolution (1 km). The results showed that the log-linear regression between MTCI and canopy $N_{1\%1}$ for the whole dataset (n = 846) was highly significant (p<0.000) and had an r² value of 0.32 and an RRMSEcv value of 18.7 % (Table 3, Fig. 5a). The relationship between MTCI and Canopy N_[%] was also investigated for each PFT individually (Fig. 5b-e). For DBF plots, the relationship between MTCI and canopy $N_{1\%1}$ had an r² value of 0.254 (n = 80) and was significant. However, although statistically significant, the r² of the relationship between MTCI and canopy $N_{1\%1}$ for EBF and ENF plots were lower and equal to 0.032 (n = 186) and 0.10 (n = 564), respectively.

The relationship between MTCI and canopy $N_{[\%]}$ was also significant for one individual species, *Fagus sylvatica*. The proportion of explained variance for this species was equal to 0.6971 (n = 15). This result, although obtained on a restricted number of

315 plots, showed that the significant relationships between MTCI and canopy $N_{[\%]}$ not only existed when all DBF plots were included but also held for one individual DBF species.

3.3.2 Relationship between MTCI and canopy N content

Significant relationships between MTCI and canopy $N_{[area]}$ were found for the whole dataset as for EBF and ENF plots (Table 3). The scatterplots between MTCI and canopy $N_{[area]}$ are presented in Figure 6. The proportion of explained variance was higher for ENF plots compared to the other PFTs and compared to the overall relationship across all plots. The relationship between MTCI and canopy $N_{[area]}$ was also investigated for 10 individual species and one of them showed significant relationships: *Quercus ilex* (r² = 0.10, p-value < 0.000, n = 160).

4 Discussion

Our aim was to explore the relationship between the MTCI vegetation index and both canopy N_{1%3} and canopy N_{fareal} in Mediterranean forests at the regional scale in Catalonia, north eastern of Spain. This was done by using the ESA spaceborne MTCI remote sensing product and canopy N data from a forest inventory. The relationship was first investigated using MTCI and canopy N data resampled to the same, lower, spatial resolution. The relationship was then investigated across all plots and by PFT at MTCI initial spatial resolution of 1 km.

4.1 Relationship between MTCI and canopy N data at lower spatial resolution

This pre-analysis was undertaken to study the MTCI-canopy N relationships when takingindependently of the discrepancy between MTCI original spatial resolution (1 km) and the size of the forest plots (diameter of 6 m) into account. By resampling both datasets to a lower spatial resolution, i.e. 5 km, 10 km, 15 km and 20 km, the obtained values were less impacted by small-scale variations because they were obtained by averaging several values over a larger area. The results showed that the relationship between MTCI and canopy N data was significant and consistent across the resampled all spatial resolutions investigated: 5 km, 10 km, 15 km and 20 km. This, however, does not give any indication about the uncertainties resulting from the initial spatial discrepancy between both datasets and about the influence of such uncertainties on the MTCI-canopy N relationship. This showed that, when the spatial discrepancy between the original datasets, i.e. 6 m and 1 km, was taken into account, MTCI and canopy N data were linked and that the MTCI canopy N relationship was not strongly affected by the resampled spatial resolution.

340 4.2 Relationship between MTCI and canopy N data at <u>original higher</u> spatial resolution (1 km)

4.2.1 Canopy N concentration detection

The overall relationship between MTCI and canopy $N_{[\%]}$ at 1 km spatial resolution for all the forest plots (n = 846) was significant and the r² value was equal to 0.32 (Table 3, Fig. 5). This result showed that canopy N_[%] could be related to MTCI in Mediterranean forests. The performance of the MTCI vegetation index to detect canopy $N_{[\%]}$ in Mediterranean vegetation 345 was similar to the results obtained from previous studies using spaceborne MTCI at higher spatial resolution. For example, using MTCI computed from the spaceborne RapidEye sensor at 5 m spatial resolution, it was possible to detect canopy $N_{1\%1}$ in grassland savannah and sub-tropical forest with similar coefficients of determination, $r^2 = 0.35$ and $r^2 = 0.52$. respectively (Ramoelo et al., 2012; Cho et al., 2013). However, while there is a consensus regarding MTCI ability for in situ leaf or canopy N_{1%1} detection in a variety of crops using handheld spectrometers (Tian et al., 2011;Li et al., 2014), there is no general agreement 350 about MTCI ability for canopy $N_{1\%1}$ detection across vegetation and sensor types at larger scales. For example, MTCI computed from airborne data at 3 m spatial resolution could not be related to canopy $N_{[\%]}$ from a mixed temperate forest (Wang et al., 2016). In this context our finding brings new insight into MTCI $N_{[\%]}$ sensing capabilities at a much coarser spatial resolution (1 km) compared to what has been done before. In these comparisons, it should be taken into account that most previous studies were based on a short sampling campaign while our study incorporates canopy N data from a forest inventory that was carried 355 out during the entire growing season and therefore includes differences in phenology.

Investigating the influence of the PFTS on the overall relationship highlighted the difference between DBF, EBF and ENF types of vegetation regarding canopy $N_{[\%]}$ detection by spaceborne MTCI. The relationships between MTCI and canopy $N_{[\%]}$ were significant for all the PFT taken separately (p-value<0.05). However, a higher proportion of variance was explained for DBF and mixed plots ($r^2 = 0.245$ and $r^2 = 0.447$ for DBF and mixed plots, respectively) compared to the other plant functional types $(r^2 = 0.10 \text{ and } r^2 = 0.032 \text{ for ENF and EBF trees, respectively})$ and the relationship between MTCI and canopy N_[%] was especially 360 weaker for EBF plots. This indicates that Therefore, the relationship observed for all the forest plots wasmay be mainly driven by DBF and mixed plots. This result is different from what was observed by Ollinger et al. (2008) in boreal forests, where canopy $N_{1\%1}$ was related to NIR reflectance for both broadleaf and needleleaf plots taken separately. Moreover, the results obtained for ENF tree species are surprising as previous studies investigating the relationship between foliar $N_{[\%]}$ and in situ measured spectra reported higher r² values, $r^2 = 0.59$ and $r^2 = 0.81$ in spruce and pine forest, respectively (Stein et al., 365 2014; Schlerf et al., 2010). The differences in scale and methodology might explain the divergent results compared to previous findings. Indeed, in our study, the analysis is carried out at a much coarser spatial resolution using spaceborne data compared to the fine spatial scale obtained with in situ devices. Moreover, most of these studies were carried out in temperate forests and studies investigating canopy $N_{[\%]}$ detection in Mediterranean regions are scarce. When investigating the relationship between canopy $N_{[\%]}$ and MTCI at the species level, we also found that it was significant for *Fagus sylvatica* plots ($r^2 = 0.6974$). 370

In the literature, the relationship between MTCI and canopy $N_{[\%]}$ is often not stratified by PFT or species (Sullivan et al., 2013; Wang et al., 2016). In this study, we showed that investigating this relationship for each PFT taken separately yielded

additional insight. Indeed, to our knowledge the difference in explained variance between DBF and other PFTs in MTCI and canopy $N_{[\%]}$ relationship has not been observed before. Moreover, the results observed for *Fagus sylvatica* plots (n = 15) were consistent with the stronger relationship observed for DBF plots.

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4.2.2 Canopy N content detection

The relationship between MTCI and canopy N_[area], which was obtained by combining canopy N concentration values with biomass data, was significant across all plots (n = 841) (Table 4, Fig. 6). Although the r² value was lower for the relationship between MTCI and canopy N_[area] (r² = 0.17) than for the relationship between MTCI and canopy N_[area] (r² = 0.32), it is interesting to note that canopy N_[area] can be related to spaceborne MTCI as remotely sensed detection of canopy N_[area] is rarely investigated in forest environments (Mirik et al., 2005). In comparison, previous studies conducted in grasslands reported higher prediction accuracy e.g. by using spaceborne MTCI at 300 m spatial resolution or a simple ratio-type vegetation index computed from airborne imagery at 1 m spatial resolution, canopy N_[area] was detected with r² values equal to 0.29 and 0.66, respectively (Mirik et al., 2005;Ullah et al., 2012).

The relationship between MTCI and canopy $N_{[area]}$ was only significant for ENF and EBF plots (Fig. 6b-e), with a higher proportion of explained variance for ENF plots ($r^2 = 0.1920$). However, when this relationship was investigated at the species scale, significant results were found for *Quercus ilex* (EBF) plots. This is accordance with a previous study examining the <u>remote sensingdetection</u> of canopy $N_{[area]}$ in *Quercus ilex* trees by MTCI computed from in situ spectra ($r^2 = 0.43$) (Pacheco-Labrador et al., 2014).

390 4.3 Comparing results obtained for canopy N concentration and canopy N content detection

This analysis highlighted the difference between canopy N expressed as a percentage of leaf dry matter (canopy $N_{[\%]}$) and on an area basis (canopy $N_{[area]}$) regarding the log-linear relationship withdetection by spaceborne_MTCI for the different PFTs. Canopy $N_{[\%]}$ of DBF and mixed plots showed higher correlation with MTCI compared to EBF and ENF plots while the relationship between canopy $N_{[area]}$ of ENF plots with MTCI was stronger than for any other PFTs. These differences between the log-linear relationship between MTCI and either canopy $N_{[\%]}$ and canopy $N_{[area]}$ detection by remote sensing can be related to previous findings showing that canopy $N_{[area]}$ but not canopy $N_{[\%]}$ could be detected by MTCI in grassland (Ullah et al., 2012) and by a simple ratio index in heterogeneous rangelands (Mirik et al., 2005) at various spatial scales, 300 m and 1 m, respectively. In the literature, canopy $N_{[\%]}$ is more often used to detect N state of foliage in forest while canopy $N_{[area]}$ is regularly employed in grasslands but also in crops (Clevers and Gitelson, 2013;Schlemmer et al., 2013). Our results showed that, for ENF plots, when biomass was accounted for, as in canopy $N_{[area]}$, the relationship between MTCI and canopy $N_{[area]}$ was stronger compared to canopy $N_{[\%]}$.-This suggests that biomass had an influence on and was a confounder of the MTCI-canopy N loglinear relationship.

4.4 Possible confounding factors of the MTCI canopy N relationship

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The relationships between MTCI and both canopy $N_{1\%}$ and canopy N_{fareal} were influenced by the PFT of the plots. The relationship between MTCI and canopy N_[%] was stronger for DBF and mixed plots compared to EBF and ENF plots while the opposite was true for the MTCI-canopy $N_{\text{[area]}}$ relationship. In the ongoing discussion about the mechanisms underlying the remote sensingdetection of canopy N, some authors argued that the difference in structural properties between different PFTs was a confounding factor of the observed relationship between canopy N and remote sensing data, rendering it spurious (Knyazikhin et al., 2013). Other authors suggested that the role of canopy structure as confounding factor can be explained by 410 an indirect association between canopy N and canopy structure resulting from convergent adaptive processes (Ollinger et al., 2013; Townsend et al., 2013). In this context, our analysis showed that the PFTs of the plots and the biomass had an influence on the MTCI canopy N relationship in a specific type of ecosystem, namely Mediterranean forests. Other confounding factors associated with N availability that might affect the observed relationship possibly include biomass, biomass allocation, leaf area index (LAI), water availability, soil type. The data from the forest inventory used in this analysis, i.e. the Catalonian National 415 Forest Inventory, were extensively studied, showing that water availability was the most limiting factor in this region. Water availability was positively correlated with both the $N_{\text{[area]}}$ and $N_{[\%]}$ in leaves, as well as with foliar and total above-ground biomass through MAP (Sardans et al., 2011; Sardans and Peñuelas, 2013). The MAP also influenced the PFT distribution as DBF plots were located in wetter areas than EBF plots, which were found in wetter sites than ENF plots. Regarding the influence of PFT on the foliar biomass, DBF plots had on average 45% less foliar biomass than EBF or ENF plots (Sardans and Peñuelas, 2013). This shows that canopy N_{1%1} and canopy N_{1areal} were interrelated to biomass, PFT and MAP.

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4.5 **Perspectives for future applications**

The methodology applied in this paper is different from the usual methodology implemented to detect canopy N concentration in forests. Remote sensing of N in forest canopies by hyperspectral sensors is often coupled with intensive forest sampling measurements. This method has been effective at detecting canopy N concentration locally in a vast range of environments 425 (Serrano et al., 2002;Smith et al., 2002;Townsend et al., 2003;Ollinger et al., 2008;Wang et al., 2016). Applying this technique at larger scales has already been explored. For example, Martin et al., (2008) compiled 137 field plots data from previous studies in various forest types and investigated the possibility to find a common detection algorithm. However, due to the different treatments required as well as the limited swath width associated with the high spatial resolution (from 3 m to 30 m for Hyspex airborne and Hyperion spaceborne sensors, respectively, Wang et al., 2016; Smith et al., 2003), applying imaging spectrometry 430 at a broader scale, although feasible, <u>might reveal laborious</u> might reveal time-consuming. Depending on the sensors as well as on the extent of the study area, this might involve correcting the acquired images for atmospheric influences and cloud cover as well as combining several images into a larger scale image. A recent study in northern temperate forests explored the effect of spatial resolution on canopy $N_{1\%1}$ estimation. The results showed that, although the prediction accuracy was reduced compared to what was achieved using PLS regression at higher spatial resolution, it was still possible to estimate canopy $N_{[\%]}$ with r²

435 between 0.34 and 0.81 using various vegetation indices computed from MODIS reflectance data at 500 m spatial resolution (Lepine et al., 2016). In this context, the methodology applied in this article could be a valuable alternative to explore remote sensing of canopy N detection at larger scale. Using published data from an extensive field plot inventory, we were able to relate both canopy $N_{[\%]}$ and canopy $N_{[area]}$ to MTCI at <u>1 km spatial resolution different spatial resolutions</u>. Although the relationships found were modest, our study contributes to the ongoing discussion about how to remotely sense canopy N over larger area. As 440 MTCI time series (1 km) are readily and almost globally available, it could eventually be possible to assess our approach at a broader scale in different types of biomes. The results obtained for DBF species and *Fagus sylvatica* in particular suggest that this method may be efficient at estimating canopy N in temperate forests. If the strength of the relationship between MTCI and canopy N can further be improved, this could lead to canopy N monitoring possibilities at regional scale. In this context, the new sensors OLCI, onboard Sentinel--3 satellite, and especially MSI, onboard Sentinel--2 satellite might be promising due to 445 their higher spatial resolution, from 10 to 60 m for Sentinel-Sentinel-2. They have bands well positioned to compute the MTCI vegetation index. Although the OLCI Terrestrial Chlorophyll Index (OTCI), i.e. the successor of the MTCI for the OLCI sensor, is already included in the OLCI level 2b reflectance image, no level 3 product (mosaicked over larger areas and temporally averaged hence similar to the MTCI time series used in this analysis) is available yet. In addition to more detailed remote sensing data, adding supplementary additional ground based canopy N observations could better constrain to the regression models could 450 better constrain these models as well. It would in particular be promising to use canopy N data over larger scale areas and for more diverse and globally distributed vegetation types. Obtaining reliable ground based canopy N data over larger areas and for diverse and globally distributed vegetation types would also be necessary to calibrate and validate global vegetation models, as the model performance will depend on the ground data availability and distribution. Remotely sensed canopy N estimates would also the calibration of such models. In a recent study, the global vegetation model LPJ-Guess was able to simulate the 455 differences in foliar nitrogen between different PFTs but not within one PFT (Fleischer et al., 2015). In this context, improving remotely sensed canopy N estimates for homogeneous vegetation types would be a beneficial development for such models.

5 Conclusion

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In this study, we investigated the relationship between spaceborne MTCI from ENVISAT and both canopy N_{1%1} and canopy N_[area] at regional scale in Mediterranean forests. We found significant results across all plots both when the original data were resampled to 5 km, 10 km, 15 km and 20 km and for the original spatial resolution of 1 km. The relationship between MTCI and canopy N data was also significant for some individual PFTs and species. The r^2 values were 0.32 and 0.17 for the overall relationships between MTCI and either canopy $N_{[\%]}$ or canopy $N_{[area]}$, respectively. We highlighted the differences between PFTs and both canopy $N_{[\%]}$ and canopy $N_{[area]}$: the relationship between MTCI and canopy $N_{[\%]}$ was stronger for DBF and mixed plots while canopy N_[area] was more linked to MTCI for ENF plots. Such differences in relationships between MTCI and either 465 canopy $N_{[\%]}$ or canopy $N_{[area]}$ were already observed in grasslands ecosystem. Our results showed that MTCI could be related to canopy N for some individual PFTs, indicating an influence of the PFTs on the MTCI-canopy N relationship. The methodology developed in this study could be investigated at larger scales in different types of ecosystem. While this could already be undertaken using the ENVISAT MTCI 10 years time series as it is almost globally available, ESA new Sentinel-2 satellite launched on 23 June 2015 yields reflectance data at improved spatial and temporal resolution than ENVISAT-MERIS.

470 Canopy N estimates collected through larger scales applications could be exploited in vegetation modelling studies including both the C and N cycles.

6 Data availability

The canopy data used in this study can be obtained from the TRY Plant Trait Database (https://www.try-db.org/TryWeb/Home.php, dataset 91) or by directly contacting the authors.

475 **7** Appendix A

This appendix presents the inter-annual variation of canopy $N_{[\%]}$ (Fig. A 1), the correlation matrices for DBF (Fig. A2), EBF (Fig. A3) and ENF plots (Fig. A3) as well as the tables representing the allometric relationships between foliar biomass and diameter at breast height (DBH, Table A 1), the number of PFT (Table A 2), the number of species (Tables A 3) and the number of sampling years (Table A 4) per resampled pixel, by pixel spatial resolution

480 8 Competing interest

The authors declare that they have no conflict of interest.

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10 References

495

500

515

Bierkens, M. F. P., Finke, P. A., and De Willigen, P.: Upscalling and Downscalling Methods for Environmental Research, Developments in Plant and Soil Sciences, Kluwer Academic Publishers, Dordrecht, 2000.

Bontemps, S., Defourny, P., Van Bogaert, E., Arino, O., Kalogirou, V., and Ramos Perez, J.: Globcover 2009 Products Description and Validation Report, UCLouvain & ESA Team, 53 pp., available at: http://due.esrin.esa.int/files/GLOBCOVER2009 Validation Report 2002.2002.pdf, last access 2011 November 2016, 2011.

Boyd, D. S., Almond, S., Dash, J., Curran, P. J., and Hill, R. A.: Phenology of vegetation in southern england from envisat meris terrestrial chlorophyll index (MTCI) data, International Journal of Remote Sensing, 32, 8421 8447, doi:10.1080/01431161.2010.542194. 2011.

Boyd, D. S., Almond, S., Dash, J., Curran, P. J., Hill, R. A., and Foody, G. M.: Evaluation of envisat MERIS terrestrial chlorophyll index based models for the estimation of terrestrial gross primary productivity, IEEE Geoscience and Remote Sensing Letters, 9, 457–461, 2012.

Cho, M. A., Ramoelo, A., Debba, P., Mutanga, O., Mathieu, R., van Deventer, H., and Ndlovu, N.: Assessing the effects
of subtropical forest fragmentation on leaf nitrogen distribution using remote sensing data, Landscape Ecology, 28, 1479-1491, doi:10.1007/s10980-013-9908-7, 2013.

Clevers, J. G. P. W., and Gitelson, A. A.: Remote estimation of crop and grass chlorophyll and nitrogen content using rededge bands on sentinel-2 and-3, International Journal of Applied Earth Observation and Geoinformation, 23, 344-351, doi:10.1016/j.jag.2012.10.008, 2013.

510 Crabbe, R. A., Dash, J., Rodriguez Galiano, V. F., Janous, D., Pavelka, M., and Marek, M. V.: Extreme warm temperatures alter forest phenology and productivity in Europe, Science of The Total Environment, 563-564, 486-495, http://dx.doi.org/10.1016/j.scitotenv.2016.04.124, 2016.

Curran, P. J., and Dash, J.: Algorithm Theoretical basis document ATBD 2.22 Chlorophyll Index, University of Southampton, Southampton, 41 pp., available at: https://earth.esa.int/documents/700255/2042855/MERIS_ATBD_2.22_v1.2++2005.pdf, last access 11 November 16, 2005.

Curran, P. J., Dash, J., Lankester, T., and Hubbard, S.: Global composites of the MERIS Terrestrial Chlorophyll Index, International Journal of Remote Sensing, 28, 3757-3758, 10.1080/01431160600639685, 2007.

Dash, J., and Curran, P. J.: The MERIS terrestrial chlorophyll index, International Journal of Remote Sensing, 25, 5403-5413, doi:10.1080/0143116042000274015, 2004.

Dash, J., and Curran, P. J.: Evaluation of the MERIS terrestrial chlorophyll index (MTCI), Advances in Space Research,
39, 100-104, doi:10.1016/j.asr.2006.02.034, 2007.

Dash, J., and Vuolo, F.: Algorithm Theoretical Basis Document OLCI Terrestrial Chlorophyll Index (OTCI), UniversityofSouthampton,21pp.,availableat:

https://sentinel.esa.int/documents/247904/349589/OLCI L247902 ATBD OLCI Terrestrial Chlorophyll Index.pdf, last access 8 March 2018, 2010.

Drusch, M., Del Bello, U., Carlier, S., Colin, O., Fernandez, V., Gascon, F., Hoersch, B., Isola, C., Laberinti, P., Martimort, P., Meygret, A., Spoto, F., Sy, O., Marchese, F., and Bargellini, P.: Sentinel-2: ESA's Optical High-Resolution Mission for GMES Operational Services, Remote Sensing of Environment, 120, 25-36, doi:10.1016/j.rse.2011.11.026, 2012.

Evans, J. R.: Photosynthesis and nitrogen relationships in leaves of C3 plants, Oecologia, 78, 9-19, doi:10.1007/BF00377192, 1989.

ESA: GlobCover 2009 (Global Land Cover Map), ESA and UCLouvain, available at: http://due.esrin.esa.int/page_globcover.php, last access: 11 November 2016, 2010.

Ferwerda, J. G., Skidmore, A. K., and Mutanga, O.: Nitrogen detection with hyperspectral normalized ratio indices across multiple plant species, International Journal of Remote Sensing, 26, 4083-4095, 10.1080/01431160500181044, 2005.

535 Filella, I., and Penuelas, J.: The red edge position and shape as indicators of plant chlorophyll content, biomass and hydric status, International Journal of Remote Sensing, 15, 1459-1470, http://dx.doi.org/10.1080/01431169408954177, 1994.

Filella, I., Serrano, L., Serra, J., and Penuelas, J.: Evaluating wheat nitrogen status with canopy reflectance indices and discriminant analysis, Crop Science, 35, 1400-1405, doi:10.2135/cropsci1995.0011183X003500050023x, 1995.

Fleischer, K., Wårlind, D., van der Molen, M. K., Rebel, K. T., Arneth, A., Erisman, J. W., Wassen, M. J., Smith, B.,
Gough, C. M., Margolis, H. A., Cescatti, A., Montagnani, L., Arain, A., and Dolman, A. J.: Low historical nitrogen deposition effect on carbon sequestration in the boreal zone, Journal of Geophysical Research: Biogeosciences, 120, 2542-2561, 10.1002/2015JG002988, 2015.

Foody, G. M., and Dash, J.: Discriminating and mapping the C3 and C4 composition of grasslands in the northern Great Plains, USA, Ecological Informatics, 2, 89–93, doi:10.1016/j.ecoinf.2007.03.009, 2007.

Glenn, E. P., Huete, A. R., Nagler, P. L., and Nelson, S. G.: Relationship between remotely-sensed vegetation indices, canopy attributes and plant physiological processes: What vegetation indices can and cannot tell us about the landscape, Sensors, 8, 2136-2160, doi:10.3390/s8042136, 2008.

González, J. R., and Pukkala, T.: Characterization of forest fires in Catalonia (north-east Spain), European Journal of Forest Research, 126, 421-429, doi:10.1007/s10342-006-0164-0, 2007.

Gracia, C., Ibàñez, J. J., Burriel, J. A., Mata, T., and Vayreda, J.: Inventari Ecològic i Forestal de Catalunya. Mètodes, CREAF, Bellaterra, available at: http://www.creaf.uab.es/iefc/pub/Metodes/index.htm, last access 11 November 2016, 2004.

Green, D. S., Erickson, J. E., and Kruger, E. L.: Foliar morphology and canopy nitrogen as predictors of light-use efficiency in terrestrial vegetation, Agricultural and Forest Meteorology, 115, 163-171, http://dx.doi.org/10.1016/S0168-1923(02)00210-1, 2003.

555

550

Hansen, P. M., and Schjoerring, J. K.: Reflectance measurement of canopy biomass and nitrogen status in wheat crops using normalized difference vegetation indices and partial least squares regression, Remote Sensing of Environment, 86, 542-553, doi:10.1016/S0034-4257(03)00131-7, 2003.

Harris, A., and Dash, J.: The potential of the MERIS Terrestrial Chlorophyll Index for carbon flux estimation, Remote Sensing of Environment, 114, 1856–1862, doi:10.1016/j.rse.2010.03.010, 2010.

Harris, A., and Dash, J.: A new approach for estimating northern peatland gross primary productivity using a satellitesensor derived chlorophyll index, Journal of Geophysical Research: Biogeosciences, 116, doi:10.1029/2011JG001662, 2011.

Huber, S., Kneubühler, M., Psomas, A., Itten, K., and Zimmermann, N. E.: Estimating foliar biochemistry from hyperspectral data in mixed forest canopy, Forest Ecology and Management, 256, 491-501, doi:10.1016/j.foreco.2008.05.011, 2008.

Karssenberg, D., Schmitz, O., Salamon, P., de Jong, K., and Bierkens, M. F. P.: A software framework for construction of process-based stochastic spatio-temporal models and data assimilation, Environmental Modelling and Software, 25, 489-502, doi:10.1016/j.envsoft.2009.10.004, 2010.

Kergoat, L., Lafont, S., Arneth, A., Le Dantec, V., and Saugier, B.: Nitrogen controls plant canopy light-use efficiency in temperate and boreal ecosystems, Journal of Geophysical Research: Biogeosciences, 113, doi:10.1029/2007JG000676, 2008.

Knyazikhin, Y., Schull, M. A., Stenberg, P., Mõttus, M., Rautiainen, M., Yang, Y., Marshak, A., Latorre Carmona, P.,
Kaufmann, R. K., Lewis, P., Disney, M. I., Vanderbilt, V., Davis, A. B., Baret, F., Jacquemoud, S., Lyapustin, A., and Myneni,
R. B.: Hyperspectral remote sensing of foliar nitrogen content, Proceedings of the National Academy of Sciences, 110, E185–
E192, 10.1073/pnas.1210196109, 2013.

575

560

565

570

Kumar, L., Schmidt, K., Dury, S., and Skidmore, A.: Imaging Spectrometry and Vegetation Science, in: Imaging Spectrometry: Basic Principles and Prospective Applications, edited by: Meer, F. D. v. d., and Jong, S. M. D., Springer Netherlands, Dordrecht, 111-155, 2006.

Lana, X., and Burgueño, A.: Spatial and temporal characterization of annual extreme droughts in Catalonia (Northeast Spain), International Journal of Climatology, 18, 93-110, 1998.

Lepine, L. C., Ollinger, S. V., Ouimette, A. P., and Martin, M. E.: Examining spectral reflectance features related to foliar
 nitrogen in forests: Implications for broad-scale nitrogen mapping, Remote Sensing of Environment, 173, 174-186, doi:10.1016/j.rse.2015.11.028, 2016.

Li, F., Miao, Y., Feng, G., Yuan, F., Yue, S., Gao, X., Liu, Y., Liu, B., Ustin, S. L., and Chen, X.: Improving estimation of summer maize nitrogen status with red edge-based spectral vegetation indices, Field Crops Research, 157, 111-123, doi:10.1016/j.fcr.2013.12.018, 2014.

585 Liu, D., Ogaya, R., Barbeta, A., Yang, X., and Peñuelas, J.: Contrasting impacts of continuous moderate drought and episodic severe droughts on the aboveground-biomass increment and litterfall of three coexisting Mediterranean woody species, Global Change Biology, 21, 4196-4209, doi:10.1111/gcb.13029, 2015.

Martin, M. E., Plourde, L. C., Ollinger, S. V., Smith, M. L., and McNeil, B. E.: A generalizable method for remote sensing of canopy nitrogen across a wide range of forest ecosystems, Remote Sensing of Environment, 112, 3511-3519, doi:10.1016/j.rse.2008.04.008, 2008.

Mirik, M., Norland, J. E., Crabtree, R. L., and Biondini, M. E.: Hyperspectral one-meter-resolution remote sensing in Yellowstone National Park, Wyoming: I. Forage nutritional values, Rangeland Ecology and Management, 58, 452-458, doi:10.2111/04-17.1, 2005.

NEODC - NERC Earth Observation Data Centre: http://neodc.nerc.ac.uk/, access: 06/02/2015, 2015.

595 Ollinger, S. V., and Smith, M. L.: Net primary production and canopy nitrogen in a temperate forest landscape: An analysis using imaging spectroscopy, modeling and field data, Ecosystems, 8, 760-778, doi:10.1007/s10021-005-0079-5, 2005.

Ollinger, S. V., Richardson, A. D., Martin, M. E., Hollinger, D. Y., Frolking, S. E., Reich, P. B., Plourde, L. C., Katul, G. G., Munger, J. W., Oren, R., Smith, M. L., Paw U, K. T., Bolsta, P. V., Cook, B. D., Day, M. C., Martin, T. A., Monson, R. K., and Schmid, H. P.: Canopy nitrogen, carbon assimilation, and albedo in temperate and boreal forests: Functional relations and

600

610

615

potential climate feedbacks, Proceedings of the National Academy of Sciences of the United States of America, 105, 19336-19341, doi:10.1073/pnas.0810021105., 2008.

Ollinger, S. V.: Sources of variability in canopy reflectance and the convergent properties of plants, New Phytologist, 189, 375-394, doi:10.1111/j.1469-8137.2010.03536.x, 2011.

Ollinger, S. V., Reich, P. B., Frolking, S., Lepine, L. C., Hollinger, D. Y., and Richardson, A. D.: Nitrogen cycling, forest
 canopy reflectance, and emergent properties of ecosystems, Proceedings of the National Academy of Sciences, 110, E2437, 10.1073/pnas.1304176110, 2013.

Pacheco-Labrador, J., González-Cascón, R., Pilar Martín, M., and Riaño, D.: Understanding the optical responses of leaf nitrogen in mediterranean holm oak (Quercus ilex) using field spectroscopy, International Journal of Applied Earth Observation and Geoinformation, 26, 105-118, doi:10.1016/j.jag.2013.05.013, 2014.

Peng, Y., and Gitelson, A. A.: Application of chlorophyll related vegetation indices for remote estimation of maize productivity, Agricultural and Forest Meteorology, 151, 1267–1276, doi:10.1016/j.agrformet.2011.05.005, 2011.

Peñuelas, J., Gamon, J. A., Fredeen, A. L., Merino, J., and Field, C. B.: Reflectance indices associated with physiological changes in nitrogen- and water-limited sunflower leaves, Remote Sensing of Environment, 48, 135-146, doi:10.1016/0034-4257(94)90136-8, 1994.

<u>R Development Core Team: R: A Language and Environment for Statistical Computing, computer program, R Foundation</u> for Statistical Computing, Vienna, Austria, 2014.

Ramoelo, A., Skidmore, A. K., Cho, M. A., Schlerf, M., Mathieu, R., and Heitkönig, I. M. A.: Regional estimation of savanna grass nitrogen using the red-edge band of the spaceborne rapideye sensor, International Journal of Applied Earth Observation and Geoinformation, 19, 151-162, doi:10.1016/j.jag.2012.05.009, 2012.

620

Reich, P. B., Walters, M. B., Kloeppel, B. D., and Ellsworth, D. S.: Different photosynthesis-nitrogen relations in deciduous hardwood and evergreen coniferous tree species, Oecologia, 104, 24-30, doi:10.1007/BF00365558, 1995.

Reich, P. B., Walters, M. B., and Ellsworth, D. S.: From tropics to tundra: Global convergence in plant functioning, Proceedings of the National Academy of Sciences of the United States of America, 94, 13730-13734, doi:10.1073/pnas.94.25.13730, 1997. 625 Reich, P. B., Ellsworth, D. S., Walters, M. B., Vose, J. M., Gresham, C., Volin, J. C., and Bowman, W. D.: Generality of leaf trait relationships: A test across six biomes, Ecology, 80, 1955-1969, doi:10.2307/176671 1999.

Reich, P. B.: Key canopy traits drive forest productivity, Proceedings of the Royal Society B: Biological Sciences, 279, 2128-2134, doi:10.1098/rspb.2011.2270, 2012.

Rodriguez-Galiano, V. F., Dash, J., and Atkinson, P. M.: Characterising the land surface phenology of Europe using
decadal MERIS data, Remote Sensing, 7, 9390-9409, doi:10.3390/rs70709390, 2015.

Sardans, J., Rivas-Ubach, A., and Peñuelas, J.: Factors affecting nutrient concentration and stoichiometry of forest trees in Catalonia (NE Spain), Forest Ecology and Management, 262, 2024-2034, doi:10.1016/j.foreco.2011.08.019, 2011.

Sardans, J., and Peñuelas, J.: Tree growth changes with climate and forest type are associated with relative allocation of nutrients, especially phosphorus, to leaves and wood, Global Ecology and Biogeography, 22, 494-507, doi:10.1111/geb.12015, 2013.

635

640

Sardans, J., and Peñuelas, J.: Trees increase their P: N ratio with size, Global Ecology and Biogeography, 24, 147-156, doi:10.1111/geb.12231, 2015.

Schlemmer, M., Gitelson, A., Schepers, J., Ferguson, R., Peng, Y., Shanahan, J., and Rundquist, D.: Remote estimation of nitrogen and chlorophyll contents in maize at leaf and canopy levels, International Journal of Applied Earth Observation and Geoinformation, 25, 47-54, doi:10.1016/j.jag.2013.04.003, 2013.

Schlerf, M., Atzberger, C., Hill, J., Buddenbaum, H., Werner, W., and Schüler, G.: Retrieval of chlorophyll and nitrogen in Norway spruce (Picea abies L. Karst.) using imaging spectroscopy, International Journal of Applied Earth Observation and Geoinformation, 12, 17-26, doi:10.1016/j.jag.2009.08.006, 2010.

Serrano, L., Peñuelas, J., and Ustin, S. L.: Remote sensing of nitrogen and lignin in Mediterranean vegetation from AVIRIS
 data: Decomposing biochemical from structural signals, Remote Sensing of Environment, 81, 355-364, doi:10.1016/S0034-4257(02)00011-1, 2002.

Smith, B., Wårlind, D., Arneth, A., Hickler, T., Leadley, P., Siltberg, J., and Zaehle, S.: Implications of incorporating N cycling and N limitations on primary production in an individual-based dynamic vegetation model, Biogeosciences, 11, 2027-2054, doi:10.5194/bg-11-2027-2014, 2014.

650 Smith, M. L., and Martin, M. E.: A plot-based method for rapid estimation of forest canopy chemistry, Canadian Journal of Forest Research, 31, 549-555, doi:10.1139/x00-187, 2001.

Smith, M. L., Ollinger, S. V., Martin, M. E., Aber, J. D., Hallett, R. A., and Goodale, C. L.: Direct estimation of aboveground forest productivity through hyperspectral remote sensing of canopy nitrogen, Ecological Applications, 12, 1286-1302, doi:10.2307/3099972, 2002.

655 Smith, M. L., Martin, M. E., Plourde, L., and Ollinger, S. V.: Analysis of hyperspectral data for estimation of temperate forest canopy nitrogen concentration: Comparison between an airborne (AVIRIS) and a spaceborne (Hyperion) sensor, IEEE Transactions on Geoscience and Remote Sensing, 41, 1332-1337, doi:10.1109/TGRS.2003.813128, 2003. Stein, B. R., Thomas, V. A., Lorentz, L. J., and Strahm, B. D.: Predicting macronutrient concentrations from loblolly pine leaf reflectance across local and regional scales, GIScience and Remote Sensing, 51, 269-287, doi:10.1080/15481603.2014.912875, 2014.

660

Sullivan, F. B., Ollinger, S. V., Martin, M. E., Ducey, M. J., Lepine, L. C., and Wicklein, H. F.: Foliar nitrogen in relation to plant traits and reflectance properties of New Hampshire forests, Canadian Journal of Forest Research, 43, 18-27, doi:10.1139/cjfr-2012-0324, 2013.

Tian, Y. C., Yao, X., Yang, J., Cao, W. X., Hannaway, D. B., and Zhu, Y.: Assessing newly developed and published
 vegetation indices for estimating rice leaf nitrogen concentration with ground- and space-based hyperspectral reflectance, Field
 Crops Research, 120, 299-310, doi:10.1016/j.fcr.2010.11.002, 2011.

Townsend, P. A., Foster, J. R., Chastain Jr, R. A., and Currie, W. S.: Application of imaging spectroscopy to mapping canopy nitrogen in the forest of the central Appalachian mountains using hyperion and AVIRIS, IEEE Transactions on Geoscience and Remote Sensing, 41, 1347-1354, doi:10.1109/TGRS.2003.813205, 2003.

670

675

685

Townsend, P. A., Serbin, S. P., Kruger, E. L., and Gamon, J. A.: Disentangling the contribution of biological and physical properties of leaves and canopies in imaging spectroscopy data, Proceedings of the National Academy of Sciences of the United States of America, 110, 10.1073/pnas.1300952110, 2013.

Ullah, S., Si, Y., Schlerf, M., Skidmore, A. K., Shafique, M., and Iqbal, I. A.: Estimation of grassland biomass and nitrogen using MERIS data, International Journal of Applied Earth Observation and Geoinformation, 19, 196-204, doi:10.1016/j.jag.2012.05.008, 2012.

Vilà-Cabrera, A., Coll, L., Martínez-Vilalta, J., and Retana, J.: Forest management for adaptation to climate change in the Mediterranean basin: A synthesis of evidence, Forest Ecology and Management, 407, 16-22, doi:10.1016/j.foreco.2017.10.021, 2018.

Vuolo, F., Dash, J., Curran, P. J., Lajas, D., and Kwiatkowska, E.: Methodologies and uncertainties in the use of the
 terrestrial chlorophyll index for the sentinel-3 mission, Remote Sensing, 4, 1112-1133, 10.3390/rs4051112, 2012.

Wang, Z., Wang, T., Darvishzadeh, R., Skidmore, A. K., Jones, S., Suarez, L., Woodgate, W., Heiden, U., Heurich, M., and Hearne, J.: Vegetation indices for mapping canopy foliar nitrogen in a mixed temperate forest, Remote Sensing, 8, doi:10.3390/rs8060491, 2016.

Wickham, H.: ggplot2: Elegant Graphics for Data Analysis, Springer-Verlag, New York, 2009.

Wright, I. J., Reich, P. B., Westoby, M., Ackerly, D. D., Baruch, Z., Bongers, F., Cavender-Bares, J., Chapin, T.,
Cornelissen, J. H. C., Diemer, M., Flexas, J., Garnier, E., Groom, P. K., Gulias, J., Hikosaka, K., Lamont, B. B., Lee, T., Lee,
W., Lusk, C., Midgley, J. J., Navas, M.-L., Niinemets, U., Oleksyn, J., Osada, N., Poorter, H., Poot, P., Prior, L., Pyankov, V.
I., Roumet, C., Thomas, S. C., Tjoelker, M. G., Veneklaas, E. J., and Villar, R.: The worldwide leaf economics spectrum, Nature,
428, 821-827, doi:10.1038/nature02403, 2004.

690 Yao, X., Zhu, Y., Tian, Y., Feng, W., and Cao, W.: Exploring hyperspectral bands and estimation indices for leaf nitrogen accumulation in wheat, International Journal of Applied Earth Observation and Geoinformation, 12, 89-100, http://dx.doi.org/10.1016/j.jag.2009.11.008, 2010.

Zaehle, S., and Friend, A.: Carbon and nitrogen cycle dynamics in the O - CN land surface model: 1. Model description, site - scale evaluation, and sensitivity to parameter estimates, Global Biogeochemical Cycles, 24, doi:10.1029/2009GB003521,
2010.





Figure 3. Map showing the forest plots (n = 846) location in the region of Catalonia, north eastern Spain. <u>Two zoom windows are</u> included showing the density of the plots, one with high density and one with low density, relatively to the MTCI 1 km pixel grid. DBF = Deciduous Broadleaf Forest, EBF = Evergreen Broadleaf Forest, ENF = Evergreen Needleleaf Forest, mixed = mixed forest.



Figure 4. Boxplot of (a) canopy nitrogen (N) concentration (canopy N_[%], %N) for Deciduous Broadleaf Forest plots (DBF, n = 80), Evergreen Broadleaf Forest plots (EBF, n = 186), Evergreen Needleleaf Forest plots (ENF, n = 564) and mixed forest plots (mixed, n = 16); (b) canopy N content (canopy N_[area], g m²) for Deciduous Broadleaf Forest plots (DBF, n = 80), Evergreen Broadleaf Forest plots (EBF, n = 186), Evergreen Needleleaf Forest plots (ENF, n = 563) and mixed forest plots (mixed, n = 12);



Figure 5. The upper right part of this figure shows the Pearson correlation matrix between canopy $N_{[\%]}$ (%N), canopy $N_{[area]}$ (g m⁻²) and foliar biomass (g m⁻²) variables for the whole dataset, n = 841. The diagonal presents the histograms of the variables on the x-axis, while the y-axis represents the number of counts. The lower left part of this figure represents the scatterplots between the variables. PFT = Plant Functional Type, DBF = Deciduous Broadleaf Forest, EBF = Evergreen Broadleaf Forest, ENF = Evergreen Needleleaf Forest, mixed = mixed forest.



Figure 6. Scatterplot between the MERIS Terrestrial Chlorophyll Index (MTCI) (-) and canopy nitrogen concentration (canopy $N_{[\%]}$, %N) after resampling the datasets to 20 km-spatial resolution (n = 204).



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Figure 7. Scatterplot and <u>log-</u>linear regression line between the MERIS Terrestrial Chlorophyll Index (MTCI) (-) and canopy nitrogen (N) concentration (canopy $N_{[\%]}$, %N) for (a) whole dataset (n = 846); (b) Deciduous Broadleaf Forest plots (DBF, n = 80); (c) Evergreen Broadleaf Forest plots (EBF, n = 186); (d) Evergreen Needleleaf Forest plots (ENF, n = 564); (e) mixed forest plots (n = 16). PFT = Plant functional type. The grey shading represents the prediction intervals (95 %). Canopy $N_{[\%]}$ variable was log transformed to fulfil linear model assumptions.



Figure 8. Scatterplot and log-linear regression line between the MERIS Terrestrial Chlorophyll Index (MTCI) (-) and canopy N content (canopy $N_{\text{[area]}}$, g m⁻²) for (a) whole dataset (n = 841); (b) Deciduous Broadleaf Forest plots (DBF, n = 80); (c) Evergreen Broadleaf Forest plots (EBF, n = 186); (d) Evergreen Needleleaf Forest plots (ENF, n = 563); (e) mixed forest plots (n = 12). PFT = Plant functional type. The grey shading represents the prediction intervals (95 %). Canopy $N_{\text{[area]}}$ variable was log transformed to fulfil linear models assumptions.

Table 1. Descriptive analysis of canopy nitrogen (N) concentration ($N_{[\%]}$, g 100g⁻¹), foliar biomass (g m⁻²) and canopy N content ($N_{[area]}$, g m⁻²) by tree species. PFT = Plant Functional Type, DBF = Deciduous Broadleaf Forest, EBF = Evergreen Broadleaf Forest, ENF = Evergreen Needleleaf Forest, mixed = mixed forest, min = minimum, max = maximum, mean = average, sd = standard deviation, ^a codominant plots refer to the plots where two tree species were dominant in the canopy, ^b foliar biomass data was lacking for five of the plots. Foliar biomass and canopy N content statistics are thus measured on a restricted number of plots.

		Abundanaa	Canopy $N_{[\%]}$ (g 100g ⁻¹)				Foliar biomass (g m ⁻²)			Canopy N _[area] (g m ⁻²)					
Species	PFT	Number of plots	Abundance (% of total number of plots)	min	max	mean	sd	min	max	mean	sd	min	max	mean	sd
Castanea sativa	DBF	14	1.7	1.62	2.81	2.08	0.36	18.13	425.90	203.46	123.49	0.40	11.99	4.25	2.89
Fagus sylvatica	DBF	15	1.8	1.22	3.13	2.28	0.61	49.94	279.86	173.54	68.70	1.21	7.40	3.96	1.95
Pinus halepensis	ENF	240	28.4	0.56	1.57	0.90	0.19	9.58	827.80	197.23	145.54	0.09	7.29	1.77	1.33
Pinus nigra	ENF	37	4.4	0.56	1.28	0.89	0.19	32.25	923.98	294.29	224.32	0.23	8.87	2.67	2.18
Pinus pinaster	ENF	5	0.6	0.82	1.08	0.93	0.13	271.75	718.87	501.67	211.53	2.30	7.69	4.75	2.25
Pinus pinea	ENF	5	0.6	0.75	1.06	0.95	0.14	103.28	275.50	179.74	66.80	1.08	2.91	1.71	0.75
Pinus sylvestris	ENF	198	23.4	0.67	2.14	1.11	0.20	10.48	828.63	326.44	181.20	0.10	12.86	3.65	2.22
Pinus uncinata	ENF	69	8.2	0.46	1.33	0.87	0.19	183.59	1744.50	687.22	345.21	1.41	16.97	5.92	3.25
Quercus canariensis	DBF	3	0.4	1.97	2.78	2.25	0.46	122.11	197.85	160.32	37.87	2.41	5.51	3.71	1.61
Quercus faginea	DBF	4	0.5	1.49	2.11	1.82	0.31	10.34	419.14	233.47	187.01	0.17	8.83	4.64	4.09
Quercus humilis	DBF	9	1.1	1.53	3.11	2.41	0.42	56.12	337.33	142.65	92.11	1.21	8.64	3.33	2.19
Quercus cerriodes	DBF	17	2.0	1.44	2.80	2.07	0.37	12.97	834.68	262.24	237.49	0.29	15.42	5.06	4.31
Quercus ilex	EBF	160	18.9	0.81	2.87	1.32	0.26	16.63	1033.31	378.23	238.61	0.22	16.61	4.95	3.23
Quercus petraea	DBF	17	2.0	1.37	2.70	2.21	0.41	20.45	741.42	279.96	229.78	0.32	15.37	5.98	4.66
Quercus suber	EBF	23	2.7	1.25	2.08	1.55	0.21	26.26	219.05	110.49	55.65	0.40	4.34	1.72	0.96
Codominant ^a	mixed	30 (25) ^b	3.5	0.92	2.54	1.45	0.41	23.45	342.58	153.70	77.39	0.33	5.74	2.06	1.02

<u> </u>	Number of plots per pixel					
Spatial resolution (km)	min	max	mean	sd		
5	1	6	1.44	0.77		
10	1	11	2.19	1.53		
15	1	15	3.11	2.59		
20	1	22	4.09	3.74		

Table 2. Descriptive statistics of the number of plots per pixel, for different spatial resolution (km, pixel length). min = minimum, max = maximum, mean = average, sd = standard deviation.

Table 3. Observed log-linear regression equations between the MERIS Terrestrial Chlorophyll Index (MTCI) (-) and canopy nitrogen concentration (CN_[%], %N) for different subgroups. Number of plots (n), determination coefficient (r²), p-value and Relative Root Mean Square Error of cross-validation (RRMEcv). PFT = Plant Functional type, DBF = Deciduous Broadleaf Forest, EBF = Evergreen Broadleaf Forest, ENF = Evergreen Needleleaf Forest, mixed = mixed forest.

group	n	log-linear regression	95% confidence interval intercept	95% confidence interval slope	\mathbf{r}^2	p-value	RRMSEcv
overall	846	$MTCI = 2.18 + 0.79 \log(CN_{[\%]})$	[2.15, 2.20]	[0.71, 0.87]	0.32	< 0.000	17.0
DBF	80	$MTCI = 2.07 + 0.95 \log(CN_{[\%]})$	[1.78, 2.36]	[0.59, 1.32]	0.2 <mark>54</mark>	< 0.000	12.7
EBF	186	$MTCI = 2.39 + 0.29 \log(CN_{[\%]})$	[2.31, 2.48]	[0.04, 0.54]	0.0 <u>32</u>	0.021	12.4
ENF	564	$MTCI = 2.13 + 0.61 \log(CN_{[\%]})$	[2.10, 2.17]	[0.46, 0.76]	0.10	< 0.000	19.2
mixed	16	$MTCI = 2.05 + 1.35 \log(CN_{[\%]})$	[1.63, 2.46]	[0.53, 2.17]	0.4 7<u>4</u>	0.003	12.4

Table 4. Observed log-linear regressions equations between the MERIS Terrestrial Chlorophyll Index (MTCI) (-) and canopy nitrogen content (CN_[area], g m⁻²) for different subgroups. Number of plots (n), determination coefficient (r²), p-value and Relative Root Mean Square Error of cross-validation (RRMEcv). PFT = Plant Functional type, DBF = Deciduous Broadleaf Forest, EBF = Evergreen Broadleaf Forest, ENF = Evergreen Needleleaf Forest, mixed = mixed forest.

group	n	log-linear regression	95% confidence interval intercept	95% confidence interval slope	\mathbf{r}^2	p-value	RRMSEcv
Overall	841	$MTCI = 2.08 + 0.20 \log(CN_{[area]})$	[2.04, 2.12]	[0.17, 0.23]	0.17	< 0.000	18.7
DBF	80	$MTCI = 2.72 + 0.06 \log(CN_{[area]})$	[2.58, 2.87]	[-0.04, 0.15]	0.0 <mark>2<u>03</u></mark>	0.263	14.7
EBF	186	$MTCI = 2.39 + 0.07 \log(CN_{[area]})$	[2.32, 2.46]	[0.02, 0.12]	0.04	0.005	12.4
ENF	563	$MTCI = 1.94 + 0.20 \log(CN_{[area]})$	[1.91, 1.99]	[0.17, 0.24]	0. <u>219</u>	< 0.000	18.2
mixed	12	$MTCI = 2.43 + 0.34 \log(CN_{[area]})$	[2.05, 2.82]	[-0.26, 0.95]	0. 14<u>05</u>	0.236	12.8

11 Appendix



Figure A1. Inter-annual variation of canopy N_[%] (%N) for each month included in the analysis. The numbers 6 – 10 (right side of the figure, row numbers) refer to the month of June, July, August, September and October, respectively. DBF = Deciduous Broadleaf Forest, EBF = Evergreen Broadleaf Forest, ENF = Evergreen Needleleaf Forest, mixed = mixed forest. Each point represents an observation at a forest plot. Note that the forest plots were not sampled multiple times, hence the inter-annual variation encompasses both temporal variation and spatial variation.



Figure A 2. The upper right part of this figure shows the Pearson correlation matrix between canopy $N_{[\%]}$ (%N), canopy $N_{[area]}$ (g m-2) and foliar biomass (g m-2) variables for deciduous broadleaf forest plots (DBF), n = 80. The diagonal presents the histogram of the variable on the x-axis, while the y-axis represents the number of counts. The lower left part of this figure represents the scatterplots between the variables.



Figure A 3. The upper right part of this figure shows the Pearson correlation matrix between canopy $N_{[\%]}$ (%N), canopy $N_{[area]}$ (g m⁻²) and foliar biomass (g m⁻²) variables for evergreen broadleaf forest (EBF) plots, n = 186. The diagonal presents the histogram of the variable on the x-axis, while the y-axis represents the number of counts. The lower left part of this figure represents the scatterplots between the variables.



Figure A 4. The upper right part of this figure shows the Pearson correlation matrix between canopy $N_{[\%]}$ (%N), canopy $N_{[area]}$ (g m⁻²) and foliar biomass (g m⁻²) variables for evergreen needleleaf forest (ENF) plots, n = 563. The diagonal presents the histogram of the variable on the x-axis, while the y-axis represents the number of counts. The lower left part of this figure represents the scatterplots between the variables.

Species	Foli	iar biomas	$as = a \cdot D$	BH⁵
	а	b	n	r ²
Castanea sativa	0.032	1.669	86	0.49
Fagus sylvatica	0.026	1.546	285	0.66
Pinus halepensis	0.037	1.656	2420	0.65
Pinus nigra	0.022	1.870	1641	0.65
Pinus pinaster	0.034	1.848	169	0.67
Pinus pinea	0.014	2.029	335	0.72
Pinus sylvestris	0.036	1.651	2755	0.66
Pinus uncinata	0.087	1.410	770	0.62
Quercus canariensis	0.120	1.322	36	0.57
Quercus faginea	0.197	0.943	170	0.40
Quercus humilis	0.047	1.462	595	0.59
Quercus cerrioides	0.023	1.805	138	0.73
Quercus ilex	0.063	1.576	2151	0.60
Quercus petraea	0.014	1.888	121	0.73
Quercus suber	0.026	1.446	314	0.55

Table A1. Allometric relationships between foliar biomass and DBH for the different species included in this analysis. DBH = Diameter at breast height (cm). Adapted from (Sardans and Peñuelas, 2015).

805 Table A 2. Descriptive statistics of the number of plant functional types (PFT) per pixel, by pixel spatial resolution (km). min = minimum, max = maximum, mean = average, sd = standard deviation.

Spotial resolution (Irm)		Number of I	PFT per pixel	
Spatial resolution (kin)	min	max	mean	sd
5	1	3	1.08	0.29
10	1	4	1.22	0.48
15	1	4	1.34	0.61
20	1	4	1.45	0.69

Table A 3. Descriptive statistics of the number of species per pixel, by pixel spatial resolution (km). min = minimum,810max = maximum, mean = average, sd = standard deviation.

Spatial resolution (Irm)	Number of species per pixel						
Spatial resolution (km)	min	max	mean	sd			
5	1	4	1.14	0.41			
10	1	4	1.38	0.67			
15	1	4	1.58	0.85			
20	1	6	1.79	1.07			

Table A 4. Descriptive statistics of the number of sampling years per pixel, by pixel spatial resolution (km). min = minimum,815max = maximum, mean = average, sd = standard deviation.

Spatial resolution (Irm)	Number of sampling years per pixel						
Spatial resolution (km)	min	max	mean	sd			
5	1	2	1.02	0.15			
10	1	3	1.07	0.26			
15	1	3	1.10	0.33			
20	1	3	1.14	0.40			