

1 Litter quality outweighs climate as a driver of 2 decomposition across the tundra biome

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79 **Abstract**

80 Considerable uncertainty exists regarding the strength, direction and relative importance of
81 the drivers of decomposition in the tundra biome, partly due to a lack of coordinated
82 decomposition field studies in this remote environment. Here, we analysed 3717 incubations
83 of two uniform litter types, green and rooibos tea, buried at 330 circum-Arctic and alpine sites
84 to quantify the effects of temperature, moisture and litter quality on decomposition. We found
85 a surprisingly linear positive relationship between decomposition and soil temperature across
86 all sites, counter to theory and previous model estimates. Litter mass loss was greater at
87 wetter sites, even where soils reached almost full water saturation. However, litter quality was
88 the strongest driver of litter mass loss across the tundra biome, explaining six times more
89 variation in summer decomposition than soil temperature. Our results indicate that climate
90 warming will directly increase decomposition across tundra environments. Yet, the indirect
91 effects of climate change on vegetation communities, and thus plant litter inputs and quality,
92 could have a more profound impact than direct effects on the balance of this globally important
93 carbon store.

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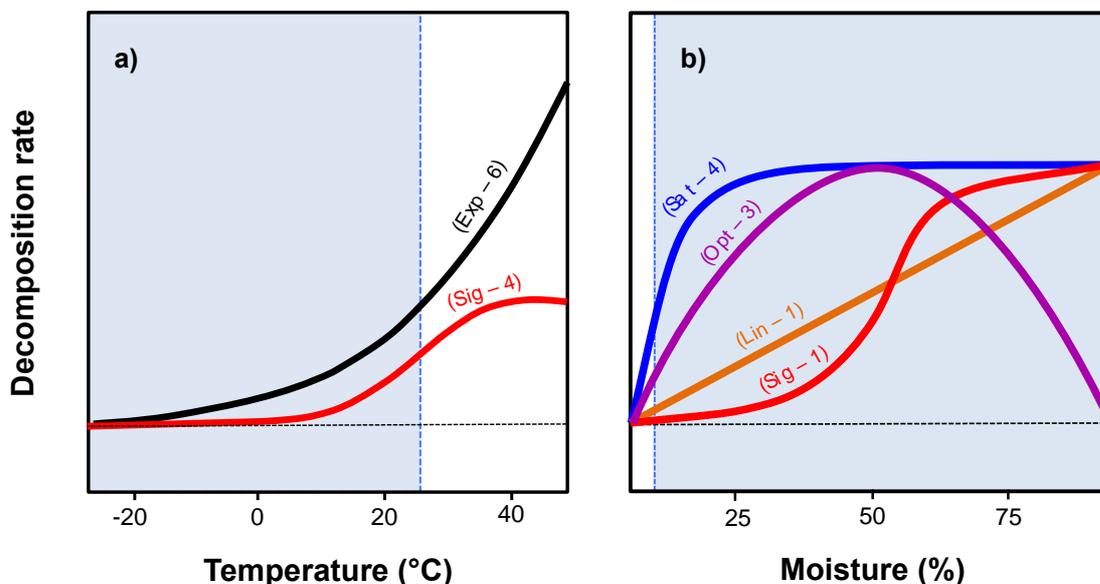
95 **Introduction**

96 The decomposition of terrestrial carbon pools is a vital component of the global carbon cycle^{1,2}
97 and is sensitive to temperature³. Climate warming is predicted to accelerate both the
98 decomposition process and carbon emissions^{2,4,5}. Quantifying changes in decomposition is
99 thus critical for identifying major feedbacks to climate change⁶. Perhaps nowhere is this more
100 true than in high-latitude ecosystems, which contain over a third of global soil carbon⁷⁻⁹, more
101 than double the current atmospheric stocks¹⁰. Decomposition in the tundra is currently
102 constrained by cold temperatures, frozen soils and recalcitrant litter, encouraging the build-up
103 of organic matter in soils¹¹. Tundra ecosystems are warming at up to four times the global
104 average rate¹², with annual temperatures in the Arctic predicted to increase by 2-10°C by the
105 end of the century relative to the period from 1850 to 1900¹³. As a result, decomposition rates
106 are expected to increase in the tundra¹¹, potentially releasing 37 to 174 Pg of carbon by 2100,
107 equivalent to an additional 17 to 82 ppm CO₂ in the Earth's atmosphere¹⁴. Climate warming
108 impacts can either directly reduce carbon stores by accelerating decomposition¹¹, or indirectly
109 by changing plant litter inputs^{15,16}. In addition, warming impacts on decomposition are not
110 occurring in isolation from other environmental change including changes to soil moisture^{17,18}.
111 Arctic carbon emissions could determine whether soils globally are a sink or source of carbon
112 under accelerating global change⁶. Thus, there is an urgent need to explore the drivers of
113 decomposition across the tundra biome.

114

115 Despite the potential substantial impact of climate change on carbon cycling in Arctic terrestrial
 116 ecosystems, the relative influence of environmental drivers of decomposition have yet to be
 117 experimentally tested at the tundra biome scale. Temperature and soil moisture are
 118 considered to be the primary drivers of decomposition¹¹, and together explain approximately
 119 70% of variation in decomposition rates globally^{2,19,20}. However, biogeochemical models
 120 incorporate substantially different relationships between decomposition, temperature and soil
 121 moisture, particularly at climatic extremes²¹ (Fig. 1). Earth system model relationships
 122 between temperature and decomposition rate are either assumed to exponentially decline or
 123 saturate near zero at sites with colder temperatures and relationships with moisture vary
 124 between saturating, optimal, linear or sigmoidal relationships²² (Fig. 1). This lack of
 125 consistency in the assumed relationships between both soil temperature and moisture and
 126 decomposition is partly driven by a lack of *in situ* data from high-latitude regions^{5,22}, and
 127 contributes to the large uncertainty surrounding predictions of global soil carbon losses^{4,5}.
 128 Thus, reducing this uncertainty requires *in situ* decomposition data across a range of
 129 temperature and moisture conditions within the tundra biome.

130



131

132 **Figure 1.** Biogeochemical models include a number of different shapes of relationships
 133 between decomposition rate of soil organic matter and temperature (a) and moisture (b).
 134 Summary of 19 biogeochemical model functions included in Sierra et al. 2015. The
 135 relationship between decomposition and temperature is modelled as exponential (Exp: black,
 136 six models) or sigmoidal (Sig: red, four models). The relationship between decomposition and
 137 moisture is modelled as saturating (Sat: blue, four models), optimal (Opt: purple, three
 138 models), linear (Lin: orange, one model) or sigmoidal (Sig: red, one model). The blue shaded
 139 area indicates the range of temperatures and soil moisture values for the 330 sites included
 140 in this study. Note that temperature conditions generally do not exceed ~25°C in the tundra
 141 biome.

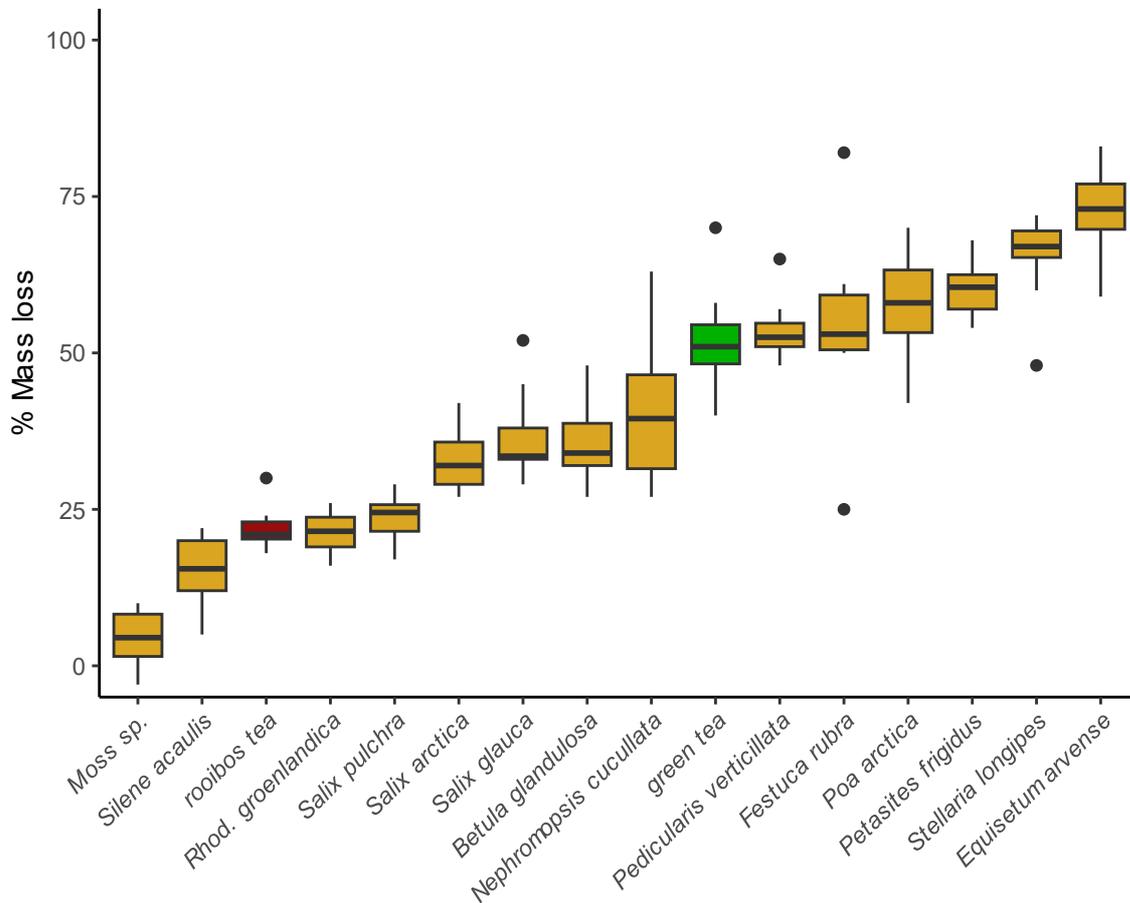
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143 Soil organic matter has many forms, one of which is leaf litter deposited on the soil surface
144 and incorporated into the soil profile over time including through cryoturbation processes in
145 tundra ecosystems²³. Decomposition of leaf litter is dependent on litter quality, and thus the
146 structural and chemical traits of plant tissues of different species across ecological
147 communities^{24–28}. Plant traits and litter characteristics may be the dominant control on litter
148 decomposition worldwide, outweighing environmental drivers even across biomes^{25,29,30}. As
149 plant community composition changes with warming, so too will the litter inputs to the soil and
150 decomposition rates of soil organic matter¹⁵. Many tundra plant communities are undergoing
151 widespread changes^{31,32}, notably an expansion of shrub species^{33,34}, that could dramatically
152 alter litter inputs to soils¹⁶. Site-scale experiments indicate that litter quality explains more
153 variation in litter decomposition than environmental variables^{26,35,36}. Cross-site studies of
154 decomposition using common substrates have been conducted for other global
155 biomes^{20,22,25,29,30,37–39}. However, the relative influence of litter quality versus environmental
156 controls on decomposition has not yet been tested across the tundra biome, primarily due to
157 difficulties of controlling for litter homogeneity.

158

159 In this study, we quantify the drivers of litter decomposition at 330 sites across the circum-
160 Arctic and alpine tundra (Table S1) and 3717 incubations using the Tea Bag Index¹⁹. The Tea
161 Bag Index is a standardised protocol that employs two commercially available types of tea
162 (labile green and recalcitrant rooibos tea) to estimate stabilisation factor (S) and
163 decomposition rate (k) and provide a chemically validated⁴⁰ and highly replicable method for
164 measuring leaf litter decomposition across sites^{19,29,41–43}. Decomposability of the two tea types
165 is also representative of leaf litters for a range of tundra species (Fig. 2, Fig. S1) and thus
166 provides an analogue for the potential impact of plant community change on litter
167 decomposability in tundra ecosystems^{15,28,40,44–46}. Due to relationships described in theoretical
168 and experimental studies^{2,21,22,26}, we predict that decomposition will increase exponentially
169 with temperature, and that temperature will be the strongest driver of decomposition across
170 the broad biogeographical gradients of the tundra biome.

171



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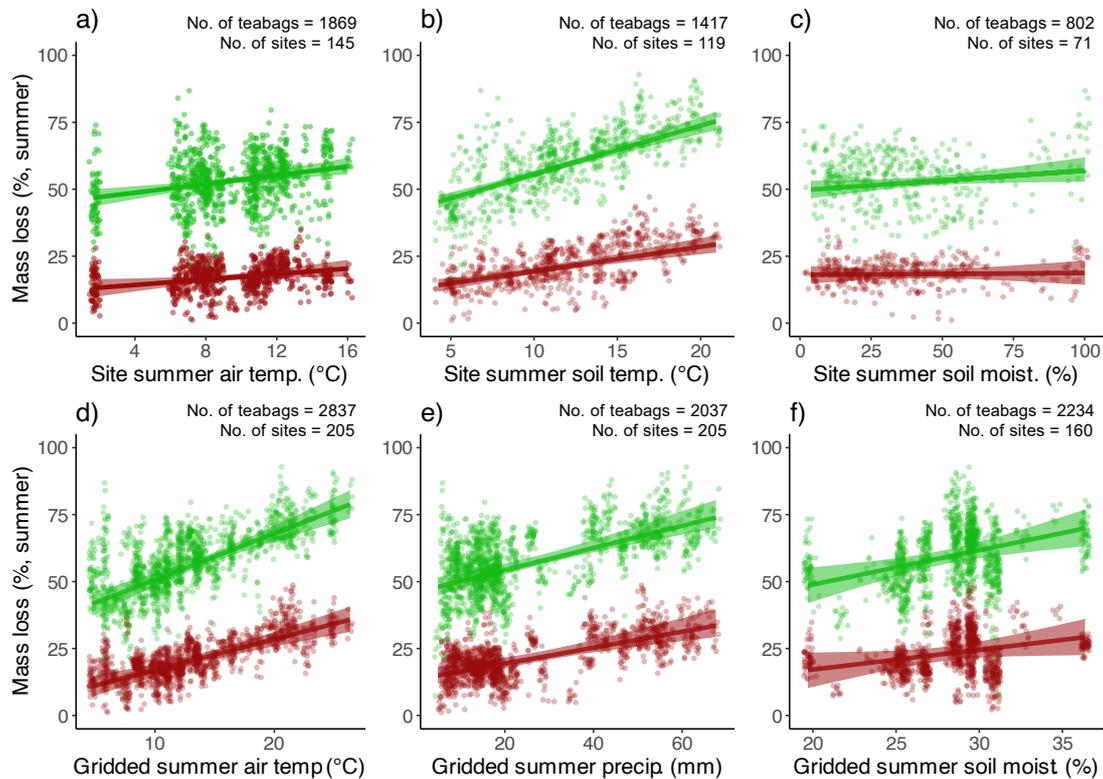
173 **Figure 2.** Annual mass loss of green and rooibos tea compared to mass loss of a range of
 174 representative tundra species. Tea types are indicated by red (rooibos tea) and green (green
 175 tea) boxplots. Tundra species were collected from two sites: the Kluane Range Mountains,
 176 Yukon, Canada (62°N) and Qikiqtaruk-Herschel Island, Yukon, Canada (70°N). All litter and
 177 tea were decomposed for one year in a common litter bed at 5-8cm depth at Kluane Lake
 178 following methods outlined in Cornelissen et al. (2007), with 10 replicates per species' litterbag
 179 and tea type. Methods are described in the Decomposition experiment section.

180

181 **Results**

182 We found that summer mass loss increased linearly with soil temperature (Fig. 3b) across
 183 tundra sites by $1.94\% \pm 0.31\%$ per °C for labile green tea and $1.09\% \pm 0.29\%$ per °C for
 184 recalcitrant rooibos tea, even when including a polynomial term (that allows for exponential
 185 relationships) in the models (Fig. S2). Relationships were consistent across incubation periods
 186 (Figs. S3-S4) and decomposition metrics (Figs. S5-S6), with higher temperatures associated
 187 with a lower stabilisation factor (S) and a faster decomposition rate (k). Within-site mass loss
 188 also increased with soil temperature (Fig. S7).

189



190

191 **Figure 3.** Relationships between litter decomposition, climate and environmental variables
 192 were linear across the range of environmental conditions found across study sites (see also
 193 Figs. S2 and S8 for non-linear models). Plotted relationships are between litter decomposition
 194 (mass loss), locally-measured environmental variables (a-c) and gridded climate data (d-f) for
 195 the summer incubation period (see also Fig. S3 for winter incubations and Fig. S4 for year-
 196 long incubations). Points indicate individual tea bag replicates across all sites. Lines indicate
 197 hierarchical Bayesian model fit with 95% credible intervals. Colours indicate tea type (red =
 198 roibos tea, green = green tea). See Table S2 for model outputs, and Table S3 and Fig. S2
 199 for results assuming polynomial relationships. Methods are detailed in the Environmental
 200 relationships section.

201

202 Summer mass loss increased with locally measured soil moisture across tundra sites (green
 203 tea: $0.07\% \pm 0.06\%$ per % moisture, roibos tea $0.01\% \pm 0.06\%$ per % moisture, Fig. 3c).

204

205 Relationships for winter and year-long incubations were weaker than for summer incubations
 206 (Figs. S3-4). Mass loss showed a weak positive relationship with soil moisture within sites
 207 (Fig. S9). Soil moisture did not influence the relationships between soil temperature and mass
 208 loss, but litter mass loss was higher at wetter versus drier sites at any given temperature (Fig.
 209 S9).

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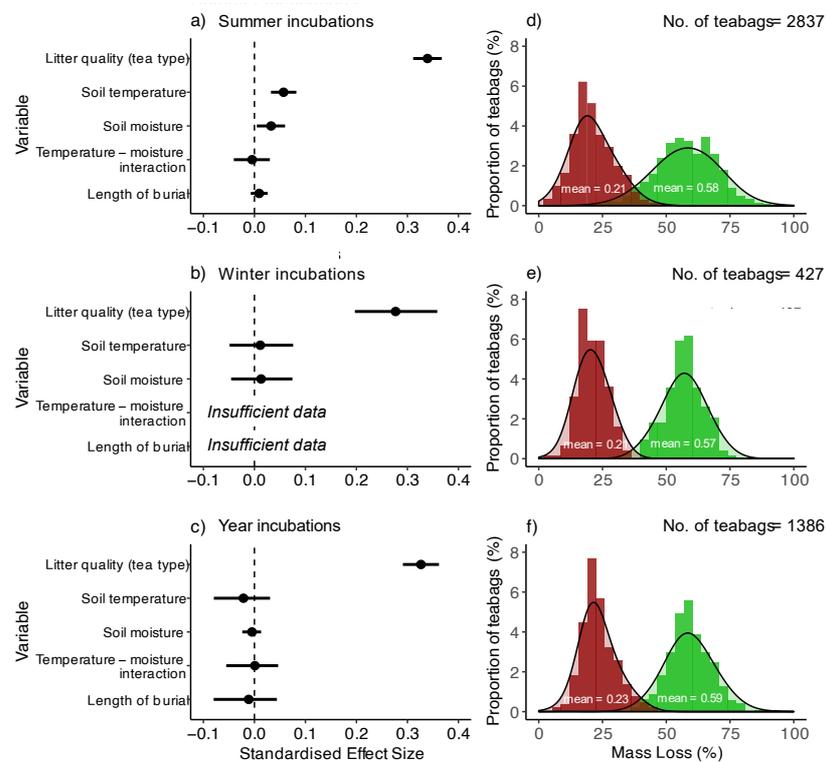
211 Relationships across sites were best explained by linear, rather than polynomial relationships
 212 for temperature (Fig. S2), although relationships of green tea summer mass loss with site air
 213 temperature and soil moisture showed weak signs of saturation and an exponential

214 relationship, respectively (Fig. S2a,c). In the Eastern Hemisphere, we found an exponential
 215 relationship with soil moisture that was driven by high mass loss despite high soil moisture at
 216 sites on Svalbard, while the relationship was best explained by a linear model in the Western
 217 Hemisphere (Fig. S8).

218

219 Litter quality was the strongest predictor of litter decomposition (Fig. 4a-c), explaining six and
 220 ten times more variation in summer mass loss than soil temperature and soil moisture,
 221 respectively. This strong effect of litter quality was maintained across incubation periods (Fig.
 222 4d-f) and mass loss of the two tea types did not converge after two years (Fig. S10).

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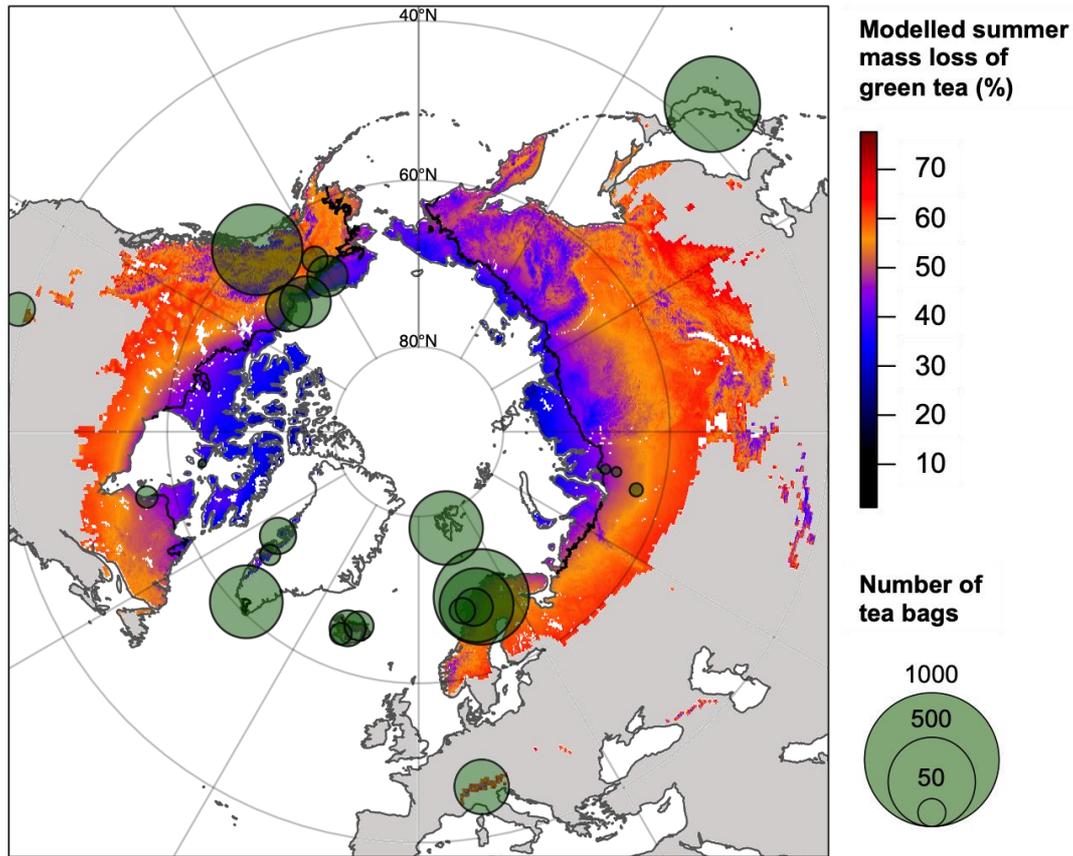
225 **Figure 4.** Litter quality explained greater variation in litter mass loss than environmental
 226 variables. Standardised effect sizes of locally measured environmental variables and litter
 227 substrate for summer incubations, winter incubations and year incubations (a-c). We did not
 228 have enough variation in incubation period to calculate the effect of length of burial (days
 229 within each incubation period) and the dataset was not large enough to test for a temperature-
 230 moisture interaction for winter incubations. Distribution of mass loss values for the two litter
 231 substrates (rooibos tea – red, green tea – green) for summer, winter and year incubations (d-
 232 f). Methods are detailed in the Environmental relationships – temperature moisture interaction
 233 section.

234

235 Relationships using gridded climate data were consistent with site-level climate data (Figs. 3,
 236 S3 and S4). We found strong linear relationships between decomposition and all gridded
 237 climate variables (Fig. 3d-f) and the interaction between gridded temperature and moisture

238 similarly suggested greater mass loss in wetter sites (Fig. S9). Extrapolating relationships
239 across tundra and subarctic regions based on gridded climate and soil moisture revealed
240 strong spatial variation in decomposition along biogeographic gradients (Fig. 5).

241



242

243 **Figure 5.** Modelled summer decomposition (percent mass loss) of green tea for tundra and
244 sub-Arctic regions based on 1979 to 2013 mean summer air temperature (Climatologies at
245 high resolution for the Earth's land surface, CHELSA) and soil moisture (European Space
246 Agency data, ESA) from 1979 to 2013. Field collection locations are illustrated by green
247 circles, grouped by geographic region (Table S1, figure excludes Australian alpine region).
248 Circle size indicates the number of tea bag replicates within each geographic region. Tundra
249 and sub-Arctic classifications are based on Köppen-Geiger classification⁴⁷. Ice-covered areas
250 are excluded. The circum-Arctic treeline is indicated with a black line⁴⁸. Methods in the
251 Mapping decomposition section.

252

253 Discussion

254 Contrasting with theory and model estimates, we find strong linear relationships, rather than
255 exponential, between decomposition and summer soil temperature and moisture across the
256 tundra biome (Fig. 3). Our findings provide comprehensive evidence that climate-driven
257 changes to plant communities, and thus litter quality, could have a greater impact on litter
258 decomposition than the direct effects of warming in the tundra (Fig. 4). Our results align with
259 site-specific studies that find that decomposition is more strongly influenced by litter quality

260 than climatic variability^{15,26–29,35,36,46,49}. Many sites across the tundra are currently undergoing
261 rapid vegetation change^{31,32,50,51}, notably an increase in shrub species with relatively
262 recalcitrant litter, which in many cases are out-competing graminoids with relatively labile
263 litter^{33,34}. This vegetation change has been hypothesised to partly counteract the effects of
264 warming on litter decomposition¹⁶. Our results suggest that the biotic effects of vegetation
265 change could outweigh the direct effects of warming on tundra litter decomposition, though
266 biotic changes will likely occur more slowly and lag behind warming^{15,50}.

267

268 We found positive linear relationships between decomposition, soil temperature and soil
269 moisture, with the greatest decomposition occurring in warmer and wetter sites (Fig. 3, Fig.
270 S9). Our 330 study sites encompass the linear range of global biogeochemical modelled
271 relationships of temperature and decomposition, but the non-linear range of soil moisture (Fig.
272 1, ²¹). Thus, our findings indicate that soil moisture may play a key role in mediating the effects
273 of warming on litter decomposition across the temperature-limited tundra biome. Based on
274 these relationships, we estimate that predicted Arctic warming of 2 to 10°C over the 21st
275 century could double summer litter mass loss at the coldest tundra sites. However, changes
276 are highly contingent upon site-specific factors, including moisture availability, substrate
277 quality and decomposer community^{15,24,25,27–30,44,52,53}. Although we focus on short-term
278 decomposition processes, greater early-stage decomposition could accelerate
279 biogeochemical cycling⁵⁴ and stimulate the loss of older organic carbon^{16,55} through nitrogen
280 mining^{56–58} or priming of microbial communities^{23,59,60}.

281

282 Contrary to the relationships assumed in many Earth system models²¹, we observed neither
283 an exponential increase in decomposition with temperature²⁰, nor a decrease in
284 decomposition at the highest moisture values (Fig. 3). However, we observed considerable
285 within-site variation in decomposition, emphasising the importance of site-specific factors⁵²
286 such as microbial community⁶¹ and soil chemistry⁶². We explored the site-level relationships
287 with general additive models as well as polynomial Bayesian models, and found that overall
288 relationships between environmental variables and decomposition were best fit by linear
289 relationships across variation in temperature and soil moisture (Figs.S2 and S8). However,
290 we did find that for the Eastern Hemisphere, there was an exponential relationship between
291 mass loss and soil moisture driven by data from Svalbard (Figs.S2 and Fig. S8). Overall, our
292 findings could indicate that decomposition is underestimated at colder or wetter tundra sites,
293 but overestimated at warmer sites in current model simulations.

294

295 Discrepancy between field observations and modelled decomposition could be caused by
296 environmental interactions. Environmental drivers such as warming and freeze-thaw

297 dynamics may have different influences across the year³⁵. With warming, higher temperatures
298 dry surface soils and reduce decomposer activity⁶³, as has been observed in warming
299 experiments^{17,64} and long-term monitoring⁶⁵. Biotic changes to either plant^{15,27,29} or
300 decomposer communities^{57,66,67} may also respond in complex ways to environmental change.
301 In addition, spatial patterning of landforms and environmental change such as permafrost thaw
302 can create wetter and drier microclimates within the same landscapes that can alter
303 decomposition across scales^{44,46,68}. Accounting for real-world biotic and abiotic patterns and
304 interactions among the drivers of decomposition in Earth system models will be critical to more
305 accurately projecting the effects of warming on decomposition and resulting losses to carbon
306 stores^{18,20,21,63}.

307

308 Our tundra-wide decomposition experiment has a number of caveats. Green and rooibos tea
309 are not tundra plant species, but they do encompass the decomposability of tundra plant
310 species (Fig. 2, Fig. S1) and thus provide an excellent common substrate for decomposition
311 studies. Although tea undergoes leaching processes, losing mass due to the loss of water-
312 soluble compounds during *in situ* decomposition⁶⁹, so too do tundra plants⁴¹. We tested
313 leaching rates in our study, finding ~20% greater mass loss for green tea and ~7% greater
314 mass loss for rooibos tea in two-month incubations rather than in 24-hour incubations in liquid
315 water. We found no substantial difference in mass loss with replacement of water across
316 incubations (Fig. S11), suggesting that leaching processes with lateral water flow is likely not
317 a major driver of mass loss in Tea Bag Index studies. Our study only encompasses short-term
318 decomposition with incubation lengths from three months to two years. Litter quality may have
319 a weaker effect on decomposition over longer time periods, and climate or other environmental
320 influences may become stronger over time⁷⁰⁻⁷³.

321

322 We used gridded climate data for our tundra-wide extrapolation and for climate data at sites
323 where *in situ* measurements were not recorded. Gridded climate data at high latitudes are
324 extrapolated from more limited meteorological data than at lower latitudes, and at high
325 latitudes, precipitation data are particularly limited⁷⁴. Thus, extrapolations of our statistical
326 results across the tundra biome contain substantial inherent uncertainty (Fig. 5). Our results
327 suggest that decomposition can nonetheless be mapped across large scales⁷⁵, and potentially
328 facilitate prediction of decomposition under future climatic conditions. Surprisingly,
329 relationships fitted with *in situ* or the corresponding gridded data displayed consistent trends
330 (Figs. 3, S3-7) although gridded data displayed steeper curves. This suggests that gridded
331 data are fit to represent macroclimatic control over decomposition but could lead to
332 overestimations of decomposition under future climatic conditions. An open question is how the

333 more nuanced view of decomposition offered by *in situ* data scales up in explaining global
334 patterns of decomposition ⁷⁶.

335

336 Changing decomposition rates will have profound implications for the global carbon cycle as
337 the climate is warming ². Warming-induced acceleration of litter decomposition could greatly
338 increase carbon losses in the tundra and other high-latitude ecosystems⁶, which have
339 historically acted as long-term carbon sinks^{14,77}. Tundra regions are also predicted to undergo
340 some of the greatest carbon losses over the coming century⁴, although predictions are highly
341 sensitive to data availability⁵. Our study provides well-resolved statistical relationships from
342 standardised field observations that can be used to parameterise Earth system models and
343 refine estimates of this critical feedback to the global carbon cycle. Ultimately, our findings
344 indicate that climate change is likely to increase decomposition across the tundra biome, but
345 that warming-induced vegetation change could have even more pronounced repercussions
346 for this globally important high-latitude carbon store.

347

348 **Methods**

349 We buried 5647 tea bags *in situ* at 5-8 cm depth at 330 sites across the tundra biome (Fig. 2,
350 Table S1). Our database has a hierarchical structure with plots (geographic areas including
351 multiple tea bag incubations) within sites (unique grid referenced locations of multiple plots)
352 within grid cells (the pixels of the gridded climate data, Table S1).

353

354 We recovered tea after three- (summer), nine- (winter), twelve-month (year) and two-year
355 incubations and calculated three metrics of decomposition: (1) percent mass loss, indicating
356 the proportion of initial mass decomposed, (2) stabilisation factor (*S*), indicating the proportion
357 of labile material remaining when initial decomposition has stabilised, and thus long-term
358 carbon storage potential and (3) decomposition rate (*k*), indicating the rate at which labile
359 material is lost, and thus short-term turnover¹⁹. We removed tea bags with experimental
360 treatments, that increased in mass due to fungal growth, got lost, split during extraction, where
361 labels were no longer legible or when only one site or plot was included per grid cell, resulting
362 in a sample size of 3717 tea bags in the final analysed dataset.

363

364 We examined relationships among the three decomposition metrics, three locally-measured
365 environmental variables (air temperature, soil temperature and soil moisture), and three
366 gridded climate variables: air temperature and precipitation from Climatologies at High
367 Resolution for the Earth's Land Surface (CHELSA)⁷⁸ and European Space Agency (ESA) soil
368 moisture data⁷⁹ using hierarchical Bayesian models. We also modelled decomposition across
369 tundra and sub-Arctic regions⁴⁷ by extrapolating relationships using CHELSA and ESA soil

370 moisture data from 1979 to 2013.

371

372 **Site Descriptions**

373 We established 330 decomposition sites encompassing 26 geographic regions across the
374 circum-Arctic and alpine tundra (Table S1). Mean annual air temperatures ranged from -
375 10.2°C to 12.7°C, with mean summer temperatures of 24.9°C at the warmest site (Alpine
376 Japan, Site SSJ) and 3.3°C at the coldest site (Svalbard, Endalen Cassiope heath). Sites were
377 largely above treeline though some subarctic and alpine sites extended below treeline.

378

379 **Decomposition experiment**

380 We measured decomposition using two types of tea in woven nylon mesh bags – a labile
381 green tea and a recalcitrant rooibos tea – following the Tea Bag Index method¹⁹. The two tea
382 types represent dried leaves of two shrub species (*Camellia sinensis* – green tea and
383 *Aspalathus linearis* – rooibos tea), which strongly differ in their leaf structural and chemical
384 traits^{19,80,81}. Although these two species are not native to the tundra, their mass losses are
385 comparable with a range of tundra species (Fig. 2), and allow comparison across sites
386 globally^{19,82}. Mass loss via leaching of these tea bags is also comparable with previous studies
387 employing the common litter bag method (24-hour mass loss: 14% for rooibos tea and 37%
388 for green tea, compared to 8 - 32% in litter leaching studies⁸³).

389

390 We buried tea bags *in situ* at 5-8 cm depth during 2015-2017. We incubated tea for three
391 approximate time periods – three months (summer: on average 81 days across all sites from
392 late spring – late summer), nine months (winter: late summer to late spring) and twelve months
393 (year: late spring to late spring). Due to the logistical constraints of accessing some field sites,
394 not all incubations were carried out at all sites. We buried a minimum of three tea bag pairwise
395 replicates at each site for each given period. Tea bags were buried, rather than placed on the
396 surface, for consistency with the global standardised Tea Bag Index protocol¹⁹. Moreover, this
397 increased the likelihood of recovery across the time periods covered in this study. Surface
398 litter likely undergo greater fluctuations in temperature and moisture that may reduce
399 decomposition⁸³. Within a common site, we found that annual mass loss was greater for buried
400 teabags compared to those in the litter layer for green tea, but not rooibos tea (Fig. S12).
401 However, litter is commonly mixed into tundra soils through cryoturbation processes²³. Thus,
402 using a buried litter substrate serves as a proxy for both leaf litter decomposition when
403 incorporated into the soil and soil organic matter decomposition⁸⁴.

404

405 We weighed tea bags prior to burial, including both the bag and tag. Upon recovery, we dried
406 bags at 70°C for at least 48 hours, removed any attached soil or roots, and reweighed tea

407 bags. We subtracted the mass of the bag and label to determine the mass of the tea only, and
408 the initial masses were corrected to account for initial moisture and loss of material in transit
409 to field sites (approximately $5.6 \pm 0.8\%$ of mass for rooibos tea and $3.8 \pm 0.4\%$ for green tea,
410 measured using 10 unused tea bags at three different field sites).

411

412 ***Decomposition variables***

413 We calculated three metrics of decomposition: (1) overall mass loss (final tea mass divided by
414 initial tea mass) for each tea type.

415

416 *Equation 1:*

$$417 \text{ mass loss} = 1 - \left(\frac{M_t}{M_0}\right)$$

418 where M_t is equal to the mass of tea at time point t (days) and M_0 is the initial mass.

419

420 (2) The stabilisation factor (S), which describes the proportion of potentially decomposable
421 compounds (the hydrolysable or acid-digestible fraction, H) remaining upon stabilisation of
422 decomposition. S is calculated using green tea, for which mass loss has stabilised within three
423 months of burial¹⁹ (Fig. S10), whereby:

424

425 *Equation 2:*

$$426 S = 1 - \left(\frac{a_g}{H_g}\right)$$

427

428 where a_g is the decomposable fraction (mass loss) of green tea and H_g is the hydrolysable
429 fraction of green tea.

430

431 (3) The decomposition rate (k), which represents the rate at which decomposable compounds
432 are lost during decomposition. This two-pool decomposition constant was calculated based
433 on the methods outlined in Keuskamp et al. (2013), and is calculated using rooibos tea, for
434 which decomposition has not yet stabilised during the incubation periods covered by this
435 analysis¹⁹ (Fig. S10).

436

437 *Equation 3:*

$$438 k = \ln\left(\frac{a_r}{M_{t(r)} - a_r}\right) \times \frac{1}{t}$$

439 where M is equal to the mass of rooibos tea at time point t (days) and a_r is the decomposable
440 fraction of rooibos tea. a_r is calculated from the hydrolysable fraction of rooibos tea (H_r) and
441 stabilisation factor (S), whereby $a_r = H_r(1 - S)$.

442

443 ***Environmental variables***

444 Where possible, we measured local environmental variables at each site for the duration of
445 the incubation period. Soil temperatures were measured using digital iButtons (DS1921G
446 Thermochron iButtons, Maxim Integrated, San Jose, CA, US) or data loggers (HOBO RX3000
447 Remote Monitoring Station Data Logger, Onset Computer Corporation, Pocasset, MA; HOBO
448 Pendant temperature and light data loggers, Part # UA-002-64, Onset Computer Corporation,
449 Pocasset, MA; Lascar EL USB-1 temperature loggers, Lascar electronics, Salisbury, UK;
450 Theta Probe ML3 attached to a HH2 Moisture Meter Logger, DELTA-T-DEVICES, Cambridge,
451 UK). Soil moisture (percent volumetric water content) was measured using hand-held moisture
452 probes (Spectrum (SM100); HydroSense II; Stevens POGO probe, Stevens Water Monitoring
453 Systems Inc., Portland, OR, USA) at 5 cm depth. Where site-measured data were not
454 available, notably for air temperature, we used local weather station data, provided either by
455 the authors or additional contributors⁸⁵ and unpublished data (Annika Kristoffersson pers.
456 comm. 2017, Phil Marsh, pers. comm. 2017). All environmental data were trimmed to the
457 corresponding incubation period for analyses. Sites that did not have available local
458 environmental data were excluded from relevant analyses.

459

460 ***Gridded climate variables***

461 We used 'Climatologies at high resolution for the Earth's land surface areas' data (CHELSA,
462 0.0083×0.0083 degree resolution⁷⁸, <http://chelsa-climate.org>) to provide gridded temperature
463 and precipitation data for all sites, and to extrapolate decomposition across the tundra biome.
464 We extracted climatologies (covering the time period 1979 to 2013) for summer (June-July-
465 August), winter (December-January-February) and annual temperature and precipitation. We
466 used European Space Agency (ESA) Climate Change Initiative combined soil moisture data
467 product (0.25×0.25 -degree resolution⁷⁹, <https://www.esa-soilmoisture-cci.org>) to provide
468 modelled soil moisture for all sites and to extrapolate decomposition across the tundra. We
469 used daily data for the period 1979 to 2013 to build climatologies (summer, winter, year) to
470 align with CHELSA data.

471

472 We compared site-measured environmental data to gridded climate data using hierarchical
473 Bayesian models with grid cell and site as nested random effects using the R package
474 *MCMCglmm*⁸⁶ (Fig. 3, Figs. S2-S7 and S9). Site-measured temperature variables correlated

475 closely with gridded temperature data, exhibiting a near 1:1 relationship (Fig. S13). Site-
476 measured moisture was not correlated with average ESA soil moisture data or long-term
477 CHELSA precipitation data (Fig. S13). This discrepancy may result from greater spatial and
478 inter-annual variability in moisture or precipitation compared to temperature⁸⁷, or high within-
479 site variation in soil moisture that is not captured by spatially variable and data-poor high-
480 latitude precipitation records at the grid cell scale.

481

482 ***Environmental Relationships***

483 We conducted three analyses of the relationships among decomposition metrics and
484 environmental variables: (i) relationships between each individual decomposition metric and
485 each environmental variable across all sites (Fig. 3, Figs. S3-S6), (ii) relationships between
486 mass loss and environmental variables within each grid cell (Fig. S7) and (iii) relationships
487 between mass loss and environmental variables accounting for interactions between
488 temperature and moisture (Fig. S9).

489

490 Analyses of environmental relationships were conducted in the statistical programming
491 language *Stan* run through R (v. 3.3.3 to 4.2.3) using packages *rjags*⁸⁸ (v. 4.6) and *rstan*⁸⁹ (v.
492 2.17.3). In all cases, models were run until convergence was reached, which was assessed
493 both visually in trace plots and by ensuring that all Gelman–Rubin convergence diagnostic
494 values (\hat{R})⁹⁰ were less than 1.1. Code is available at:

495 <https://github.com/ShrubHub/TundraTeaHub>

496

497 *Environmental Relationships – individual variables*

498 The relationship between each decomposition metric (decomp) and environmental variable
499 (EV) was estimated from a hierarchical Bayesian model, with climatic variables as the
500 predictor variable and decomposition as the predictor variable, with grid cell (g), site (s, unique
501 grid referenced location) and plot (p, replicate plots within each location) as random effects,
502 varying by tea type (t):

503

504 *Equation 4:*

$$505 \text{decomp}_{p,t} \sim \text{Normal}(\alpha_{p,t} + \alpha_{s,t} + \alpha_{g,t}, \sigma)$$

506

507 We estimated relationships with decomposition metrics over space at the level at which
508 environmental variables were measured, including incubation length (days) as a fixed effect.
509 For example, relationships for gridded climate data were estimated at the level of the grid cell
510 (g), with site (s) and plot (p) as nested hierarchical random effects. Relationships for site-

511 measured variables were estimated at the site level, with plot (p) as a random effect. If
512 environmental variables were measured at the plot level, we summarised variables to the site
513 level and carried forward the standard deviation among plots into models. If there was only
514 one teabag per plot, one plot per site or one site per grid cell, $\alpha_{p,t}$ or $\alpha_{s,t}$ was set to zero. Note
515 that data availability differs for each environmental variable. For stabilisation factor (S) and
516 decomposition rate (k) models, we did not vary effects by tea type (t), since only one tea type
517 is used for each of these variables.

518

519 *Equation 5:*

$$520 \quad \alpha_{g,t} \sim \text{Normal}(\gamma_0_t + \gamma_1_t * EV_{g,t} + \gamma_2_t * days_{g,t}, \theta)$$

$$521 \quad \alpha_{p,t} \sim \text{Normal}(0, \sigma_1)$$

$$522 \quad \alpha_{s,t} \sim \text{Normal}(0, \sigma_2)$$

523

524 We modelled all incubation periods separately due to large differences in the availability of
525 environmental data and qualitative differences between conditions in different seasons such
526 as frozen ground during the winter.

527

528 Decomposition relationship with temperature and moisture can also be exponential, saturating
529 or sigmoidal ^{22,91,92}. To account for this, we also tested a potential non-linear relationship
530 between summer mass loss and the environmental variables by including a fourth fixed
531 parameter in equation 5, multiplying the square of the environmental variable, hence allowing
532 the relationship to be polynomial.

533

534 *Environmental Relationships – within grid cells*

535 We modelled the relationship between decomposition metrics and environmental variables
536 (single variables only) within grid cells using the same model structure, but by standardising
537 all environmental variables within a grid cell using mean zero and unit-variance scaling.

538

539 *Environmental Relationships – temperature and moisture interactions*

540 We modelled the relationships between mass loss and environmental variables over space
541 accounting for both temperature and moisture within the same model (both for site-measured
542 soil temperature and soil moisture, and for gridded air temperature and soil moisture). We
543 used the same model structure as for individual variables, but also included an interaction
544 term between these two environmental variables.

545

546 *Equation 6:*

547 $\alpha_{g,t} \sim \text{Normal}(\gamma_0_t + \gamma_1_t * \text{temp}_{g,t} + \gamma_2_t * \text{moisture}_{g,t} + \gamma_3_t * \text{temp}_{g,t} * \text{moisture}_{g,t} + \gamma_4_t$
548 $* \text{days}_{g,t}, \theta)$

549

550 We ran models with environmental data in original units, and also using standardised
551 environmental variables and incubation length using mean zero and unit-variance scaling to
552 allow comparison across environmental variables.

553

554 ***Mapping decomposition***

555 We used model estimates from the gridded climate variable model (Equation 6) to map
556 decomposition over space based on summer temperature and moisture for tundra and
557 subarctic climate regions. We mapped gridded temperature of the warmest quarter (CHELSA
558 bio10) and gridded summer soil moisture (ESA, June-July-August) as environmental
559 variables. We used the coefficients for green tea (Fig. 5) and rooibos tea (Fig. S14), and
560 assumed the mean incubation length across summer treatments (81 days). We masked
561 estimates to tundra and subarctic climate regions based on the Köppen-Geiger climate
562 classification⁹³ (regions ET, Dsc, Dsc, Dwc, Dwd, Dfc, Dfd). We included an estimation of
563 global treeline based on the Circum-Arctic Vegetation Map (CAVM) classification⁴⁸.

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789 **Supplementary Tables**

790 **Table S1.** Summary of geographic locations used in main study, indicating number of sites
 791 and plots (the base study unit), number of tea bag replicates used in study, and mean
 792 temperatures (CHELSA data 1979-2013, summer = warmest quarter, winter = coldest
 793 quarter), mean elevation of the sites, and a general description of the sites.

Geographic Region	Number of sites	Number of plots	Number of tea bags	Mean temperature (°C) (year / summer / winter)			Elevation (m a.s.l.)	Sites description
Alpine Japan	45	46	776	6.7	18.7	-4.5	1576	Alpine tundra
Auðkúluheiði, Iceland	3	26	110	0.9	8.6	-5.0	486	Subarctic shrub tundra dominated by <i>Betula nana</i>
Australian Alps	1	18	191	4.7	11.4	-1.5	1880	Alpine tundra
Disko Island, Greenland	7	7	112	-4.0	7.1	-15.4	8	Prostrate/hemi-prostrate dwarf-shrub, lichen tundra
Fairbanks, Alaska	7	14	56	-4.8	14.3	-22.0	501	Erect dwarf-shrub, moss tundra
Kilpisjärvi, Finland	82	120	751	-2.1	10.2	-13.2	672	Subarctic tundra from birch forest to birch shrubs
Gothic Mountain, Colorado, USA	5	5	95	2.2	13.4	-8.1	2922	Alpine tundra
Italian Alps	2	14	116	-1.6	7.5	-10.1	2569	Alpine tundra dominated by <i>Salix herbacea</i> and mosses
Kangerlussuaq, Greenland	2	2	36	-5.6	7.5	-16.6	280	<i>Salix glauca</i> and <i>Betula nana</i> tundra and steppe tundra
Khanymey, western Siberia	2	2	15	-3.6	15.4	-21.1	66	Subarctic pine and larch forests
Kluane, Yukon, Canada	15	72	757	-3.1	8.8	-14.1	1414	Alpine tundra
Lofoten Islands, Norway	1	16	55	5.8	12.6	0.8	15	Subarctic grass and forb tundra
Narsarsuaq, Greenland	10	49	450	-3.3	6.6	-12.0	207	Prostrate dwarf-shrub, herb, lichen tundra
Northern Norway	35	62	119	0.5	11.9	-9.7	356	Gradient from birch forests to alpine tundra (shrubs, meadows).
Northern Sweden	56	122	467	-2.1	9.9	-12.8	796	Gradient from birch forests to alpine tundra (heathland, shrubs, tundra meadows).
Qikiqtaruk-Herschel Island, Yukon, Canada	9	14	224	-9.4	7.6	-24.3	38	Coastal tundra with shrub, sedge, grass and forb.
Southampton Island, Nunavut, Canada	1	1	5	-9.4	6.4	-26.6	12	Cryptogam, barren complex
Svalbard	25	109	468	-6.3	4.2	-14.7	40	Graminoid, forb, cryptogam tundra and dwarf-shrub, lichen tundra
Swiss Alps	3	61	256	-1.00	8.0	-9.4	2180	Alpine tundra
Peistareykir, Iceland	2	12	72	1.7	8.9	-3.5	340	Subarctic shrub tundra dominated by <i>Betula nana</i>
Pingvellir, Iceland	1	10	40	4.0	10.7	-1.1	120	Subarctic tundra dominated by birch trees and dwarf birches
Tazovsky, western Siberia	1	1	8	-7.4	12.5	-25.1	24	Subarctic shrub tundra with sporadic trees
Toolik Lake, Alaska, USA	2	7	140	-10.2	10.4	-26.6	760	Alpine tundra
Trail Valley, NWT, Canada	10	30	180	-9.1	12.0	-27.2	95	From sporadic boreal tree species to shrub tundra
Umiujaq, Québec, Canada	2	2	40	-3.9	9.9	-20.8	81	Subarctic shrub tundra with maritime influence
Urengoy, western Siberia	1	1	8	-6.3	13.7	-24.0	26	Subarctic tundra with sporadic trees

794 **Table S2.** Model outputs for individual environmental variable – decomposition relationships.
795 Bold rows designate relationships (slope parameter) for which the credible interval does not
796 cross zero (i.e., the relationship is “significant”). Sample size indicates number of tea samples
797 available to test relationships. Effective sample size indicates number of convergent model
798 runs. G and R indicate green and rooibos tea, respectively.

Environ. Variable	Decomp. variable	Time period	Tea Type	Parameter	Mean	SD	2.50%	97.50%	Sample size	Effective sample size
Air temp. (measured)	Mass loss	Summer	G	Intercept	0.531	0.006	0.519	0.543	1913	15 000
Air temp. (measured)	Mass loss	Summer	R	Intercept	0.17	0.006	0.158	0.182	1913	15 000
Air temp. (measured)	Mass loss	Summer	G	Slope	0.008	0.002	0.004	0.012	1913	15 000
Air temp. (measured)	Mass loss	Summer	R	Slope	0.005	0.002	0.001	0.004	1913	15 000
Soil temp. (measured)	Mass loss	Summer	G	Intercept	0.605	0.006	0.593	0.616	1560	15 000
Soil temp. (measured)	Mass loss	Summer	R	Intercept	0.22	0.006	0.208	0.231	1560	15 000
Soil temp. (measured)	Mass loss	Summer	G	Slope	0.02	0.002	0.017	0.023	1560	15 000
Soil temp. (measured)	Mass loss	Summer	R	Slope	0.011	0.002	0.008	0.014	1560	15 000
Moisture (measured)	Mass loss	Summer	G	Intercept	0.523	0.009	0.504	0.541	917	15 000
Moisture (measured)	Mass loss	Summer	R	Intercept	0.183	0.009	0.165	0.201	917	15 000
Moisture (measured)	Mass loss	Summer	G	Slope	7.36E-04	3.75E-04	1.45E-05	1.47E-03	917	14 142
Moisture (measured)	Mass loss	Summer	R	Slope	7.16E-05	3.68E-04	-6.57E-04	8.09E-04	917	15 000
Air temp. (CHELSA)	Mass loss	Summer	G	Intercept	0.559	0.009	0.541	0.577	2837	15 000
Air temp. (CHELSA)	Mass loss	Summer	R	Intercept	0.204	0.009	0.187	0.222	2837	15 000
Air temp. (CHELSA)	Mass loss	Summer	G	Slope	0.017	0.002	0.013	0.021	2837	7178
Air temp. (CHELSA)	Mass loss	Summer	R	Slope	0.012	0.002	0.008	0.015	2837	15 000
Precip. (CHELSA)	Mass loss	Summer	G	Intercept	0.565	0.01	0.545	0.585	2837	15 000
Precip. (CHELSA)	Mass loss	Summer	R	Intercept	0.209	0.01	0.19	0.229	2837	15 000
Precip. (CHELSA)	Mass loss	Summer	G	Slope	0.004	0.001	0.003	0.005	2837	15 000
Precip. (CHELSA)	Mass loss	Summer	R	Slope	0.003	0.001	0.003	0.004	2837	15 000
Moisture (ESA)	Mass loss	Summer	G	Intercept	0.595	0.013	0.569	0.621	2234	15 000
Moisture (ESA)	Mass loss	Summer	R	Intercept	0.232	0.013	0.206	0.258	2234	15 000
Moisture (ESA)	Mass loss	Summer	G	Slope	0.013	0.004	0.005	0.02	2234	15 000
Moisture (ESA)	Mass loss	Summer	R	Slope	0.007	0.004	-9.38E-05	0.015	2234	15 000
Air temp. (measured)	Mass loss	Winter	G	Intercept	0.561	0.03	0.503	0.621	176	15 000
Air temp. (measured)	Mass loss	Winter	R	Intercept	0.226	0.031	0.165	0.287	176	15 000
Air temp. (measured)	Mass loss	Winter	G	Slope	0.044	0.016	0.012	0.074	176	15 000
Air temp. (measured)	Mass loss	Winter	R	Slope	-0.002	0.016	-0.033	0.029	176	15 000
Soil temp. (measured)	Mass loss	Winter	G	Intercept	0.498	0.041	0.416	0.58	71	5842

Soil temp. (measured)	Mass loss	Winter	R	Intercept	0.195	0.113	0.113	0.278	71	15 000
Soil temp. (measured)	Mass loss	Winter	G	Slope	0.003	0.008	-0.013	0.019	71	7964
Soil temp. (measured)	Mass loss	Winter	R	Slope	0.004	0.008	-0.012	0.02	71	15 000
Moisture (measured)	Mass loss	Winter	G	Intercept	0.553	0.007	0.538	0.567	206	9488
Moisture (measured)	Mass loss	Winter	R	Intercept	0.19	0.007	0.175	0.204	206	8980
Moisture (measured)	Mass loss	Winter	G	Slope	0.001	0.001	-1.51E-05	0.002	206	12 291
Moisture (measured)	Mass loss	Winter	R	Slope	4.42E-04	0.001	-5.93E-04	0.001	206	15 000
Air temp. (CHELSA)	Mass loss	Winter	G	Intercept	0.542	0.005	0.451	0.637	427	15 000
Air temp. (CHELSA)	Mass loss	Winter	R	Intercept	0.208	0.005	0.116	0.3	427	15 000
Air temp. (CHELSA)	Mass loss	Winter	G	Slope	0.019	0.012	-0.006	0.043	427	15 000
Air temp. (CHELSA)	Mass loss	Winter	R	Slope	-3.47E-05	0.012	-0.024	0.024	427	15 000
Precip. (CHELSA)	Mass loss	Winter	G	Intercept	0.54	0.046	0.453	0.634	427	15 000
Precip. (CHELSA)	Mass loss	Winter	R	Intercept	0.206	0.046	0.114	0.299	427	15 000
Precip. (CHELSA)	Mass loss	Winter	G	Slope	0.005	0.003	-0.001	0.011	427	15 000
Precip. (CHELSA)	Mass loss	Winter	R	Slope	-4.88E-04	0.003	-0.007	0.006	427	15 000
Moisture (ESA)	Mass loss	Winter	G	Intercept	0.541	0.045	0.451	0.633	309	15 000
Moisture (ESA)	Mass loss	Winter	R	Intercept	0.207	0.046	0.118	0.298	309	15 000
Moisture (ESA)	Mass loss	Winter	G	Slope	0.073	0.032	0.008	0.137	309	15 000
Moisture (ESA)	Mass loss	Winter	R	Slope	-0.01	0.033	-0.075	0.054	309	15 000
Air temp. (measured)	Mass loss	Year	G	Intercept	0.581	0.006	0.578	0.6	1251	15 000
Air temp. (measured)	Mass loss	Year	R	Intercept	0.228	0.006	0.217	0.24	1251	15 000
Air temp. (measured)	Mass loss	Year	G	Slope	0.011	0.002	0.007	0.015	1251	15 000
Air temp. (measured)	Mass loss	Year	R	Slope	0.014	0.002	0.01	0.018	1251	15 000
Soil temp. (measured)	Mass loss	Year	G	Intercept	0.591	0.014	0.564	0.619	342	15 000
Soil temp. (measured)	Mass loss	Year	R	Intercept	0.263	0.014	0.237	0.29	342	15 000
Soil temp. (measured)	Mass loss	Year	G	Slope	0.018	0.004	0.011	0.027	342	15 000
Soil temp. (measured)	Mass loss	Year	R	Slope	0.019	0.004	0.011	0.027	342	15 000
Moisture (measured)	Mass loss	Year	G	Intercept	0.614	0.005	0.604	0.625	760	15 000
Moisture (measured)	Mass loss	Year	R	Intercept	0.255	0.006	0.245	0.266	760	15 000
Moisture (measured)	Mass loss	Year	G	Slope	-0.001	3.89E-04	-0.001	1.54E-04	760	15 000
Moisture (measured)	Mass loss	Year	R	Slope	-0.001	4.06E-04	-0.002	1.06E-04	760	15 000
Air temp. (CHELSA)	Mass loss	Year	G	Intercept	0.606	0.011	0.585	0.628	1377	15 000
Air temp. (CHELSA)	Mass loss	Year	R	Intercept	0.236	0.011	0.215	0.258	1377	15 000
Air temp. (CHELSA)	Mass loss	Year	G	Slope	0.015	0.003	0.009	0.02	1377	10 775
Air temp. (CHELSA)	Mass loss	Year	R	Slope	0.01	0.003	0.004	0.015	1377	9975

Precip. (CHELSA)	Mass loss	Year	G	Intercept	0.601	0.015	0.572	0.631	1377	15 000
Precip. (CHELSA)	Mass loss	Year	R	Intercept	0.236	0.015	0.207	0.265	1377	15 000
Precip. (CHELSA)	Mass loss	Year	G	Slope	0.001	4.15E-04	2.48E-04	0.002	1377	15 000
Precip. (CHELSA)	Mass loss	Year	R	Slope	3.06E-04	4.05E-04	-5.02E-04	0.001	1377	15 000
Moisture (ESA)	Mass loss	Year	G	Intercept	0.62	0.017	0.588	0.655	1098	15 000
Moisture (ESA)	Mass loss	Year	R	Intercept	0.252	0.017	0.219	0.285	1098	15 000
Moisture (ESA)	Mass loss	Year	G	Slope	0.008	0.004	-4.67E-05	0.015	1098	15 000
Moisture (ESA)	Mass loss	Year	R	Slope	0.005	0.004	-2.84E-03	0.007	1098	15 000
Air temp. (measured)	k	Summer	R	Intercept	0.011	3.89E-04	0.01	0.012	927	15 000
Air temp. (measured)	k	Summer	R	Slope	4.28E-04	1.36E-04	1.59E-04	6.97E-04	927	15 000
Soil temp. (measured)	k	Summer	R	Intercept	0.011	3.29E-04	0.01	0.011	704	15 000
Soil temp. (measured)	k	Summer	R	Slope	1.46E-04	8.92E-05	-2.85E-05	3.20E-04	704	15,000
Moisture (measured)	k	Summer	R	Intercept	0.012	0.001	0.011	0.013	398	15 000
Moisture (measured)	k	Summer	R	Slope	-3.97E-05	2.54E-05	-9.01E-05	9.95E-06	398	15 000
Air temp. (CHELSA)	k	Summer	R	Intercept	0.011	0.001	0.01	0.012	1403	15 000
Air temp. (CHELSA)	k	Summer	R	Slope	1.72E-04	1.20E-04	-6.58E-05	4.08E-04	1403	15 000
Precip. (CHELSA)	k	Summer	R	Intercept	0.011	0.001	0.01	0.012	1403	15 000
Precip. (CHELSA)	k	Summer	R	Slope	1.07E-05	3.22E-05	-5.23E-05	7.35E-05	1403	15 000
Moisture (ESA)	k	Summer	R	Intercept	0.011	0.001	0.01	0.012	1108	15 000
Moisture (ESA)	k	Summer	R	Slope	-7.21E-07	1.23E-06	-2.90E-04	3.02E-04	1108	15 000
Air temp. (measured)	S	Summer	G	Intercept	0.372	0.009	0.366	0.39	944	15 000
Air temp. (measured)	S	Summer	G	Slope	-0.007	0.003	-0.014	-9.41E-04	944	15 000
Soil temp. (measured)	S	Summer	G	Intercept	0.327	0.009	0.309	0.346	715	15 000
Soil temp. (measured)	S	Summer	G	Slope	-0.026	0.002	-0.031	-0.021	715	15 000
Moisture (measured)	S	Summer	G	Intercept	0.373	0.015	0.344	0.403	408	15 000
Moisture (measured)	S	Summer	G	Slope	-0.001	4.67E-06	-0.002	8.67E-06	408	15 000
Air temp. (CHELSA)	S	Summer	G	Intercept	0.364	0.013	0.338	0.39	1436	8376
Air temp. (CHELSA)	S	Summer	G	Slope	-0.021	0.003	-0.026	-0.016	1436	8560
Precip. (CHELSA)	S	Summer	G	Intercept	0.355	0.015	0.326	0.384	1436	15 000
Precip. (CHELSA)	S	Summer	G	Slope	-0.005	0.001	-0.007	-0.004	1436	15 000
Moisture (ESA)	S	Summer	G	Intercept	0.297	0.019	0.258	0.334	1128	15 000
Moisture (ESA)	S	Summer	G	Slope	-0.019	0.005	-0.029	-0.008	1128	15 000

800 **Table S3.** Model outputs for individual environmental variable – summer mass loss polynomial
801 relationships. Bold rows designate polynomial relationships (polynomial parameter) for which
802 the credible interval does not cross zero (i.e., the relationship is “significant”). Sample size
803 indicates number of tea samples available to test relationships. Effective sample size indicates
804 number of convergent models runs. G and R indicate green and roibos tea, respectively.

Environ. Variable	Decomp. variable	Time period	Tea Type	Parameter	Mean	SD	2.50%	97.50%	Sample size	Effective sample size
Air temp. (measured)	Mass loss	Summer	G	Intercept	0.535	0.00624	0.523	0.547	1913	31625
Air temp. (measured)	Mass loss	Summer	R	Intercept	0.172	0.00634	0.159	0.184	1913	30383
Air temp. (measured)	Mass loss	Summer	G	Slope	0.0078	0.0021	0.00374	0.0119	1913	25671
Air temp. (measured)	Mass loss	Summer	R	Slope	0.00516	0.0021	0.0011	0.00929	1913	27935
Air temp. (measured)	Mass loss	Summer	G	Polynomial	-5.48E-05	2.31E-05	-0.000101	-9.29E-06	1913	15544
Air temp. (measured)	Mass loss	Summer	R	Polynomial	-2.48E-05	3.37E-05	-8.99E-05	4.17E-05	1913	15676
Soil temp. (measured)	Mass loss	Summer	G	Intercept	0.617	0.00856	0.6	0.633	1560	17804
Soil temp. (measured)	Mass loss	Summer	R	Intercept	0.215	0.00824	0.199	0.232	1560	18677
Soil temp. (measured)	Mass loss	Summer	G	Slope	0.0201	0.00157	0.017	0.0231	1560	22592
Soil temp. (measured)	Mass loss	Summer	R	Slope	0.0109	0.00153	0.00797	0.0139	1560	21301
Soil temp. (measured)	Mass loss	Summer	G	Polynomial	-0.000642	0.000329	-0.00129	6.70E-06	1560	15355
Soil temp. (measured)	Mass loss	Summer	R	Polynomial	0.000259	0.00032	-0.000375	0.000897	1560	15769
Moisture (measured)	Mass loss	Summer	G	Intercept	0.495	0.0116	0.473	0.518	917	24515
Moisture (measured)	Mass loss	Summer	R	Intercept	0.178	0.0113	0.155	0.2	917	26840
Moisture (measured)	Mass loss	Summer	G	Slope	-0.000848	0.000587	-0.00202	0.000274	917	15676
Moisture (measured)	Mass loss	Summer	R	Slope	-0.000198	0.000499	-0.00117	0.000775	917	16094
Moisture (measured)	Mass loss	Summer	G	Polynomial	5.16E-05	1.43E-05	2.38E-05	8.02E-05	917	16802
Moisture (measured)	Mass loss	Summer	R	Polynomial	9.66E-06	1.27E-05	-1.51E-05	3.52E-05	917	17865
Air temp. (CHELSA)	Mass loss	Summer	G	Intercept	0.555	0.0127	0.53	0.58	2837	6177
Air temp. (CHELSA)	Mass loss	Summer	R	Intercept	0.197	0.0119	0.174	0.22	2837	5970
Air temp. (CHELSA)	Mass loss	Summer	G	Slope	0.0166	0.00245	0.0119	0.0214	2837	5759
Air temp. (CHELSA)	Mass loss	Summer	R	Slope	0.0107	0.00216	0.00651	0.015	2837	7225
Air temp. (CHELSA)	Mass loss	Summer	G	Polynomial	0.000197	0.000363	-5.00E-04	0.000934	2837	5128
Air temp. (CHELSA)	Mass loss	Summer	R	Polynomial	0.00031	0.000343	-0.000363	0.000979	2837	6156
Precip. (CHELSA)	Mass loss	Summer	G	Intercept	0.56	0.0165	0.527	0.592	2837	11978
Precip. (CHELSA)	Mass loss	Summer	R	Intercept	0.189	0.0164	0.157	0.221	2837	13451
Precip. (CHELSA)	Mass loss	Summer	G	Slope	0.00384	0.00103	0.00179	0.00585	2837	10612
Precip. (CHELSA)	Mass loss	Summer	R	Slope	0.00174	0.000988	-0.000212	0.00365	2837	12732
Precip. (CHELSA)	Mass loss	Summer	G	Polynomial	1.38E-05	3.94E-05	-6.38E-05	9.10E-05	2837	10820

Precip. (CHELSA)	Mass loss	Summer	R	Polynomial	6.33E-05	3.91E-05	-1.26E-05	0.00014	2837	12939
Moisture (ESA)	Mass loss	Summer	G	Intercept	0.601	0.016	0.569	0.632	2234	35778
Moisture (ESA)	Mass loss	Summer	R	Intercept	0.23	0.0162	0.198	0.261	2234	35783
Moisture (ESA)	Mass loss	Summer	G	Slope	0.012	0.00402	0.00413	0.0199	2234	36885
Moisture (ESA)	Mass loss	Summer	R	Slope	0.00781	0.00406	-0.000131	0.0158	2234	37415
Moisture (ESA)	Mass loss	Summer	G	Polynomial	-0.000486	0.00073	-0.0019	0.000944	2234	32330
Moisture (ESA)	Mass loss	Summer	R	Polynomial	0.000236	0.000743	-0.00121	0.00171	2234	30986

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806 **Table S4.** Model outputs for environmental variable – decomposition relationships within grid
807 cells. Bold rows designate relationships (slope parameter) for which the credible interval does
808 not cross zero (i.e., the relationship is “significant”). Sample size indicates number of tea
809 samples available to test relationships. Effective sample size indicates number of convergent
810 model runs. Variables are standardised within grid cells using mean zero and unit-variance
811 scaling. All models are for summer incubations only. G and R indicate green and rooibos tea,
812 respectively.

Environmental variable	Decomp variable	Tea Type	Parameter	Mean	SD	2.5%	97.5%	Sample size	Effective sample size
Air temp.	Mass loss	G	Intercept	0.815	0.112	0.595	1.033	1504	4865
Air temp.	Mass loss	R	Intercept	-0.996	0.111	-1.215	-0.782	1504	5131
Air temp.	Mass loss	G	Slope	-0.029	0.036	-0.100	0.041	1504	1964
Air temp.	Mass loss	R	Slope	0.033	0.026	-0.034	0.066	1504	4794
Soil temp.	Mass loss	G	Intercept	0.856	0.122	0.616	1.088	1311	230
Soil temp.	Mass loss	R	Intercept	-0.822	0.120	-1.061	-0.588	1311	137
Soil temp.	Mass loss	G	Slope	0.144	0.027	0.091	0.197	1446	261
Soil temp.	Mass loss	R	Slope	0.073	0.026	0.021	0.122	1446	420
Moisture	Mass loss	G	Intercept	0.816	0.164	0.492	1.132	802	597
Moisture	Mass loss	R	Intercept	-0.877	0.161	-1.191	-0.565	802	817
Moisture	Mass loss	G	Slope	0.049	0.052	-0.054	0.153	802	368
Moisture	Mass loss	R	Slope	0.059	0.046	-0.034	0.151	802	657

813

814 **Table S5.** Model outputs for temperature – decomposition relationships, including an
815 interaction with soil moisture. Bold rows designate relationships (slope parameter) for which
816 the credible interval does not cross zero (i.e., the relationship is “significant”). Sample size
817 indicates number of tea samples available to test relationships. Effective sample size indicates
818 number of convergent model runs. Environmental variables are unscaled and in original units.
819 All models are for summer incubations only. G and R indicate green and rooibos tea,
820 respectively.

Environmental variable	Decomp. variable	Tea Type	Parameter	Mean	SD	2.5%	97.5%	Sample size	Effective sample size
Measured soil temp. × moisture	Mass loss	G	Intercept	0.522	0.018	0.487	0.558	624	5009
Measured soil temp. × moisture	Mass loss	G	Temp. slope	0.029	0.008	0.014	0.044	624	4470
Measured soil temp. × moisture	Mass loss	G	Moisture slope	0.002	0.001	0.001	0.004	624	6430
Measured soil temp. × moisture	Mass loss	G	Interaction	-1.38e-04	3.32e-04	-5.35e-04	7.86e-04	624	6277
Measured soil temp. × moisture	Mass loss	R	Intercept	0.176	0.016	0.144	0.209	624	6799
Measured soil temp. × moisture	Mass loss	R	Temp. slope	0.008	0.007	-0.004	0.022	624	6349
Measured soil temp. × moisture	Mass loss	R	Moisture slope	0.001	0.001	-0.001	0.002	624	8261
Measured soil temp. × moisture	Mass loss	R	Interaction	4.97e-05	2.81e-04	-5.07e-04	6.05e-04	624	7942
Gridded temp. × moisture	Mass loss	G	Intercept	0.581	0.009	0.563	0.599	2,234	15 000
Gridded temp. × moisture	Mass loss	G	Temp. slope	0.019	0.002	0.015	0.023	2,234	15 000
Gridded temp. × moisture	Mass loss	G	Moisture slope	0.009	0.003	0.004	0.014	2,234	15 000
Gridded temp. × moisture	Mass loss	G	Interaction	0.001	0.001	-6.39e-04	0.002	2,234	15 000
Gridded temp. × moisture	Mass loss	R	Intercept	0.218	0.009	0.201	0.234	2,234	15 000
Gridded temp. × moisture	Mass loss	R	Temp. slope	0.012	0.002	0.008	0.015	2,234	15 000
Gridded temp. × moisture	Mass loss	R	Moisture slope	0.004	0.002	-0.001	0.009	2,234	15 000
Gridded temp. × moisture	Mass loss	R	Interaction	0.001	0.001	2.67e-05	0.003	2,234	15 000

821

822 **Table S6.** Model outputs for relationships between measured environmental variables and
823 gridded environmental variables. Bold rows designate relationships (slope parameter) for
824 which the credible interval does not cross zero (i.e., the relationship is “significant”). Sample
825 size indicates number of sites available to test relationships. Effective sample size indicates
826 number of convergent model runs.

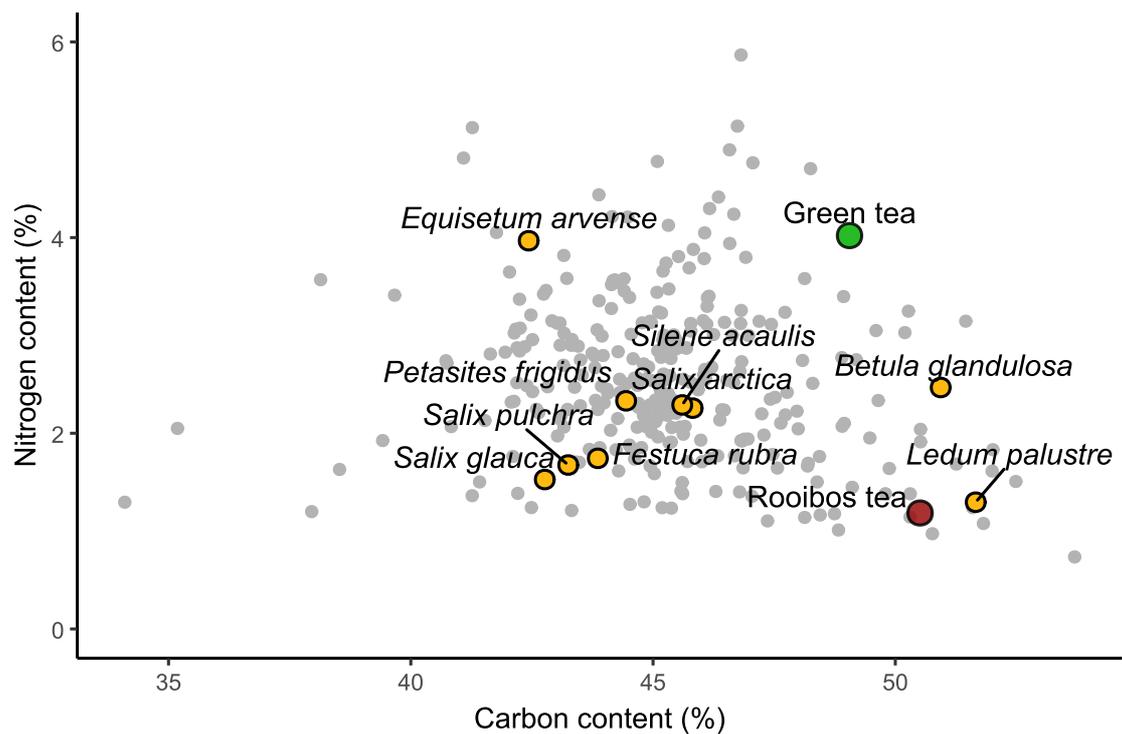
Measured variable	Gridded variable	Parameter	Mean	2.5%	97.5%	Sample size	Effective sample size
Air temperature	CHELSA air temperature	Intercept	-0.225	-1.596	1.194	151	15 000
Air temperature	CHELSA air temperature	Slope	0.877	0.474	1.013	151	15 000
Soil temperature	CHELSA air temperature	Intercept	-2.259	-3.507	-1.013	134	15 000
Soil temperature	CHELSA air temperature	Slope	1.24	1.130	1.352	134	15 000
Soil moisture	CHELSA precipitation	Intercept	16.876	14.625	19.197	79	15 000
Soil moisture	CHELSA precipitation	Slope	-0.120	-0.170	-0.074	79	11 445
Soil moisture	ESA soil moisture	Intercept	24.405	22.237	26.612	39	15 000
Soil moisture	ESA soil moisture	Slope	0.061	-0.004	0.128	39	15 000

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828 **Supplementary Figures**

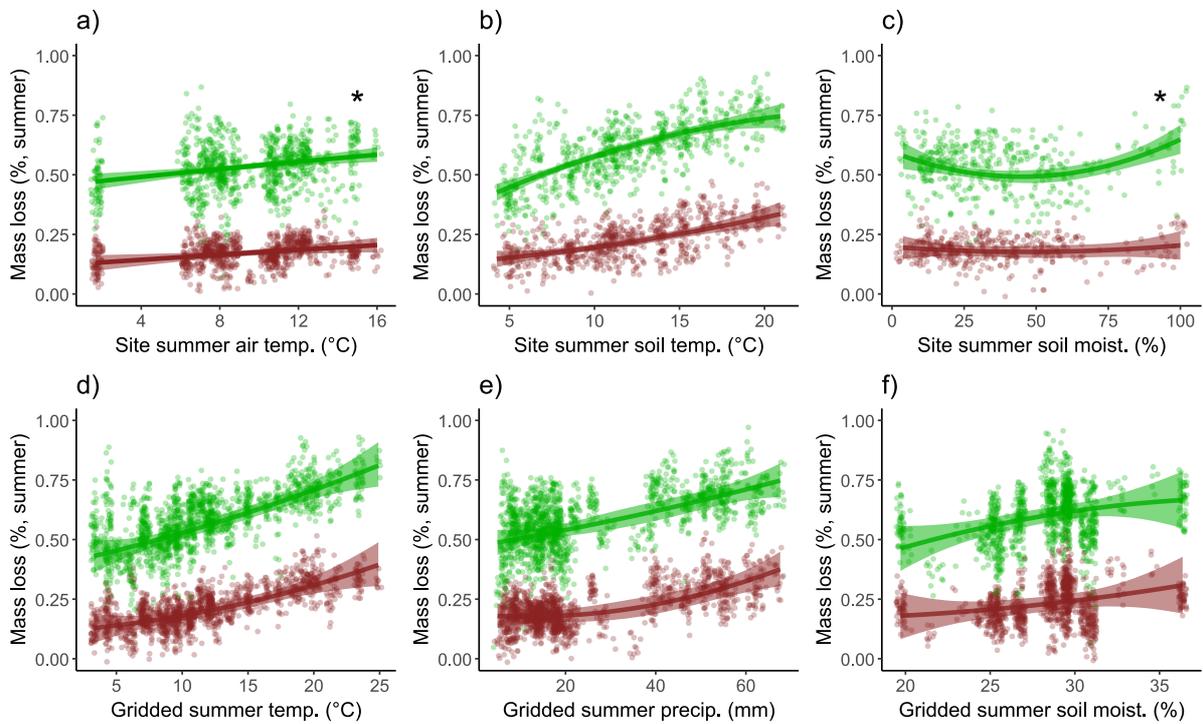
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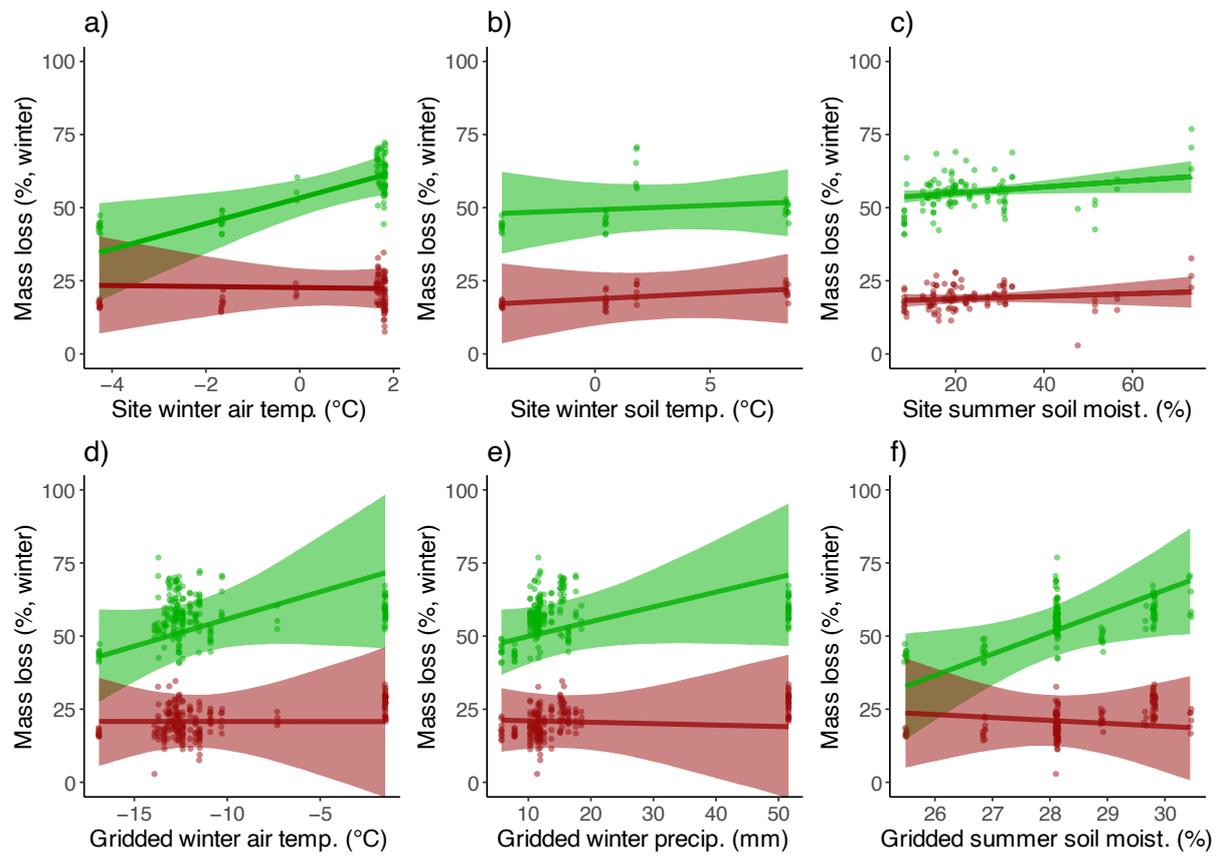
832 **Figure S1.** Carbon and nitrogen content of green and rooibos tea compared a range of
833 representative tundra species. Tea types are indicated by red (rooibos tea) and green (green
834 tea) plots. Yellow dots and corresponding names indicate tundra species for which we
835 collected and run the incubation experiment presented in Fig. 2. Grey dots represent a tundra
836 species with available data from the Tundra Trait Team Database^{94,95}.



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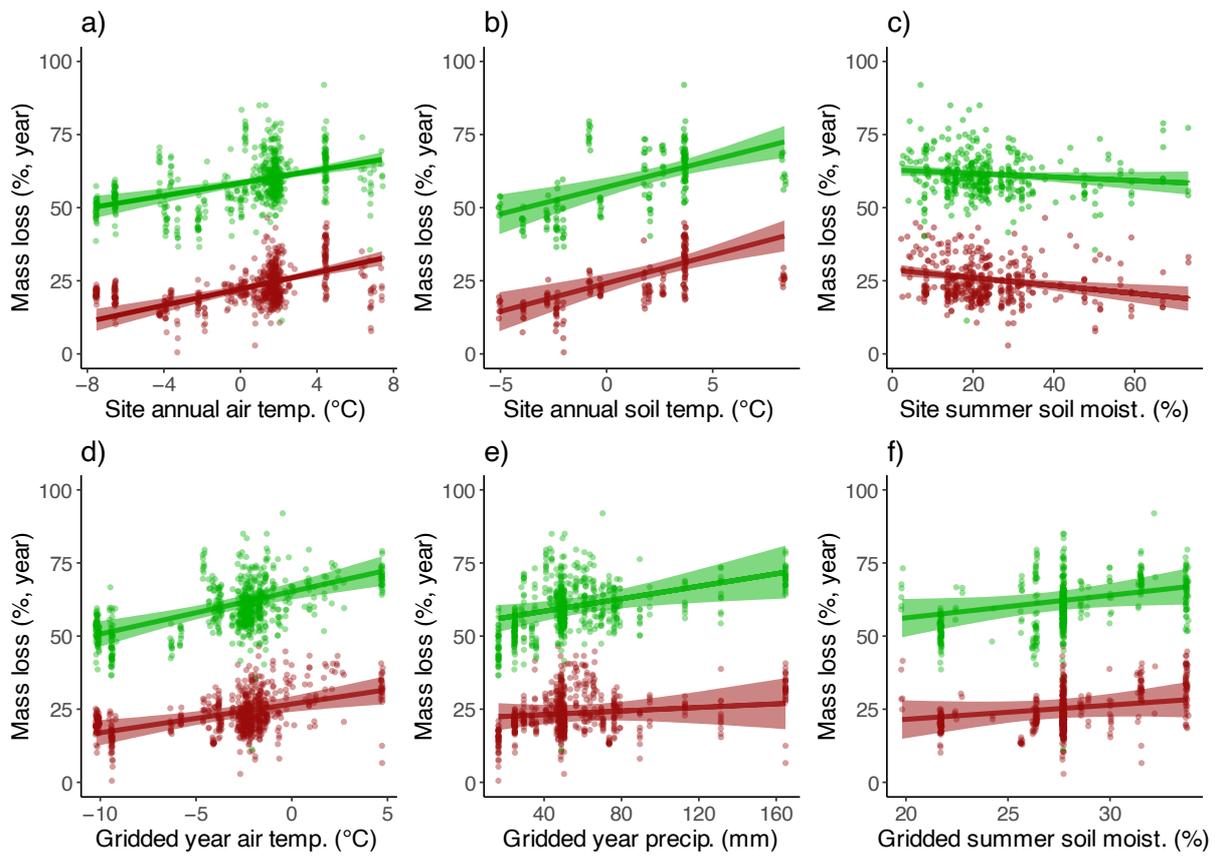
839 **Figure S2.** Polynomial relationships between decomposition (mass loss), measured
 840 environmental variables (a-c) and gridded climate data (d-f) for summer tea
 841 incubations. Points indicate individual tea bag replicates across all sites. Lines
 842 indicate hierarchical Bayesian polynomial model fits with 95% credible intervals.
 843 Colours indicate tea type (red = rooibos tea, green = green tea). See Table S3 for
 844 model outputs. Polynomial parameters with 95% credible interval not including 0 are
 845 highlighted with (*).



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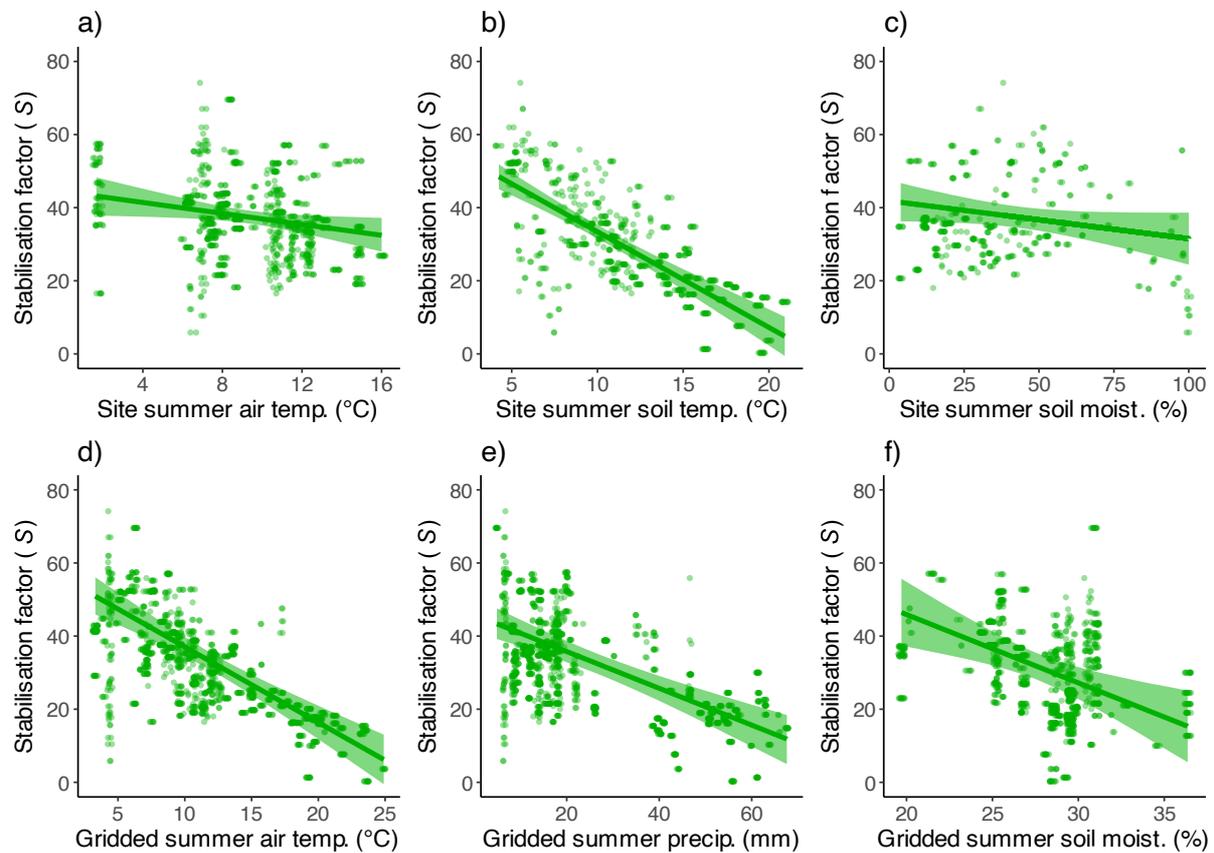
848 **Figure S3.** Relationships between decomposition (mass loss), measured environmental
 849 variables (a-c) and gridded climate data (d-f) for winter tea incubations, as opposed to summer
 850 incubations in main text (Fig. 3) or year-long incubations (Fig. S4). Points indicate individual
 851 tea bag replicates across all sites. Lines indicate hierarchical Bayesian model fits with 95%
 852 credible intervals. Colours indicate tea type (red = rooibos tea, green = green tea). See Table
 853 S2 for model outputs.



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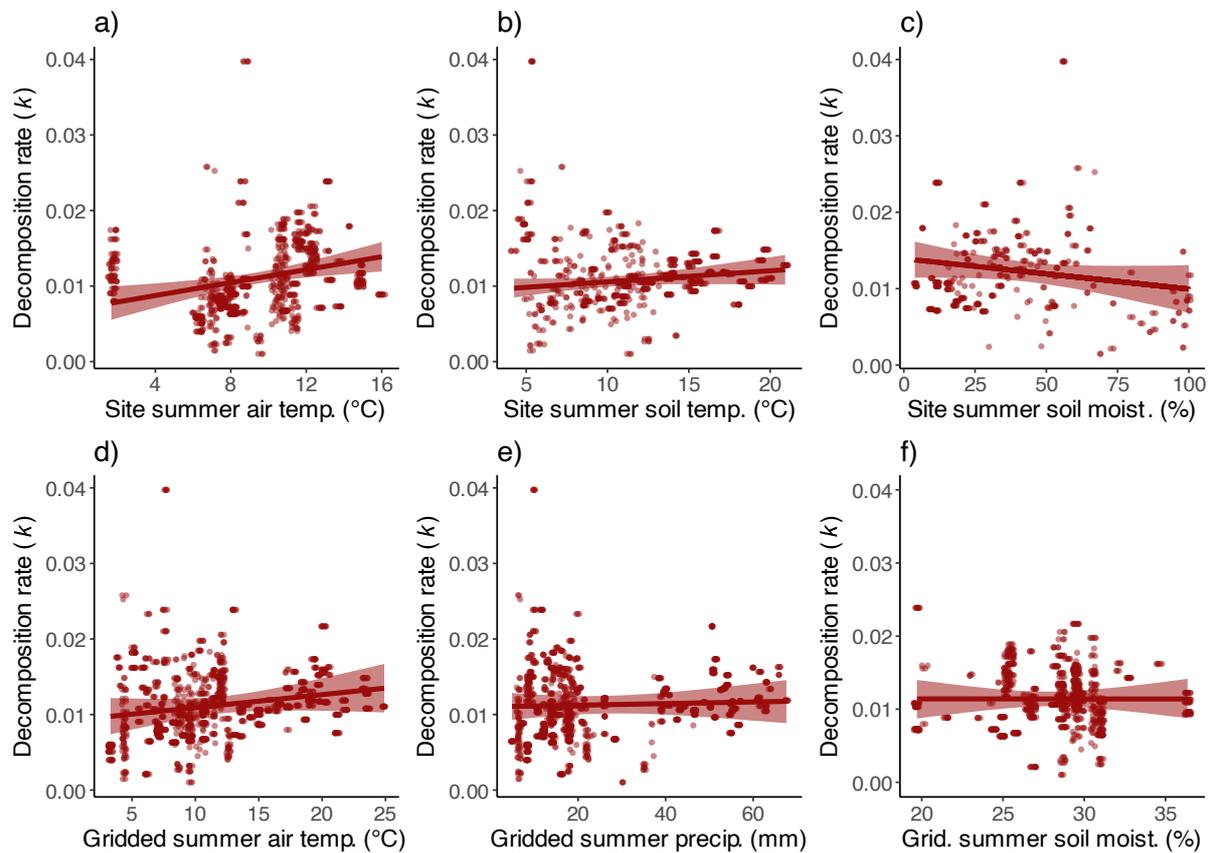
856 **Figure S4.** Relationships between decomposition (mass loss), measured environmental
 857 variables (a-c) and gridded climate data (d-f) for year-long tea incubations, as opposed to
 858 summer incubations in main text (Fig. 3) or winter incubations (Fig. S3). Points indicate
 859 individual tea bag replicates across all sites. Lines indicate hierarchical Bayesian model fits
 860 with 95% credible intervals. Colours indicate tea type (red = rooibos tea, green = green tea).
 861 See Table S2 for model outputs.



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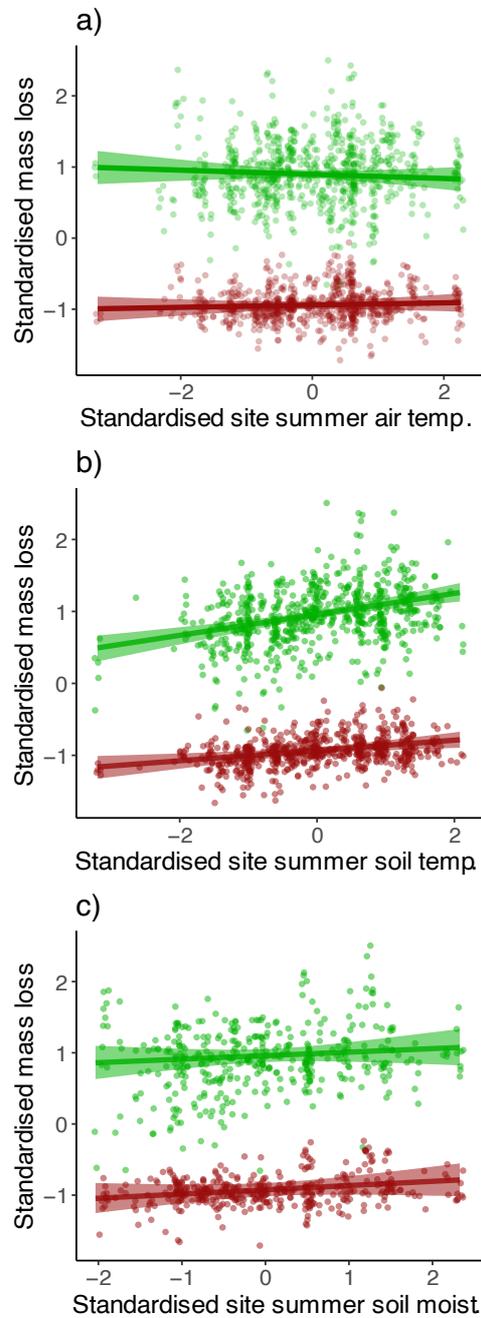
864 **Figure S5.** Relationships between stabilisation factor (S), measured environmental variables
 865 (a-c) and gridded climate data (d-f) for summer tea incubations, as opposed to summer mass
 866 loss in main text (Fig. 3). S is calculated based on decomposition of green tea, and is assumed
 867 to be consistent across tea types¹⁹. S represents the proportion of labile material remaining
 868 once decomposition has stabilised, and thus long-term carbon storage. Points indicate
 869 individual tea bag replicates across all sites. Lines indicate hierarchical Bayesian model fits
 870 with 95% credible intervals. See Table S2 for model outputs.



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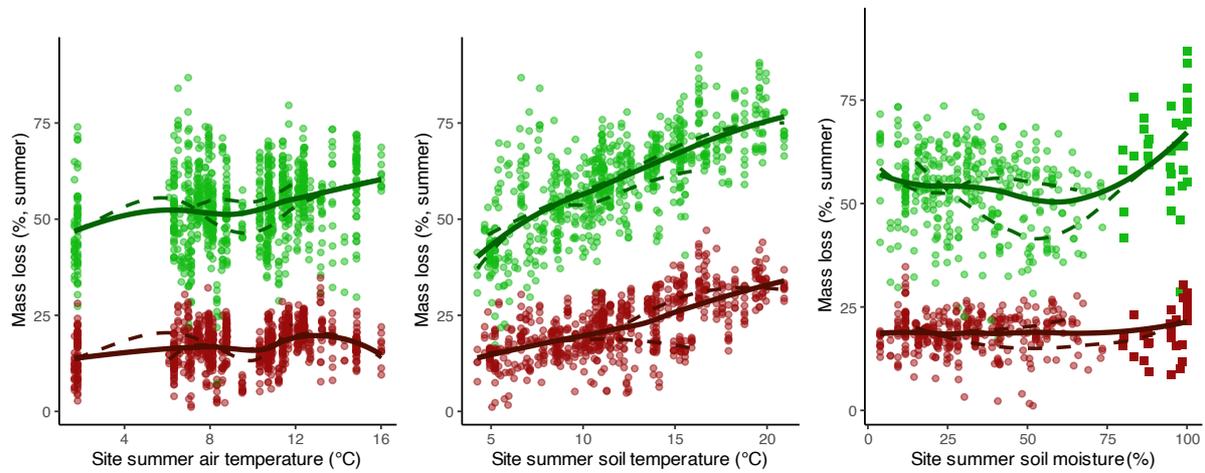
873 **Figure S6.** Relationships between decomposition rate (k), measured environmental variables
 874 (a-c) and gridded climate data (d-f) for summer tea incubations, as opposed to summer mass
 875 loss in main text (Fig. 3). k is calculated based on decomposition of rooibos tea, and is
 876 assumed to be consistent across tea types¹⁹. k represents the rate of loss of labile material,
 877 and thus short-term decomposition dynamics and biogeochemical cycling. Points indicate
 878 individual tea bag replicates across all sites. Lines indicate hierarchical Bayesian model fits
 879 with 95% credible intervals. See Table S2 for model outputs.



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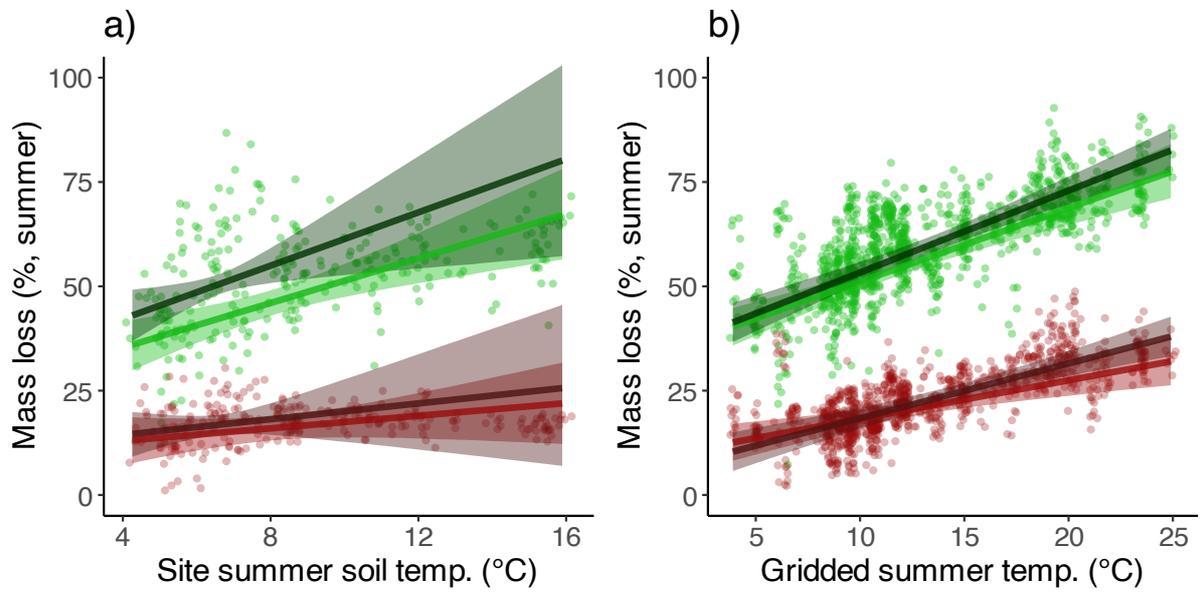
882 **Figure S7.** Within-grid cell relationships reflect among-site relationships between
 883 environmental variables and mass loss, but with greater variability. Within-grid cell
 884 relationships between summer decomposition (mass loss) and measured environmental
 885 variables, as opposed to among sites in main text (Fig. 3). Environmental and decomposition
 886 variables are standardised within 0.25×0.25 -degree resolution grid cells using mean zero
 887 and unit-variance scaling. Points indicate individual tea bag replicates. Lines indicate
 888 hierarchical Bayesian model fits with 95% credible intervals. Colours indicate tea type (red =
 889 rooibos tea, green = green tea). See Table S3 for model outputs.



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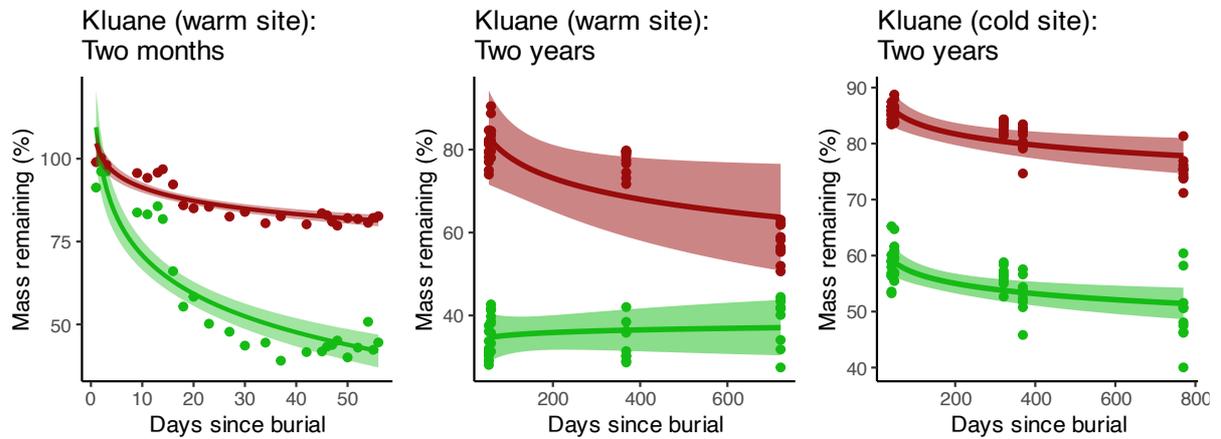
892 **Figure S8.** Overall, relationships between climate variables and mass loss were best
 893 described by linear rather than exponential models. In order to test the linearity of the
 894 relationships between climate variables and mass loss of tea types, we fit general additive
 895 models with a loess fit to the overall dataset (solid lines) and to the Western and Eastern
 896 hemispheres as two subsets of the data (dashed lines). The relationship between site summer
 897 soil moisture and green tea mass loss was more exponential, but this was driven by data from
 898 Svalbard located at particularly wet sites (square points) and thus we do not have confidence
 899 that the exponential relationships can be generalised to the tundra biome, which were better
 900 fit by hierarchical linear models (Fig. 3).



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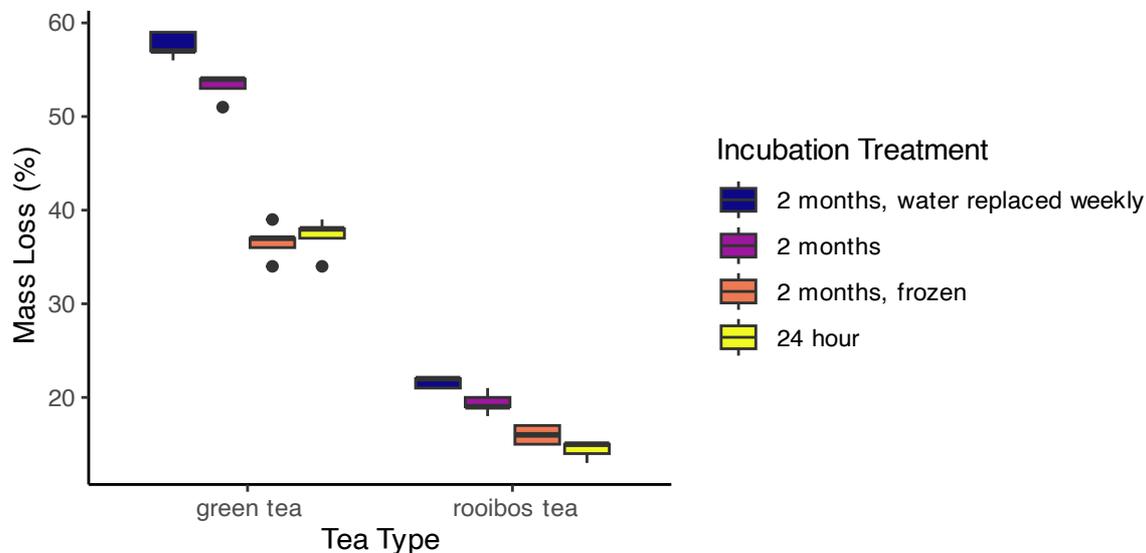
903 **Figure S9.** Soil moisture did not influence the relationships between soil temperature and
 904 mass loss, but decomposition was higher at wetter versus drier sites at any given temperature.
 905 Relationships between summer decomposition (mass loss), (a) measured soil temperature
 906 and soil moisture, and (b) gridded temperature (CHELSA) and soil moisture (ESA). Models
 907 incorporate the interaction between soil temperature and soil moisture. Lines indicate
 908 predicted decomposition at upper (dark) and lower (light) quartiles of soil moisture,
 909 representing wet and dry sites respectively, based on hierarchical Bayesian model fits with
 910 95% credible intervals. Points indicate individual tea bag replicates. Colours indicate tea type
 911 (red = rooibos tea, green = green tea). See Table S4 for model outputs.



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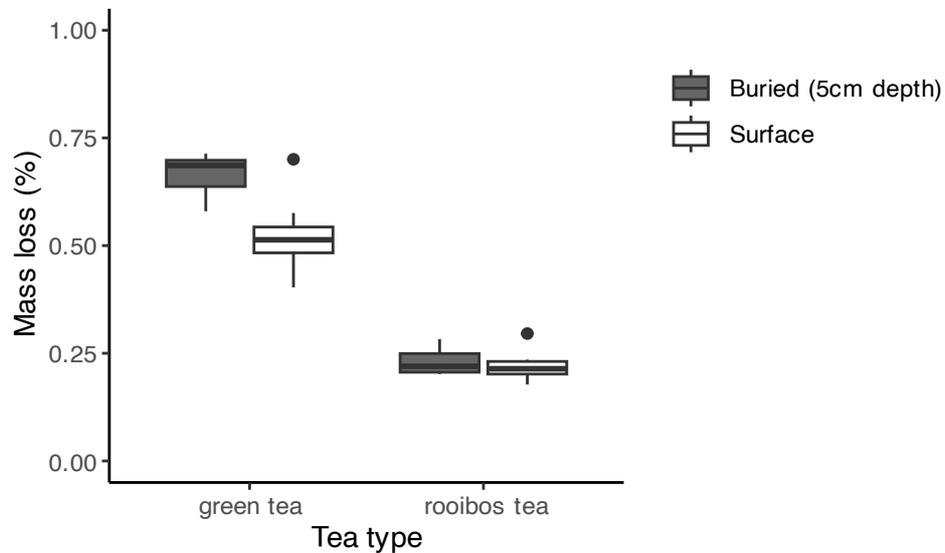
914 **Figure S10.** Mass loss of tea types did not converge after two years and stabilised after
 915 approximately 30 days. Mass remaining over time of roibos and green tea at warm and cold
 916 tundra sites at the Kluane Lake location (see Table S1). Mass loss is calculated using a single-
 917 phase exponential decay decomposition model. (a) Mass remaining at the warm experimental
 918 site, with tea extracted every two days over a two-month summer period; (b) mass remaining
 919 at the warm experimental site with summer, one-year and two-year incubation lengths; and
 920 (c) mass remaining at the cold experimental site with summer, two-month, one-year and two-
 921 year incubation lengths.



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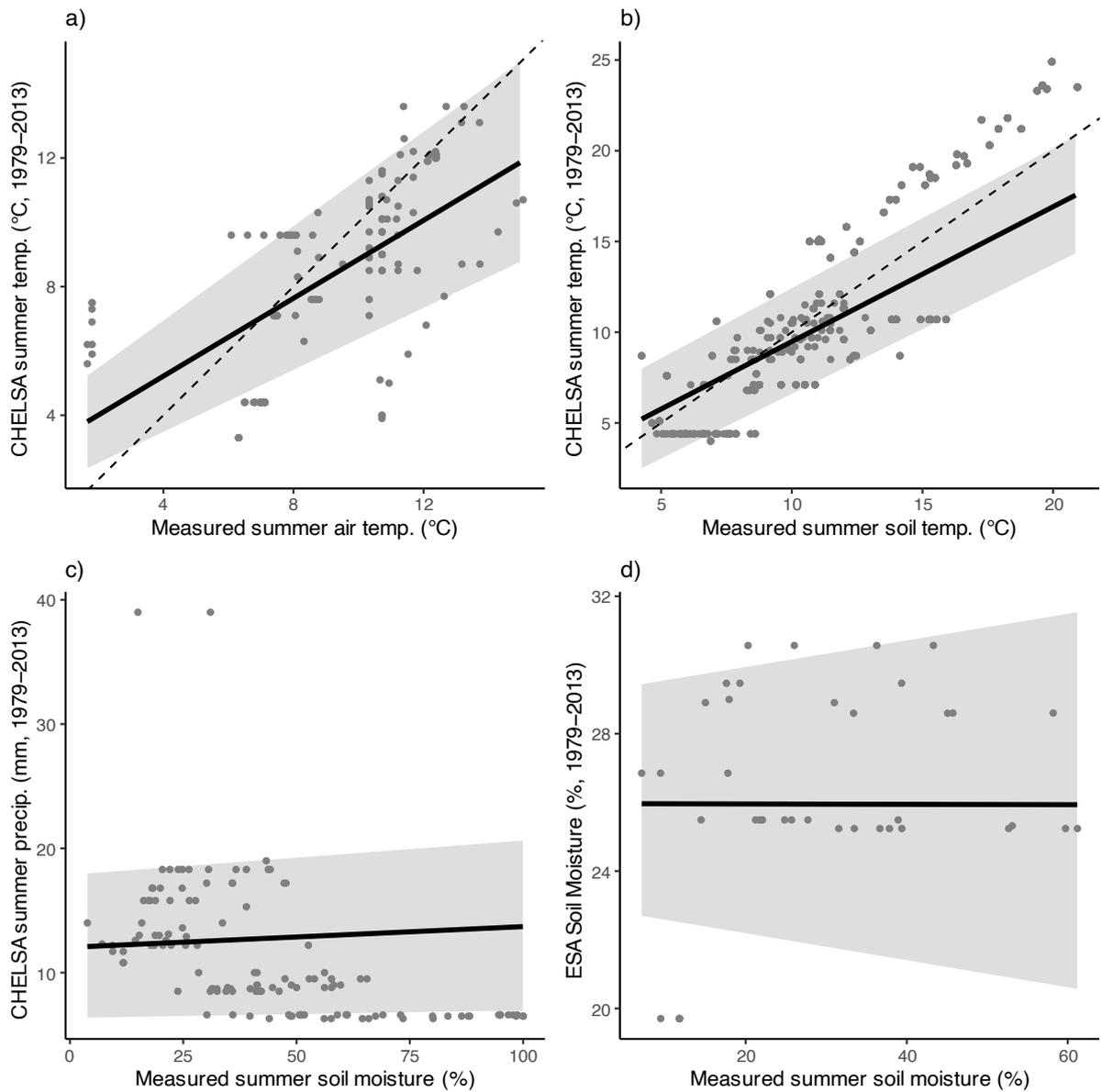
924 **Figure S11.** In order to test the influence of leaching, we conducted 2-month and 24-hour
 925 incubations of green and rooibos tea in a laboratory environment at room temperature, in a
 926 4°C fridge and a 20°C freezer. We found ~20% greater mass loss for green tea and ~7%
 927 greater mass loss for rooibos tea in two-month incubations rather than in 24-hour incubations
 928 in liquid water. Leaching was not strongly influenced by replacement of water and was slower
 929 in frozen conditions for green and rooibos tea.



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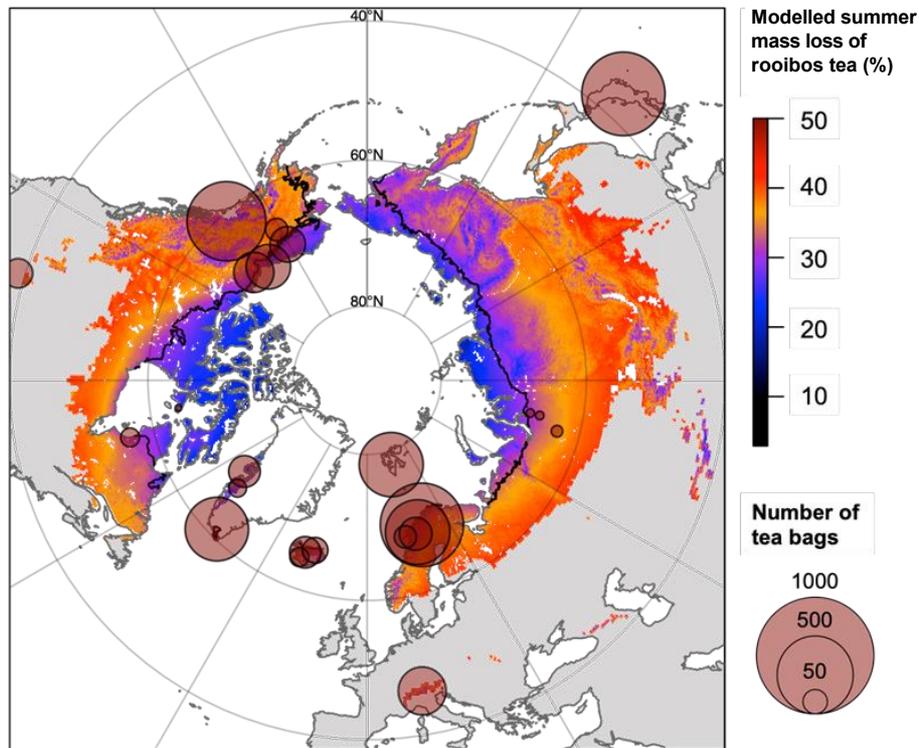
932 **Figure S12.** Mass loss of buried tea bags was significantly greater than tea placed on the
 933 ground surface for green tea, but not different for rooibos tea. Teabags were incubated in a
 934 common site (Kluane Lake, see Table S1) and were either buried at 5 cm depth directly in the
 935 soil (grey) or placed within a litter bed and covered in a local litter medium (white), following
 936 protocols outlined in Cornelissen et al. (2007). Teabags were incubated for one year, though
 937 the time periods of incubation differed between the two treatment types (buried: June – June,
 938 surface: August – August) as tea bags are taken from two different, but adjacent, experiments.
 939 Stars indicate significance (***, $P < 0.001$, ns, $P > 0.05$, t-test).



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941

942 **Figure S13.** Site-measured environmental variables aligned with gridded climate variables for
 943 summer temperature, but not summer soil moisture. Relationships between site-measured
 944 environmental variables and gridded climate data for all tea bag sites with available data. Lines
 945 indicate hierarchical Bayesian model fits and errors are 95% credible intervals. See Table S5
 946 for model outputs.



947

948 **Figure S14.** Modelled summer decomposition (percent mass loss) of rooibos tea for tundra
 949 and sub-Arctic regions based on 1979 to 2013 mean summer air temperature (Climatologies
 950 at high resolution for the Earth's land surface, CHELSA) and soil moisture (European Space
 951 Agency data, ESA) from 1979 to 2013. Field collection locations are illustrated by red circles,
 952 grouped by geographic region (Table S1, figure excludes Australian alpine region). Circle size
 953 indicates the number of tea bag replicates within each geographic region. Tundra and sub-
 954 Arctic classifications are based on Köppen-Geiger classification⁴⁷. Ice-covered areas are
 955 excluded. The circum-Arctic treeline is indicated with a black line⁴⁸.