

**Response to referee comments and suggestions on bg-2018-521 by N. Löbs et al.:
“Microclimatic conditions and water content fluctuations experienced by epiphytic bryophytes in an Amazonian rain forest”**

Dear Professor Bahn,

we would like to thank you and the reviewer for the manuscript evaluation and the comments, which helped to improve it once again. We indeed highly appreciate the efforts made by Maaïke Bader and have already included her in the acknowledgement section. Below we respond with a point-by-point explanation to the comments from the peer-reviewer with our responses in blue color following every comment. At the end of the comments we provide the manuscript and the supplement with all changes being marked.

Sincerely,

Nina Löbs, on behalf of the co-authors.

Comments on the text:

Black text shows the original referee comment, and blue text shows the response of the authors and the explicit changes in the revised text. The figure and table numbers refer to the revised manuscript.

Referee report #1

Maaïke Bader, 04 August 2020

Dear authors,

Thank you for once again reviewing your manuscript and for seriously and effectively addressing my concerns. I think that you have now solved the main problem in the previous versions, the plausibility of the water content values. I think that the calibration method and the orders of magnitude of the results now make sense, and I am really pleased about this. Of course, it still does not imply that they necessarily present the correct absolute values, but together with the other microclimatic data they do show nicely how the dynamics of the water contents in these bryophytes may respond to environmental conditions. I think this is an important contribution of this paper.

Now that the main issue is solved, I think the manuscript just needs one more round of polishing, highlighting and discussing the exciting findings more, correcting some mistakes, optimising some graphs and formulations and checking for style and grammar.

The WC values now look more plausible to me and in the examples shown (Figs 2 and S6) they show a wonderful reaction to rain events. Could you corroborate this impression by a correlation between the WC and the rain amount in the preceding 12 h (or something like that)?

Dear Maaïke Bader, thank you very much for your positive review! We are glad that you also consider the main issue to be solved now. As suggested by you, we plotted a correlation graph between the WC and the rain amount of the preceding 24 h, and a linear regression line shows that over 30% of the variability of the mean water content could be explained by the precipitation within the preceding 24 hours. These results are now shown in Fig. S8. The figure is referred to in the manuscript (P 12, L31-32: “The WC of the bryophytes reacted quite reliably upon rain that had fallen in the preceding hours (Fig. S5; Fig. S8) with some differences between the different height levels. ...”).

It is interesting that the WC of the samples at 23 m sometimes do (especially on 10-09-2016) and sometimes do not at all (March 2015) react to rain. Apart from possible umbrella effects of the canopy, could it also be due to a reinstallation and recalibration of the sensors around June 2015? There seems to be a large and systematic difference between the EC (and WC) values before and after this maintenance period for the sensors at 1.5 m (Fig S5). I guess this may be partly due to the drier season after this period, but I am not convinced that this explains the pattern completely. It may be worth elaborating a bit on this point.

As mentioned by you, some of the EC data indeed look different after as compared to before the maintenance period around June 2015. This definitely is partly caused by the different climatic conditions and the resulting growth of new foliage. But we indeed cannot completely exclude that also the repositioning of the sensors had some effect. We now explain that in the discussion in the section on the sensors at 23 m height (P16, L12-14: “Apart from the effect of growing foliage, one has to keep in mind that there was a re-installation of the sensors around July 2015, which could cause differences in the sensor readings before and after this event.”) and in the section on the understory sensors (P16, L22-23: “Also here, a potential effect of the sensor re-installation around July 2015 has to be kept in mind.”).

The diel fluctuations (mentioned P17, L1) in WC are not mentioned clearly enough in the results. They can be seen a little bit in Fig S8 and better in S5c, but the former figure is referred to only in the context of condensation and the latter not at all in results or discussion. I think these daily fluctuations deserve more attention (especially as you use it as a main conclusion), so I would discuss them with reference to figure S5c, or add another figure similar to Fig S8 but showing more days (only on days without much rain so that the scale of the WC can be precise enough to show the fluctuations - these are hard to see in Fig 2 and S6).

Following your suggestion, information on the diel fluctuations in the results section was made clearer and a reference to figure S5c was added. In addition, we also created a new Fig. S11, which shows these fluctuations during the dry and the wet season. Indeed, we observed that the fluctuations occur during all seasons, but are more regular and thus obvious during the dry season, due to the regular climate with only rare precipitation events. This information is now presented in the results section (P13, L2-5: “Overall, the bryophytes at 8 m and 23 m showed a regular and pronounced daily fluctuation of the WC, which occurred during all seasons, but was particularly regular during the dry season, due to the rare interfering rain events (Fig. 2, Fig. S5c, Fig. S11).”).

In the discussion section, we also added the following text and a link to figure S5c: (P16 L33 - P17L3: “The water content data at 8 m and 23 m height showed diel fluctuations, which were particularly regular during the dry season, due to rare interfering rain events (Fig. S5c). They showed a parallel behavior to the RH data with the highest values reached during the morning hours (Fig. S11). It is well known from the

literature that moist bryophytes and many cryptogams could utilize high air humidity as a source of water (Lange et al., 2001; Raggio et al., 2017) and this likely also occurs here.”)

We also included this information in the conclusions (P19L15-17: “The bryophytes at 8 m and 23 m height showed regular daily fluctuations of the WC contents, which went in parallel to RH and reached highest values during the morning hours.”).

I like Figure S9, but it does not show whether reaching the dew point is related to the fluctuations in the WC. An interesting analysis would be to calculate a correlation between the distance below the dew point reached and the response in WC on that morning. The same could be done for the fog events. The difference in daily fluctuations between the seasons (mentioned in the conclusions) is also interesting and worth some more discussion (P15 L2-3: is this phenomenon more frequent in the dry season?). In fact, I think this point could perhaps make for an additional publication by itself!

As suggested by you, we looked at the events when the dew point was reached in some more detail. We plotted a characteristic sequence during the wet and the dry season (Fig. S11), and here we saw, that the temperature in the bryophytes dropped below the dewpoint of the ambient air mostly during the morning hours. This happened more frequently during the wet as compared to the dry season, which might be caused by a shading effect of the foliage, leading to a lower bryophyte temperature and causing a drop below dewpoint temperature. We also show exemplary dewpoint events at 1.5 and 23 m, where negative dewpoint spread values seem to cause increased water content values (Fig. 12). To analyze this potential effect, we extracted the minimum dewfall spread value per day, and for all negative daily values we calculated the slope of the water content data of the last 4 hours prior the negative dewpoint spread. Relating these parameters with each other, we obtained small negative Pearson’s R values for the three WC sensors at 23 m height (i.e., R_{WC21} : -0.071; R_{WC23} : -0.076; R_{WC24} : -0.040), suggesting that bryophyte temperatures below the dewpoint temperature of the ambient air cause an increase in water content and thus a condensation of water. For fog, such a relationship could not be observed.

We adapted the text accordingly (P13L5-17: “A potential condensation, when the temperature of the bryophytes drops below the dewpoint of the ambient air, was mostly reached during the morning hours (Fig. S9-S12). This occurred during ~50 % of the wet season and ~30% of the dry season days at the surface of bryophytes at 23 m height (Fig. S10, Fig. S11). Contrastingly, at 1.5 m height dew point temperatures were only surpassed during ~9 % of the days, independently of season. Plots of exemplary dewpoint events at 1.5 m and 23 m height suggest that negative dewpoint spread values (i.e., bryophytes temperature below dewpoint of ambient air) cause increased water content values (Fig. 12). To analyze this potential effect, we extracted the largest dewfall spread value per day, and for all negative daily values we calculated the slope of the water content data of the last 4 hours prior the negative dewpoint spread. Relating these parameters with each other, we obtained small negative Pearson’s R values for the three WC sensors at 23 m height (i.e., R_{WC21} : -0.071; R_{WC23} : -0.076; R_{WC24} : -0.040), suggesting that bryophyte temperatures below the dewpoint temperature of the ambient air caused an increase in water content and thus a condensation of water.”).

The new analysis is described in the methods section (P9L12-17:” To analyze if condensation might influence the WC in the bryophytes, a correlation between events with negative ΔT_d and the change of the WC in the 4h before these events were calculated. The potential effect of fog was analyzed by calculating

the average change in water content (ΔWC) from the beginning of the fog event until 1h later. The dew point and fog calculations and correlations were performed with R version 3.6.1 (2019-07-05)...”).

The discussion section was adapted accordingly (P17L3-10: “In addition, also condensation and fog need to be considered as potential additional sources of water for epiphytic covers as well as for near-stem vegetation at the forest floor (Lakatos et al., 2012; van Stan and Gordon, 2018; León-Vargas et al., 2006). Our data show that the necessary conditions for condensation were regularly met and occurred most frequently during the wet season at 23 m height (Fig. S11). Cases with a negative dewpoint spread, when condensation could occur, were related to increasing water contents of the bryophytes, supporting the calculated condensation data. During the occurrence of fog, an increase in water contents could not be directly proven, suggesting that fog does not represent a major water source for the bryophytes.”).

We also looked in the daily fluctuations in some more detail and found that they indeed occur in all seasons but are just more obvious during the dry season, as the daily patterns are more regular, due to the rare rain events. We treated this topic in the previous comment.

I think Figure S5 is useful, but I recommend making the graphs larger. In the supplement I do not think that there is the need to save space, so I would go for clarity here. I also like the magnification shown in S5c, though judging from the rain amounts it does not really correspond to the box in the coarse-scale graph, which I would recommend stretching vertically to make clear that the entire graph (all parameters) are shown in the magnification.

Many thanks for your suggestion; Figure S5 was magnified to show all the parameters with clarity, and the box shown in S5c was changed accordingly.

The way in which stemflow is discussed could be improved. In my view, this is not ‘additional’ water in the sense that moisture input from condensation may be (so I would not merge these two points into one sentence like you now do, P16 L 30), but it is redistributed rain water (mostly, as condensation and fog would hardly lead to stemflow) that benefits epiphytes on the tree stem (i.e. in your case those at 1.5 m in particular). As stemflow concentrates rainfall from a larger area to the stem, this could explain why the bryophytes on the stem respond so reliably to rainfall. For a recent review about this see Mendieta-Leiva, G., P. Porada and M.Y. Bader, 2020. Interactions of epiphytes with precipitation partitioning. In: Precipitation partitioning by vegetation. A global synthesis, J. T. Van Stan IIE. Gutmann and J. Friesen, (Eds.). Springer Nature Switzerland: pp: 133-145.

The text was altered following your suggestion: (P16 L24-32: “In this rain forest environment, epiphytes growing in different parts of the tree and along the stem can benefit from different sources of water. The gross precipitation, as the main water source, can be converted into throughfall, stemflow, water storage and water vapor (Mendieta-Leiva et al., 2020). Thus, rainwater can influence the bryophytes in variable ways depending on its redistribution and the microenvironmental conditions: at the canopy level, direct interception of precipitation can be used for water storage, whereas in the understory, stemflow is more pronounced and contributes to the water supply of the bryophytes. It has been estimated that in tropical forests up to 4 % of the annual rainfall amount could be converted into stemflow (Lloyd and Marques F, 1988; Marin et al., 2000; van Stan and Gordon, 2018), corresponding to maximum values of 68 and 75 mm in the years 2015 and 2016, respectively, at the ATTO site.”).

And I think I found one important mistake in your calculation of activity times, or possibly two (sorry I did not notice this in the previous round):

1. You assume that “For the net photosynthesis (NP) it is required that ... $T > TCP$ ”. However, the TCP is an upper limit, so for a positive NP you need T to be below this compensation point!

This is correct, the calculation was done considering $T < TCP$, it was only a typo and it is now correct: (P31 L6-7) “For the net photosynthesis (NP) it is required that $WC > WCP$, $PAR > LCP_i$ and $T < TCP...$ ”

2. The range for T_{opt} can be considered a range within (not above or below!) which NP is optimal. Therefore I would expect only one value for the % of time where the bryophytes at a particular height in the tree are within this range, not a range of %. This is in contrast to the other ranges which are ranges of estimates for which we do not know which applies so that indeed it is useful to report on the estimated % of time if we assume one or the other value.

Thank you for this valuable comment. We adopted the data accordingly.

Small typos / grammar problems (not exhaustively reviewed) and needs for short clarifications:

P3 L31 ...communities were conducted ...

Following your suggestion, this alteration was done (P3 L26).

P4 L25: add the tree species and some information about its phenology (it seems from the discussion that it is a semideciduous tree..?). This is quite relevant for extrapolating the results.

The required information was added to the text: (P4 L19-23) “The parameters temperature and light within/on top of the bryophyte communities and their WC were measured with a microclimate station installed along one evergreen tree of the species *Buchenavia parvifolia* Ducke (Combretaceae) in September 2014 (Fig. S1). The family regularly occurs in the Amazon rainforest, and represents a common genus in tropical America, growing on clayey soil in plateau environments. It presents flowers during the dry and fruits during the wet season (Stace, 2007).”

P5 L12: 26-m-high

The text was modified accordingly (P5 L26).

P5 29: data that could be used...

The text was changed accordingly (P6 L8).

P5 L33: The electrical conductivity (EC) values on which the WC calculations were based (see Paragraph 2.3 below) showed some unexplained oscillation, causing ...

The text was changed accordingly: (P6 L9-10) “The electrical conductivity (EC) values, on which the WC calculations were based (see Paragraph 2.3 below) showed some unexplained oscillation, causing an inaccuracy...”

P6 L9: please explain this ‘fluctuated’

The values obtained from the light sensors varied in a range of $10 \mu\text{mol m}^{-2} \text{s}^{-1}$ photosynthetic photon flux density (PPFD), and all the 5-min data were averaged to present the same period for all the parameters (30-min). The sentence was rewritten to make this clearer: (P6 L19-24: “The average daily PAR values were calculated from the data collected during daytime, i.e., 6:00 to 18:00, while PAR_{max} represents the daily maximum value. The 5-minute readings obtained from the light sensors fluctuated by approximately $\pm 10 \mu\text{mol m}^{-2} \text{s}^{-1}$ photosynthetic photon flux density (PPFD). To smoothen the microclimate data (i.e. the PPFD values obtained from light sensors, the temperature values measured within the bryophytes and water content values obtained from electrical conductivity sensors), 30-minute averages were calculated and used for all further calculations.”)

P7 L17: I would add here that 0% was an approximation (simplification), as no moss would really reach 0% unless in a good drying oven... But they would dry to a minimum WC of a few %, so near enough to 0% for this model assumption.

The text was modified according to the suggestion: (P7 L26-32: “The minimum electrical conductivity achieved in the field was used as an approximation (simplification) of a water content of 0%. We are aware of the fact that bryophytes do not reach a water content of 0% under field conditions, but they can be expected to dry to a water content of a few %, which we assume as close enough for this model assumption in a tropical rainforest environment.”)

P8 L24: Please start this point by explaining why you calculated this (to estimate the likelihood of water input by dew?). It also would not hurt to introduce this idea in the introduction. I think it is a very important and exciting point!

A short explanation was inserted in the text: (P8 L29-31: “In order to assess the potential water input by condensation, we calculated the dew point temperature, at which saturated air humidity levels are reached.”).

The idea was also added in the Introduction: (P2 L31 – P3 L4: “In a dry state, many of them can outlast extreme weather conditions, being reactivated by water (Oliver et al., 2005; Proctor, 2000; Proctor et al., 2007; Seel et al., 1992). This water can be supplied by precipitation, either directly intercepted or taken up from stemflow. For several species, also condensation of fog and dew can serve as a source of water (Lancaster et al., 1984; Lange et al., 2006; Lange and Kilian, 1985; Reiter et al., 2008).”).

P9 L7: Fig S5 does not show PAR values...

This is correct, the Fig S5 does not show PAR values, but the text (P11 L7) refers to Table S5 where the values for monthly mean values and standard deviations of PAR (average and maximum) are shown.

P13 L8: remove in before 90%

The word was replaced by “during” (P13 L23)

P13 L10: relatively similar or similarly high

The sentence was corrected (P13 L24-25): “maximum light intensities were similarly high”

P16 L7: those at 23 m

The sentence was modified accordingly (P16 L8)

P16 L23 remove the second ‘during the dry season’

The phrase was removed accordingly (P16 L18)

P17 L8: we observed (instead of ‘observed by us’)

The sentence was removed over the course of the editing process (as suggested by you).

P17 L14-16: stating that *Leucobryum* may have a low water retention seems strange, as this genus has special water-retaining cells... I would think that it is not particularly good at capturing water (e.g. from fog), but then very good at retaining it.

This is correct, the sentence has been removed over the course of the reviewing process (as suggested by you).

P17: I would consider removing L5-32, as it does not really add much to your story. The life-form discussion seems too much discussion for the amount of data you have per life form and does not sound like you have profoundly researched it yet. Similarly, with the number of samples you studied it seems unjustified to draw any conclusions about the pattern of distributions among height zones. You could mention this in the methods section where you present the species studied, to justify your choice by showing that you chose typical species for the ecosystem.

This is correct, the life form was not profoundly researched, and the life-form description was moved to the Methods section: (P5 L3-24: “The WC sensors were placed in four different bryophyte communities being heterogeneously distributed along three height levels. At 1.5 m height, the WC sensors were installed in communities dominated by *Sematophyllum subsimplex* (5 sensors) and *Leucobryum*

martianum (1 sensor), at 8 m in *Octoblepharum cocuiense* (2 sensors) and *Symbiezidium barbiflorum* (1 sensor), and at 23 m in *Symbiezidium barbiflorum* (3 sensors; Fig. S2, Fig. S3). The communities used for a placement of the sensors reflect the distribution of bryophytes among height zones in the Amazon rainforest (Cornelissen and ter Steege, 1989; Mota de Oliveira, 2010; Mota de Oliveira and ter Steege, 2015; Pantoja et al., 2015). Studies describe that Lejeuneaceae (common liverwort family of the Amazon region comprising the genus *Symbiezidium*) are more diverse and abundant in the canopy area, while mosses are mainly concentrated at the tree base and trunk in a plateau ecosystem (Campos et al. 2019; Mota de Oliveira 2010, 2018). The species identified by us (Table S1) have also been reported as being frequent at other tropical rain forest sites (Campos et al., 2015; Dislich et al., 2018; Gradstein and Salazar Allen, 1992; Mota de Oliveira et al., 2009; Pinheiro da Costa, 1999). They show different water holding capacities, which are influenced by their life form (Lakatos et al., 2006; Romero et al., 2006; Williams and Flanagan, 1996; Proctor, 1990). The liverwort *Symbiezidium barbiflorum* (Lejeuneaceae) has been described to have the life-form of mats (Batista and Santos, 2016; Mägdefrau, 1982; Valente et al., 2017), which are characterized by a high capillarity retention of water, supporting the storage of condensed water. Mats also have an increased drought-tolerance, being more adapted to dry conditions as well as to extreme changes (Gimingham and Birse, 1957). *Sematophyllum subsimplex* (Sematophyllaceae) and *Leucobryum martianum* (Dicranaceae) belong to the life-forms of wefts and turfs, respectively (Mägdefrau, 1982, Batista and Santos, 2016; Valente et al., 2017). Turfs show high capillary water conduction and are well known for special water-retaining cells, whereas wefts show high values of capillary water conduction but lower values of water retention (Mägdefrau, 1982), being characteristic for humid areas (Gimingham and Birse, 1957).”)

P22 L 1: I would not say that the WC has “turned out to be” the key parameter controlling the overall physiological activity of the organisms, because you did not measure this activity. Better use a more cautious formulation like “appears to be”

The text was modified accordingly (P19L8)

P22 L3 remove the comma after major rain events

The comma was removed (P19 L10)

P22 L8: here also remove the speculative remark about thallus morphology. This is, by the way, not a seasonal feature anyway...

As suggested, the remark regarding the thallus morphology has been removed.

P22 L13-14: do you really know whether this nightly condensation activates physiological processes? Is it enough for photosynthesis, for example? I do not think that we know this, and I do think it would be very important to know it, so perhaps formulate it as a research need rather than a finding.

The sentence was rewritten for clarity (P 19 L18-22: “Thus, our data suggest that the relevant water source for bryophytes in the understory is rain, while for the bryophytes in the canopy RH fluctuations and dew

condensation might be relevant. With the current data at hand, however, it cannot be answered if the daily fluctuations and the dew condensation events are large enough to activate physiological processes; this topic, indeed, would deserve to be investigated in a separate in-depth study.”).

Table 1b: please help the reader by providing the number of days for which the fog sensor was operational (in addition to the sensor outage times)

The requested information was included in Table 1 and Table S3.

Table 3: please make clearer what data were used: the averages of the different samples per height zone? Or the individual samples (i.e. is the range shown a function only of the range of possible parameter values for the cardinal points, or also a function of the differences between samples?).

The data show the averages of the different samples per height zone. This is now also written in the legend (P31 Table 3: “The data show the averages of the different samples per height zone.”).

Fig 4 caption: I do not think that the ‘Estimated’ is appropriate here.

The word was removed from the caption (P39 L2): “Frequency of mean photosynthetically active radiation...”

Figure S8 caption: Exemplary

The spelling was corrected: “Exemplary daily”.

Microclimatic conditions and water content fluctuations experienced by epiphytic bryophytes in an Amazonian rain forest

Nina Löbs^{1*}, David Walter^{1,2}, Cybelli G. G. Barbosa¹, Sebastian Brill¹, Rodrigo P. Alves¹, Gabriela R. Cerqueira³,
5 Marta de Oliveira Sá³, Alessandro C. de Araújo⁴, Leonardo R. de Oliveira³, Florian Ditas¹, Daniel Moran-Zuloaga¹,
Ana Paula Pires Florentino¹, Stefan Wolff¹, Ricardo H. M. Godoi⁵, Jürgen Kesselmeier¹, Sylvia Mota de Oliveira⁶,
Meinrat O. Andreae^{1,7}, Christopher Pöhlker¹, Bettina Weber^{1,8*}

10 1 Multiphase Chemistry and Biogeochemistry Departments, Max Planck Institute for Chemistry, Mainz, 55128, Germany

2 Biogeochemical Process Department, Max Planck Institute for Biogeochemistry, Jena, 07701, Germany

3 Large Scale Biosphere-Atmosphere Experiment in Amazonia (LBA), Instituto Nacional de Pesquisas da Amazonia (INPA), Manaus-AM, CEP 69067-375, Brazil

4 Empresa Brasileira de Pesquisa Agropecuária (EMBRAPA), Belém-PA, CEP 66095-100, Brazil

15 5 Environmental Engineering Department, Federal University of Parana, Curitiba, PR, Brazil

6 Biodiversity Discovery Group, Naturalis Biodiversity Center, Leiden, 2333 CR, The Netherlands

7 Scripps Institution of Oceanography, University of California San Diego, La Jolla, CA 92037, US

8 Institute for Biology, Division of Plant Sciences, University of Graz, 8010 Graz, Austria

*Correspondence to: Nina Löbs (n.loeb@mpic.de) and Bettina Weber (bettina.weber@uni-graz.at)

20 To be submitted in journal *Biogeosciences*, *ATTO special issue*

Abstract. In the Amazonian rain forest, major parts of trees and shrubs are covered by epiphytic cryptogams of great taxonomic variety, but their relevance in biosphere-atmosphere exchange, climate processes, and nutrient cycling are largely unknown. As cryptogams are poikilohydric organisms, they are physiologically active only under moist conditions. Thus, information on their water content, as well as temperature and light conditions experienced by them are essential to analyze their impact on local, regional, and even global biogeochemical processes. In this study, we present data on the microclimatic conditions, including water content, temperature, and light conditions experienced by epiphytic bryophytes along a vertical gradient and combine these with above-canopy climate data collected at the *Amazon Tall Tower Observatory (ATTO)* in the Amazonian rain forest between October 2014 and December 2016. While the monthly average of above-canopy light intensities revealed only minor fluctuations over the course of the year, the light intensities experienced by the bryophytes varied depending on the location within the canopy, probably caused by individual shading by vegetation. In the understory (1.5 m), monthly average light intensities were similar throughout the year and individual values were extremely low, remaining below $3 \mu\text{mol m}^{-2} \text{s}^{-1}$ photosynthetic photon flux density during more than 84 % of the time. Temperatures showed only minor variations throughout the year with higher values and larger height-dependent differences during the dry season. The indirectly assessed water contents of bryophytes varied depending on precipitation, air humidity, dew condensation, and bryophyte type. Whereas bryophytes in the canopy were affected by diel fluctuations of the relative humidity and condensation, those close to the forest floor mainly responded to rainfall patterns. In general, bryophytes growing close to the forest floor were limited by light availability, while those growing in the canopy had to withstand larger variations in microclimatic conditions, especially during the dry season. For further research in this field, these data may be combined with CO_2 gas exchange measurements to investigate the role of bryophytes in various biosphere-atmosphere exchange processes, and could be a tool to understand the functioning of the epiphytic community in greater detail.

1 Introduction

Epiphytic bryophyte communities widely cover the stems and branches of trees in the tropics (Campos et al., 2015). Within that habitat, they may play a prominent role in environmental nutrient cycling (Coxson et al., 1992) and also influence the microclimate within the forest (Porada et al., 2019), thus contributing to the overall fitness of the host plants and the surrounding vegetation (Zartman, 2003). However, they are also affected by deforestation and increasing forest fragmentation (Zartman, 2003; Zotz et al., 1997).

Physiologically, cryptogamic organisms are characterized by their poikilohydric nature, as they do not actively regulate their water status but passively follow the water conditions of their surrounding environment (Walter and Stadelmann, 1968). In a dry state, many of them can outlast extreme weather conditions, being reactivated by water

(Oliver et al., 2005; Proctor, 2000; Proctor et al., 2007; Seel et al., 1992). This water can be supplied by precipitation, either directly intercepted or taken up from stemflow. For several species, also condensation of even fog and dew can serve as a source of water (Lancaster et al., 1984; Lange et al., 2006; Lange and Kilian, 1985; Reiter et al., 2008). In contrast, high water contents (WC) may cause suprasaturation, when gas diffusion is restrained, causing reduced CO₂ gas exchange rates (Cowan et al., 1992; Lange and Tenhunen, 1981; Snelgar et al., 1981) and even ethanolic fermentation, as shown for lichens (Wilske et al., 2001). Accordingly, their physiological activity is primarily regulated by the presence of water and only secondarily by light and temperature (Lange et al., 1996, 1998, 2000; Rodriguez-Iturbe et al., 1999). The amount of available water is relevant to determine the range of photosynthetic activity of these organisms; therefore, rainfall, stemflow and condensation processes, as well as morphological characteristics of bryophytes influencing water conduction or retention are important to understand the activity patterns.

In the Amazonian rain forest, cryptogamic communities mainly occur epiphytically on the stems, branches, and even leaves of trees, and in open forest fractions they may also occur on the soil (Richards, 1954). By 2013, 800 species of mosses and liverworts, 250 lichen species, and 1,800 fungal species have been reported for the Amazon region (Campos et al., 2015; Gradstein et al., 2001; Komposch and Hafellner, 2000; Normann et al., 2010; Piepenbring, 2007). Whereas studies in temperate zones address the importance of cryptogamic communities for the ecosystem (Gimeno et al., 2017; Rastogi et al., 2018), only few reports for the tropical region can be found in the literature.

The Amazonian rain forest has been described to play important roles in the water cycle, as well as in carbon, nitrogen, and phosphorus fluxes on regional and global scales (Andreae et al., 2015). Up to now, the relevance of cryptogamic communities in these regional cycling processes are largely unknown (Hargis et al., 2019). This data are urgently needed, as this ecosystem is under severe pressure and it is hard to predict to which extent the ongoing and envisioned environmental changes will still ensure its ecological services as the “green lung” and carbon sink of planet Earth (Soepadmo, 1993).

In the current study, long-term continuous measurements of temperature, light and water content inside bryophyte communities ~~have been~~ were conducted along a vertical gradient. To our knowledge, our study is the first one measuring microclimatic parameters and the water status inside bryophyte communities in a rainforest environment. With these data on the microclimate along a vertical profile and during different seasons, we believe to provide a unique dataset, combined with an estimation of the activity patterns of bryophyte communities in a tropical rainforest.

2 Material and Methods

2.1 Study site

The study site is located within a *terra firme* (plateau) forest area in the Amazonian rain forest, approx. 150 km northeast of Manaus, Brazil. The average annual rainfall is 2,540 mm a⁻¹ (de Ribeiro, 1984), reaching its monthly maximum of ~ 335 mm in the wet (February to May) and its minimum of ~ 47 mm in the dry season (August to November) (Andreae et al., 2015). These main seasons are linked by transitional periods covering June and July after the wet and December and January after the dry season (Andreae et al., 2015; Martin et al., 2010; Pöhlker et al., 2016). The *terra firme* forest has an average growth height of ~ 21 meters, a tree density of ~ 598 trees ha⁻¹, and harbors around 4,590 tree species on an area of ~ 3,784,000 km², thus comprising a very high species richness compared to other forest types (McWilliam et al., 1993; ter Steege et al., 2013). Measurements were conducted at the research site ATTO (*Amazon Tall Tower Observatory*; S 02° 08.602', W 59° 00.033', 130 m a. s. l.), which has been fully described by Andreae and co-authors (2015). It comprises one walk-up tower and one mast of 80 m each, being operational since 2012, and a 325 m tower, which has been erected in 2015. The ATTO research platform has been established to investigate the functioning of tropical forests within the Earth system. It is operated to conduct basic research on greenhouse gas as well as reactive gas exchange between forests and the atmosphere and contributes to our understanding of climate interactions driven by carbon exchange, atmospheric chemistry, aerosol production, and cloud condensation.

2.2 Microclimatic conditions within epiphytic habitat

The parameters temperature and light within/on top of the bryophyte communities and their WC were measured with a microclimate station installed along one evergreen tree of the species *Buchenavia parvifolia* Ducke (Combrretaceae) in September 2014 (Fig. S1). The family regularly occurs in the Amazon rainforest, and represents a common genus in tropical America, growing on clayey soil in plateau environments. It presents flowers during the dry and fruits during the wet season (Stace, 2007). The sensors were placed along a vertical gradient at— 1.5, 8, and 23 m above the ground on one tree, corresponding to the zones 1, 2 and 4 (i.e., at the base, the lower trunk, and at the base of the crown) described by Mota de Oliveira and ter Steege (2015), to investigate the variation within the story structure of the forest.

It needs to be mentioned, that not only one single species was measured by one sensor, but usually several bryophyte species and also other cryptogams, such as lichenized and non-lichenized fungi and algae, as well as heterotrophic fungi, bacteria and archaea, which grow together forming a cryptogamic community. Thus, the organisms mentioned throughout this paper were the dominating but not solitarily living species. The restriction of the measurements to one individual tree needs to be considered, as a complete independence of the replicate sensors could not be assured. However, due to the large effort of such an installation within the rain forest, it was not possible to

equip more trees with additional instruments. Thus, the data obtained from the measurements on this individual tree should be considered as exemplary.

~~For that reason, the~~ The WC sensors were placed in four different bryophyte communities being heterogeneously distributed along three height levels. At 1.5 m height, the WC sensors were installed in communities dominated by *Sematophyllum subsimplex* (5 sensors) and *Leucobryum martianum* (1 sensor), at 8 m in *Octoblepharum cocu-
5 iense* (2 sensors) and *Symbiezidium barbiflorum* (1 sensor), and at 23 m in *Symbiezidium barbiflorum* (3 sensors; Fig. S2, Fig. S3). The communities used for a placement of the sensors reflect the distribution of bryophytes among height zones in the Amazon rainforest (Cornelissen and ter Steege, 1989; Mota de Oliveira, 2010; Mota de Oliveira and ter Steege, 2015; Pantoja et al., 2015). Studies describe that Lejeuneaceae (common liverwort family of the
10 Amazon region comprising the genus *Symbiezidium*) are more diverse and abundant in the canopy area, while mosses are mainly concentrated at the tree base and trunk in a plateau ecosystem (Campos et al. 2019; Mota de Oliveira 2010, 2018). The species identified by us (Table S1) have also been reported as being frequent at other tropical rain forest sites (Campos et al., 2015; Dislich et al., 2018; Gradstein and Salazar Allen, 1992; Mota de
15 Oliveira et al., 2009; Pinheiro da Costa, 1999). They show different water holding capacities, which are influenced by their life form (Lakatos et al., 2006; Romero et al., 2006; Williams and Flanagan, 1996; Proctor, 1990). The liverwort *Symbiezidium barbiflorum* (Lejeuneaceae) has been described to have the life-form of mats (Batista and Santos, 2016; Mägdefrau, 1982; Valente et al., 2017), which are characterized by a high capillarity retention of water, supporting the storage of condensed water. Mats also have an increased drought-tolerance, being more adapted to dry conditions as well as to extreme changes (Gimingham and Birse, 1957). *Sematophyllum subsimplex*
20 (*Sematophyllaceae*) and *Leucobryum martianum* (*Dicranaceae*) belong to the life-forms of wefts and turfs, respectively (Mägdefrau, 1982, Batista and Santos, 2016; Valente et al., 2017). Turfs show high capillary water conduction and are well known for special water-retaining cells, whereas wefts show high values of capillary water conduction but lower values of water retention (Mägdefrau, 1982), being characteristic for humid areas (Gimingham and Birse, 1957).

25 Additionally, at each height level two temperature and two light sensors (except for 1.5 m with only one light sensor) were installed in/on top of the bryophyte communities located on the approximately ~~26-26-m-m~~-high tree (Fig. S2, Table S1). The temperature sensors were installed in the same communities as the WC sensors, and the light sensors were installed adjacent to them on ~ 5 cm long sticks (Fig. S1). As the morphology of the different species affects their overall WC, different maximum WC contents were observed (Tab. S1, S2). At 1.5 and 8 m
30 the sensors were installed vertically along the trunk, whereas at 23 m they were placed at the upper side of a horizontal branch. Thus, also the orientation at the stem may influence the WC of the bryophyte communities, not only the species and the height above ground. Furthermore, sample properties as their thickness and density might play a relevant role for their WC, as samples at 1.5 m height tended to be more loose and thinner as compared to the ones at the upper height levels (Fig. S4). Since the installation, automatic measurements at 5-minute intervals

were taken with a data logger (CR1000; Campbell Scientific, Logan, Utah, USA) equipped with a relay multiplexer (AM16/32; Campbell Scientific, Bremen, Germany) and two interfaces.

The WC sensors, initially developed for biological soil crust research (Tucker et al., 2017; Weber et al., 2016), were optimized for measurements in epiphytic bryophyte communities by a straight-lined construction and with outer pins of 25 mm length, serving as an effective holdfast. However, during stormy episodes and/or physical friction, some WC and temperature sensors fell out of the moss samples and required a reinstallation. Additionally, during some episodes the sensors showed unreliable data, which had to be removed from the overall dataset. All data, ~~which that~~ could be used for data analysis, are shown in Fig. S5.

The electrical conductivity (EC) values, on which the WC calculations were based (see Paragraph 2.3 below) showed some unexplained oscillation, causing an inaccuracy corresponding to approximately 20 mV. Besides the specific position in the substrate, the EC also depends on the texture of the sample material, its ion concentration, and the temperature. Because of all these factors influencing the sensor readings, the provided values of the WC should be considered as the best possible estimates and not as exact values. For the temperature measurements, thermocouples (Conatex, St. Wendel, Germany) with a tip length of 80 mm and a measurement accuracy of ± 0.5 °C were used. For the light sensors, GaAsP-photodiodes (G1118, Hamamatsu Photonics Deutschland GmbH, Herrsching, Germany) were placed in a housing covered by a convex translucent polytetrafluoroethylene (PTFE) cap and calibrated against a PAR (photosynthetically active radiation) quantum sensor (SKP215; Skye Instruments, Llandrindod Wells, Powys, UK).

The average daily PAR values were calculated from the data collected during daytime, i.e., 6:00 to 18:00, while PAR_{max} represents the daily maximum value. The values-5-minute readings obtained from the light sensors fluctuated by approximately $\pm 10 \mu\text{mol m}^{-2} \text{s}^{-1}$ photosynthetic photon flux density (PPFD). To smoothen the microclimate data, (i.e. the PPFD values obtained from light sensors, the temperature values measured within the bryophytes and water content values obtained from electrical conductivity sensors), 30-minute averages were calculated and used for all further calculations, as well photosynthetic photon flux density (PPFD) obtained from light sensors, and water content obtained from electrical conductivity sensors. During measurements, the light sensors were regularly checked for algal growth and cleaned accordingly.

2.3 Calculation of the water content (WC)

The WC sensors measure the electrical conductivity in the field (EC_t), which is influenced by temperature; consequently, a temperature correction was performed according to Eq. (1), analogous to Weber et al. (2016):

$$EC_{25} = f_T * EC_t, \quad (1)$$

with EC_{25} as EC at 25 °C, T as bryophyte temperature [°C] and the temperature conversion factor f_T :

$$f_T = 0.447 + 1.4034 e^{-T/26.815}. \quad (2)$$

The WC sensor has a fixed distance between the sensor pins, which ensures that in all sensors the resistance is equal. This guarantees that the electrical voltage, being the inverse resistance, is proportional to the electrical conductivity. The values of the sensors were recorded as electrical voltage in mV and by calibration transformed into the WC of the samples, given as dry weight percentage, as explained below.

5 To determine the maximum water content of the different bryophyte communities, samples of them were collected in the forest area surrounding the ATTO site. They were removed from the stem with a pocket knife and stored in paper bags in an air conditioned lab container until calibration (few hours after collection). Prior to the calibration, the samples were cleaned from adhering material using forceps. The weight of the bryophytes was determined when they were moistened until saturation (temperature 30° C, RH 100 %) and again after drying in a dryer over-
10 night (temperature 40° C, RH 30 %) to simulate the natural range of the WC under controlled temperature and RH conditions. The dry weight (*DW*) was determined after drying at 60° C until weight consistency was reached (Caesar et al., 2018). The WC of the sample was calculated according to the formula in Weber et al. (2016):

$$WC [\% DW] = \frac{(FW-DW)}{DW} * 100 \%, \quad (3)$$

with *FW* as sample fresh weight [g] and *DW* as sample dry weight [g].

15 In a previous approach, calibration curves were established under controlled conditions, logging the electrical conductivity values and the corresponding weight/water content of samples of the different bryophyte species during drying, analogous to (Weber et al., 2016). However, the variability of electrical conductivity values between samples and even at different spots within one sample turned out to be too large and thus this was not a feasible approach to calibrate the sensors. On the other hand, the electrical conductivity values decreased in a linear way
20 with decreasing sample weight, demonstrating that a linear relationship between both factors could be assumed (except for water contents close to saturation).

In the current approach, the calibration of the water content was performed, based on the maximum and minimum values of electrical conductivity reached in the field and the maximum WCs reached during the laboratory measurements. We assumed, that the maximum electrical conductivity achieved in the field corresponds to the maxi-
25 mum WC achieved in the laboratory, due to water saturation of the samples during the laboratory measurement.

The minimum electrical conductivity achieved in the field was used as an approximation (simplification) of ~~corresponds to~~ a water content of 0%. We are aware of the fact that bryophytes do not reach a water content of 0% under field conditions, but they can be expected to dry to a water content of a few %, which we assume as close enough for this model assumption which is a simplification for the minimal WC values achieved (few % and close
30 enough to 0%). This is an approximation, to represent ~~thas we assumed that all samples dried state during at least once over the whole measurement period for this model assumption at the~~ in a tropical rainforest environment.

Accordingly, the water content (WC) was calculated as follows:

$$WC [\% DW] = \frac{WC_{max}}{(EC_{max}-EC_{min})} * (EC_i - EC_{min}), \quad (4)$$

with WC_{max} corresponding to the maximum WC measured in the laboratory, EC_i as electrical conductivity, EC_{min} as the minimum electrical conductivity-, and EC_{max} as the maximum electrical conductivity measured in the field. The measured electrical conductivity values showed short-time oscillations, which might be caused by the fact that the bryophytes cushions have some air spaces in-between, as we observed that these oscillations are less pronounced in denser substrate. Nevertheless, the overall functionality of the sensors is still ensured also in less dense material, and the short-term fluctuations could be removed with a 30-minute smoothing algorithm. Thus, for all calculations the 30-minute averages were used. The electrical conductivity data of replicate samples at the same height were combined to obtain average values for each height.

2.4 Meteorology

For the purpose of long-term monitoring, a set of meteorological parameters is being measured within the ATTO project since 2012. In our study we used rainfall data measured at 81 m [$mm\ min^{-1}$] (Rain gauge TB4, Hydrological Services Pty. Ltd., Australia), relative humidity (RH) measured at 26 m [%], air temperature measured at 26 m [$^{\circ}C$] (Termohygrometer CS215, Rotronic Measurement Solutions,UK), and photosynthetically active radiation (PAR) measured at 75 m height above the ground [$\mu mol\ m^{-2}\ s^{-1}\ PPF$] (Quantum sensor PAR LITE, Kipp & Zonen, Netherlands). All data were recorded at 1-minute intervals with data loggers (CR3000 and CR1000, Campbell Scientific, Logan, Utah, USA) on the walk-up tower (Andreae et al., 2015).

For calculation of the average light intensities per month, season or year (PAR_{avg} month, PAR_{avg} season, PAR_{avg} year) only values during daytime (6:00 – 18:00 local time) were considered. Rainfall data are presented as accumulated values in millimeters per month, season, or year, which were calculated by an integration of 30-minute intervals. As there were gaps in the data record of the rain detection, additional information from the WC sensors were used to calculate the number of days with rain events. The sensors at 1.5 m height were found to react reliably to rain events. Thus, the gaps in the rain detection were corrected with the information received from these sensors. Furthermore, the amount of rain within each month was corrected by assuming that during the missing days there were the same amounts as during the rest of the month. Overall, a malfunction of the rain detection was observed on only 6 % of the days (Table S4).

The information on fog events was provided by visibility measurements using an optical fog sensor installed at 50 m height (OFS, Eigenbrodt GmbH, Königsmoor, Germany). Fog was defined to occur at visibility values below 2,000 m.

~~In order to assess the potential water input by condensation, we calculated the dew point temperature, at which was calculated. Based on the climate data, we calculated the dew point temperature, to understand at which temperature the saturated air humidity levels are reached. If the temperature drops below the dew point, condensation might occur and form liquid water. The dew point spread is the temperature difference between a surface temper-~~

ature and the dew point of the surrounding air and can be used to assess potential condensation processes at surfaces. If the difference between the surface temperature T_s and the dew point of the surrounding air T_d is negative, water vapor is able to condense at the colder surface. The calculations were performed according to the following equations:

$$5 \quad \Delta T_d = T_s - T_d \quad (5)$$

and

$$T_d = \frac{241.2 \ln\left(\frac{\varphi}{100}\right) + \frac{4222.03716 \vartheta}{241.2 + \vartheta}}{17.5043 - \ln\left(\frac{\varphi}{100}\right) - \frac{17.5043}{241.2 + \vartheta}} \quad (6)$$

10 With ΔT_d = dew point spread [°C], T_s = surface temperature (bryophytes sensors, 23 m or 1.5 m) [°C], T_d = dew point temperature air [°C], ϑ = temperature [°C], φ = relative humidity [%]. The dew point spread calculation was performed for the levels of 23 m and 1.5 m height. As a data source for air temperature and humidity, meteorology data assessed at the walk-up tower at 26 m and 1.5 m were used. For surface temperature (T_s), the temperature data measured within the bryophyte communities at 23 m and 1.5 m height were applied. To analyze if condensation might influence the WC in the bryophytes, a correlation between events with negative ΔT_d and the change of the WC in the 4h before these events were calculated. The potential effect of fog was analyzed by calculating the average change in water content (ΔWC) from the beginning of the fog event until 1h later. The dew point and fog calculations and correlations were performed with R version 3.6.1 (2019-07-05).

15 Time readings are always presented as UTC (universal coordinated time) values, except for diurnal cycles, where local time (LT, i.e., UTC-4) is shown, as labeled in the figures.

20 **2.5 Potential physiological activity of bryophytes**

The physiological activity of bryophytes – and of cryptogams in general – is primarily controlled by water and light, whereas temperature plays a secondary role, at least in the environment of the central Amazon (Lösch et al., 1994; Wagner et al., 2013). While the availability of water determines the overall time of physiological activity, the light intensity regulates whether net photosynthesis (NP) or dark respiration (DR) dominates the overall metabolic balance. Furthermore, high nighttime temperatures cause increased carbon losses due to high respiration rates, as previously shown for lichens (Lange et al., 1998, 2000). For tropical bryophytes along an altitudinal gradient in Panama, however, it has been shown that respiration loss during the night might not play the determining role for an overall positive net carbon balance, as species acclimatized to elevated temperatures, but that the restricted time for photosynthesis was a decisive factor (Wagner et al., 2013).

25 To assess the potential physiological activity of bryophyte communities, the water and light conditions as major drivers of the metabolism were investigated in somewhat greater detail. The lower water compensation point (WCP) presents the minimum WC that allows positive net photosynthesis. For the tropical liverwort *Symbiezidium*

spp., occurring in the lowlands near sea level in Panama, WCP values in the range between ~ 30 and ~ 80 % have been determined (Wagner et al., 2013) (Table S4).

The lower light compensation point (LCP_1) represents the minimum light intensity that allows a positive primary production; it ranges between ~ 3 and ~ 12 $\mu\text{mol m}^{-2} \text{s}^{-1}$ for bryophytes (based on measurements of *Ectropothecium* cf. *perrotii*, *Frullania* spec., *Neckera* spec., *Plagiochila divergens*, *Plagiochila squamulosa*, *Porothamnium stipiatum*, *Porotrichum molliculum*, *Racopilum tomentosum*, *Radula boryana*, *Rhizogonium spiniforme*) occurring in African tropical lowland rain forests (Lösch et al., 1994). The epiphytic bryophytes grew in an upper lowland rain forest in the Kahuzi-Biega National Park (Zaire) at about 800 m a. s. l.. Microclimatic conditions inside the forest were similar to the conditions at the ATTO site, as RH ranged around 60–70 % during sunny days and temperatures remained above 20 °C during night and day. At light intensities below the LCP_1 and WCs below the WCP, respiration rates are higher than NP rates, causing overall net respiration to occur.

With regard to temperature, a range for optimum NP (T_{opt}) and an upper compensation point, where NP equals DR (TCP) can be defined. For tropical bryophytes (i. e., the species *Octoblepharum pulvinatum*, *Orthostichopsis te-tragona*, *Plagiochila* sp. 1, *Stictolejeunea squamata*, *Symbiezidium* spp., *Zelometeorium patulum*), T_{opt} ranges between 24 and 27 °C and the TCP ranges between 30 and 36 °C, as described by Wagner and coauthors (Wagner et al., 2013). For long-term survival and growth, the bryophytes need to be predominantly exposed to temperatures below the upper compensation point, at least under humid conditions. The measurements performed by Wagner et al. (2013) were conducted at a study site (BT) in a lowland rainforest in Western Panama on the Bocas del Toro archipelago, located approximately at sea level. The mean temperature was 25 °C (26 °C during day, 24 °C during night), thus slightly warmer than the temperatures measured at ATTO. With 3,300 mm a^{-1} of rain, BT is in a similar range as the ATTO site (2,540 mm a^{-1}). Unfortunately, literature data on the compensation points are rare, facilitating only a first approximate assessment of the physiological processes (Lösch et al., 1994; Wagner et al., 2013). A WC above the compensation point allows NP if both light intensity and temperature are above the lower compensation point. If WCs are above the compensation point but light intensities are too low, or if temperatures are above the upper compensation point, net DR occurs. There is also a narrow span of low WCs, when samples are activated already but despite sufficient light intensities only net respiration can be measured. As this span of WCs is narrow and respiration rates are low, it has been neglected in the current calculations. The compensation points for the different parameters are also to some extent interrelated, e.g., the water compensation point of lichens has been shown to slightly increase with increasing temperature (Lange, 1980), but this can be neglected in such a first qualitative approach. Finally, also inter- and intraspecific variation of compensation points could not be considered in the current study.

2.6 Data analysis

All data processing steps and analyses were performed with the software IGOR Pro (Igor Pro 6.37, WaveMetrics, Inc, Lake Oswego, Oregon, USA). For the average values obtained at the different height levels, the data of the individual sensors were pooled.

5 3 Results

3.1 Microclimatic conditions

3.1.1 Annual fluctuation of monthly mean values

Over the course of the two years of measurements, the monthly mean values of the WC, temperature, and light conditions experienced by the epiphytic bryophyte communities, as well as the above-canopy meteorological conditions, varied between seasons and years. Comparing the two consecutive years, the effect of an El Niño event was clearly detectable, as rainfall amounts were 35 % lower (525 mm versus 805 mm) and relative air humidity 11 % lower (81 % versus 92 %) between October 2015 and February 2016 as compared to the same time-span in the previous year (Fig. 1, Table S3).

The monthly mean values of above-canopy PAR (PAR_{avg}) were rather stable throughout the years and did not differ between the years 2015 and 2016, ranging between 635 and 1150 $\mu\text{mol m}^{-2} \text{s}^{-1}$ during the daytime (Fig. 1, Table S3). The PAR_{avg} values in the understory at 1.5 m also showed only minor seasonal variation, whereas those at higher levels revealed larger variations (Fig. 1, Table. S5). However, the light conditions observed at one individual tree are strongly influenced by its canopy structure and foliage and thus could not be considered as data representative for the canopy in general.

Over the course of the years, the monthly mean temperatures at all heights as well as above-canopy temperatures showed a parallel behavior (Fig. 1). The temperatures decreased in a stepwise manner from the canopy to the understory, and temperatures within bryophytes at 23 m height were frequently higher than the temperatures measured above the canopy (Fig. 1, Fig. S67). Overall, temperatures at all height levels were lower and more similar during the wet than the dry seasons.

Over the course of the years, the monthly WCs of epiphytic bryophytes showed similar patterns corresponding to the increasing and decreasing values of rain and RH. During the dry season 2015, it rained on 25 % of the days, while in the previous and subsequent years rain occurred at a higher frequency (58 % and 31 % of the days, respectively; Fig. 1, Table S3). Monthly rain amounts varied from 9 mm during the dry to 341 mm during the wet season. In 2016, the rain increased from January to March and decreased from March to August, while in 2015 the monthly rain amounts were more variable but still lower throughout the year. The lowest monthly average of the RH was observed during the dry season 2015 with 74 ± 15 %. The monthly WC values of epiphytic bryophyte

communities at 1.5 and 8 m varied between seasons in parallel to the monthly rain amounts, whereas at 23 m the values remained relatively stable over the complete measurement time. During the dry seasons, the WC of the mosses at 1.5 m tended to be lowest, whereas during the wet season this was the case for the liverworts at 23 m height (Fig. 1, Tab. 2).

5 3.1.2 Seasonal changes between wet and dry season

The wet seasons were characterized by a high frequency of precipitation events, large amounts of rain per event, the frequent appearance of fog, and high RH values, ranging mostly above 70 % (Fig. 2, Table 2, Fig. S6S7). In contrast, during the dry season the precipitation events were much rarer and smaller, there was hardly any occurrence of fog, and the RH regularly had values below 60 %. Comparing environmental conditions of the seasons, 10 the diel amplitudes of ambient light, temperature, and RH were larger in the dry compared to the wet season (Fig. 3). While the microclimatic temperature and light conditions within and on top of the epiphytic bryophyte communities mostly followed the above-canopy conditions, modified by canopy shading, the WC of bryophytes did not present a clear pattern (Fig. 3).

The above-canopy light intensity (PAR_{avg} daytime) tended to be higher and to show somewhat stronger fluctuations 15 in the dry season than in the wet season ($950 \pm 93 \mu\text{mol m}^{-2} \text{s}^{-1}$ vs. $738 \pm 46 \mu\text{mol m}^{-2} \text{s}^{-1}$; Table 2). During both main seasons the average light intensity (PAR_{avg} daytime) decreased from the canopy towards the understory. During the dry season this happened in a regular stepwise manner, whereas in the wet season there were some irregularities, probably caused by the local canopy structure (Fig. 2, 3 Table 2).

The temperatures showed a decreasing gradient from the canopy (wet season: $25.7 \pm 0.7 \text{ }^\circ\text{C}$, dry season: 20 $27.2 \pm 1.0 \text{ }^\circ\text{C}$) towards the understory (wet season: $24.9 \pm 0.4 \text{ }^\circ\text{C}$, dry season: $26.0 \pm 0.8 \text{ }^\circ\text{C}$) and the differences among heights and diel amplitudes were more pronounced during the dry season (Fig. 2, 3, Table 2). During the dry season, temperatures within the bryophyte communities at 23 m height were frequently higher than the above-canopy values, and even the seasonal average temperature was 0.6°C higher, probably due to surface heating (Table 2).

25 During the wet seasons of 2015 and 2016, rain occurred on average on 84 % of the days and in the dry season on 28 % of the days (Table S3). During the wet season, an average RH of $94 \pm 2 \text{ } \%$ and frequently even full saturation was reached, while during the dry season the RH reached an average value of $84 \pm 6 \text{ } \%$ (Table 2). Fog was recorded on 60 % of the days during the wet seasons and on 20 % of the days during the dry seasons, respectively (Fig. 2, Table S3). According to our observations, fog observed above the canopy normally also occurred (at least to some 30 extent) within the forest.

The WC of the bryophytes reacted quite reliably upon rain that had fallen in the preceding hours (Fig. S5; Fig. S8) with some differences between the different height levels. The sensors at 1.5 and 8 m responded consistently to rain events, while for the liverworts at 23 m height only sometimes an immediate response was observed. During

the wet season, the bryophytes at 1.5 and 8 m height contained an increased WC over several days after a rain event, while in the dry season the samples tended to dry quickly again. Overall, the bryophytes at 8 m and 23 m showed a regular and pronounced daily fluctuation of the WC, which occurred during all seasons, but was particularly regular during the dry season, due to the rare interfering rain events especially during the dry season (Fig. 2, Fig. S5c, Fig. S11). A potential condensation of dew, when the temperature of the bryophytes drops below the dewpoint of the ambient air, was mostly reached during the morning hours (Fig. S9-S12). This occurred during condensation might serve as an additional source of water, as in ~50 % of the wet season and ~30% of the dry season days at the surface of bryophytes at 23 m height the surface temperatures of the bryophytes at 23 m height went below the calculated dew point temperature (Fig. S108, Fig. S119). Contrastingly, at 1.5 m height dew point temperatures were only surpassed during ~9 % of the days, independently of season. Plots of exemplary dewpoint events at 1.5 m and 23 m height suggest that negative dewpoint spread values (i.e., bryophytes temperature below dewpoint of ambient air) cause increased water content values (Fig. 12). To analyze this potential effect, we extracted the largest dewfall spread value per day, and for all negative daily values we calculated the slope of the water content data of the last 4 hours prior the negative dewpoint spread. Relating these parameters with each other, we obtained small negative Pearson's R values for the three WC sensors at 23 m height (i.e., R_{WC21} : -0.071; R_{WC23} : -0.076; R_{WC24} : -0.040), suggesting that bryophyte temperatures below the dewpoint temperature of the ambient air caused an increase in water content and thus a condensation of water. An effect of fog occurrence on the WC of bryophytes could not be directly shown, as the WC decreased within one hour after the fog event started (i.e., ΔWC_{21} : -1.6%; ΔWC_{23} : -0.4%; ΔWC_{24} : -3.4%), probably due to the effect of rising temperatures during the morning hours.

3.2 Potential physiological activity of bryophytes

Whereas overall light intensities at the upper two height levels were rather similar, with values below 108 and 147 $\mu\text{mol m}^{-2} \text{s}^{-1}$ (at 8 and 23 m height) during in 90 % of the time, the values at 1.5 m height remained below 10 $\mu\text{mol m}^{-2} \text{s}^{-1}$ during the same time fraction (Fig. 4). In contrast to that, maximum light intensities were relatively similarly similar high, reaching 1,550 (1.5 m), 1,500 (8 m), 1,040 (18 m), and 950 $\mu\text{mol m}^{-2} \text{s}^{-1}$ (23 m). If we assume a lower light compensation point (LCP₁) ranging between 3 and 12 $\mu\text{mol m}^{-2} \text{s}^{-1}$ (Löscher et al., 1994), the understory samples (1.5 m) exceeded that range only in less than 20 % of the time during both the wet and the dry season, whereas at the uppermost height level the bryophytes exceeded these values in during one third to almost half of the time (Table 3).

The temperatures inside the moss stands at different height levels mainly ranged between 22 °C and ~28 °C during the wet and between 23 °C and ~33 °C during the dry season (Fig. 4). For tropical lowland regions, the optimum temperatures for bryophytes (T_{opt}) have been suggested to range between 24.0 °C and 27.0 °C (Wagner et al., 2013). If we assume this range for our study, the temperatures in the understory (at 1.5 m height) remained within

these limits in ~~2-77~~91 % of the time during the wet season and in ~~9-60~~76 % of the time during the dry season (Table 3). In the canopy (at 23 m height), temperatures remained in this range in ~~4-54~~70 % of the time during the wet and in ~~6-35~~46 % of the time during the dry season (Table 3). For an upper temperature compensation point (TCP), above which respiration exceeds photosynthesis, Wagner and co-authors suggested 30.0 °C – 36.0 °C (Wagner et al., 2013). In the understory, this TCP was never surpassed during the wet season and only rarely during the dry season. Similarly, at the uppermost canopy level the upper TCP was surpassed in 0-3 % of the time during the wet and in 0-27 % of the time during the dry season. Overall, the highest temperatures were reached when the bryophytes were relatively dry and most probably inactive (Fig. S1~~30~~). The WC of bryophytes differed along the vertical profile, with smaller values in the understory at 1.5 m height than in the canopy (8 m and 23 m), particularly during the dry season (Fig. 4). Considering a lower water compensation point (WCP) between 30 and 80 % according to the literature (Wagner et al., 2013), bryophytes at the uppermost level surpassed these values during 3-80% of the time during the wet and 6-64 % of the time during the dry season. Contrastingly, at 1.5 m and 8 m height there were larger differences between seasons, as the assumed upper range of the WCP was only rarely reached during the dry but during ~ 40% of the time during the wet season (Fig. 4; Table 3).

15 4 Discussion

4.1 Microclimatic conditions

In the current study we measured the microclimatic conditions experienced by epiphytic bryophyte communities along a vertical gradient over the course of more than two years. In previous studies, microclimatic data on the light, temperature, and air humidity have been assessed, at different height levels within the forest (Chazdon and
20 Fletcher, 1984; Lösch et al., 1994; Romero et al., 2006), but long-term measurements of the water content and the light and temperature on top and inside the cryptogamic communities and have been missing up to now.

The microclimatic conditions experienced by epiphytic bryophyte communities along a height gradient at the ATTO site followed the meteorological parameters to some extent, but they also revealed microsite-specific properties regarding annual, seasonal, and diel microclimate patterns. Whereas the water content and the temperature
25 mostly followed the patterns of the meteorological parameters precipitation and temperature, the light intensities were clearly altered, particularly in the understory, due to the local canopy structure.

Within one height level, the small-scale environmental conditions, such as radiation and shading, water conditions, and wind velocity vary, depending on the specific habitat conditions, as e.g. exposition, tree foliage and inclination of the substrate (Barkman, 1958; Campos et al., 2019; Cornelissen and ter Steege, 1989; Oliveira and Oliveira,
30 2016; Sierra et al., 2018). These small-scale patterns also explain the variability observed within one height level. Over the measurement period, the monthly averages of above-canopy light conditions (PAR_{avg}) were rather stable (Fig. 1, Table S3). Within the canopy, the monthly PAR_{avg} values at 23 m height tended to be higher during the

dry seasons, whereas patterns were less clear at 8 m height and there was hardly any seasonal variation at 1.5 m height. This was most probably an effect of the canopy structure, cushion orientation, and shading. The sensors at 1.5 and 8 m were installed vertically along the trunk, whereas at 23 m they were positioned on the upper side of a horizontal branch. As the light sensors at 23 m height were located within the canopy, newly growing leaves may have periodically shaded the organisms, which may explain the lower monthly PAR_{avg} values at this height level compared to the values at the lower levels, where sunbeams could come through the canopy of neighboring trees and open space.

The diel patterns of PAR_{avg} are expected to show a decreasing gradient from the canopy to the understory, as the canopy receives most solar radiation, while the understory vegetation is expected to be shaded by foliage and branches. During the dry season this general pattern was indeed observed, whereas during the wet season mean light intensities were often higher at 8 than at 23 m, probably also caused by canopy shading effects at the upper two height levels (Fig. 2). High light intensities above $1000 \mu\text{mol s}^{-2} \text{s}^{-1}$ occurred in the understory only as small light spots of short duration and thus were only observed in 0.008 % of the time. For the understory of a rain forest in Costa Rica, light intensities were reported to range from 10 to $1000 \mu\text{mol m}^{-2} \text{s}^{-1}$, and more than in 50 % of the total amount of light resulted from sun flecks (Chazdon and Fetcher, 1984). Bryophyte and lichen taxa in the understory are known to be adapted to these low light conditions and are able to make efficient use of the rather short periods of high light intensities (Lakatos et al., 2006; Lange et al., 2000; Wagner et al., 2014).

The temperatures measured inside the bryophyte communities followed the above-canopy temperature at all height levels, with a mostly increasing gradient from the understory towards the canopy, probably caused by a reduced shading effect towards the canopy (Fig. 1, Table 1, 2). At the uppermost height level, mean temperatures inside the bryophyte communities often were even higher than the mean above-canopy temperatures. During the wet season, the overall temperature conditions were more buffered due to reduced incoming radiation caused by clouds and a frequent mixing of the air masses during rain events (von Arx et al., 2012; Gaudio et al., 2017; Thompson and Pinker, 1975).

The microclimatic mean temperature differences measured inside the bryophyte stands between the understory (1.5 m) and the canopy (23 m) were $1.5 \text{ }^\circ\text{C}$ in the dry and only $0.5 \text{ }^\circ\text{C}$ in the wet season. Compared to these results, a temperature difference of $4.0 \text{ }^\circ\text{C}$ was determined during the dry season in a tropical evergreen forest in Thailand, while in the wet season it was below $1.0 \text{ }^\circ\text{C}$, thus corresponding quite well to our results (Thompson and Pinker, 1975) (Table 2). The daily amplitude of the temperature was about twice as large in the canopy as compared to the understory (Fig. 3) This could be caused by the exposure to strong solar radiation and higher wind velocity in the canopy compared to the sheltered understory (Kruijt et al., 2000).

Rainfall amounts and relative air humidity values differed between the seasons and also between the years, as they were considerably higher between October 2014 and February 2015 as compared to the following year. This was most probably to an El Niño event, which caused air humidity and WC of bryophytes to be substantially lower

compared to the previous dry season (Fig. 1, Table 1). Generally, the moisture conditions, including rain, condensation of dew and fog, and RH, differed between seasons, resulting in different WC patterns of bryophytes. A higher frequency of rain during the wet season particularly affected the bryophyte communities at the lower levels (1.5 and 8 m), whereas those in the canopy showed similar water contents during all seasons (Fig. 2, Fig. S76a, Table 2).

The data also suggest that the position of the measured communities and the tree foliage played a crucial role for rainwater absorption and the subsequent drying process. Whereas the sensors at 1.5 m and 8 m height responded quite reliably to rain events during all seasons, ~~these those~~ at 23 m responded quite reliably during the dry but only rarely during the wet seasons. This might be caused by a dense foliage during the wet season, shading the communities at 23 m from direct inundation, whereas during the dry season the canopy foliage changes a lot and intense leaf shedding takes place before new leaves develop, which seems to allow rain to also reach the samples directly below the canopy (Lopes et al., 2016). Apart from the effect of growing foliage, one has to keep in mind that there was a re-installation of the sensors around July 2015, which could cause differences in the sensor readings before and after this event.

During the wet season, the WC of bryophytes in the understory and at 8 m height responded strongly to rain events and subsequently, the water was lost gradually with bryophytes staying wet and active over prolonged time spans, indicating that large amounts of water were taken up during prolonged rain events (Fig. 2, Fig. S76). In contrast to that, during the dry season the drying of the samples occurred quite rapidly after the rain ~~during the dry season~~. Most rain events in the Central Amazon occur in the early afternoon (12:00–14:00 LT) and more than 75 % of them are weak events of less than 10 mm (Cuartas et al., 2007), which often cause no complete water saturation of the bryophytes. Consequently, the organisms tend to dry much quicker than after strong rain events ~~occurring more frequently during the wet season~~. Also here, a potential effect of the sensor re-installation around July 2015 has to be kept in mind.

In ~~this a~~ rain-forest environment, epiphytes growing in different parts of the tree and along the stem can benefit from different sources of water. The gross precipitation, as the main water source, can be converted into through-fall, stemflow, water storage and water vapor (Mendieta-Leiva et al., 2020). Thus, rainwater can influence the bryophytes in variable ways depending on its redistribution and the microenvironmental conditions: at the canopy level, direct interception of precipitation can be used for water storage, whereas in the understory stemflow is more pronounced and contributes to the water supply of the bryophytes. It has been estimated that in tropical forests ~~the stemflow water could provide~~ up to 4 % of the annual rainfall amount could be converted into stemflow (Lloyd and Marques F, 1988; Marin et al., 2000; van Stan and Gordon, 2018), corresponding to maximum values of 68 and 75 mm in the years 2015 and 2016, respectively, at the ATTO site.

The water content data at 8 m and 23 m height showed diel fluctuations, which were particularly regular during the dry season, due to rare interfering rain events (Fig. S5c). They showed a parallel behavior to the RH data with

the highest values reached during the morning hours (Fig. 11). It is well known from the literature that moist bryophytes and many cryptogams could utilize high air humidity as a source of water (Lange et al., 2001; Raggio et al., 2017) and this likely also occurs here.

In addition, also condensation and fog need to be considered as potential additional sources of water for epiphytic covers as well as for near-stem vegetation at the forest floor (Lakatos et al., 2012; van Stan and Gordon, 2018; León-Vargas et al., 2006). Our data show that the necessary conditions for condensation were regularly met and occurred most frequently during the wet season at 23 m height (Fig. S11). Cases with a negative dewpoint spread, when condensation could occur, were related to increasing water contents of the bryophytes, supporting the calculated condensation data. During the occurrence of fog, an increase in water contents could not be directly proven, suggesting that fog does not represent a major water source for the bryophytes.

4.2 Potential physiological activity of bryophytes

The microenvironmental conditions influence the WC of epiphytic bryophyte communities, but the ability to deal with these conditions differs among species (interspecific variability), being determined by morphological and physiological features. Apart from the interspecific variability, the performance of a single species under differing microenvironmental conditions can also be modulated by short-term acclimation and long-term adaption processes, with the latter being driven by environmental exposure, genetic variation among populations, and plasticity, as, e.g., shown for bryophytes and lichens (Cornelissen et al., 2007; Marks et al., 2019; Pardow et al., 2010). These aspects help to understand the occurrence of bryophytes under widely varying microclimatic conditions within the canopy. During our study, we measured the microenvironmental conditions of epiphytic bryophytes and observed bryophyte taxa to vary depending on these. Additionally, we estimated the potential ranges of physiological activity based on the compensation points for light, temperature, and WC, which have been reported from other studies in tropical forests (Lösch et al., 1994; Wagner et al., 2013).

In the canopy it is essential for the cryptogams to be adapted to high light conditions and UV radiation in order to avoid cell damage by radiation (Green et al., 2005; Pardow and Lakatos, 2013; Sinha and Häder, 2008; Westberg and Kärnefelt, 1998). As high light conditions mainly occur as short light flecks in the understory, the organisms need to react rapidly and efficiently to changing light conditions to reach overall positive net photosynthesis rates. Furthermore, understory mosses and lichens show higher rates of net photosynthesis at low light conditions as compared to canopy species (Kangas et al., 2014; Lakatos et al., 2006; Wagner et al., 2013). Epiphytic organisms growing under low-light conditions in the understory are also known to have lower LCP₁ values compared to the ones in the canopy, as documented for epiphytic lichens in French Guiana (Lakatos et al., 2006).

The temperature regulates the overall velocity of metabolic processes. While it has a strong impact on the respiration, the photosynthetic light reaction is by far less affected by it (Elbert et al., 2012; Green and Proctor, 2016;

Lange et al., 1998). As the measured net photosynthesis rates are the sum of simultaneously occurring photosynthesis and respiration processes, positive net photosynthesis may still be reached at higher temperatures, if the photosynthetic capacity is high enough, whereas during the night, high temperatures could cause a major loss of carbon due to high respiration rates (Lange et al., 2000). In the course of our study, the lowest temperatures pre-
5 dominantly occurred during the night, contributing to lower respiration rates, and values were mostly below the upper TCP. Thus, the temperature did not seem to be a limiting factor for the physiological activity of epiphytic bryophytes in this environment (Fig. S1~~10~~). Similarly, Wagner and coauthors (Wagner et al., 2013) stated that the temperature likely was not a limiting factor for the overall carbon balance of the bryophytes investigated in a low- and highland rainforest in Panama.

10 Utilizing the compensation points of water, light, and temperature taken from the literature, one can make rough estimates of the time fractions when NP and DR occur at the different height levels (Table 3).

These data suggest that at the upper height level NP occurred in 1-30 % and DR in 2-52 % of the time during the wet season and in 1-24 % (NP) and in 4-45 % (DR) of the time during the dry season, respectively (Table 3). These estimates suggest that the duration of DR was about twice as long as that of NP. For the samples at 8 m height the
15 results were similar, whereas for those in the understory the duration of DR was about 5- to 30-fold higher than the duration of NP. The large discrepancy between the time ranges for NP and DR calculated for the bryophytes in the understory gives reason to expect the LCP_1 and the WCP to be at the lower end of the range ($3 \mu\text{mol m}^{-2} \text{s}^{-1}$, 30 %) for the bryophytes at the lowest height level and to be at the upper end of the range ($12 \mu\text{mol m}^{-2} \text{s}^{-1}$, 80 %) for the bryophytes at the two upper height levels. For other habitats, LCP_1 values as low as $1 \mu\text{mol m}^{-2} \text{s}^{-1}$ have
20 been defined for lichens (Green et al., 1991), and thus it could be possible that the bryophyte communities in the understory exhibit similarly low LCP_1 values.

In the environment being studied, the acclimation of the organisms to the environmental conditions is also crucial for their survival. Thus, the time ranges of metabolic activity are only rough estimates, depending on the actual compensation points, which are influenced by inter- and intraspecific variation. There are also some differences
25 between groups, as, e.g., lichens tend to perform photosynthesis at lower WCs than bryophytes, and chlorolichens (with green algae as photobionts) may utilize high air humidity, whereas cyanolichens (cyanobacteria as photobiont) need liquid water (Green et al., 2011; Lange and Kilian, 1985; Raggio et al., 2017). Furthermore, there are

also differences between the bryophyte divisions of mosses and liverworts, and also within one division the inter-specific variability can also be large.

5 Conclusions

The microclimatic conditions experienced by bryophytes are being assessed in long-term measurements at the ATTO site since October 2014. These measurements provide a unique data set of the micrometeorological conditions within the understory and the inner canopy of tropical rain forests and facilitate a rough estimation of the physiological activity patterns of epiphytic bryophytes along a vertical gradient. Within this tropical rain forest habitat, the WC ~~has turned out~~ appears to be the key parameter controlling the overall physiological activity of the organisms with major differences between organisms of the canopy and the understory. In the understory, the WC of the bryophytes responded reliably upon rain events, and after major rain events, the samples could stay wet for several days before they dried out again. In contrast to that, the WC of the bryophytes in the canopy responded only ~~very~~ rarely to rain events during the wet season, probably caused by the dense foliage, and kept relatively stable low water contents. During the dry season, ~~probably due to the less dense foliage,~~ they responded ~~to some~~ much more reliably to rain events, which is probably caused by a less dense foliage, but also an effect of sensor re-positioning cannot be completely excluded. The bryophytes at 8 m and 23 m height showed regular daily fluctuations of the WC contents, which went in parallel to RH and reached highest values during the morning hours and showed a regular nightly increase in water contents, which might be a combined effect of nightly condensation and thallus life form morphology (life form). Thus, our data suggest that the relevant water source for bryophytes in the understory ~~might be the~~ is rain, while for the bryophytes in the canopy ~~nightly RH fluctuations and dew condensation might be relevant for an activation of the physiological processes. With the current data at hand, however, it cannot be answered if the daily fluctuations and the dew condensation events are large enough to activate physiological processes; this topic, indeed, would deserve to be investigated in a separate in-depth study.~~ The light intensity during periods of physiological activity mainly determines whether NP dominates or carbon is lost by dominating respiration. As the temperature shows only minor spatial, diel, and seasonal variation relative to the physiological tolerance of the bryophytes, it seems to be of minor physiological relevance within the given habitat.

Data on the potential physiological activity of bryophytes and cryptogamic organisms in general are not only relevant for their potential role in carbon cycling, but may also provide new insights into their relevance as sources of bioaerosols and different trace gases. Thus, these data may form a baseline for studies investigating the overall relevance of cryptogams in the context of biogeochemical cycling in tropical habitats. However, the wide ranges of potential activity and the scarcity of literature data illustrate the necessity of CO₂ gas exchange measurements

to assess the actual diel and seasonal physiological activity and productivity of rain forest cryptogams under varying environmental conditions.

Data availability

5 All data are deposited in a data portal, which is accessible via the homepage of the ATTO project (<https://www.atto-project.org/>) upon request.

Supplement link

Author contribution

10 BW, CP, and NL designed the measurement setup. NL, CGGB, SB, RPA and APPF conducted the practical measurements. NL, DW, GRC, MS, AA, LRO, FD, and SMO compiled the data and conducted the analyses. All authors discussed the results. NL and BW prepared the manuscript with contributions from all co-authors.

Disclaimer

The authors declare that they have no conflict of interest.

Special issue statement

Acknowledgement

15 This work has been supported by the Max Planck Society (MPG). We would like to acknowledge the German Federal Ministry of Education and Research (BMBF contracts 01LB1001A and 01LK1602B), supporting this project as well as the construction and operation of the ATTO site. We also acknowledge the support of the Brazilian Ministério da Ciência, Tecnologia e Inovação (MCTI/FINEP contract 01.11.01248.00) as well as the Amazon State University (UEA), FAPEAM, LBA/INPA and SDS/CEUC/RDS-Uatumã for their support during construction and
20 operation of the ATTO site. Furthermore, we would like to thank Ulrich Pöschl for provision of the scientific infrastructure and the possibility to work in the labs. We would like to thank Reiner Ditz, Susan Trumbore, Alberto Quesada, Thomas Disper, Andrew Crozier, Hermes Braga Xavier, Feliciano de Souza Coelho, Josué Ferreira de Souza, Roberta Pereira de Souza, Holger Ritter, Henno Heintz, and Henning Braß for technical, logistical, and
25 scientific support within the ATTO project. NL would like to thank the Max Planck Graduate Center (MPGC) for its support. DW, CGGB, SB, RA, FD, DMZ, APPF, SW, JK, CP and BW appreciate the support by the Max Planck

5 Society. GRC would like to thank for the support of the Instituto Nacional de Pesquisas da Amazônia (INPA) provided by the Programa de Pós-graduação em Botânica. MS and LRO appreciate the support of INPA. AA would like to thank the Empresa Brasileira de Pesquisa Agropecuária (EMBRAPA), RHMG expresses his thanks to the Federal University of Parana, SMO would like to thank Stichting Het Kronendak, and MOA appreciates the support of the Max Planck Society and the University of San Diego. We would like to thank ~~Maaïke Bader and~~ two unknown referees and particularly also Maaïke Bader, whose comments helped to considerably improve the manuscript. This paper contains results of research conducted under the Technical/Scientific Cooperation Agreement between the National Institute for Amazonian Research, the State University of Amazonas, and the Max Planck Society. The opinions expressed are the entire responsibility of the authors and not of the participating institutions.

References

- Andreae, M. O., Acevedo, O. C., Araùjo, A., Artaxo, P., Barbosa, C. G. G., Barbosa, H. M. J., Brito, J., Carbone, S., Chi, X., Cintra, B. B. L., Da Silva, N. F., Dias, N. L., Dias-Júnior, C. Q., Ditas, F., Ditz, R., Godoi, A. F. L., Godoi, R. H. M., Heimann, M., Hoffmann, T., Kesselmeier, J., Könemann, T., Krüger, M. L., Lavric, J. V., Manzi, A. O., Lopes, A. P., Martins, D. L., Mikhailov, E. F., Moran-Zuloaga, D., Nelson, B. W., Nölscher, A. C., Santos Nogueira, D., Piedade, M. T. F., Pöhlker, C., Pöschl, U., Quesada, C. A., Rizzo, L. V., Ro, C. U., Ruckteschler, N., Sá, L. D. A., De Oliveira Sá, M., Sales, C. B., Dos Santos, R. M. N., Saturno, J., Schöngart, J., Sörgel, M., De Souza, C. M., De Souza, R. A. F., Su, H., Targhetta, N., Tóta, J., Trebs, I., Trumbore, S., Van Eijck, A., Walter, D., Wang, Z., Weber, B., Williams, J., Winderlich, J., Wittmann, F., Wolff, S. and Yáñez-Serrano, A. M.: The Amazon Tall Tower Observatory (ATTO): Overview of pilot measurements on ecosystem ecology, meteorology, trace gases, and aerosols, *Atmos. Chem. Phys.*, 15, 10723–10776, doi:10.5194/acp-15-10723-2015, 2015.
- 5 von Arx, G., Dobbertin, M. and Rebetez, M.: Spatio-temporal effects of forest canopy on understory microclimate in a long-term experiment in Switzerland, *Agric. For. Meteorol.*, 166–167, 144–155, doi:10.1016/J.AGRFORMET.2012.07.018, 2012.
- 15 Caesar, J., Tamm, A., Ruckteschler, N., Lena Leifke, A. and Weber, B.: Revisiting chlorophyll extraction methods in biological soil crusts - Methodology for determination of chlorophyll a and chlorophyll a Cb as compared to previous methods, *Biogeosciences*, 15(5), 1415–1424, doi:10.5194/bg-15-1415-2018, 2018.
- Campos, L. V., ter Steege, H. and Uribe, J.: The epiphytic bryophyte flora of the Colombian Amazon, *Caldasia*, 37(1), 47–59, doi:10.15446/caldasia.v37n1.50980, 2015.
- 20 Chazdon, R. L. and Fetcher, N.: Light Environments of Tropical Forests, in *Physiological ecology of plants of the wet tropics: Proceedings of an International Symposium Held in Oxatepec and Los Tuxtlas, Mexico, June 29 to July 6, 1983*, edited by E. Medina, H. A. Mooney, and C. Vázquez-Yánes, pp. 27–36, Springer Netherlands, Dordrecht., 1984.
- Cornelissen, J. H. C., Lang, S. I., Soudzilovskaia, N. A. and During, H. J.: Comparative cryptogam ecology: A review of bryophyte and lichen traits that drive biogeochemistry, *Ann. Bot.*, doi:10.1093/aob/mcm030, 2007.
- 25 Cowan, I. R., Lange, O. L. and Green, T. G. A.: Carbon-dioxide exchange in lichens: determination of transport and carboxylation characteristics, *Planta*, 187(2), 282–294, doi:10.1007/BF00201952, 1992.
- Coxson, D. S., McIntyre, D. D. and Vogel, H. J.: Pulse Release of Sugars and Polyols from Canopy Bryophytes in Tropical Montane Rain Forest (Guadeloupe, French West Indies), *Biotropica*, 24, 121–133, 1992.
- 30 Cuartas, L. A., Tomasella, J., Nobre, A. D., Hodnett, M. G., Waterloo, M. J. and Múnera, J. C.: Interception water-partitioning dynamics for a pristine rainforest in Central Amazonia: Marked differences between normal and dry years, *Agric. For. Meteorol.*, 145(1–2), 69–83, doi:10.1016/J.AGRFORMET.2007.04.008, 2007.

- Dislich, R., Marília, E., Pinheiro, L. and Guimarães, M.: Corticolous liverworts and mosses in a gallery forest in Central Brazil: effects of environmental variables and space on species richness and composition, *Nov. Hedwigia*, 107(3), 385–406, 2018.
- 5 Elbert, W., Weber, B., Burrows, S., Steinkamp, J., Büdel, B., Andreae, M. O. and Pöschl, U.: Contribution of cryptogamic covers to the global cycles of carbon and nitrogen, *Nat. Geosci.*, 5, 459–462, doi:10.1038/ngeo1486, 2012.
- Frahm, J.-P.: Bryophyte phytomass in tropical ecosystems, *Bot. J. Linn. Soc.*, 104(1–3), 23–33, doi:10.1111/j.1095-8339.1990.tb02209.x, 1990.
- 10 Gaudio, N., Gendre, X., Saudreau, M., Seigner, V. and Balandier, P.: Impact of tree canopy on thermal and radiative microclimates in a mixed temperate forest: A new statistical method to analyse hourly temporal dynamics, *Agric. For. Meteorol.*, 237–238, 71–79, doi:10.1016/J.AGRFORMET.2017.02.010, 2017.
- Gimeno, T. E., Ogée, J., Royles, J., Gibon, Y., West, J. B., Burlett, R., Jones, S. P., Sauze, J., Wohl, S., Benard, C., Genty, B. and Wingate, L.: Bryophyte gas-exchange dynamics along varying hydration status reveal a significant carbonyl sulphide (COS) sink in the dark and COS source in the light, *New Phytol.*, 215(3), 965–976, doi:10.1111/nph.14584, 2017.
- 15 Gradstein, R., Churchill, S. P. and Salazar Allen, N.: Guide to bryophytes of tropical America, *Mem. N. Y. Bot. Gard.*, 86, 2001.
- Gradstein, S. R. and Salazar Allen, N.: Bryophyte diversity along an altitudinal gradient in Darién National Park, Panama, *Trop. Bryol.*, 5, 61–71, 1992.
- 20 Green, T. G. A. and Proctor, M. C. F.: Physiology of Photosynthetic Organisms Within Biological Soil Crusts: Their Adaptation, Flexibility, and Plasticity, in *Biological Soil Crusts: An Organizing Principle in Drylands*, edited by B. Weber, B. Büdel, and J. Belnap, pp. 347–381, Springer International Publishing, Cham., 2016.
- Green, T. G. A., Kilian, E. and Lange, O. L.: *Pseudocyphellaria dissimilis*: a desiccation-sensitive, highly shade-adapted lichen from New Zealand, *Oecologia*, 85(4), 498–503, doi:10.1007/BF00323761, 1991.
- 25 Green, T. G. A., Kulle, D., Pannewitz, S., Sancho, L. G. and Schroeter, B.: UV-A protection in mosses growing in continental Antarctica, *Polar Biol.*, 28(11), 822–827, doi:10.1007/s00300-005-0011-7, 2005.
- Green, T. G. A., Sancho, L. G. and Pintado, A.: Ecophysiology of Desiccation/Rehydration Cycles in Mosses and Lichens, in *Plant Desiccation Tolerance*, edited by U. Lüttge, E. Beck, and D. Bartels, pp. 89–120, Springer Berlin Heidelberg, Berlin, Heidelberg., 2011.
- 30 Hargis, H., Gotsch, S. G., Porada, P., Moore, G. W., Ferguson, B. and Van Stan, J. T.: Arboreal Epiphytes in the Soil-Atmosphere Interface : How Often Are the Biggest “ Buckets ” in the Canopy Empty ?, *Geosciences*, 9(342), 1–17, 2019.
- Kangas, L., Maanavilja, L., Hájek, T., Juurola, E., Chimner, R. A., Mehtätalo, L. and Tuittila, E. S.: Photosynthetic traits of *Sphagnum* and feather moss species in undrained, drained and rewetted boreal spruce swamp forests, *Ecol.*

- Evol., 4(4), 381–396, doi:10.1002/ece3.939, 2014.
- Komposch, H. and Hafellner, J.: Diversity and vertical distribution of lichens in a venezuelan tropical lowland rain forest, *Selbyana*, 21(1,2), 11–24, 2000.
- Kruijt, B., Malhi, Y., Lloyd, J., Norbre, A. D., Miranda, A. C., Pereira, M. G. P., Culf, A. and Grace, J.: Turbulence Statistics Above And Within Two Amazon Rain Forest Canopies, *Boundary-Layer Meteorol.*, 94(2), 297–331, doi:10.1023/A:1002401829007, 2000.
- Lakatos, M., Rascher, U. and Büdel, B.: Functional characteristics of corticolous lichens in the understory of a tropical lowland rain forest, *New Phytol.*, doi:10.1111/j.1469-8137.2006.01871.x, 2006.
- Lakatos, M., Obregón, A., Büdel, B. and Bendix, J.: Midday dew - an overlooked factor enhancing photosynthetic activity of corticolous epiphytes in a wet tropical rain forest, *New Phytol.*, doi:10.1111/j.1469-8137.2011.04034.x, 2012.
- Lancaster, J., Lancaster, N. and Seely, M.: Climate of the Central Namib Desert, *Madoqua*, 14, 5–61, 1984.
- Lange, O. L.: Moisture content and CO₂ exchange of lichens, *Oecologica*, 45(1), 82–87, 1980.
- Lange, O. L. and Kilian, E.: Reaktivierung der Photosynthese trockener Flechten durch Wasserdampfaufnahme aus dem Luftraum: Artspezifisch unterschiedliches Verhalten, *Flora*, 176, 7–23, doi:10.1016/S0367-2530(17)30100-7, 1985.
- Lange, O. L. and Tenhunen, J. D.: Moisture Content and CO₂ Exchange of Lichens. II. Depression of Net Photosynthesis in *Ramalina maciformis* at High Water Content is Caused by Increased Thallus Carbon Dioxide Diffusion Resistance, *Oecologia (Berl.)*, 51, 426–429, 1981.
- Lange, O. L., Green, T. G. A., Lange, O. L. and Green, T. G. A.: High thallus water content severely limits photosynthetic carbon gain of central European epilithic lichens under natural conditions, *Oecologia*, 108, 13–20, 1996.
- Lange, O. L., Belnap, J. and Reichenberger, H.: Photosynthesis of the cyanobacterial soil-crust lichen *Collema tenax* from arid lands in southern Utah, USA: Role of water content on light and temperature responses of CO₂ exchange, *Funct. Ecol.*, doi:10.1046/j.1365-2435.1998.00192.x, 1998.
- Lange, O. L., Büdel, B., Meyer, A., Zellner, H. and Zotz, G.: Lichen carbon gain under tropical conditions : water relations and CO₂ exchange of three *Leptogium* species of a lower montane rainforest in Panama, *Flora - Morphol. Distrib. Funct. Ecol. Plants*, 195, 172–190, doi:10.1016/S0367-2530(17)30965-9, 2000.
- Lange, O. L., Green, T. G. A. and Heber, U.: Hydration-dependent photosynthetic production of lichens: what do laboratory studies tell us about field performance?, *J. experim.*, 52(303), 2033–2042, 2001.
- Lange, O. L., Allan Green, T. G., Melzer, B., Meyer, A. and Zellner, H.: Water relations and CO₂ exchange of the terrestrial lichen *Teloschistes capensis* in the Namib fog desert: Measurements during two seasons in the field and under controlled conditions, *Flora - Morphol. Distrib. Funct. Ecol. Plants*, 201(4), 268–280, doi:10.1016/J.FLORA.2005.08.003, 2006.

- León-Vargas, Y., Engwald, S. and Proctor, M. C. F.: Microclimate, light adaptation and desiccation tolerance of epiphytic bryophytes in two Venezuelan cloud forests, *J. Biogeogr.*, doi:10.1111/j.1365-2699.2006.01468.x, 2006.
- Lloyd, C. R. and Marques F, A. de O.: Spatial variability of throughfall and stemflow measurements in Amazonian rainforest, *Agric. For. Meteorol.*, 42, 63–73, 1988.
- 5 Lopes, A. P., Nelson, B. W., Wu, J., Graça, P. M. L. de A., Tavares, J. V., Prohaska, N., Martins, G. A. and Saleska, S. R.: Leaf flush drives dry season green-up of the Central Amazon, *Remote Sens. Environ.*, 182, 90–98, doi:10.1016/j.rse.2016.05.009, 2016.
- Lösch, R., Mülders, P., Fischer, E. and Frahm, J. P.: Scientific Results of the BRYOTROP Expedition to Zaire and 3 . Photosynthetic gas exchange of bryophytes from different forest types in eastern Central Africa ., *Trop. Bryol.*, 10 9, 169–185, 1994.
- Marin, C. T., Bouten, W. and Sevink, J.: Gross rainfall and its partitioning into throughfall , stemflow and evaporation of intercepted water in four forest ecosystems in western Amazonia, *J. Hydrol.*, 237, 40–57, 2000.
- Marks, R. A., Pike, B. D. and Nicholas McLetchie, D.: Water stress tolerance tracks environmental exposure and exhibits a fluctuating sexual dimorphism in a tropical liverwort, *Oecologia*, 191(4), 791–802, doi:10.1007/s00442-15 019-04538-2, 2019.
- Martin, S. T., Andreae, M. O., Artaxo, P., Baumgardner, D., Chen, Q., Goldstein, A. H., Guenther, A., Heald, C. L., Mayol-Bracero, O. L., McMurry, P. H., Pauliquevis, T., Pöschl, U., Prather, K. A., Roberts, G. C., Saleska, S. R., Dias, M. A. S., Spracklen, D. V, Swietlicki, E. and Trebs, I.: Sources and properties of Amazonian aerosol particles, *Rev. Geophys.*, 48(2), doi:10.1029/2008RG000280, 2010.
- 20 McWilliam, A.-L. C., Roberts, J. M., Cabral, O., Leitao, M. V. B. R., de Costa, A., Maitelli, G. T. and Zamparoni, C. A. G. P.: Leaf Area Index and Above-Ground Biomass of terra firme Rain Forest and Adjacent Clearings in Amazonia, *Funct. Ecol.*, 7(3), 310–317, 1993.
- Mendieta-Leiva, G., Porada, P. and Bader, M. Y.: Interactions of Epiphytes with Precipitation Partitioning, in *Precipitation Partitioning by Vegetation: A Global Synthesis*, edited by J. T. Van Stan II, E. Gutmann, and J. 25 Friesen, pp. 133–146, Springer International Publishing, Cham., 2020.
- Mota de Oliveira, S., ter Steege, H., Cornelissen, J. H. C. and Gradstein, S. R.: Niche assembly of epiphytic bryophyte communities in the Guianas: A regional approach, *J. Biogeogr.*, doi:10.1111/j.1365-2699.2009.02144.x, 2009.
- Normann, F., Weigelt, P., Gehrig-Downie, C., Gradstein, S. R., Sipman, H. J. M., Obregon, A. and Bendix, J.: 30 Diversity and vertical distribution of epiphytic macrolichens in lowland rain forest and lowland cloud forest of French Guiana, *Ecol. Indic.*, 10(6), 1111–1118, doi:10.1016/J.ECOLIND.2010.03.008, 2010.
- Oliver, M. J., Velten, J. and Mishler, B. D.: Desiccation Tolerance in Bryophytes: A Reflection of the Primitive Strategy for Plant Survival in Dehydrating Habitats?, *INTEGR. COMP. BIOL.*, 45, 788–799, 2005.
- Pardow, A. and Lakatos, M.: Desiccation Tolerance and Global Change: Implications for Tropical Bryophytes in

- Lowland Forests, *Biotropica*, 45(1), 27–36, doi:10.1111/j.1744-7429.2012.00884.x, 2013.
- Pardow, A., Hartard, B. and Lakatos, M.: Morphological, photosynthetic and water relations traits underpin the contrasting success of two tropical lichen groups at the interior and edge of forest fragments, *AoB Plants*, 2010, 1–12, doi:10.1093/aobpla/plq004, 2010.
- 5 Piepenbring, M.: Inventoring the fungi of Panama, *Biodivers. Conserv.*, 16(1), 73–84, doi:10.1007/s10531-006-9051-8, 2007.
- Pinheiro da Costa, D.: Epiphytic Bryophyte Diversity in Primary and Secondary Lowland Rainforests in Southeastern Brazil, *Bryologist*, 102(2), 320–326, 1999.
- Pöhlker, M. L., Pöhlker, C., Ditas, F., Klimach, T., De Angelis, I. H., Araújo, A., Brito, J., Carbone, S., Cheng, Y.,
10 Chi, X., Ditz, R., Gunthe, S. S., Kesselmeier, J., Könemann, T., Lavrič, J. V., Martin, S. T., Mikhailov, E., Moran-Zuloaga, D., Rose, D., Saturno, J., Su, H., Thalman, R., Walter, D., Wang, J., Wolff, S., Barbosa, H. M. J., Artaxo, P., Andreae, M. O. and Pöschl, U.: Long-term observations of cloud condensation nuclei in the Amazon rain forest - Part 1: Aerosol size distribution, hygroscopicity, and new model parametrizations for CCN prediction, *Atmos. Chem. Phys.*, 16, 15709–15740, doi:10.5194/acp-16-15709-2016, 2016.
- 15 Porada, P., Tamm, A., Raggio, J., Cheng, Y., Kleidon, A., Pöschl, U. and Weber, B.: Global NO and HONO emissions of biological soil crusts estimated by a process-based non-vascular vegetation model, *Biogeosciences*, 16(9), 2003–2031, doi:10.5194/bg-16-2003-2019, 2019.
- Proctor, M. C. F.: The bryophyte paradox: Tolerance of desiccation, evasion of drought, *Plant Ecol.*, 151, 41–49, doi:10.1023/A:1026517920852, 2000.
- 20 Proctor, M. C. F., Oliver, M. J., Wood, A. J., Alpert, P., Stark, L. R., Cleavitt, N. L. and Mishler, B. D.: Desiccation-tolerance in bryophytes: a review, *Bryologist*, 110, 595–621, 2007.
- Raggio, J., Green, T. G. A., Sancho, L. G., Pintado, A., Colesie, C., Weber, B. and Büdel, B.: Metabolic activity duration of biological soil crusts across Europe can be predicted from common climate data, *Geoderma*, 306, 10–17, doi:10.1016/j.geoderma.2017.07.001, 2017.
- 25 Rastogi, B., Berkelhammer, M., Wharton, S., Whelan, M. E., Itter, M. S., Leen, J. B., Gupta, M. X., Noone, D. and Still, C. J.: Large Uptake of Atmospheric OCS Observed at a Moist Old Growth Forest: Controls and Implications for Carbon Cycle Applications, *J. Geophys. Res. Biogeosciences*, 123(11), 3424–3438, doi:10.1029/2018JG004430, 2018.
- 30 Reiter, R., Höftberger, M., G. Allan Green, T. and Türk, R.: Photosynthesis of lichens from lichen-dominated communities in the alpine/nival belt of the Alps – II: Laboratory and field measurements of CO₂ exchange and water relations, *Flora - Morphol. Distrib. Funct. Ecol. Plants*, 203, 34–46, 2008.
- de Ribeiro, A.: Local rainfall variability - a potential bias for bioecological studies in the central Amazon, *Acta Amaz.*, 14, 159–174, 1984.
- Richards, P. W.: Notes on the Bryophyte Communities of Lowland Tropical Rain Forest, with Special Reference

- to Moraballi Creek , British Guiana, *Vegetatio*, 5(1), 319–328, 1954.
- Rodriguez-Iturbe, I., D’Odorico, P., Porporato, A. and Ridolfi, L.: On the spatial and temporal links between vegetation, climate, and soil moisture, *Water Resour. Res.*, doi:10.1029/1999WR900255, 1999.
- Romero, C., Putz, F. E. and Kitajima, K.: Ecophysiology in relation to exposure of pendant epiphytic bryophytes
5 in the canopy of a tropical montane oak forest, *Biotropica*, doi:10.1111/j.1744-7429.2006.00099.x, 2006.
- Seel, W. E., Hendry, G. A. F. and Lee, J. A.: The combined effects of desiccation and irradiance on mosses from xeric and hydric habitats, *J. Exp. Bot.*, doi:10.1093/jxb/43.8.1023, 1992.
- Sinha, R. P. and Häder, D. P.: UV-protectants in cyanobacteria, *Plant Sci.*, doi:10.1016/j.plantsci.2007.12.004, 2008.
- 10 Snelgar, W. P., Green, T. G. A. and Wilkins, A. L.: Carbon dioxide exchange in lichens: resistances to CO₂ uptake different thallus water contents, *New Phytol.*, 88(6), 353–361, 1981.
- Soepadmo, E.: Tropical rain forests as carbon sinks, *Chemosphere*, 27(6), 1025–1039, doi:10.1016/0045-6535(93)90066-E, 1993.
- Stace, C. A.: Combretaceae, in *The Families and Genera of Vascular Plants. Flowering Plants. Eudicots*, vol. 9,
15 edited by K. Kubitzki, pp. 67–82, Springer Berlin Heidelberg., 2007.
- van Stan, J. T. and Gordon, D. A.: Mini-Review: Stemflow as a Resource Limitation to Near-Stem Soils, *Frontiers Plant Sci.*, 9(February), 1–7, doi:10.3389/fpls.2018.00248, 2018.
- ter Steege, H., Pitman, N. C. A., Sabatier, D., Baraloto, C., Salomão, R. P., Guevara, J. E., Phillips, O. L., Castilho, C. V., Magnusson, W. E., Molino, J. F., Monteagudo, A., Vargas, P. N., Montero, J. C., Feldpausch, T. R.,
20 Coronado, E. N. H., Killeen, T. J., Mostacedo, B., Vasquez, R., Assis, R. L., Terborgh, J., Wittmann, F., Andrade, A., Laurance, W. F., Laurance, S. G. W., Marimon, B. S., Marimon, B. H., Vieira, I. C. G., Amaral, I. L., Brien, R., Castellanos, H., López, D. C., Duivenvoorden, J. F., Mogollón, H. F., Matos, F. D. D. A., Dávila, N., García-Villacorta, R., Diaz, P. R. S., Costa, F., Emilio, T., Levis, C., Schiatti, J., Souza, P., Alonso, A., Dallmeier, F.,
Montoya, A. J. D., Piedade, M. T. F., Araujo-Murakami, A., Arroyo, L., Gribel, R., Fine, P. V. A., Peres, C. A.,
25 Toledo, M., Aymard C., G. A., Baker, T. R., Cerón, C., Engel, J., Henkel, T. W., Maas, P., Petronelli, P., Stropp, J., Zartman, C. E., Daly, D., Neill, D., Silveira, M., Paredes, M. R., Chave, J., Lima Filho, D. D. A., Jørgensen, P. M., Fuentes, A., Schöngart, J., Valverde, F. C., Di Fiore, A., Jimenez, E. M., Mora, M. C. P., Phillips, J. F., Rivas, G., Van Andel, T. R., Von Hildebrand, P., Hoffman, B., Zent, E. L., Malhi, Y., Prieto, A., Rudas, A., Ruschell, A. R., Silva, N., Vos, V., Zent, S., Oliveira, A. A., Schutz, A. C., Gonzales, T., Nascimento, M. T., Ramirez-Angulo, H., Sierra, R., Tirado, M., Medina, M. N. U., Van Der Heijden, G., Vela, C. I. A., Torre, E. V., Vriesendorp, C., et al.: Hyperdominance in the Amazonian tree flora, *Science* (80-.), 342(1243092), doi:10.1126/science.1243092, 2013.
- Thompson, O. E. and Pinker, R. T.: Wind and temperature profile characteristics in a tropical evergreen forest in Thailand, *Tellus*, 27(6), 562–573, doi:10.1111/j.2153-3490.1975.tb01711.x, 1975.

- Tucker, C. L., McHugh, T. A., Howell, A., Gill, R., Weber, B., Belnap, J., Grote, E. and Reed, S. C.: The concurrent use of novel soil surface microclimate measurements to evaluate CO₂ pulses in biocrusted interspaces in a cool desert ecosystem, *Biogeochemistry*, 135(3), 239–249, doi:10.1007/s10533-017-0372-3, 2017.
- Wagner, S., Zotz, G., Salazar Allen, N. and Bader, M. Y.: Altitudinal changes in temperature responses of net photosynthesis and dark respiration in tropical bryophytes, *Ann. Bot.*, 111(3), 455–465, doi:10.1093/aob/mcs267, 2013.
- Wagner, S., Bader, M. Y. and Zotz, G.: Physiological Ecology of Tropical Bryophytes, in *Photosynthesis in Bryophytes and Early Land Plants*, edited by D. T. Hanson and S. K. Rice, pp. 269–289, Springer Netherlands, Dordrecht., 2014.
- 10 Walter, H. and Stadelmann, E.: The Physiological Prerequisites for the Transition of Autotrophic Plants from Water to Terrestrial Life, *Bioscience*, 18(7), 694–701, 1968.
- Weber, B., Berkemeier, T., Ruckteschler, N., Caesar, J., Heintz, H., Ritter, H. and Braß, H.: Development and calibration of a novel sensor to quantify the water content of surface soils and biological soil crusts, *Methods Ecol. Evol.*, doi:10.1111/2041-210X.12459, 2016.
- 15 Westberg, M. and Kärnefelt, I.: The Genus *Fulgensia* A. Massal. & De Not., a Diverse Group in the Teloschistaceae, *Lichenol.*, 30(06), 515–532, doi:10.1006/lich.1998.0141, 1998.
- Williams, T. G. and Flanagan, L. B.: Effect of changes in water content on photosynthesis , transpiration and discrimination against ¹³C₂ and C₁₈O₁₆ in *Pleurozium* and *Sphagnum*, *Oecologia*, 2, 38–46, doi:10.1007/BF00333212, 1996.
- 20 Wilske, B., Holzinger, R. and Kesselmeier, J.: Evidence for ethanolic fermentation in lichens during periods of high thallus water content, *Symbiosis*, 31(1–3), 95–111, 2001.
- Zartman, C. E.: Habitat fragmentation impacts on epiphyllous bryophyte communities in central Amazonia, *Ecology*, 84(4), 948–954, doi:10.1890/0012-9658(2003)084[0948:HFIOEB]2.0.CO;2, 2003.
- Zotz, G., Büdel, B., Meyer, A., Zellner, H. and Lange, L.: Water relations and CO₂ exchange of tropical bryophytes in a lower montane rain forest in Panama, *Bot. Acta*, 110, 9–17, doi:10.1111/j.1438-8677.1997.tb00605.x, 1997.
- 25 Zotz, G., Schultz, S. and Rottenberger, S.: Are tropical lowlands a marginal habitat for macrolichens? Evidence from a field study with *Parmotrema endosulphureum* in Panama, *Flora - Morphol. Distrib. Funct. Ecol. Plants*, 198(1), 71–77, doi:10.1078/0367-2530-00077, 2003.

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Tables

Table 1: Annual mean values and standard deviations (\pm SD) of mean daytime photosynthetically active radiation (PAR_{avg}), temperature, and water contents (WC) of bryophytes at the three height levels and above the canopy (a). Annual sum of rain and fog days as well as the annual sum of rain (b). Mean values were calculated from 30-minute intervals. Due to data gaps in the measured rain (shown in brackets), missing values were also extrapolated from existing data as described in methods section (values behind the brackets). Values for PAR_{max} can be found in Table S6.

(a)

Height	2015		2016	
	Mean	SD	Mean	SD
PAR_{avg} daytime [$\mu\text{mol m}^{-2} \text{s}^{-1}$]				
above-canopy	819	596	824	599
23 m	32	37	49	52
8 m	43	50	8	14
1.5 m	5	15	3	8
Temperature [$^{\circ}\text{C}$]				
above-canopy	26.6	3.4	26.4	3.1
23 m	26.1	3.0	26.8	3.3
8 m	25.8	2.3	25.8	1.9
1.5 m	25.4	1.8	25.5	1.5
Water content [%]; above canopy RH [%]				
above-canopy	86	15	90	13
23 m	39	30	47	69
8 m	70	45	73	72
1.5 m	64	85	38	53

10 (b)

Parameter	2015	2016
	Sum	Sum
Rain (days)	(199) 202	(197) 215
(mm)	(1680) 1693	(1702) 1863
Fog (days)	21* <u>(217)</u>	28* <u>(176)</u>

*: Gaps in the data record due to malfunction of fog sensor during time window of 31.05. – 20.10.2015, 30.04. – 06.07.2016, and 01.09. – 31.12.2016. Number in brackets are the operational days of the sensor.

Table 2 Seasonal mean values and standard deviations (\pm SD) of the mean photosynthetically active radiation (PAR_{avg}), the temperature, and the above-canopy relative humidity (RH) or water content (WC) of bryophytes determined at different height levels and above the canopy. Mean values for the respective seasons were calculated from 30-minute intervals from October 2014 to November 2016. Values for PAR_{max} can be found in Table S7.

Height [m]	PAR_{avg} daytime [$\mu\text{mol m}^{-2} \text{s}^{-1}$]		Temperature [$^{\circ}\text{C}$]		RH (above-canopy) [%], WC [%]	
	Mean	\pm SD	Mean	\pm SD	Mean	\pm SD
Wet season						
above-canopy	738	46	25.7	0.7	94	2
23 m	27	17	25.3	0.6	41	3
8m	41	24	24.9	0.4	93	21
1.5	3	1	24.9	0.4	83	26
Transitional season Wet-Dry						
above-canopy	860	53	25.6	0.5	91	2
23 m	38	29	25.7	0.7	49	4
8 m	63	14	24.9	0.4	72	27
1.5 m	2	1	24.6	0.2	31	6
Dry season						
above-canopy	950	93	27.2	1.0	84	6
23 m	54	21	27.8	1.2	45	10
8 m	24	17	26.6	0.9	58	20
1.5 m	5	4	26.0	0.8	30	31
Transitional season Dry-Wet						
above-canopy	784	111	26.5	1.6	87	8
23 m	52	34	27.1	2.2	37	2
8 m	23	5	26.2	1.7	58	13
1.5 m	4	1	25.9	1.4	52	53

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Table 3: The potential time fractions [%], during which ~~on average~~ the epiphytic bryophytes ~~samples~~ at the different height levels exceeded the lower compensation points of light (LCP_l), the upper compensation points for temperature (TCP), the lower compensation points for water (WCP), and reached the optimal temperature for net photosynthesis (T_{opt}). The results are shown separately for a) the wet season (February–May) and b) the dry season (August–November). Values are given for the different height levels (1.5, 8, 23 m) and bryophyte divisions (M=moss, L=liverwort). For the net photosynthesis (NP) it is required that WC > WCP, PAR > LCP_l and T > TCP, for the dark respiration (DR) it is necessary that WC > WCP and PAR < LCP_l or WC > WCP and T > TCP. Thirty-minute averages of measurements during the entire measurement period from October 2014 to November 2016 were considered. The data show the averages of the different samples per height zone. The ranges of the compensation points (CP) and the optimum temperature (opt) were reported in Lösch (1994) and Wagner et al. (2013) (see Table S4).

a) Wet season

Height	Division	LCP _l	T _{opt}	TCP	WCP	NP	DR
		≥ 3-12	24.0-27.0	≥ 30.0-36.0	≥ 30-80		
		μmol m ⁻² s ⁻¹	° C	° C	% DW		
[m]	L/M	Time fraction when cardinal points are reached/exceeded [% of time]					
23	L	33-43	4-5470	0-3	3-80	1-30	2-52
8	M & L	24-31	2-7488	0	42-94	14-35	29-59
1.5	M	2-19	2-7791	0	32-80	1-13	32-67

b) Dry season

Height	Division	LCP _l	T _{opt}	TCP	WCP	NP	DR
		≥ 3-12	24.0-27.0	≥ 30.0-36.0	≥ 30-80		
		μmol m ⁻² s ⁻¹	° C	° C	% DW		
[m]	L/M	Time fraction when cardinal points are reached/exceeded [% of time]					
23	L	40-46	6-3546	0-27	6-64	1-24	4-45
8	M & L	18-35	8-5166	0-11	5-84	2-34	7-51
1.5	M	3-16	9-5976	0-4	2-21	0-5	10-26

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Figures

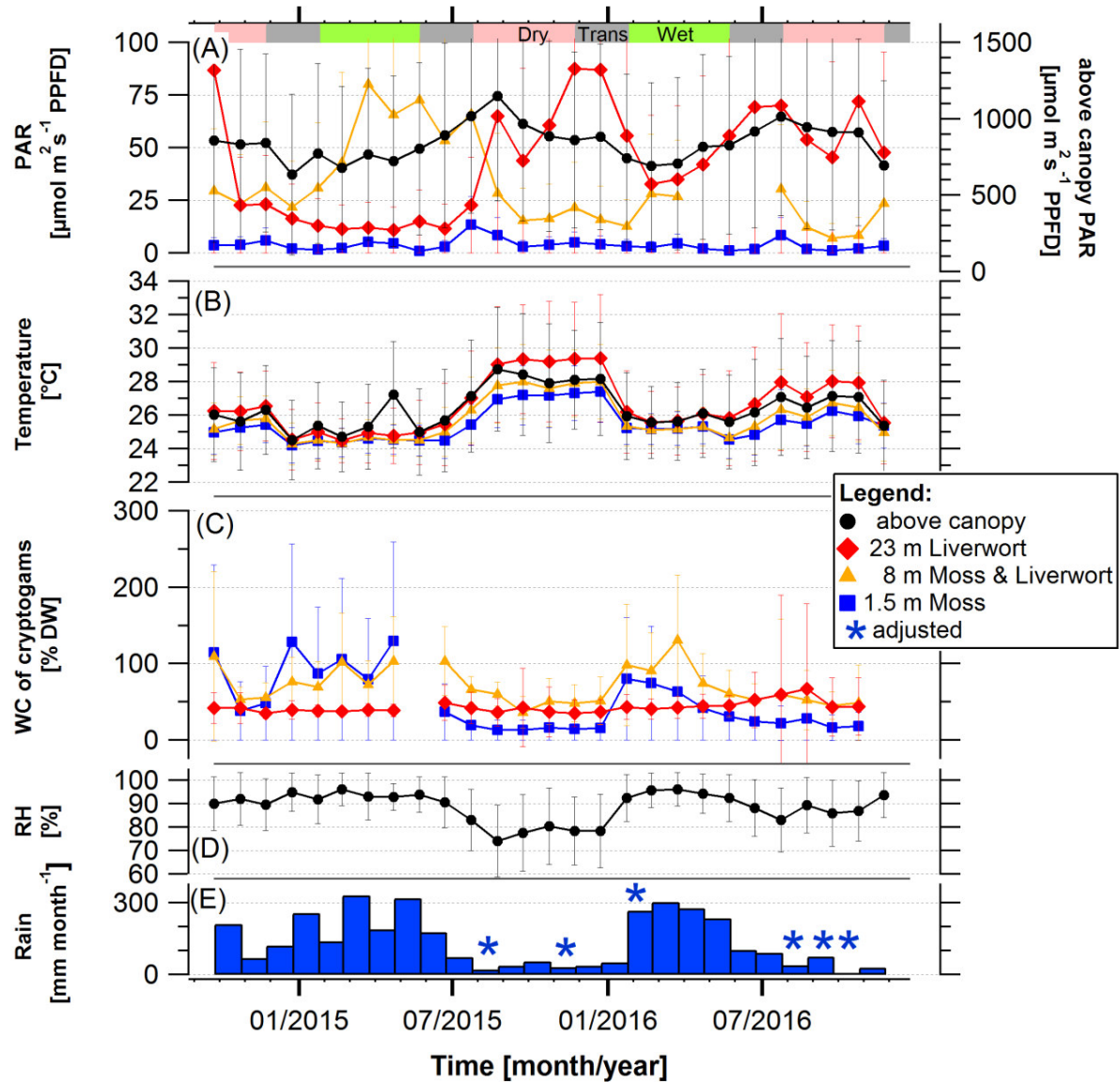
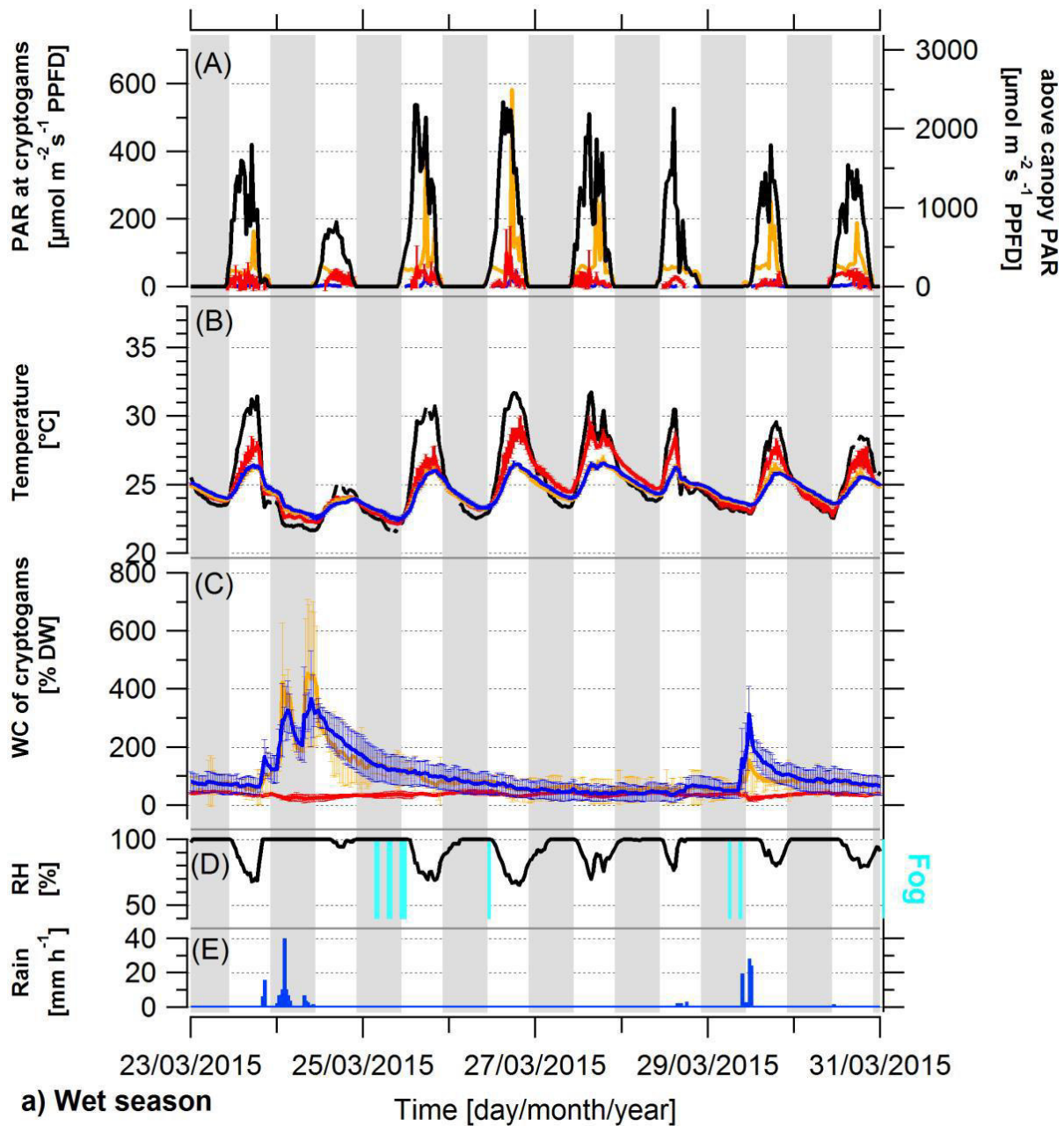


Figure 1: Mean light condition (PAR_{avg}), temperatures, and water content (WC) experienced by bryophyte communities and above-canopy meteorological conditions in the Amazonian rain forest. The micrometeorological parameters on top/within epiphytic cryptogamic communities represent monthly mean values \pm SD of (A) daily average (06:00 – 18:00 LT) of photosynthetically active radiation (PAR) on top, (B) temperature within, and (C) WC of cryptogamic communities. The above-canopy meteorological parameters comprise the (A) monthly mean value of the daily average (06:00 – 18:00 LT) of above-canopy photosynthetically active radiation (PAR at 75 m), (B) monthly mean value of above-canopy temperature (at 26 m), (D) monthly mean value of relative air humidity (RH

at 26 m height), and (E) monthly amount of rain. Data of replicate sensors installed within communities at the same height level were pooled, while above-canopy parameters were measured with one sensor each. Colored horizontal bars in the upper part of the figure indicate the seasons. Exact values and additional data are presented in Tables S3 and S5.

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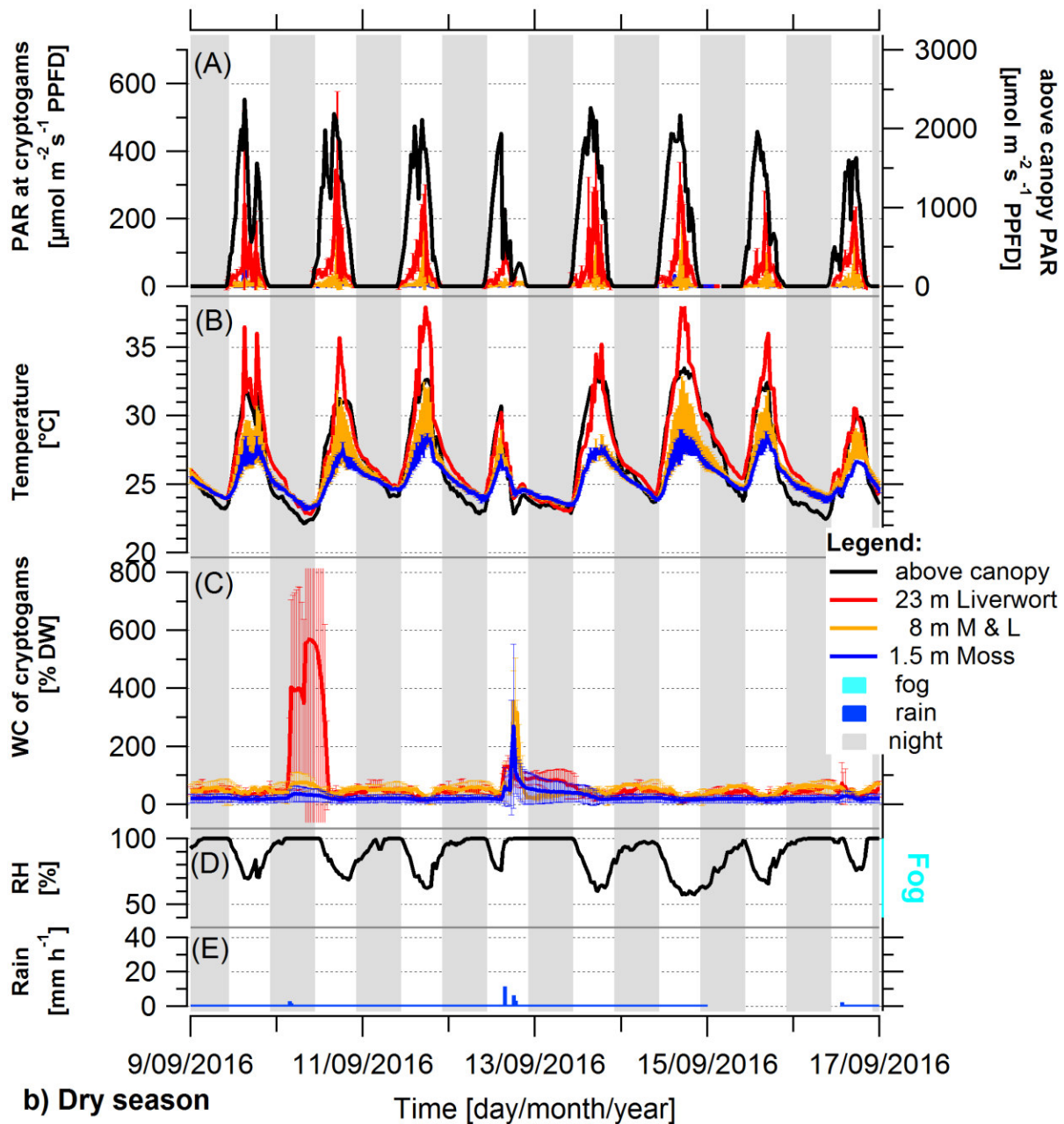


Figure 2: Representative periods during the wet and dry season under average conditions, showing light condition (PAR), temperature, and water content (WC) of bryophytes, and above-canopy meteorological conditions in the Amazonian rain forest. Shown are 8-day periods during (a) the wet season 2015 and (b) the dry season 2016. The micrometeorological parameters on top/within epiphytic cryptogamic communities represent (A) the photosynthetically active radiation (PAR) on top, (B) the temperature within, and (C) the WC of cryptogamic communities. The above-canopy meteorological parameters comprise (A) above-canopy photosynthetically active radiation

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(PAR at 75 m), (B) above-canopy temperature (at 26 m), (D) relative air humidity (RH at 26 m height), presence of fog events, and (E) rain amount. The data show 30-minute averages \pm SD except for rain, which shows hourly sums. Data of replicate sensors installed within communities at the same height level were pooled, while above-canopy parameters were measured with one sensor each. The nighttime is shaded in grey (1806:00 – 0648:00 LT).

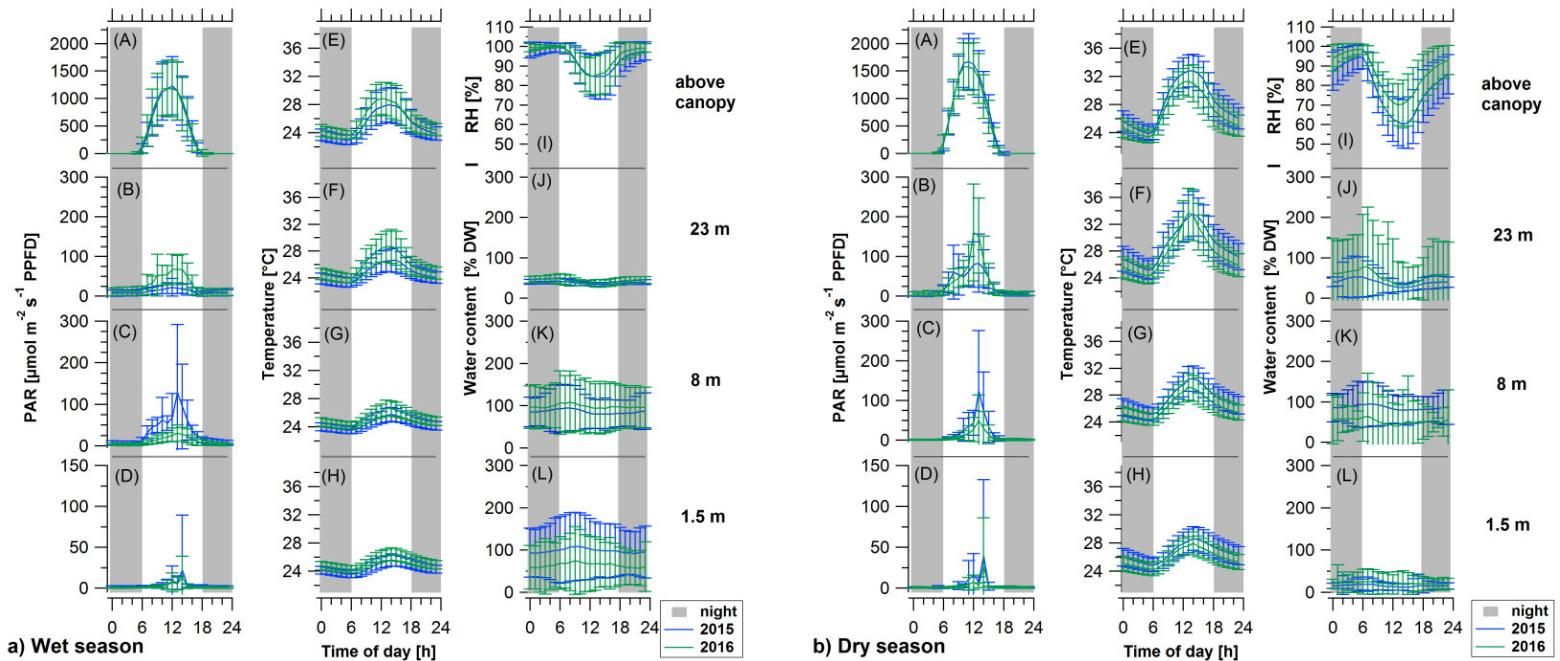
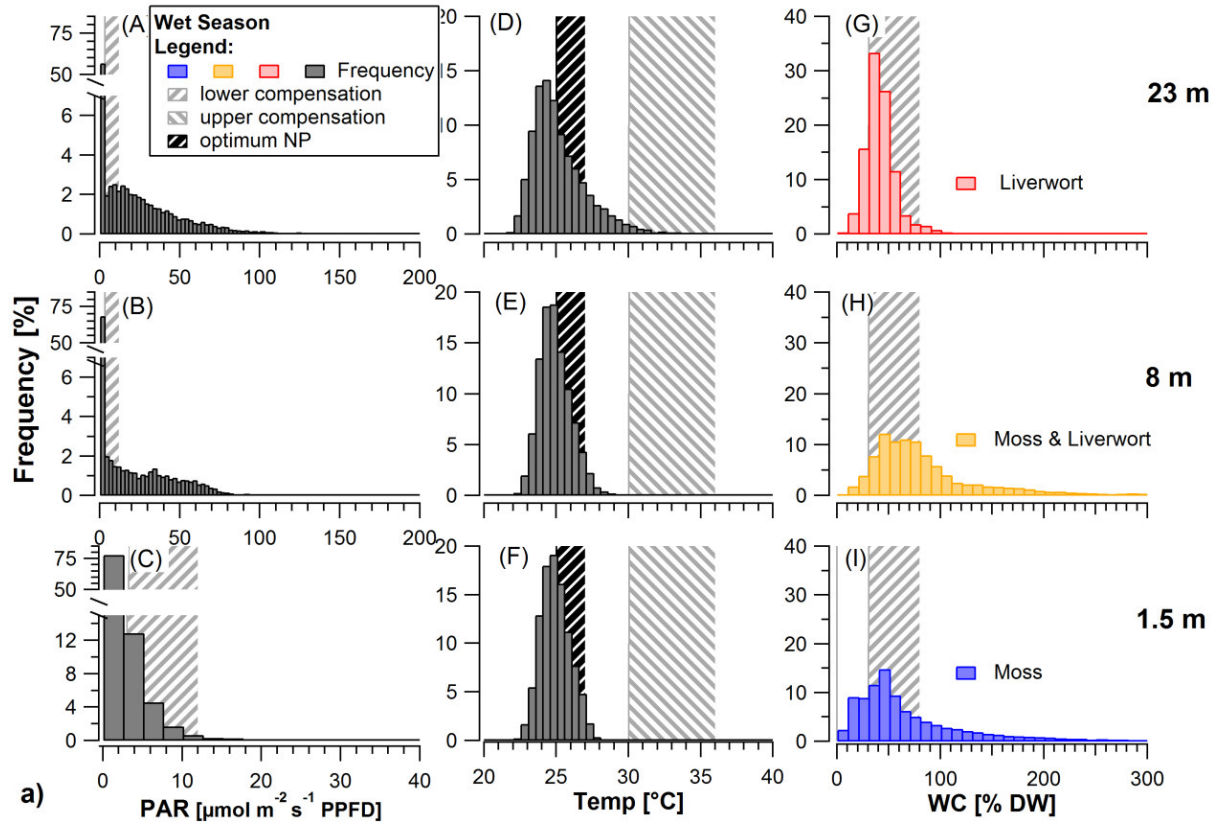


Figure 3: Mean diurnal cycles of light conditions (PAR), temperature, and water content (WC) of bryophytes, and above-canopy meteorological parameters during (a) wet season and (b) dry season of the years 2015 (blue lines) and 2016 (green lines). The above-canopy meteorological parameters comprise (A) the photosynthetically active radiation (PAR at 75 m), (E) the temperature (at 26 m), and (I) the relative air humidity (RH at 26 m height). The micrometeorological parameters measured on top/within epiphytic cryptogamic communities comprise (B – D) the photosynthetically active radiation (PAR) on top, (D – H) the temperature within, and (J – L) the WC of cryptogamic communities at different height levels. Diel cycles were calculated from 30-minute intervals of the whole seasons and show hourly mean values \pm SD. Data of the sensors installed at the same height level were pooled, while the above-canopy parameters were measured with one sensor each. Nighttime is shaded in grey (1806:00 – 1806:00 LT).



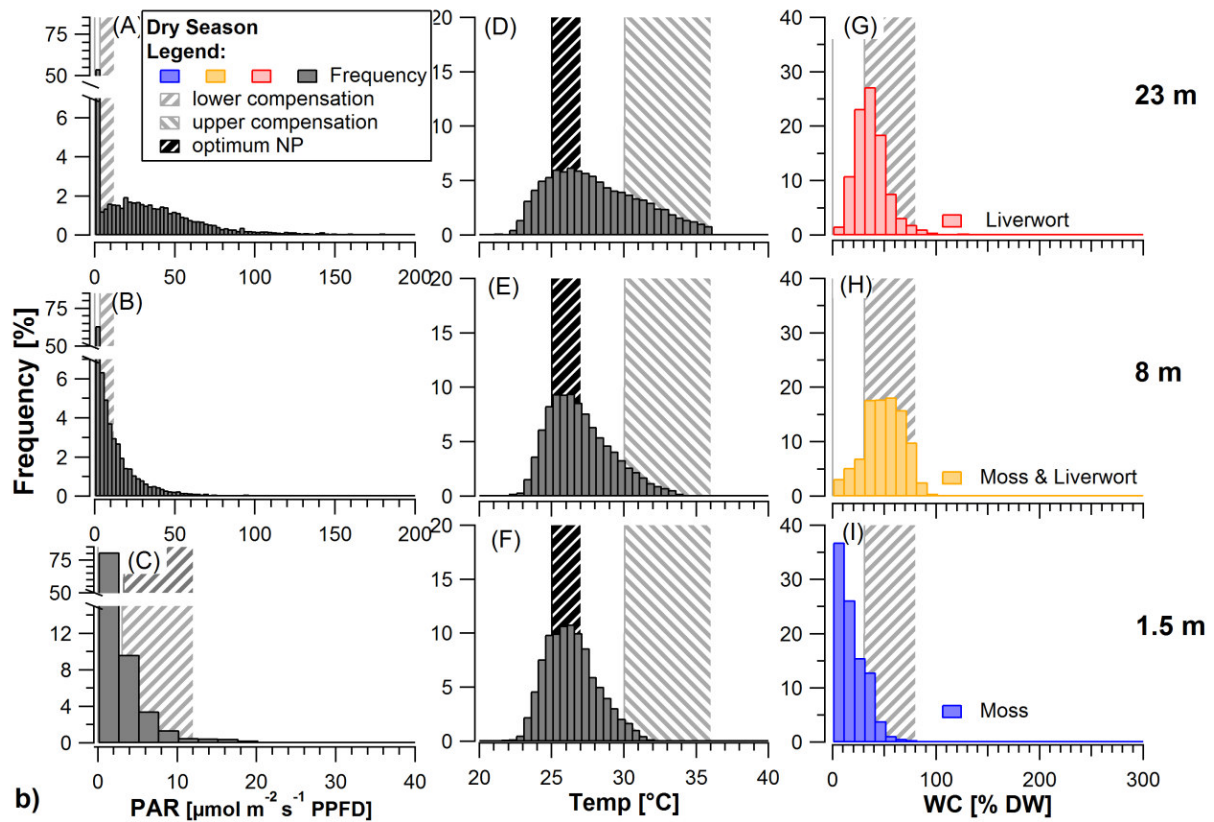
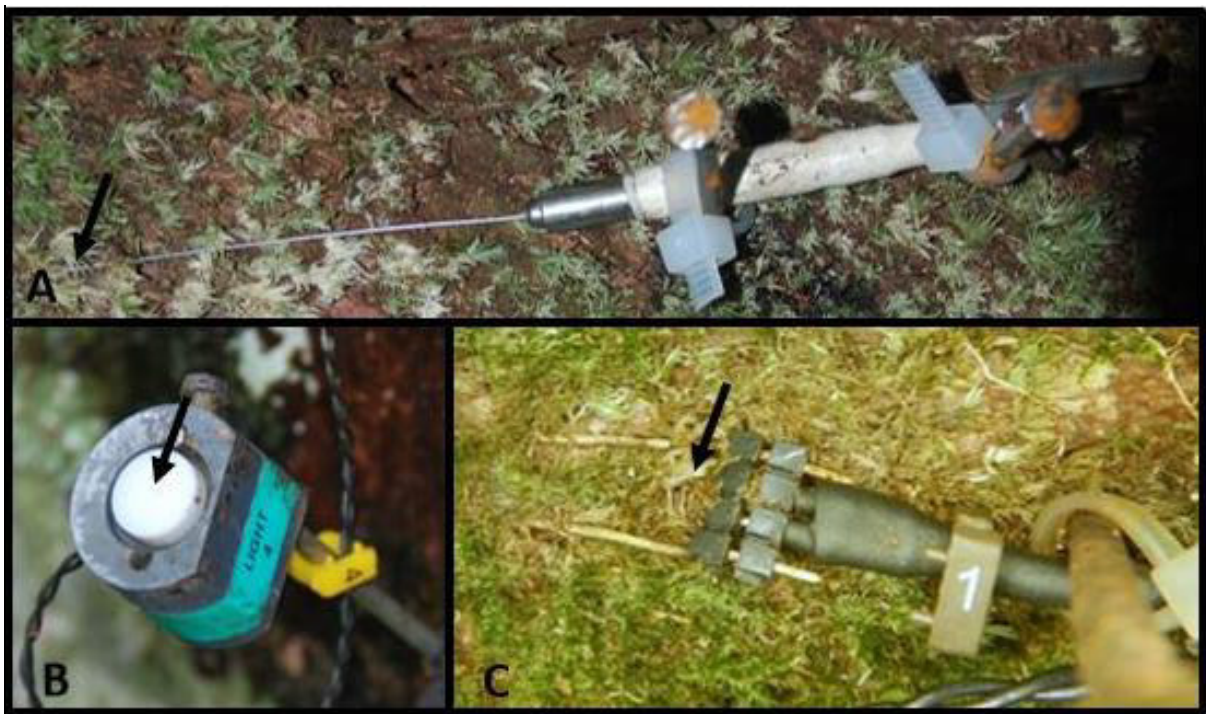


Figure 4: [Estimated frequency](#) Frequency of mean photosynthetically active radiation (PAR; A – C), temperature (Temp; D – F), and water content (WC; G – I) measured on top/within bryophytes at 1.5, 8, and 23 m height during (a) the wet and (b) the dry season. Calculation of the histograms based on 30-minute intervals. Shaded areas represent the ranges of reference values for lower compensation (PAR, WC), upper compensation (temperature), and the optimum (temperature) for net photosynthesis, as measured by Lösch (1994) and Wagner et al., (2013) (Table S4). Bin sizes: PAR: $2.5 \mu\text{mol m}^{-2} \text{s}^{-1}$; temperature: $0.5 \text{ }^\circ\text{C}$; WC: 10 %.

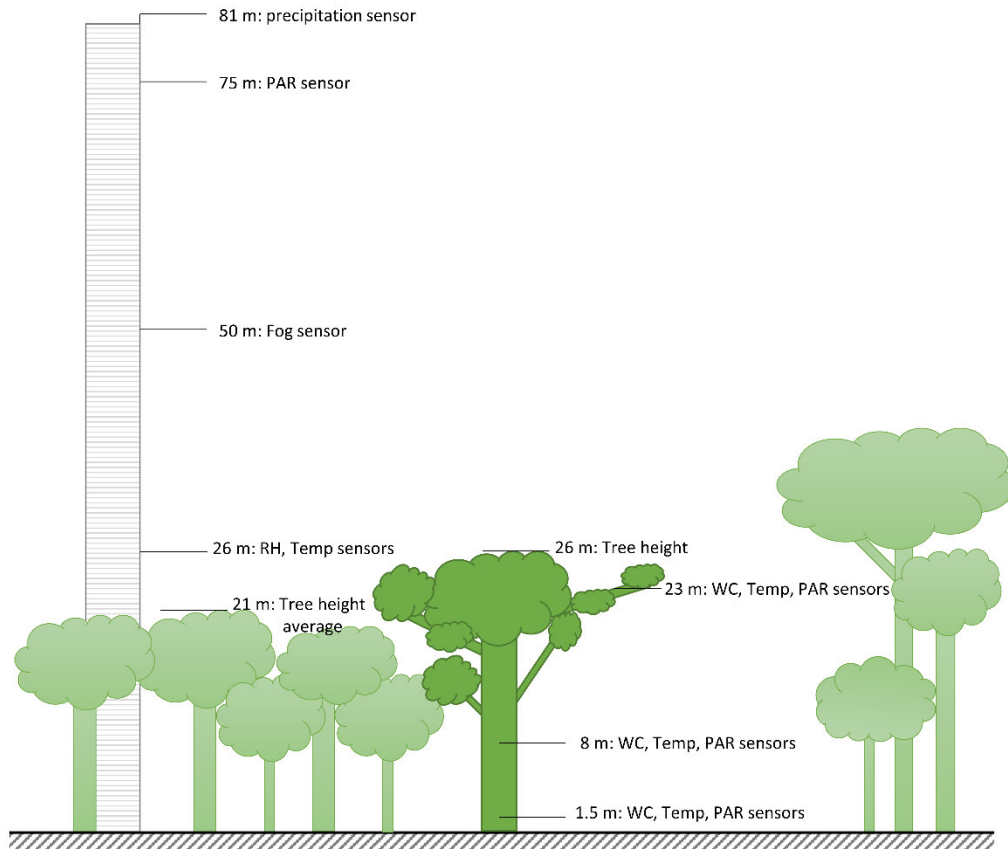
Microclimatic conditions and water content fluctuations experienced by epiphytic bryophytes in an Amazonian rain forest

5 Contents:
Figures S1 – S130
Tables S1 – S7

10



15 **Figure S1:** Examples of the temperature sensor (A), light sensor (B), and water content sensor (C) installed in epiphytic bryophytes at the ATTO site. The little arrows show the area of detection, i.e. the sensor tip of the temperature sensor, the area just below the white PTFE cap of the light sensor, and the two inner pins of the water content sensor.



5 **Figure S2:** Schematic overview of the sensors installed at different height levels below, within, and above the canopy. The parameters water content (WC) and temperature (Temp) were measured within the bryophyte samples, the light sensors (PAR) were installed directly on top of the thalli. The average tree height of 21 m was determined for the Plateau forest in general.

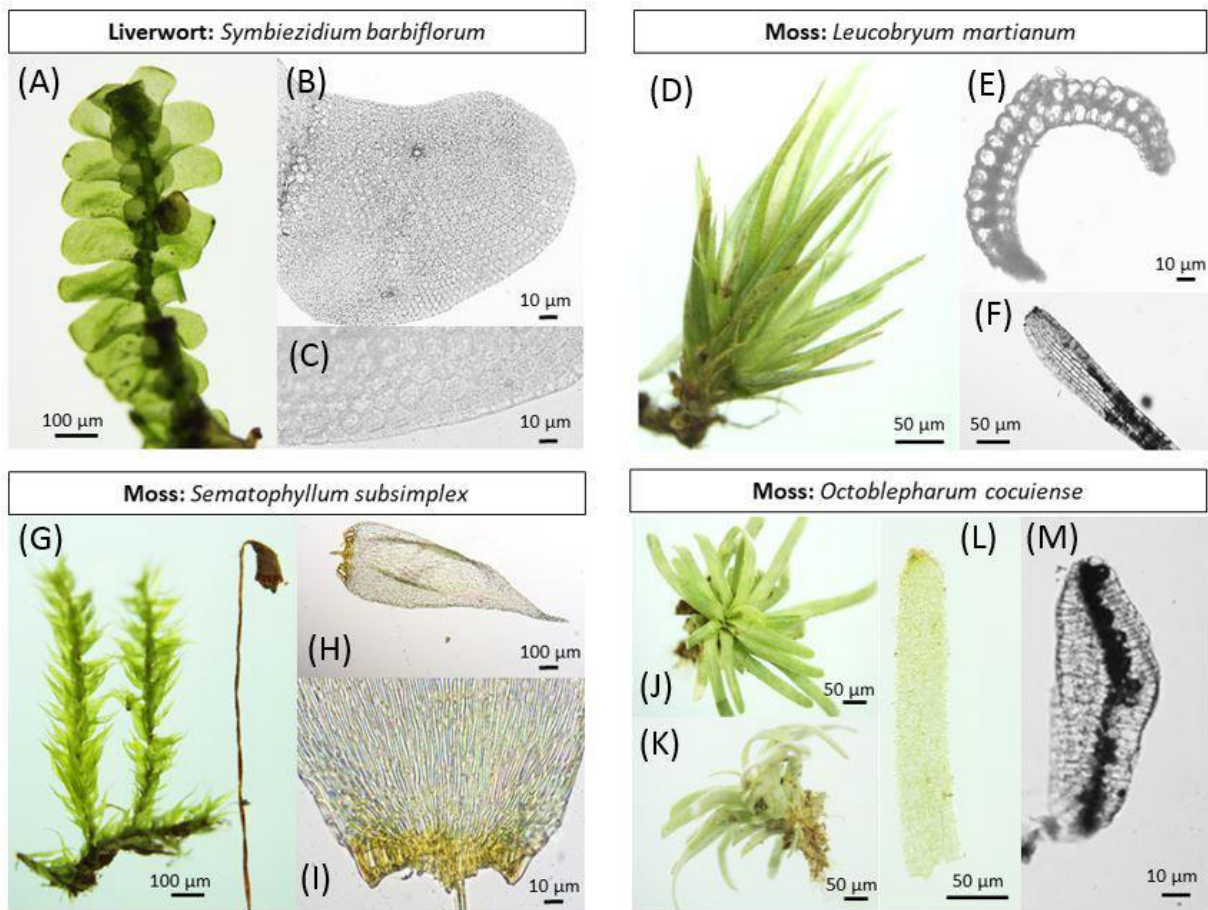


Figure S3: The four bryophyte species being used for installation of the sensors of the microclimate station. (A, D, G, J, K) overview, (B, H, L) leaf, (C, F I) cell form, and (E, M) cross section of a leaf.

5

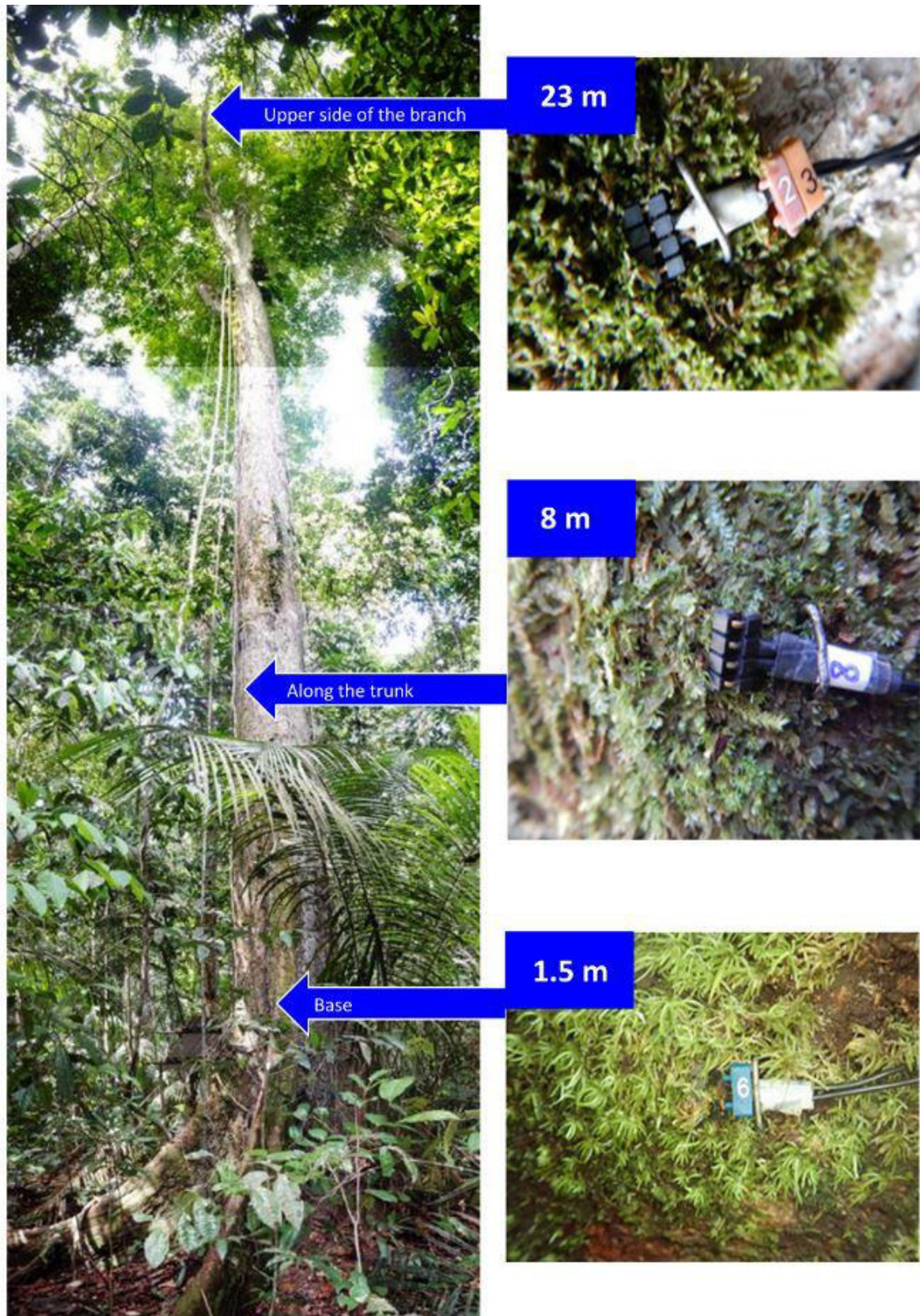
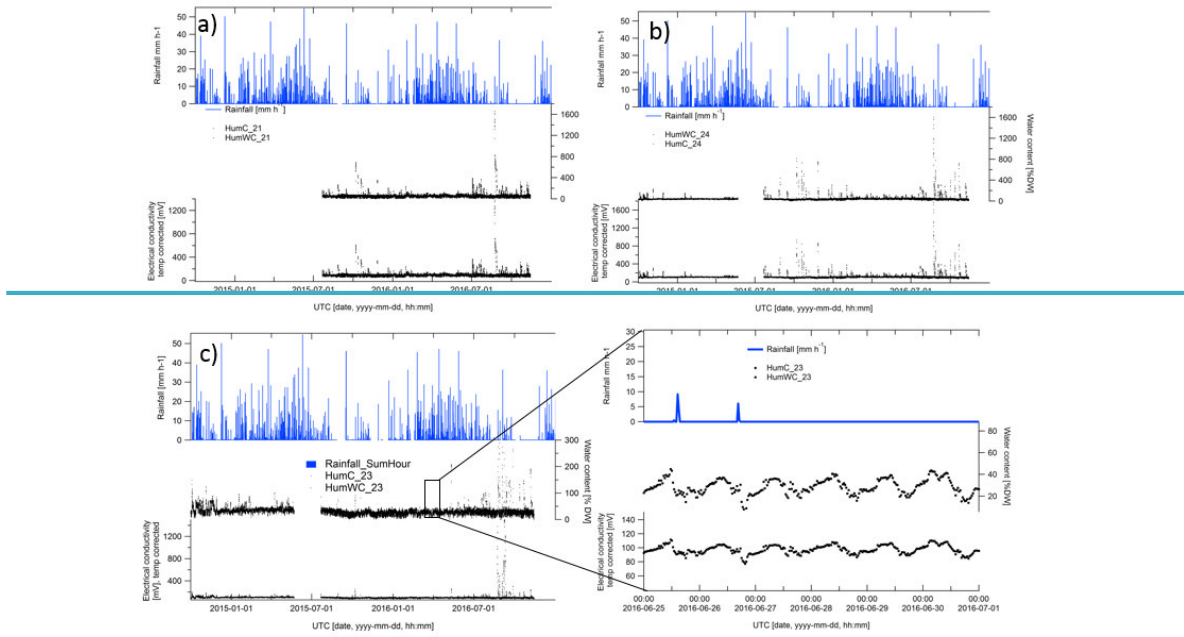
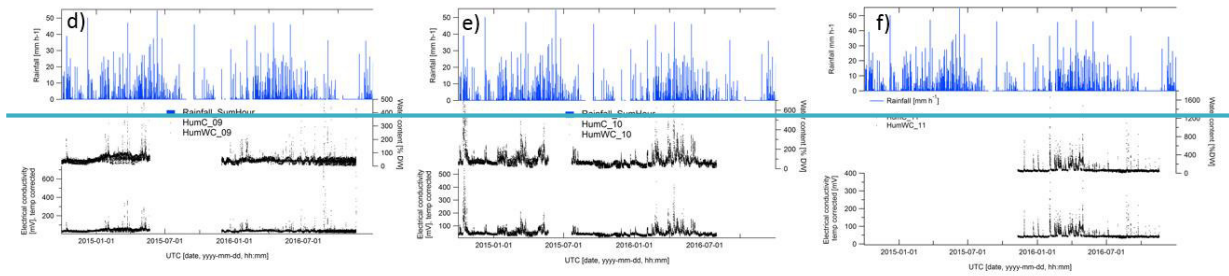


Figure S4: Overview pictures of microsensor tree and exemplary bryophyte samples with installed water content sensors at the four height levels.

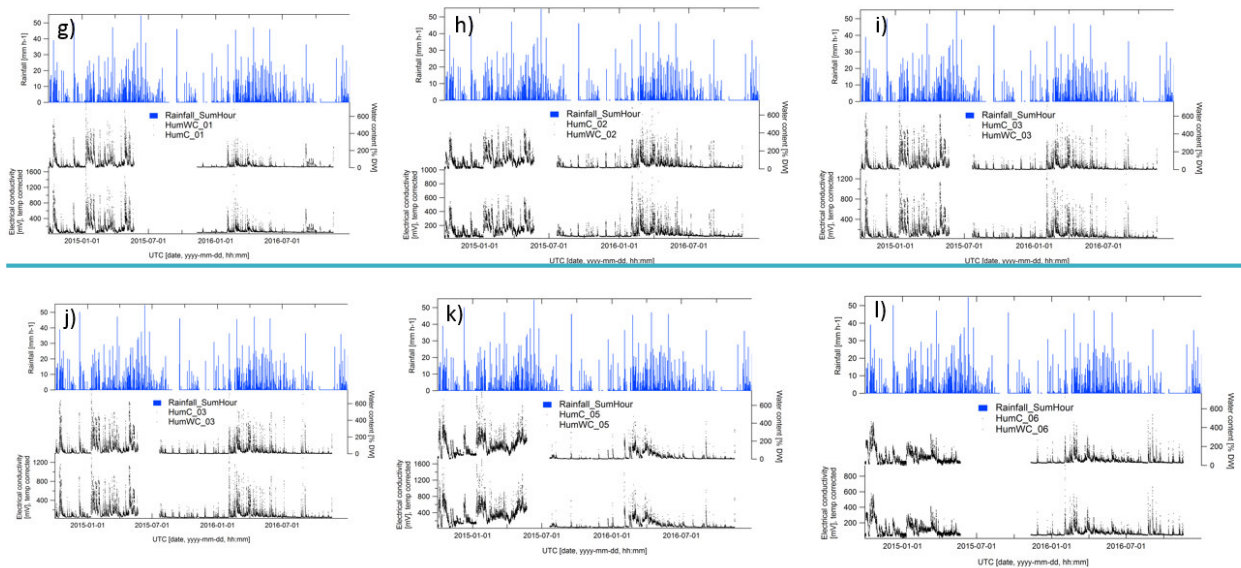
Sensors at 23 m height level (inner canopy)

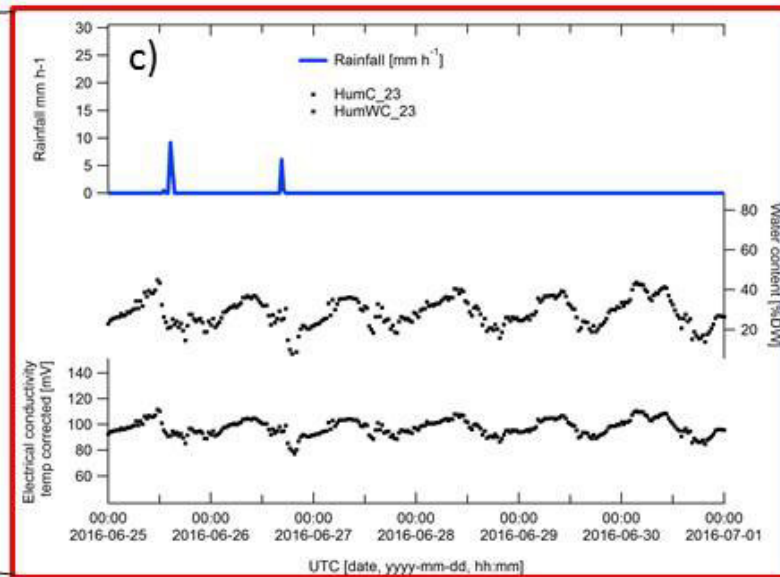
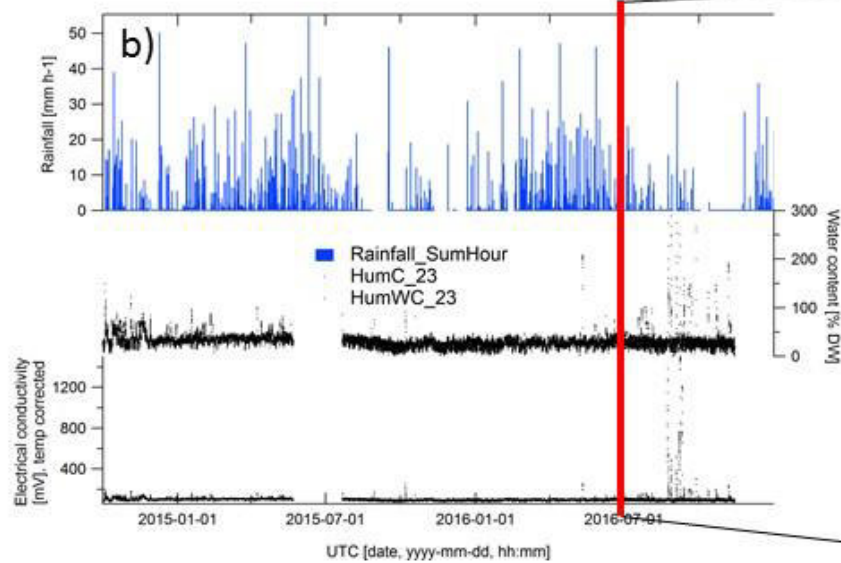
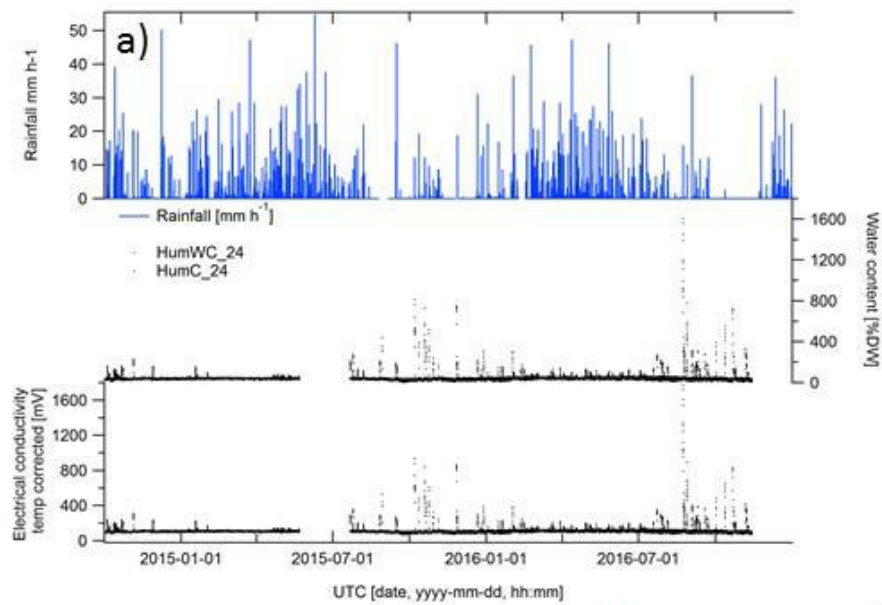


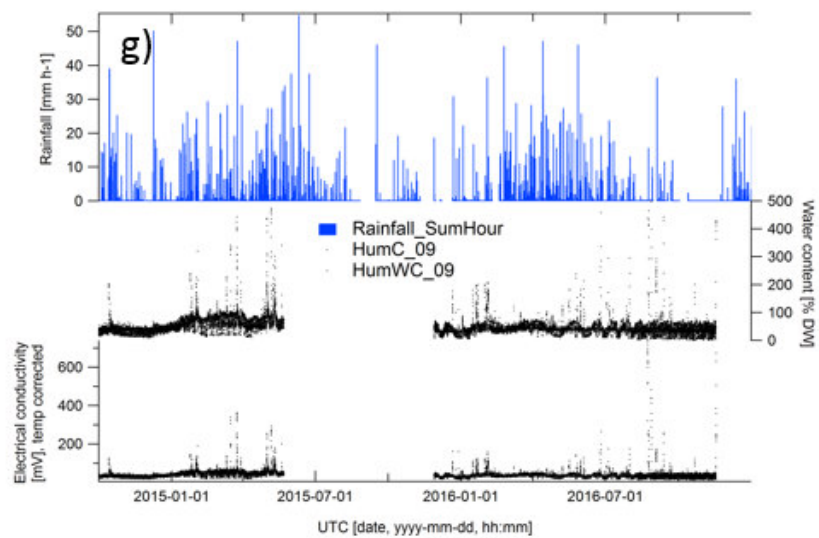
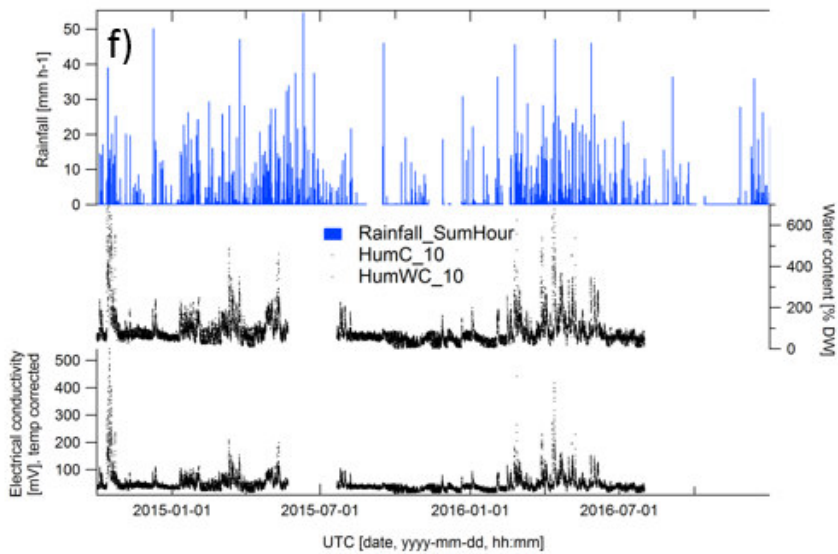
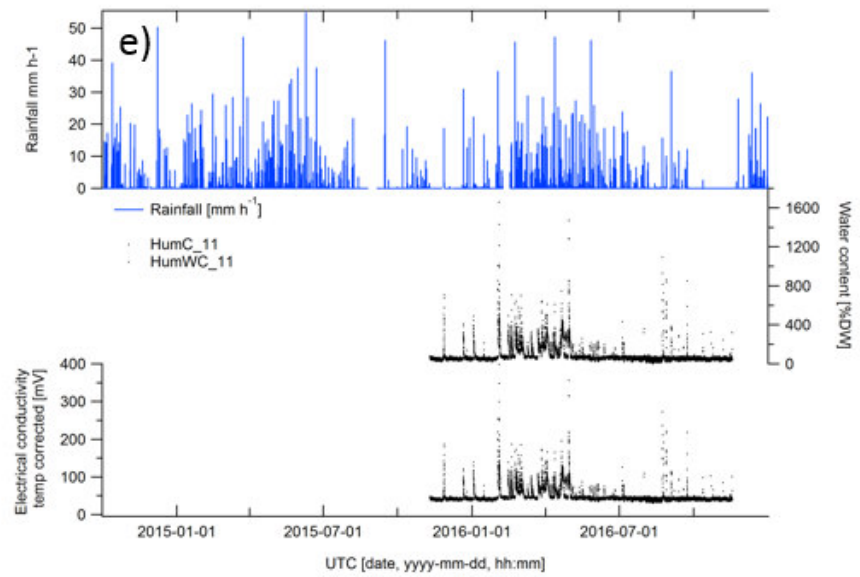
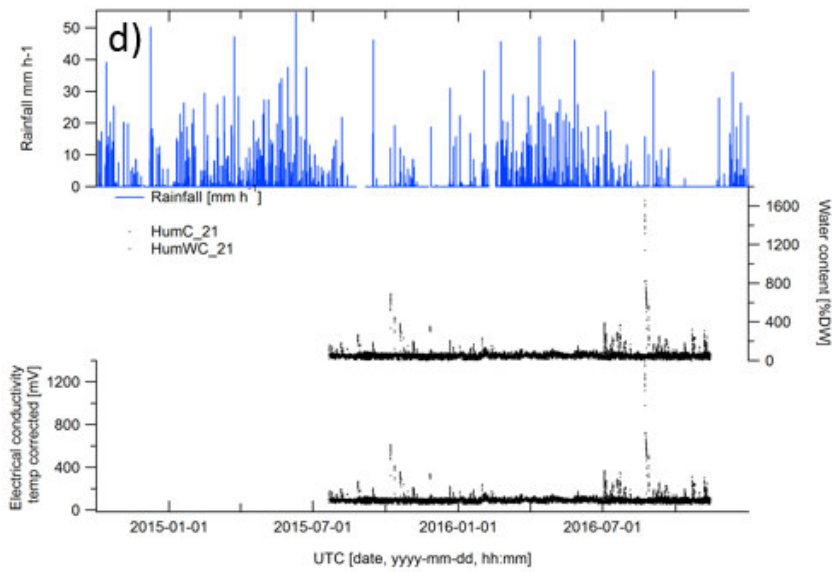
Sensors at 8 m height level (stem)

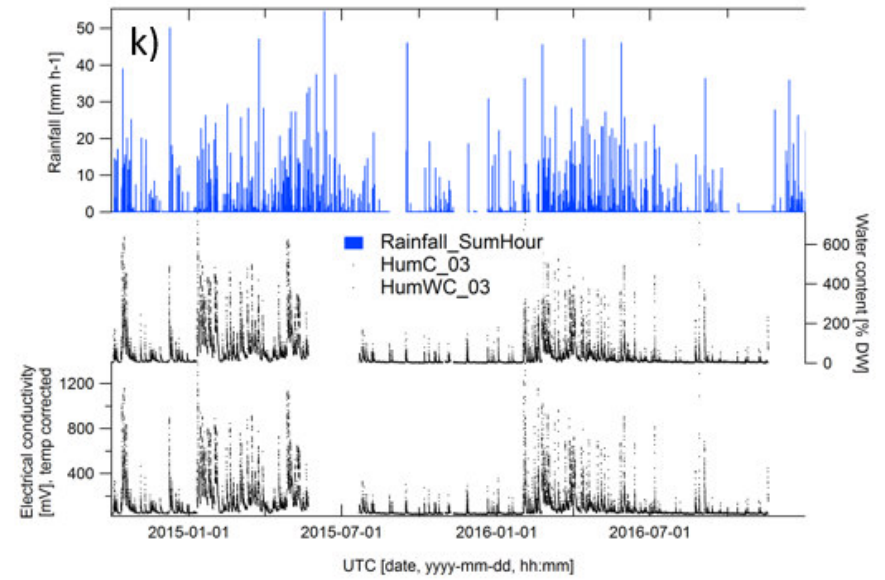
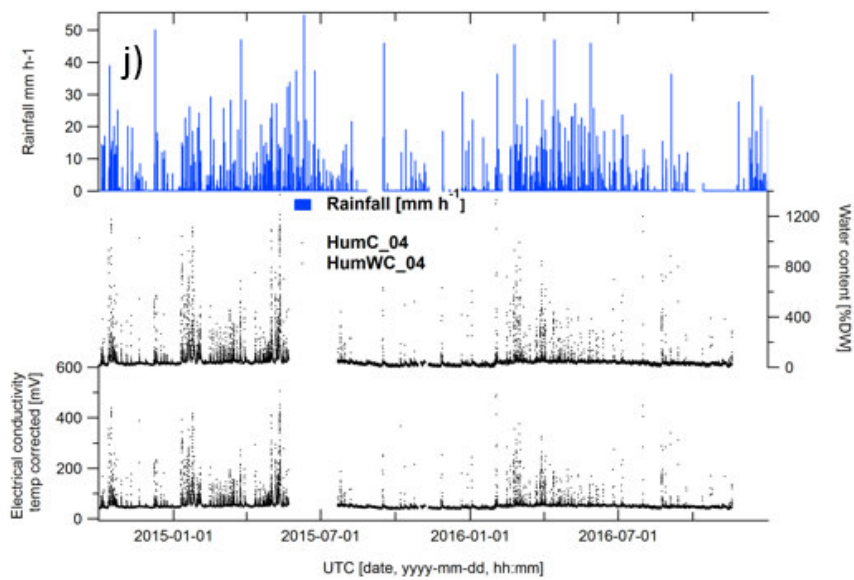
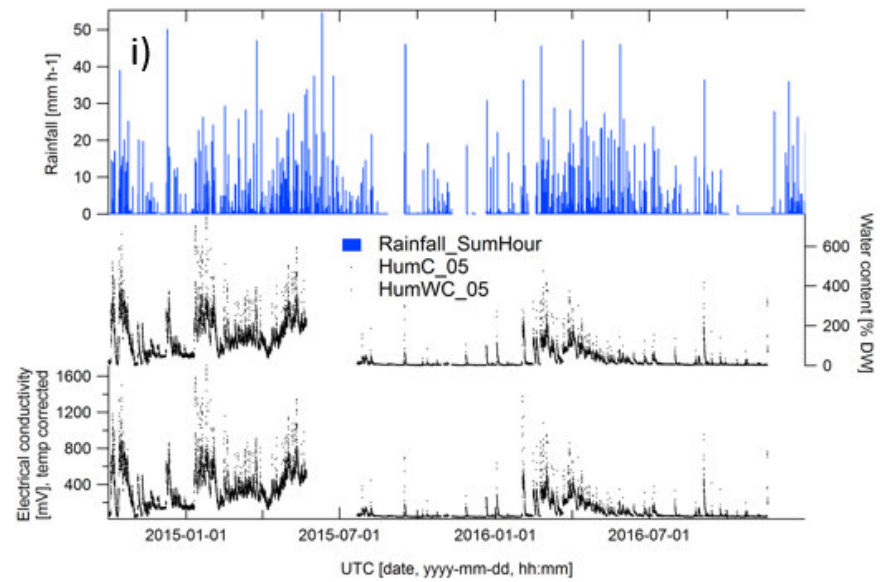
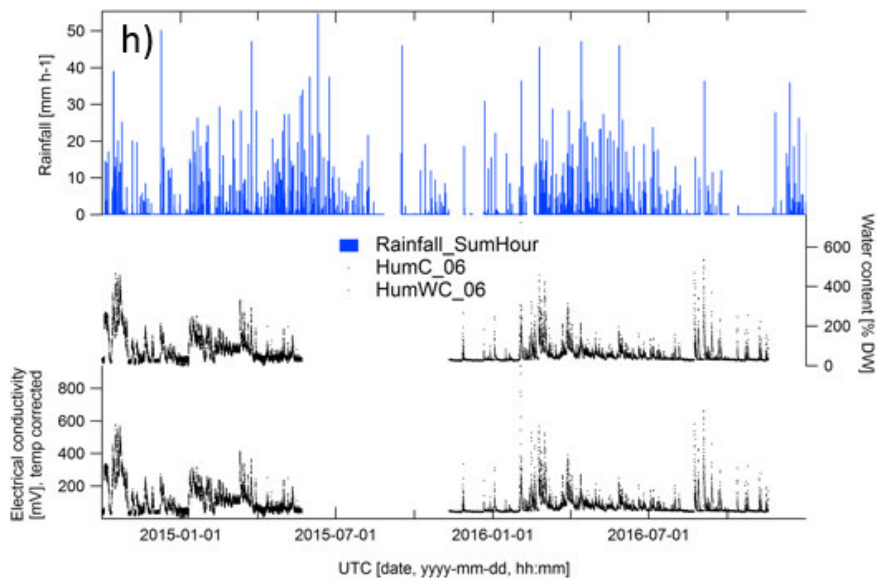


Sensors at 1.5 m height level (understory)









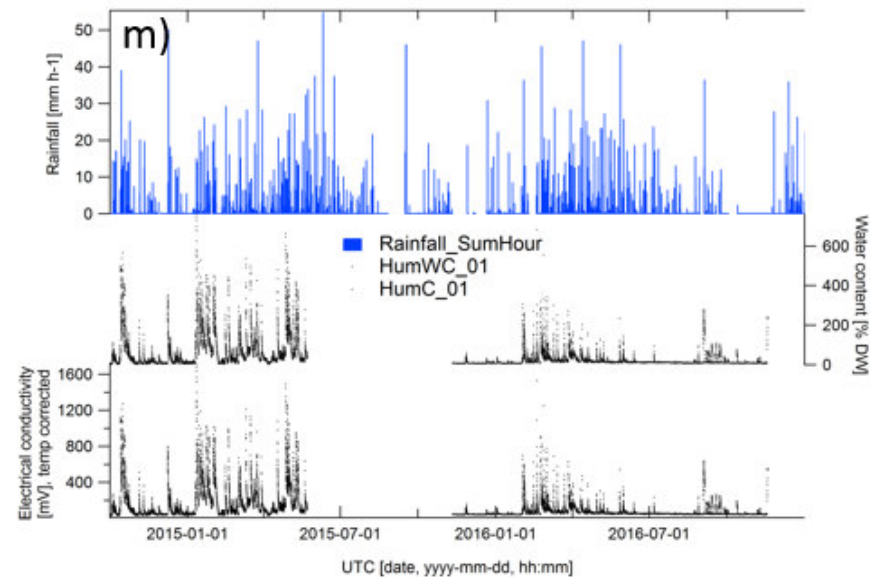
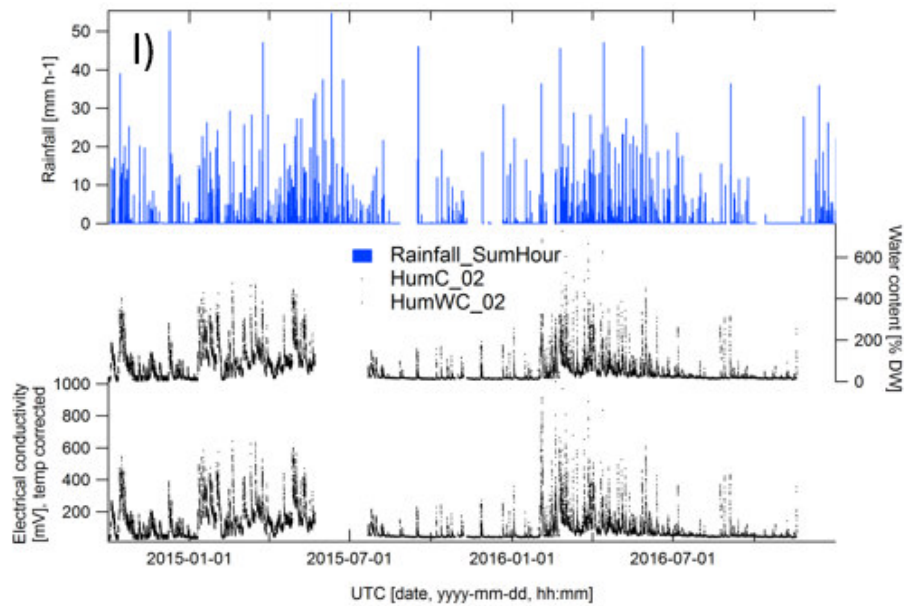
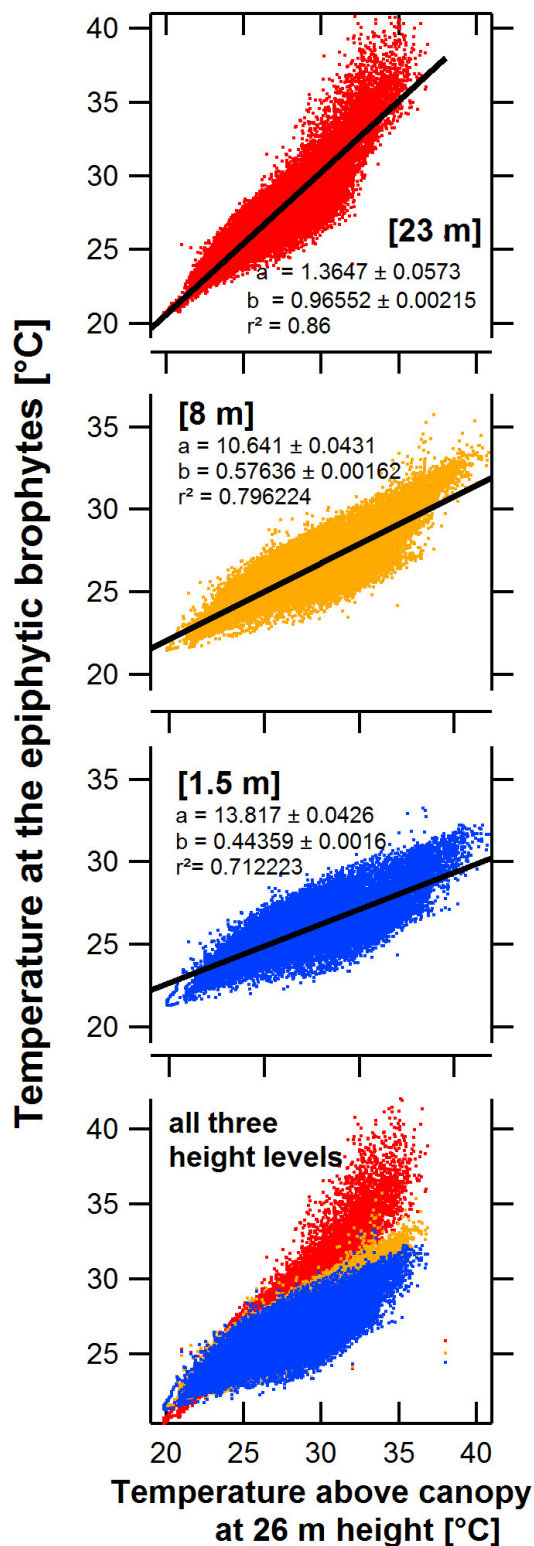
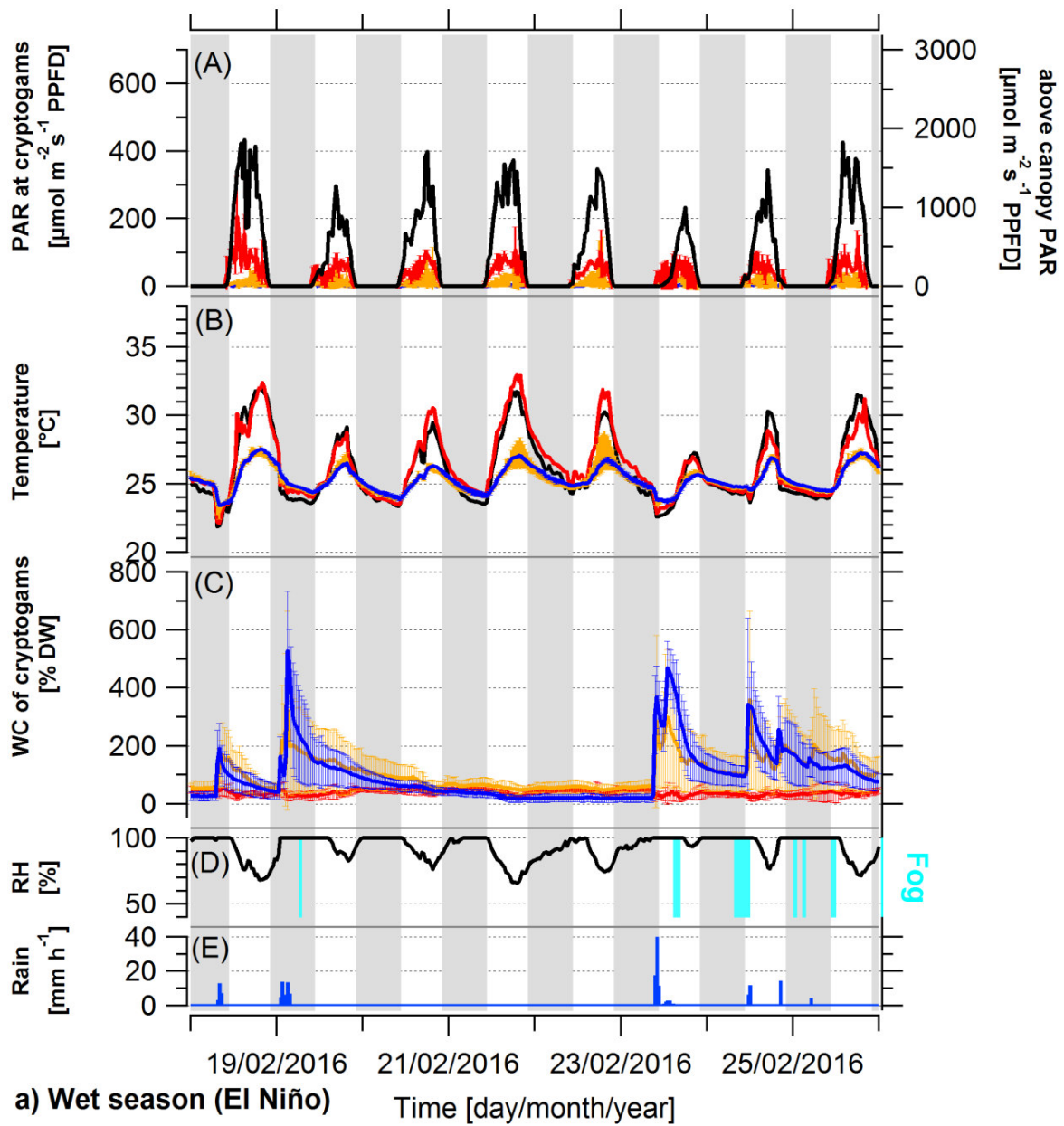


Figure S5. Long-term measurements of precipitation, electrical conductivity, and the calculated water content. All the sensors utilized for further calculations are shown: (a, b, c, d) at 23 m height, (~~e~~, e, f, g) at the 8 m height, and (~~g~~, h, i, j, k, l, m) at 1.5 m height. Gaps in the dataset correspond to maintenance periods.



5 | **Figure S6:** Temperature within bryophytes compared to the above-canopy temperature. The temperature within bryophytes was measured at 1.5 m, 8 m, and 23 m, while the above-canopy temperature was measured at 26 m height on the tower. The data are presented per height zone and also pooled together in the lowest panel. Data present 30-minute averages with linear fits, of the function $y = a + bx$, with the coefficients (± 1 std. dev.) and the R^2 are given in the figure for each height level.



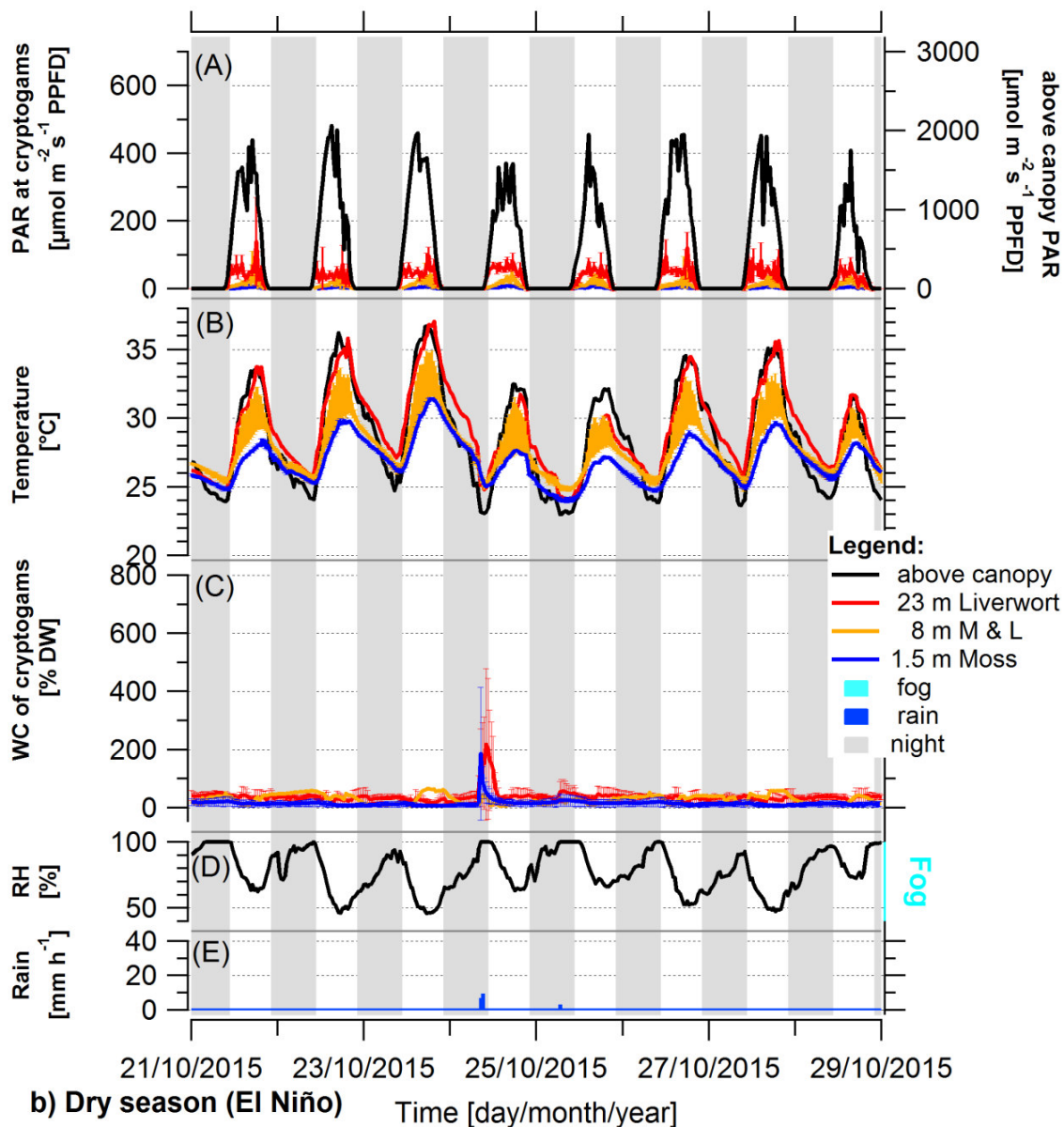


Figure S6S7: Representative periods during wet and dry season under the influence of El Niño, showing light conditions (PAR), temperature, and water content (WC) experienced by bryophytes, and above-canopy meteorological conditions in the Amazonian rain forest. Shown are 8-day periods during a) the wet season 2016 and b) the dry season 2015. The micrometeorological parameters on top/within epiphytic cryptogamic communities represent (A) the photosynthetically active radiation (PAR) on top, (B) the temperature within, and (C) the water content of cryptogamic communities. The above-canopy meteorological parameters comprise (A) the above-canopy photosynthetically active radiation (PAR at 75 m), (B) the above-canopy temperature (at 26 m), (D) the relative air humidity (RH at 26 m), the presence of fog events, and (E) the rain amount. The data show 30-minute averages \pm SD except for rain, which shows hourly sums. Data of replicate sensors installed within communities at the same height level were pooled, while above-canopy parameters were measured with one sensor each. The nighttime is shaded in grey color (18:00 – 06:18:00 LT).

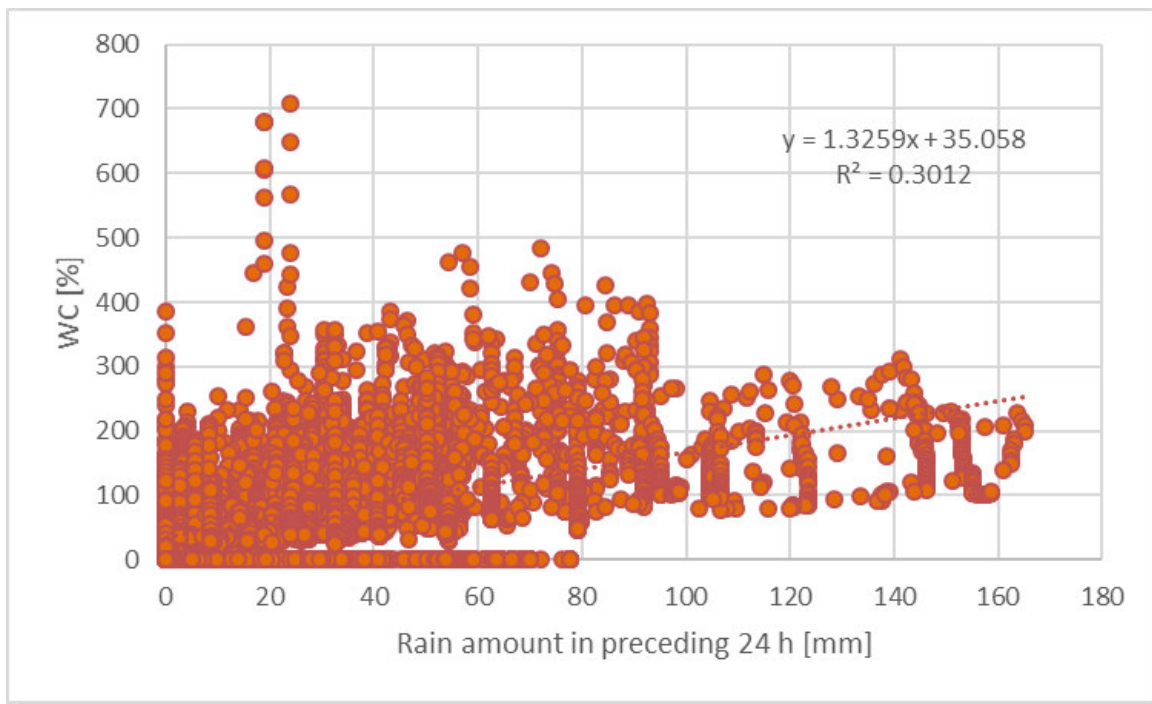
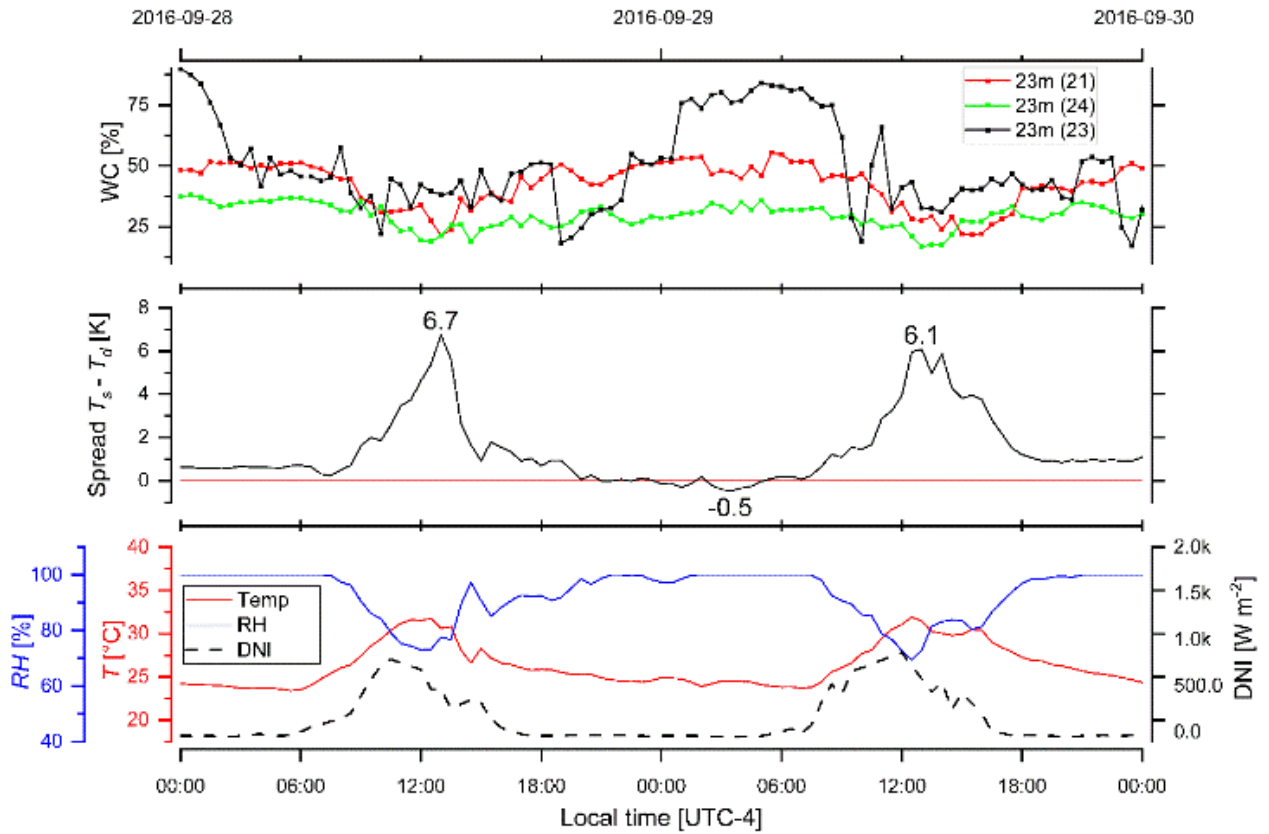


Figure S8: Correlation between the water content (WC) and the rain amount in the preceding 24 h. Linear trendline with formula added to illustrate the relationship between both parameters.



5 **Figure S98:** Exemplary daily (micro-)climatic conditions at the canopy level, showing the WC values of the 3 sensors at 23m [%] (A), the dew point spread at 23 m [°C] (B), and the environmental factors relative humidity RH[%], temperature T [°C] and direct normal irradiance DNI [W m⁻²] measured at 26 m (C).

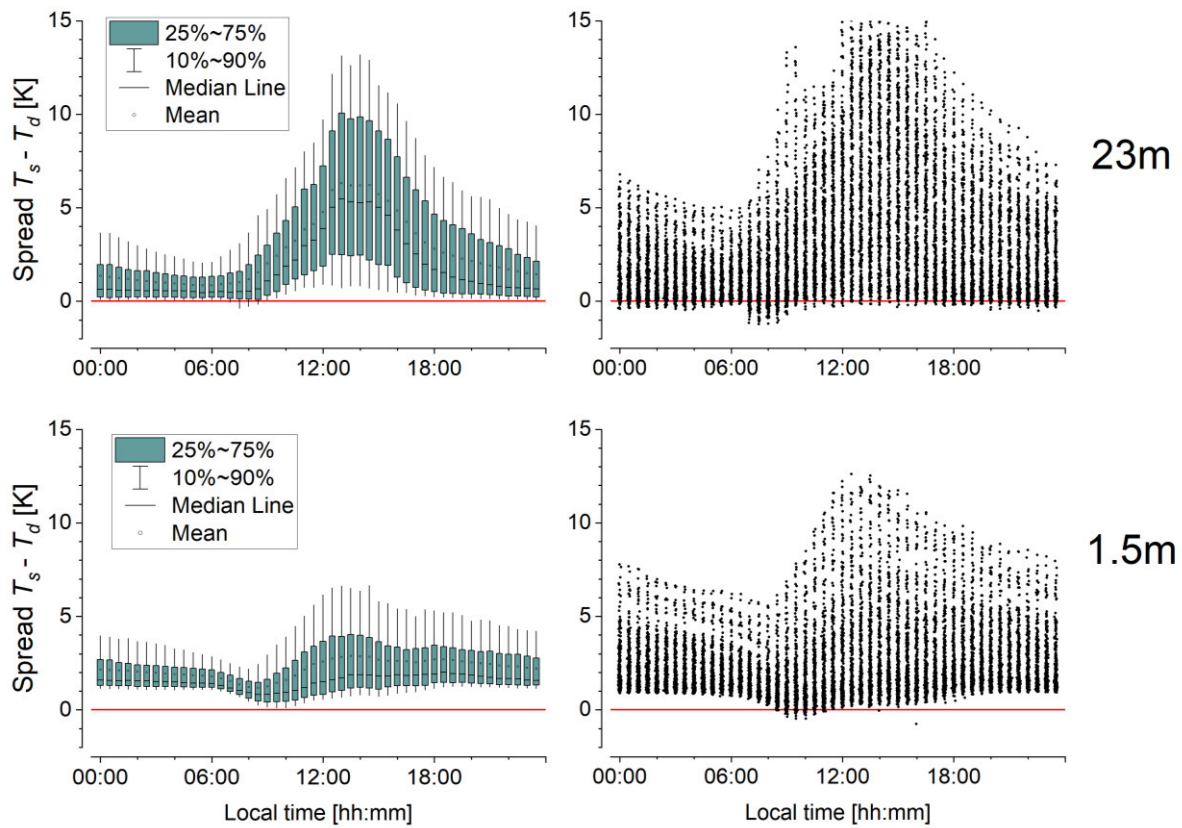


Figure S109: Diel dew point spread at 1.5 m and 23 m height levels in a 24h cycle, illustrating the difference between the temperature of the substrate (T_s) and the dew point of the surrounding air (T_d). If the surface temperature is lower than the dew point of the surrounding air (values below red line), condensation might occur.

5

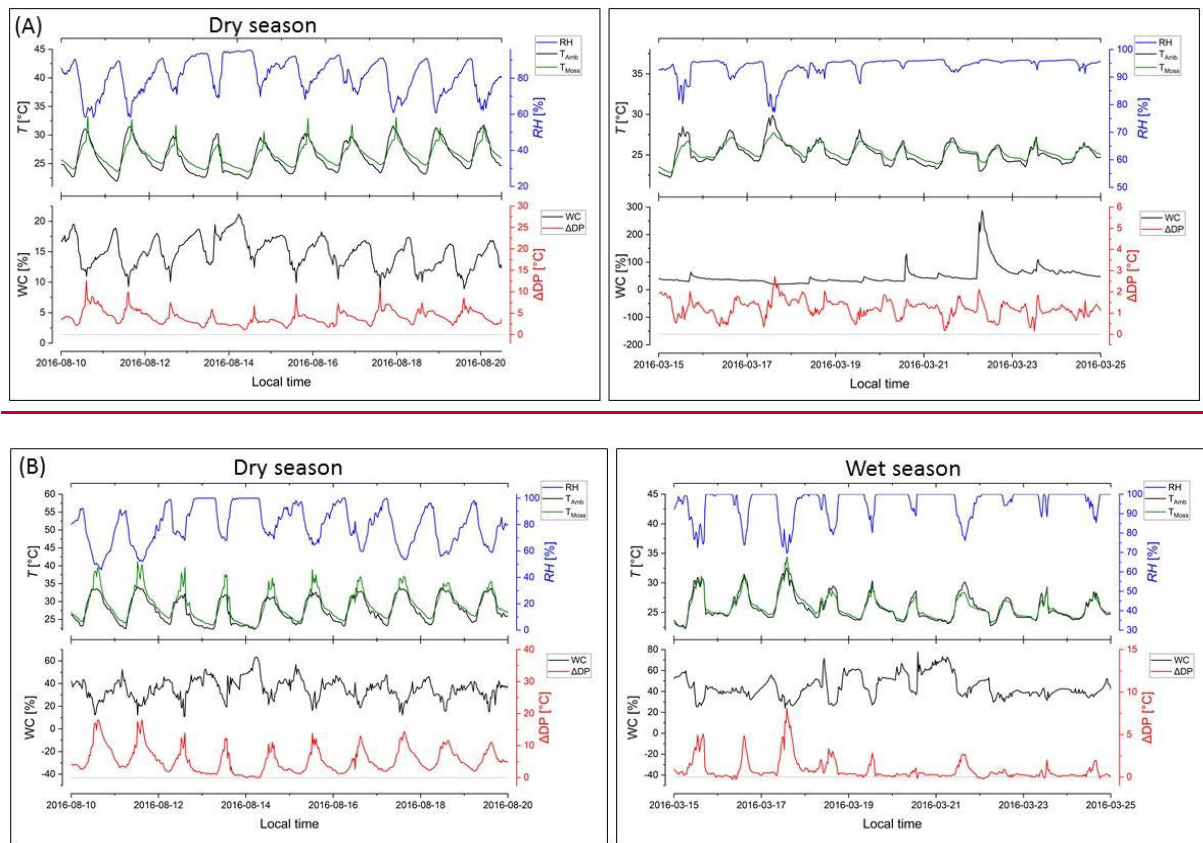


Figure S11: Characteristic sequence of (micro)climatic conditions during the dry and the wet season at (A) 23 m and (B) 1.5 m height. In the plots the relative air humidity [%], ambient and bryophyte temperature [°C], water content of the bryophytes [% dry weight], and dew point spread [°C] are shown.

5

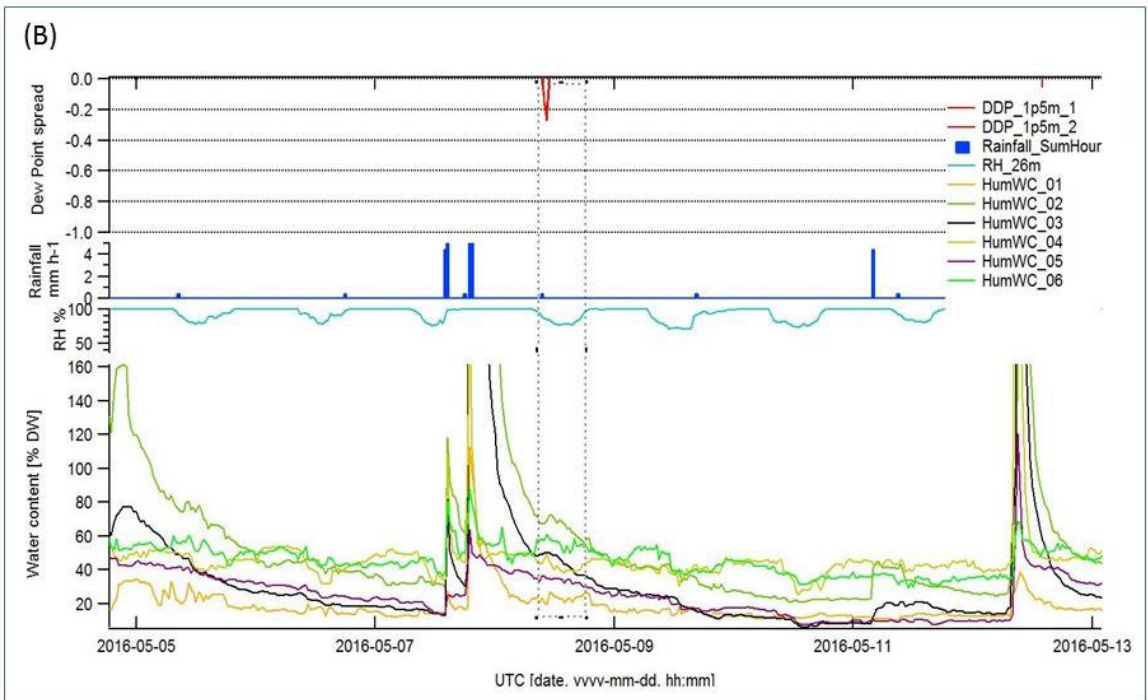
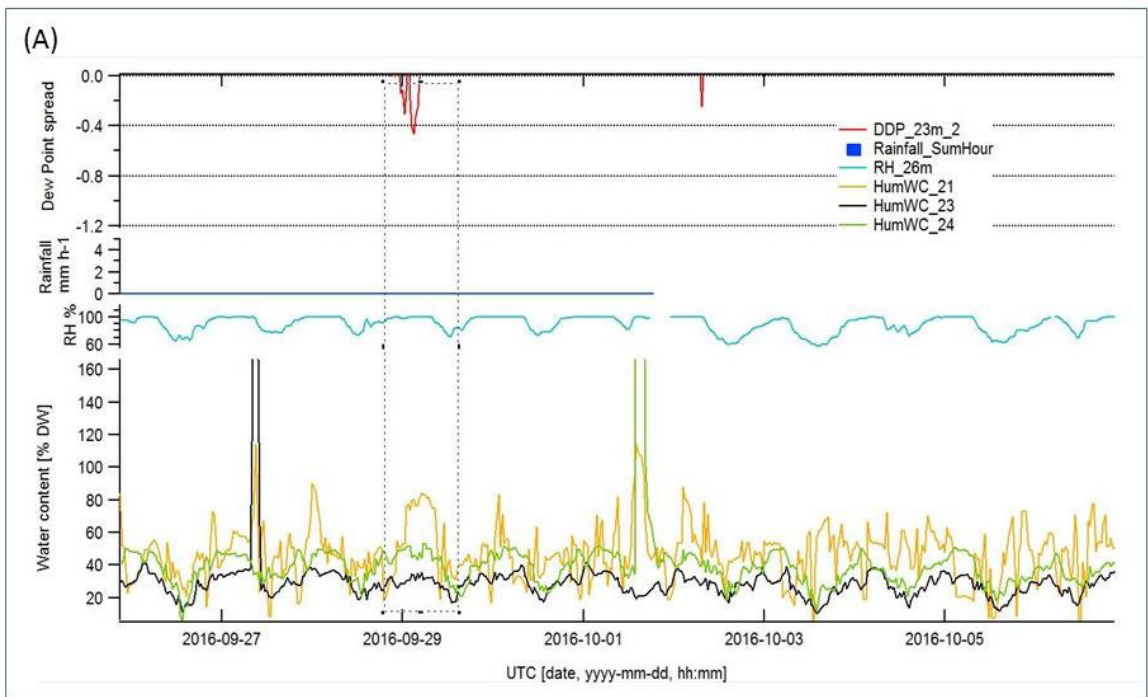


Figure S12: Exemplary (micro)meteorological data comprising dewpoint events at (A) 23 m and (B) 1.5 m height. In the plots, the dew point spread [°C], precipitation [mm * h⁻¹], relative humidity [%], and the water content [% dry weight] are shown.

5

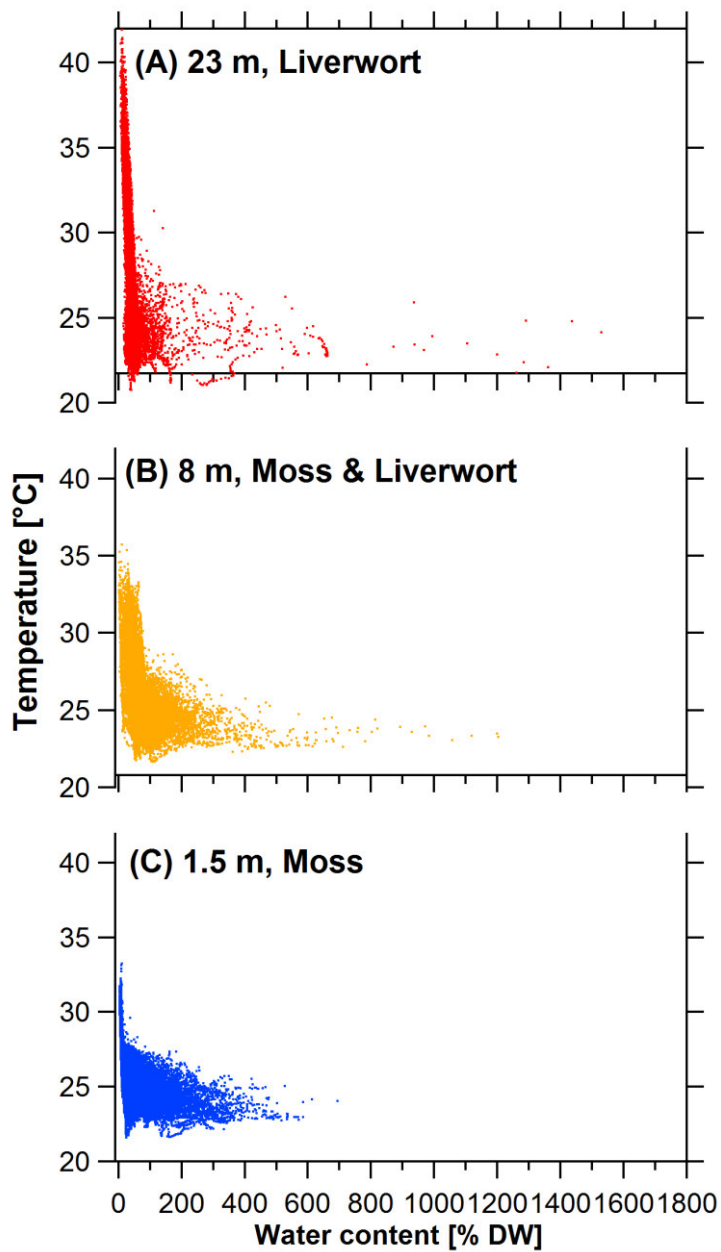


Figure S130: Temperature conditions of bryophytes related to their water content. The temperature was measured in bryophytes at different height levels along the tree. Data presented as 30-minute averages.

Table S1: Height of installation, minimum and maximum values of the individual sensors of the microclimate station measuring water content, temperature, and light. For the water content sensors, also the bryophyte species are given. Based on 30-minute averages.

Sensor No	Height [m]	Water content [% DW]		Bryophyte species	Sensor No	Height [m]	Temperature [°C]	
		min	max				min	max
Sensor 01	1.5	0	763	<i>Sematophyllum subsimplex</i>	Sensor 01	1.5	21.1	36.3
Sensor 02	1.5	0	763	<i>Sematophyllum subsimplex</i>	Sensor 02	1.5	21.4	39.4
Sensor 03	1.5	0	763	<i>Sematophyllum subsimplex</i>	Sensor 03	8	21.6	34.7
Sensor 04	1.5	0	1373	<i>Leucobryum martianum</i>	Sensor 04	8	20.9	46.3
Sensor 05	1.5	0	763	<i>Sematophyllum subsimplex</i>	Sensor 07	23	20.8	41.2
Sensor 06	1.5	0	763	<i>Sematophyllum subsimplex</i>	Sensor 08	23	20.3	48.7
Sensor 09	8	0	1318	<i>Octoblepharum cocuiense</i>		Height	PAR	
Sensor 10	8	0	1318	<i>Octoblepharum cocuiense</i>		[m]	[$\mu\text{mol m}^{-2} \text{s}^{-1}$]	
Sensor 11	8	0	1658	<i>Symbiezidium barbiflorum</i>			min	max
Sensor 21	23	0	1658	<i>Symbiezidium barbiflorum</i>	Sensor 01	1.5	0	634
Sensor 23	23	0	1658	<i>Symbiezidium barbiflorum</i>	Sensor 02	8	0	569
Sensor 24	23	0	1658	<i>Symbiezidium barbiflorum</i>	Sensor 03	8	0	1121
					Sensor 06	23	0	654
					Sensor 07	23	0	767

5 **Table S2:** Water content range measured during the calibration in the laboratory for the different replicates of the four bryophyte species. Listed are the minimum and maximum water content values (WC) measured at full water saturation (WC_{max}) and in the end of drying when weight stability was reached over more than 5 minutes (WC_{min}). Data shown for each replicate (1–4) and the species average (all).

Species	Replicate sample	WC_{min}	WC_{max}
<i>Leucobryum martianum</i>	1	32	1487
<i>Leucobryum martianum</i>	2	10	931
<i>Leucobryum martianum</i>	3	10	1241
<i>Leucobryum martianum</i>	4	7	1834
<i>Sematophyllum subsimplex</i>	1	14	614
<i>Sematophyllum subsimplex</i>	2	14	698
<i>Sematophyllum subsimplex</i>	3	14	468
<i>Sematophyllum subsimplex</i>	4	14	459
<i>Sematophyllum subsimplex</i>	5	7	1576
<i>Symbiezidium barbiflorum</i>	1	15	1657
<i>Symbiezidium barbiflorum</i>	2	15	1982
<i>Symbiezidium barbiflorum</i>	3	15	1581
<i>Symbiezidium barbiflorum</i>	4	22	1412
<i>Octoblepharum cocuiense</i>	1	23	742
<i>Octoblepharum cocuiense</i>	2	16	870
<i>Octoblepharum cocuiense</i>	3	6	2342
<i>Leucobryum martianum</i>	all	15	1373
<i>Sematophyllum subsimplex</i>	all	13	763
<i>Symbiezidium barbiflorum</i>	all	16	1658
<i>Octoblepharum cocuiense</i>	all	15	1318

Table S3: Monthly mean values and standard deviations (\pm SD) of photosynthetically active radiation (PAR_{avg} daytime, measured at 75 m), daily maxima of photosynthetically active radiation (PAR_{max}), temperature (measured at 26 m), and relative humidity (RH, measured at 26 m). Rainfall is presented as the monthly amounts and the percentage of days with rain (measured at 81 m), and also the percentage of days when rain detection malfunctioned are listed. Fog events are given as the percentage of days. Dry season data are shaded in red, wet season data in blue and transitional periods are unshaded. Due to data gaps in the measured rain data (shown in brackets) values for 21 days of rain were also extrapolated from existing data as described in methods section (values behind data in brackets). Values were calculated from 30-minute intervals. Fog has not being recorded in the time ranges of 31.05. – 20.10.2015, 30.04. – 06.07.2016, 01.09. – 31.12.2016 due to malfunction of the device, thus the last column provides data about the number of operational days per month.

Month	PAR_{avg} daytime [$\mu\text{mol m}^{-2} \text{s}^{-1}$]		PAR_{max} [$\mu\text{mol m}^{-2} \text{s}^{-1}$]		Temperature [$^{\circ}\text{C}$]		RH [%]		Rain [mm month $^{-1}$]	Rain [% days]	Defect on rain detection [% days]	Fog [% days]	<u>Days with fog data</u>
	Mean	\pm SD	Mean	\pm SD	Mean	\pm SD	Mean	\pm SD					
Oct 2014	857	668	2201	509	26.0	2.8	90	11	212	58	0	55	30
Nov 2014	832	624	2082	423	25.6	2.9	92	11	70	57	0	53	30
Dec 2014	843	582	2140	346	26.3	2.7	90	11	123	42	0	42	30
Jan 2015	637	525	1747	735	24.5	2.4	95	8	259	71	0	71	29
Feb 2015	774	589	2058	600	25.4	2.6	92	10	140	64	0	46	28
Mar 2015	680	534	2038	575	24.7	2.1	96	7	331	87	0	77	31
Apr 2015	766	564	2155	463	25.3	2.5	93	10	189	80	0	40	29
May 2015	725	559	2103	425	27.2	n.a.	93	6	320	90	0	58	30
Jun 2015	804	562	2237	128	25.0	2.3	94	8	178	80	0	0*	0
Jul 2015	892	605	2238	188	25.7	3.0	91	11	74	65	0	0*	0
Aug 2015	1017	636	1722	957	27.1	3.3	83	13	(23) 32*	23	23	0*	0
Sep 2015	1148	687	2242	467	28.7	3.7	74	15	38	13	20	0*	0
Oct 2015	968	635	2072	514	28.4	3.6	78	16	55	35	3	13*	11
Nov 2015	887	624	1859	769	27.9	3.5	81	16	(33) 37*	30	17	23	28
Dec 2015	862	575	2074	304	28.1	3.0	78	14	38	26	3	6	31
Jan 2016	882	606	2175	270	28.2	3.4	78	16	52	48	0	13	31
Feb 2016	743	550	1928	679	25.9	2.6	93	10	(267) 341*	79	52	48	29
Mar 2016	692	545	2041	545	25.6	2.1	96	7	304	90	0	77	31
Apr 2016	709	564	2088	443	25.6	2.3	96	7	277	87	0	73	28
May 2016	817	603	2230	405	26.1	2.6	94	8	236	90	0	0*	0
Jun 2016	828	584	2178	261	25.6	2.8	92	10	105	57	0	0*	0
Jul 2016	917	629	2253	118	26.2	3.2	88	12	92	58	0	26*	26
Aug 2016	1016	648	2146	593	27.1	3.5	83	14	40	32	3	16	31
Sep 2016	947	662	2230	543	26.5	3.1	89	12	(77) 96*	50	17	0*	0
Oct 2016	915	641	2323	192	27.1	3.3	86	14	(1) 9*	23	23	0*	0
Nov 2016	911	610	2227	217	27.1	3.3	87	13	(30) 89*	20	13	0*	0
Dec 2016	694	553	1955	503	25.4	2.7	94	10	223	71	0	0*	0

(*) Gaps in the data record due to malfunction of the device.

Table S4: Parameters determining the time range of photosynthesis and respiration. The ranges of values defining the lower water compensation point (WCP), the lower light compensation point (LCP_l), the temperature for optimal net photosynthesis (T_{opt}), and the upper temperature compensation point (TCP) as relevant parameters have been extracted from published studies conducted at various study sites in the tropical rain forest.

Parameter	Range of values	Reference	Study site
WCP	30–80 % DW	Wagner et al 2013	Panama, lowland rain forest, 0 m
LCP _l	3–12 $\mu\text{mol m}^{-2} \text{s}^{-1}$	Lösch et al. 1994	Zaire, lowland rain forest, 800 m
T _{opt}	24–27 °C	Wagner et al 2013	Panama, lowland rain forest, 0 m
TCP	30–36 °C	Wagner et al 2013	Panama, lowland rain forest, 0 m

Table S5: Monthly mean values and standard deviations (\pm SD) of the photosynthetically active radiation (PAR_{avg} daytime), the daily maxima of photosynthetically active radiation (PAR_{max}), temperature, and water content of bryophytes at four height levels. Dry season data are shaded in red, wet season data in blue and transitional periods are unshaded. Values were calculated from 30-minute intervals. N.a.: data not available.

Month	PAR_{avg} daytime [$\mu\text{mol m}^{-2} \text{s}^{-1}$]						PAR_{max} [$\mu\text{mol m}^{-2} \text{s}^{-1}$]					
	1.5 m		8 m		23 m		1.5 m		8 m		23 m	
	Mean	\pm SD	Mean	\pm SD	Mean	\pm SD	Mean	\pm SD	Mean	\pm SD	Mean	\pm SD
Oct 2014	4	8	30	31	88	90	75	105	285	231	624	286
Nov 2014	4	11	23	32	24	37	142	131	396	321	378	275
Dec 2014	6	18	31	50	25	33	236	172	435	228	346	235
Jan 2015	3	8	22	28	20	27	155	96	341	219	341	246
Feb 2015	2	3	31	21	16	17	46	33	173	183	234	244
Mar 2015	3	4	43	35	16	15	45	55	292	159	128	117
Apr 2015	6	20	80	105	16	18	346	310	480	231	241	231
May 2015	6	32	66	71	16	17	634	428	282	236	146	137
Jun 2015	2	3	73	64	18	20	42	51	214	125	177	141
Jul 2015	3	12	54	73	15	18	168	178	727	301	152	144
Aug 2015	13	56	66	115	24	23	601	414	746	193	227	170
Sep 2015	9	21	28	47	65	66	248	204	403	224	492	229
Oct 2015	3	4	15	15	44	30	53	47	128	99	221	157
Nov 2015	4	7	16	25	61	64	82	95	315	151	475	208
Dec 2015	5	11	22	35	88	103	112	116	308	171	645	250
Jan 2016	4	7	16	21	88	103	72	91	177	143	692	294
Feb 2016	3	4	13	11	57	46	46	54	79	76	388	237
Mar 2016	3	7	28	15	37	33	102	125	107	80	268	215
Apr 2016	5	15	27	19	38	31	192	199	59	27	270	203
May 2016	3	7	n.a.	n.a.	45	41	114	109	n.a.	n.a.	286	209
Jun 2016	2	2	n.a.	n.a.	58	68	25	34	n.a.	n.a.	416	199
Jul 2016	2	4	n.a.	n.a.	72	86	30	44	n.a.	n.a.	527	204
Aug 2016	9	34	31	52	71	94	319	216	340	241	614	256
Sep 2016	3	7	13	24	55	69	102	84	250	137	508	244
Oct 2016	2	3	7	9	47	54	35	28	106	71	421	219
Nov 2016	3	5	9	13	73	85	59	51	172	114	606	251
Dec 2016	4	12	24	38	52	56	156	131	361	282	457	274

Continuation of Table S5

Month	Temperature [°C]						Water content [% DW]					
	1.5 m		8 m		23 m		1.5 m Moss		8 m M&L		23 m Liverwort	
	Mean	± SD	Mean	± SD	Mean	± SD	Mean	± SD	Mean	± SD	Mean	± SD
Oct 14	25.0	1.3	25.2	1.6	26.3	2.9	115	107	110	111	42	20
Nov 14	25.3	1.2	25.7	1.4	26.2	2.3	38	30	53	16	42	20
Dec 14	25.4	1.1	25.8	1.3	26.6	2.1	49	48	56	20	35	7
Jan 15	24.2	1.1	24.3	1.3	24.6	1.8	129	113	76	32	39	12
Feb 15	24.5	1.0	24.5	1.1	25.0	1.8	87	67	69	33	38	8
Mar 15	24.4	0.9	24.3	0.9	24.5	1.3	106	65	102	64	38	6
Apr 15	24.6	0.9	24.7	1.1	24.9	1.8	79	65	73	31	39	8
May 15	24.6	0.9	24.5	0.9	24.8	1.7	130	90	103	58	39	8
Jun 15	24.5	0.9	24.5	1.0	25.0	1.9						
Jul 15	24.5	1.1	25.0	1.5	25.5	2.5	37	24	103	45	49	23
Aug 15	25.4	1.2	26.3	2.0	27.0	2.8	20	10	67	17	42	21
Sep 15	27.0	1.7	27.8	2.2	29.0	3.4	13	17	60	15	36	18
Oct 15	27.2	1.8	28.0	2.2	29.4	3.2	13	16	36	21	43	52
Nov 15	27.2	1.9	27.6	2.3	29.2	3.6	16	14	51	30	37	32
Dec 15	27.3	1.6	27.9	2.0	29.4	3.4	15	11	48	24	35	18
Jan 16	27.4	1.8	28.0	2.2	29.4	3.8	16	14	51	31	37	16
Feb 16	25.2	1.0	25.4	1.2	26.2	2.5	80	93	99	80	43	16
Mar 16	25.2	0.9	25.1	0.9	25.6	1.8	74	68	91	49	41	13
Apr 16	25.2	1.0	25.2	1.1	25.7	2.0	63	45	131	85	43	14
May 16	25.3	1.0	25.3	1.2	26.1	2.3	42	33	75	39	44	16
Jun 16	24.6	1.1	24.6	1.3	25.8	2.8	31	18	61	30	45	13
Jul 16	24.8	1.2	25.3	1.7	26.7	3.4	24	22	52	21	53	36
Aug 16	25.7	1.8	26.3	2.4	28.0	4.1	22	28	59	99	59	130
Sep 16	25.5	1.3	25.9	1.7	27.1	3.3	28	40	52	39	67	111
Oct 16	26.2	1.6	26.8	1.9	28.0	3.4	17	9	45	18	43	38
Nov 16	25.9	1.7	26.5	2.1	28.0	3.4	18	20	49	49	44	37
Dec 16	25.4	1.3	25.0	1.7	25.6	2.5						

Table S6: Mean values and standard deviations (\pm SD) of the daily maxima of photosynthetically active radiation (PAR_{max}) for each height level shown for 2015 and 2016, considering that 2015 was an El Niño year (additional information to Table 1).

Height	PAR_{max} [$\mu\text{mol m}^{-2} \text{s}^{-1}$]			
	2015		2016	
	Mean	\pm SD	Mean	\pm SD
above canopy	1766	415	1842	364
23 m	125	123	226	140
8 m	186	195	68	90
1.5 m	49	89	29	45

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Table S7: Mean values and standard deviations (\pm SD) of the daily maxima of photosynthetically active radiation (PAR_{max}) for each height level shown for the different seasons (additional information to Table 2).

Height [m]	PAR_{max} [$\mu\text{mol m}^{-2} \text{s}^{-1}$]	
	Mean	\pm SD
Wet season		
above-canopy	1687	431
23 m	245	82
8 m	210	151
1.5 m	191	206
Transitional season Wet-Dry		
above-canopy	1855	233
23 m	318	183
8 m	471	363
1.5 m	66	68
Dry season		
above-canopy	1924	370
23 m	457	147
8 m	314	184
1.5 m	172	177
Transitional season Dry-Wet		
above-canopy	1691	407
23 m	496	165
8 m	324	95
1.5 m	146	61