

# 1 **Technical Note: Semi-rigid chambers for methane gas flux**

## 2 **measurements on tree stems**

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9

### 10 **Abstract**

11 There is increasing interest in the measurement of methane (CH<sub>4</sub>) emissions from tree stems  
12 in a wide range of ecosystems so as to determine how they contribute to the total ecosystem  
13 flux. To date, tree CH<sub>4</sub> fluxes are commonly measured using rigid closed chambers (static or  
14 dynamic), which often pose challenges as these are bulky and limit measurement of CH<sub>4</sub>  
15 fluxes to only a very narrow range of tree stem sizes and shapes. To overcome these  
16 challenges we aimed to design and test new semi-rigid stem-flux chambers (or sleeves). We  
17 compared the CH<sub>4</sub> permeability of the new semi-rigid chambers with that of the traditional  
18 rigid chamber approach, in the laboratory and in the field, with continuous flow or syringe  
19 injections. We found that the semi-rigid chambers had reduced gas permeability and optimal  
20 stem gas exchange surface to total chamber volume ratio ( $S_c/V_{tot}$ ) allowing better headspace  
21 mixing, especially when connected in a dynamic mode to a continuous flow gas analyser.  
22 Semi-rigid sleeves can easily be constructed and transported in multiple sizes, are extremely

23 light, cheap to build and fast to deploy. This makes them ideal for use in remote ecosystems  
24 where access logistics is complicated.

25

## 26 **1 Introduction**

27 Recent research into ecosystem greenhouse gas fluxes has shown that tree stems emit  
28 significant amounts of methane (CH<sub>4</sub>) (Terazawa et al., 2007; Rusch and Rennenberg, 1998;  
29 Gauci et al., 2010; Pangala et al., 2013; Rice et al., 2010; Terazawa et al., 2015) although the  
30 transport mechanisms and global importance of tree-mediated emissions remain largely  
31 unknown. These past investigations have used a variety of closed chambers adapted to  
32 various tree-stem sizes. Presently, the most common chambers used to measure CH<sub>4</sub>  
33 emissions from tree-stems are closed rigid chambers in the form of either a vertical cylinder, a  
34 horizontal cylinder or a cube fitted around tree-stems (e.g. Gauci et al., 2010; Pangala et al.,  
35 2013; Terazawa et al., 2007; Hari et al., 1991; Rusch and Rennenberg, 1998). These chambers  
36 can be deployed either vertically by enclosing the whole stem or, alternatively when the stems  
37 are too large, laterally on the stem, covering only a small fraction of the stem surface (e.g.  
38 Levy et al., 1999; Teskey and McGuire, 2005; Ryan, 1990; Hari et al., 1991). These  
39 techniques were originally designed to measure CH<sub>4</sub> and carbon dioxide (CO<sub>2</sub>) from samples  
40 manually taken with syringes and analysed by gas chromatography. The ratio between the gas  
41 exchange surface and the chamber volume ( $S_c/V_{tot}$ ) was transposed from soil chambers and  
42 were not necessarily adapted to the lower fluxes found in tree-stems, and are therefore often  
43 too high (Hutchinson and Livingston, 2001). In other words, if the chambers are too large for  
44 a given exchange surface, mixing problems may occur, making it important to circulate the air  
45 in their headspace (Hutchinson and Livingston, 1993; Rusch and Rennenberg, 1998).

46 With the advent of continuous flow analytical techniques and increasing precision of  
47 instruments (e.g. cavity ring-down spectroscopy, infrared and photo-acoustic gas analysers),  
48 the need for longer accumulation periods to detect significant concentration changes has  
49 become obsolete. The tendency is to reduce the accumulation period as much as possible in  
50 order to be able to use more straightforward linear regressions to determine fluxes closest to  
51 the point of chamber closure. Unlike open chamber techniques which allow steady state  
52 measurements (e.g. Bortoluzzi et al., 2006; Norman et al., 1997; Subke et al., 2003;  
53 Pumpanen et al., 2004), closed chambers are non-steady state systems; the diffusive laws  
54 advocate the use of non-linear regressions of gas concentrations as a function of time to  
55 determine rates, as these decrease with increasing gas saturation (Hutchinson and Livingston,  
56 2001; Pihlatie et al., 2013; Pumpanen et al., 2004; Kutzbach et al., 2007).

57 With continuous flow gas analysers there are three main advantages: 1) they are non-  
58 dispersive as no gas needs to be taken out of the measurement system and irreversibly  
59 “consumed”, 2) they circulate air between the chamber and the gas detectors, which for small  
60 chamber volumes could represent enough mixing to avoid underestimations of fluxes by as  
61 much as 36 to 58% in non-mixed soil-atmosphere exchanges (Christiansen et al., 2011), and  
62 3) with measurement frequencies of up to 10 Hertz and precisions of  $\pm 2$  ppb the closure time  
63 needed to get a representative accumulation slope has been dramatically reduced using these  
64 devices (excluding the equilibration period) and therefore it also avoids underestimations due  
65 to regressions made over longer periods of time (Hutchinson and Livingston, 2001; Pihlatie et  
66 al., 2013). In addition, recent work has focused on trace gases (e.g. CH<sub>4</sub> and N<sub>2</sub>O) which have  
67 lower accumulation rates compared to the more frequently measured CO<sub>2</sub> (IPCC, 2007),  
68 moderating the saturation issue inherent to non-steady state setups (Hutchinson and  
69 Livingston, 2001). Altogether, these point towards the use of a smaller stem chamber with  
70 larger gas exchange surface per chamber volume proportion (Sc-to-Vtot ratio).

71 A further complicating factor is field access. Stem-methane emissions have recently begun to  
72 be investigated in remote areas such as in forested tropical wetlands with often no road access.  
73 In those areas it is a logistical challenge to carry large/heavy loads. Moreover, because of the  
74 great variety of stem sizes/shapes, a whole collection of rigid chambers is usually needed to  
75 cover most of the ecosystem tree species thus creating further logistical and cost issues.

76 In order to meet the new challenges presented by the growing interest in measuring  
77 greenhouse gas fluxes from tree-stems we aimed to design, describe and test/deploy new  
78 semi-rigid stem-emission chambers in the laboratory and in the field, and to compare their  
79 permeability to CH<sub>4</sub> (gas conductance) with previously described rigid chambers. Thus far,  
80 semi-rigid sleeve chambers have been used effectively in several of our measurement  
81 campaigns. We therefore consider their detailed reporting to be of interest to a broader  
82 constituency of eco-physiologists and biogeochemists. We also examine various  
83 methodological benefits and logistical advantages of using this new approach.

84

## 85 **2 Materials and methods**

### 86 **2.1 Chamber designs: semi-rigid sleeve and rigid chamber**

87 Our approach to measure stem CH<sub>4</sub> emissions, which could also include other greenhouse  
88 gases produced in anaerobic conditions such as N<sub>2</sub>O, uses a semi-rigid chamber (or sleeve).  
89 The preferred material was a pre-shaped and gas impermeable PET (polyethylene  
90 terephthalate) or PC (polycarbonate) plastic sheet with a natural tendency to curve induced by  
91 3-4 vertically distributed imprinted rims on the periphery. These rims ensured good stability  
92 and helped maintain the desired natural curvature of the sleeve that proved to be very helpful  
93 for the deployment of the sleeves on the stems as the sleeve could hold in place without straps.

94 To investigate permeability changes due to both the size and the approach, we used two semi-  
95 rigid sleeves together with a rigid chamber. As this was straightforward, for the smaller semi-  
96 rigid sleeve we sourced the pre-shaped material from a cylindrical 3 L soft drink bottle, which  
97 already had the desired imprinted rims. The 0.1 mm thick bottle was truncated above and  
98 below the cylindrical section, and opened vertically on the side. For the larger sleeve we  
99 sourced the material from 0.2 mm thick not pre-shaped semi-rigid PC sheets. Both types of  
100 plastic sheets have very low gas permeabilities under experimental standard ambient  
101 temperature and pressure (SATP from UIPAC) conditions and short chamber enclosure times  
102 (McKeen, 2012).

103 The edges of the sheets were framed with 1.5 cm thick and 3 cm wide adhesive backed  
104 expanded Neoprene strips (Seals+Direct Ltd, Hampshire, UK); closed cell neoprene foam  
105 that is gas tight and can be bent, but is hardly compressible ( $\leq 3\%$  with 200 N). This  
106 Neoprene strip was placed as a frame around the rectangular sheet to provide a seal and to  
107 ensure a constant volume between the sheet and the tree stem (Fig. 1). The adhesive was  
108 provided on one side of the expanded Neoprene strips. Inside this framed volume we placed  
109 two Neoprene vertical wedges (1.5 cm thick and 3 cm wide) to keep the sheet equidistant  
110 from the stem all along the radial periphery of the sleeve. The sleeve was also equipped with  
111 two snap-on rubber caps with inserted three-way Luer-lock stopcocks (BBraun, Bethlehem,  
112 USA) that permitted connection to the Ultraportable Greenhouse Gas Analyser (UGGA, Los  
113 Gatos Research Inc., Mountain View, USA) via two 4.6 m long and 5 mm inside diameter  
114 PTFE (polytetrafluoroethylene) coated PVC (polyvinyl chloride) parallel tubes (Nalgene,  
115 Rochester, USA). As venting was recommended (Hutchinson and Livingston, 2001;  
116 Christiansen et al., 2011) both sleeves were equipped with a coiled vent tube (18 cm long, 1.2  
117 mm inner diameter). We downscaled the vent described by Hutchinson and Livingston (2001)  
118 by a factor 48 in terms of vent volume whereas the sleeves were a factor 10 to 20 less

119 voluminous as compared to the authors' chamber (14 L). Their study showed that in a  
120 perfectly sealed chamber, after 30 minutes of deployment the gas mass loss through the sole  
121 vent represented 0.038% of the target gas.

122 We tested all the components of the semi-rigid sleeves independently for unwanted  
123 background contaminations that could interfere with CH<sub>4</sub> emissions from the stems by  
124 incubating them for two hours in 500 mL borosilicate glass beakers filled with air and  
125 connected in continuous flow with the UGGA. The selected raw material was inert and did  
126 not interfere with measurements from the environment. We also tested the compressibility of  
127 sleeves by pulling the straps with a 200 N force (twice 100N) and measuring the thickness of  
128 the Neoprene frame before and after pulling (Fig. 2, see also chamber deployment section).

129 We also compared the CH<sub>4</sub> losses from our new semi-rigid sleeves with a previously used  
130 rigid chamber design, similar to the ones constructed and described in other studies (Rusch  
131 and Rennenberg, 1998; Gauci et al., 2010; Pangala et al., 2013). The closed rigid chamber  
132 was constructed from cylindrical Perspex® (Perspex, Tamworth, UK) of inner diameter of 28  
133 cm and had an inner height of 30 cm. The cylinder was cut into two halves, which were held  
134 together with a metal hinge. The two half-cylinders were framed within a 5 cm wide and 1 cm  
135 thick frame made of flat Perspex® that was fitted with Neoprene strips. The cylindrical  
136 chamber had a central opening to enclose the tree stem. Two smaller cylinders (18 cm  
137 diameter x 5 cm height) were attached on either side of that opening (Fig. 3). The chamber  
138 was equipped with a gas sampling port and a small vent tube (12 cm long; 6 mm diameter).

139

## 140 **2.2 Enclosed chamber volume and gas exchange surface determinations**

141 The volume of the semi-rigid sleeves could be determined precisely in two different ways.  
142 Firstly, we extrapolated the empirical total chamber volume ( $V'_{tot}$ ) from the CH<sub>4</sub>

143 concentration dilution factor after having inserted a known volume ( $V_{standard}$ ) of a 2000 ppmv  
 144  $CH_4$  standard (Air Liquide, Paris, France) into the sleeve's enclosed volume and measuring  
 145 the end concentration ( $C_0$ ) after dilution, and subtracting the atmospheric  $CH_4$  concentration  
 146 ( $C_{atm}$ ) originally in the chamber. The two semi-rigid sleeves and a rigid chamber were  
 147 attached to an inert stainless steel cylinder (see chamber deployment). The dilution was done  
 148 in 90 seconds so that the losses through gas permeability of the chambers remained negligible.  
 149 This extrapolation was formalised as:

$$150 \quad V'_{tot} = V_{standard} * \frac{(C_{standard})}{(C_0 - C_{atm})} \quad (1)$$

151 Secondly, we also calculated the theoretical volume of the sleeves ( $V_c$ ) by subtracting a sector  
 152 (K) of both, a smaller cylinder volume ( $V_{stem}$ ) from a larger cylinder volume ( $V_{ext}$ ), minus the  
 153 volume taken by the vertical wedges ( $V_{wedges}$ ) (Fig. 4). The sector (K) was determined from a  
 154 ratio between the sleeve length (L) and the circumference at the external edge of the sleeve  
 155 ( $\pi D_{ext}$ ). The sleeve length (L) is the length of the incompressible external edge of the chamber  
 156 and represents a fraction of the total circumference given by  $\pi D_{ext}$ . The diameter of the  
 157 smaller cylinder (the compressible internal foamy edge) is given by the diameter of the stem  
 158 ( $D_{stem}$ ). The larger cylinder diameter ( $D_{ext}$ ) is the diameter given by the stem ( $D_{stem}$ ) plus the  
 159 thickness (T) of the sleeve. Both cylinders have the same height (H). Thereafter, we have:

$$160 \quad D_{ext} = D_{stem} + 2T \quad (2)$$

$$161 \quad K = \frac{L}{\pi D_{ext}} = \frac{L}{\pi(D_{stem} + 2T)} \quad (3)$$

$$162 \quad V_c = K(V_{ext} - V_{stem}) - V_{wedges} = \frac{HL}{(D_{stem} + 2T)} * \left[ \left( \frac{D_{stem} + 2T}{2} \right)^2 - \left( \frac{D_{stem}}{2} \right)^2 \right] - V_{wedges} \quad (4)$$

163 However, the total volume ( $V_{tot}$ ) is the sum of the chamber volume ( $V_c$ ) plus the dead volume  
 164 enclosed in the gas analyser and the tubes ( $V_{dead}$ ):

165  $V_{tot} = V_C + V_{dead}$  (5)

166 Similarly, the gas exchange surface of the sleeves ( $S_c$ ) was calculated by considering the  
167 sector ( $K$ ) of the stem surface ( $S_{stem}$ ) covered by the chamber at the circumference of the stem  
168 ( $\pi D_{stem}$ ) and the height of the sleeve ( $H$ ), minus the small surface covered by the vertical  
169 wedges ( $S_{wedges}$ ):

170  $S_c = K * S_{stem} - S_{wedges} = \frac{HL}{(D_{stem}+2T)} * D_{stem} - S_{wedges}$  (6)

171

### 172 **2.3 Chamber deployment**

173 The three types of chambers (two semi-rigid sleeves and one rigid chamber) were deployed  
174 on a gas-inert stainless steel cylinder of diameter 15 cm. The semi-rigid chambers were  
175 flattened around the cylinder and subsequently attached and tightened with two metal cam  
176 straps at the top and bottom of the frame (Figure 2). The straps were 1.5 m long and 3 cm  
177 wide. An additional strap was necessary at mid-height of the bigger sleeve to ensure a good  
178 cohesion of the vertical Neoprene frames and vertical wedges with the stem (steel cylinder in  
179 this case).

180 Before installing the rigid acrylic chamber, closed cell Neoprene foam bands (7 cm wide and  
181 4 cm thick) were attached at the bottom of the inert stainless steel cylinder and also at 35 cm  
182 height using double-sided Scotch tape (3M, St-Paul, USA) to append the extremities of the  
183 band as well as packing brown tape (5 cm wide) to tighten the band firmly against the  
184 metallic cylinder. The two mobile panels of the chamber were opened and the upper and  
185 lower half-necks of one panel were lodged around the two foam bands by compressing the  
186 foam so as to ensure gas tightness. Finally, all open-end flanges surrounding the cylindrical  
187 volume were progressively closed with Handy-grips (Irwin, Vernier, Switzerland).



188 We used the larger semi-rigid chamber to exemplify the field deployment (Table 1). We  
189 deployed it on twelve tree-stems (diameter at breast height: 25-45 cm) located in the northern  
190 boreal zone (*Pinus sylvestris* and *Betula pendula*, Degerö mire, Sweden) as well as in a  
191 tropical lowland forest (*Heisteria concinna*, Barro Colorado Island, Panama). The sleeves  
192 were placed at mid-height on the stems at 35 cm of height. We shaded the sleeves with a  
193 plasticised aluminum foil to minimise any changes to chamber temperature which might  
194 otherwise affect stem-gas exchange processes. In the lab this measure was unnecessary. We  
195 tested the sleeve's CH<sub>4</sub> concentration change on both, very smooth birch stems and very  
196 rough pine-tree stems to contrast the concentration readings as much as possible. To ensure  
197 optimal gas tightness it was important to distribute the pressure of each strap all around the  
198 surface of the sleeve. We visually checked for gaps between the stem and the Neoprene strips.  
199 Monitoring the CH<sub>4</sub> concentration change in a continuous flow mode made an optimal gas  
200 tightness test. A leaking chamber (mainly pressure-driven bulk flow following Hagen-  
201 Poiseuille's law) typically displayed fluctuating concentrations with concentration build-up  
202 being recurrently drawn down. Finally, we also used the larger semi-rigid sleeve together with  
203 a manual syringe sampling. For that purpose we used a 30 mL plastic syringe fitted with a  
204 Luer-lock three-way stopcock (BBraun, Bethlehem, USA) and connected it to one of the two  
205 stopcocks on the sleeve. At t=0, t=5, t=10 and t=15 minutes we collected 12 mL of gas  
206 sample from the sleeve and transferred it into pre-evacuated glass Exetainers (Labco Ltd,  
207 Ceredigion, UK) before analysing CH<sub>4</sub> concentrations on a Fast Methane Analyser (Los  
208 Gatos Research Inc., Mountain View, USA) equipped with a sampling loop as described in  
209 Baird et al. (2010).

210

## 211 **2.4 Gas Analyses**

212 For the permeability tests, the CH<sub>4</sub> concentration change was analysed in the laboratory under  
213 SATP conditions for three types of chamber (Table 1, Supplement S1); a rigid chamber and  
214 two semi-rigid sleeves. We injected 50 mL of a 2000 ppmv methane standard (Air Liquide,  
215 Paris) into these chambers after which the CH<sub>4</sub> concentration decline was measured over 20  
216 minutes in continuous flow mode. Each chamber type was tested in triplicate. For the blanks,  
217 we injected ambient air. The slopes were measured from a linear regression of declining  
218 concentrations starting after an equilibration time of 90 seconds (dead band) and running for  
219 20 min. This dead band represents a maximum time for the continuous flow circuit to mix the  
220 entire headspace ( $V_{tot}$ ).

221 In the field, the CH<sub>4</sub> concentration changes of a larger sleeve were monitored when deployed  
222 on various tree-stem species (see chamber deployment). In order to have a set of contrasting  
223 responses we selected, a posteriori, measurement runs with both high and low rates, and also  
224 included runs where leakages of the sleeve were present (Figs. 5 and 6, Supplements S3 and  
225 S4). Methane concentration accumulations were measured as in the laboratory although with  
226 shorter runs of approximately 420 seconds. In the manual sampling mode with syringe, the  
227 accumulation period was 900 seconds. The slopes were measured from linear, quadratic and  
228 exponential regressions of increasing concentrations starting after a dead band of 90 seconds.  
229 The gas pressure, temperature and humidity inside the stem sleeve were measured from the  
230 circulated gas running through the UGGA's flow-cell and we used temperature, pressure and  
231 humidity compensated CH<sub>4</sub> concentrations for the slope calculations. The advantage of using  
232 the cell temperature is the perfect synchronicity of the airflow with the temperature  
233 measurement. In previous tests we showed that the cell temperature was strongly correlated  
234 ( $R^2 = 0.994$ ) to the chamber temperature measured with a small data logger (ST-171, Clas  
235 Ohlson, Insjön, Sweden). Besides, the analytical laser did not significantly increase the  
236 temperature of the closed circuit (cell, connection tubes and chamber), as the temperature drift

237 over 20 minutes of enclosure was only +0.7 % under lab conditions (SATP). The chamber  
238 pressure was equilibrated to the outside monitored atmospheric pressure (Gas pressure sensor,  
239 Vernier, Beaverton, USA) via the vent tube.

240 All chambers were connected to an UGGA via two flexible tubes (see chamber designs  
241 section) set in parallel in a continuous flow mode; one tube bringing air from the gas analyser  
242 towards the chambers and the other tube pumping air from the headspaces towards the gas  
243 analyser. The tubes were connected to the gas analyser via two ¼ inch push-connect fittings.  
244 The UGGA's pump ensured a continuous flow of 2-4 L min<sup>-1</sup>. The UGGA measured CH<sub>4</sub>  
245 with the Off-Axis Integrated Cavity Output Spectroscopy (OA-ICOS) at a frequency of 0.33  
246 Hz. The analyser's uncertainty in the range of 0.01 ppmv to 100 ppmv of methane is <1%  
247 without calibration and the precision is ±0.6 ppb over a period of 100 seconds (LGR, 2013).

248

## 249 **2.5 Methane permeability calculations**

250 In order to quantify and compare CH<sub>4</sub> losses from the three types of chambers (two semi-rigid  
251 sleeves and one rigid chamber) attached to an inert stainless steel cylinder we corrected the  
252 loss rates by taking into account both the stem exchange surface covered by each sleeve (or  
253 chamber) as well as the concentration gradient between inside and outside of each chamber.  
254 To express this we calculated the permeability as a function of the effluxes (outgoing fluxes)  
255 and the concentration gradient between inside and outside the chambers.

256 In the first step we multiplied the slope (mg m<sup>-3</sup> s<sup>-1</sup>) by the total volume of the chamber ( $V_{\text{tot}}$ )  
257 to get the loss rates (mg s<sup>-1</sup>). We then divided the loss rates from each sleeve (or chamber) by  
258 the stem exchange surface ( $S_c$ ) covered by each sleeve (or chamber) to express the methane  
259 flux (J) which can be used for both the permeability experiment on the metallic cylinder and  
260 the methane accumulation runs from tree-stems in the field:

261  $Loss\ rate = slope * V_{tot} = \frac{dC}{dt} * V_{tot} \left[ \frac{mg}{s} \right]$  (7)

262  $Flux\ (J) = \frac{Loss\ rate}{S_c} = \frac{dC}{dt} * \frac{V_{tot}}{S_c} \left[ \frac{mg}{m^2s} \right]$  (8)

263 In the second step, from Fick's first law (Fick, 1855) we could apply the general equation  
 264 often used in cell biological or textile fabric applications (Ogulata and Mavruz, 2010) to  
 265 calculate, for each sleeve (or chamber), the CH<sub>4</sub> permeability (P) through a porous medium by  
 266 dividing the CH<sub>4</sub> flux (J) by the CH<sub>4</sub> concentration gradient (ΔC) between inside (C<sub>chamber</sub>)  
 267 and outside of the sleeve (C<sub>atm</sub>). We assume that the diffusive CH<sub>4</sub> losses (including dilutions)  
 268 through the rigid and semi-rigid material are negligible at SATP conditions (McKeen, 2012).  
 269 Thereafter the equation was:

270  $J = -P * \Delta C \rightarrow Permeability\ (P) = -\frac{J}{(C_{chamber} - C_{atm})} \left[ \frac{m^3}{m^2s} \right]$  (9)

271

## 272 **2.6 Numerical analyses**

273 We used linear, quadratic and exponential regressions to fit the CH<sub>4</sub> concentrations as a  
 274 function of the accumulation time in the chambers. The fitting was based on sum of squares'  
 275 minimisation of the errors. The frequency distribution, homogeneity and homoscedacity of  
 276 the residuals were previously checked using normal quartile plots, residual versus predicted  
 277 plots, and box plots. The coefficient of determination (R<sup>2</sup>) was used to quantify the level of fit.  
 278 All the data was analysed with the SAS software (SAS Institute Inc., Toronto, Canada).

279

## 280 **3 Results**

### 281 **3.1 Calibration of the semi-rigid sleeves**

282 The compared predicted (theoretical) and the mean observed empirical  $V'_{tot}$  (Eqs. 1 and 4)  
283 were respectively: 966 and 933 cm<sup>3</sup> for the small sleeve, 1406 and 1439 cm<sup>3</sup> for the large  
284 sleeve, and 13581 and 13026 cm<sup>3</sup> for the rigid chamber (Supplement S1). The observed  $V'_{tot}$   
285 values included variability due to the possible but very tiny compaction of the Neoprene foam  
286 over the whole frame. This compaction was less than 3% of  $V_{tot}$ , which was a maximum  
287 considering the pulling force of 200 N applied on the straps (twice 100N).

288 The difference between the mean observed  $V'_{tot}$  and the predicted  $V_{tot}$  values gave us an  
289 estimate of the bias in size. By dividing the absolute value of the bias through the predicted  
290 value we get an estimate of the inaccuracy of  $V_{tot}$  (chamber, tubes and detector's cell). As all  
291 terms of the fraction (Eq. 4) are linearly dependent, the inaccuracy of the permeability (P) is  
292 the quadratic mean of all other terms (Table 1, footnote). The gas exchange surface ( $S_c$ ) could  
293 be precisely determined and we assume that there is no error associated with it. The  
294 inaccuracies in the concentration measurements are dependent on the uncertainty of the  
295 UGGA, which in our case was <1% for the un-calibrated device.

296 The precision of our measurement system, related to repeatability, is the level to which  
297 repeated measurements show the same results under the same conditions. For each sleeve or  
298 chamber we repeatedly injected 50 mL of a 200 ppmv standard and measured the initial  
299 concentration ( $C_0$ , Table 1, Supplement S1) in the enclosed volume. We used the relative  
300 standard error (RSE) of the initial concentration ( $C_0$ ) to express the level of precision between  
301 different types of chambers. Thereafter, precision is of  $\pm 1.82\%$  for the small sleeve,  $\pm 1.59\%$   
302 for the large sleeve and  $\pm 1.68\%$  for the rigid chamber.

303

### 304 **3.2 Chamber permeability comparisons**

305 The comparison of permeability (Table 1, Supplement S1) of the three types of chamber  
306 shows that the semi-rigid sleeves are on average less permeable than the rigid chamber, and  
307 that the smaller semi-rigid sleeve had a higher permeability compared to the larger one. It was  
308 also interesting to note that the CH<sub>4</sub> loss (negative slope) is lower for the rigid chamber  
309 compared to semi-rigid sleeves. The contrasting higher permeability of the rigid chamber was  
310 counterbalanced by the much greater V<sub>tot</sub> as well as a much lower initial concentration  
311 gradient between inside and outside of the chamber ( $dC = C_0 - C_{atm}$ ). The rigid chamber was  
312 14.1 times that of the small sleeve in volume and 9.7 times that of a large sleeve, and the  
313 initial concentration gradient in the rigid chamber was only 1/14 of that in the smaller sleeve  
314 and 1/9 of that in the bigger sleeve. Moreover, the larger sleeve had a larger S<sub>c</sub>-to-V<sub>tot</sub> ratio  
315 (0.42) compared to the smaller sleeve (0.34).

316 In order to understand why the permeability of the semi-rigid sleeves was lower than that of  
317 the rigid chamber we compared and calculated the potential contact distances between air  
318 from inside and outside of the chamber volumes (Fig. 3, Supplement 2). Those contact zones  
319 represented the paths where gas effusion could occur, which were driven by the architecture  
320 of the chamber. For that purpose we distinguished two types of contact lines: 1) mobile lines  
321 that needed to be sealed properly every time the chambers were deployed and from which  
322 most of the losses were likely to occur, and 2) fixed lines that resulted from the manufacture  
323 which could be cracked and leak as a result of twisting forces on the rigid joints. The result  
324 was that for the same theoretical stem gas exchange surface (S<sub>c</sub>) between the two chambers  
325 (same length and height), the ratio between the length of the mobile lines and the stem gas  
326 exchange surface (S<sub>c</sub>) was 2.17x smaller for the semi-rigid as compared to the rigid approach.

327

### 328 **3.3 Stem-methane emissions and field deployments**

329 In the field, the manual sampling by syringe showed steady concentration changes with the  
330 sleeve technique (Fig. 5, Supplement S3), and the linear fitting of those concentration changes  
331 was always high ( $R^2 \geq 0.924$ ). When applying a quadratic fit the coefficient of determination  
332 improved substantially ( $R^2 \geq 0.995$ ). In continuous flow mode the concentration changes were  
333 also consistent with the sleeve technique (Fig. 6, Supplement S4), and the linear fitting was  
334 very high ( $R^2 \geq 0.989$ ) for all the runs not displaying leakages. Equally to the manual sampling  
335 mode, in continuous flow mode, the fitting improved slightly when applying a quadratic  
336 function or an exponential function ( $R^2 \geq 0.998$ ).

337 The two modes also distinguish themselves by the fact that with the continuous flow mode the  
338 runs are shorter compared to the manual mode. The runs were set to 15 minutes closure for  
339 the manual mode and to 7 minutes closure for the continuous flow mode. These times  
340 included a maximum of 90 seconds equilibration time just after the sleeve was deployed to  
341 allow the headspace to mix properly.

342 Runs 3 and 6 of the continuous flow mode were deliberately presented to display situations  
343 where leakages from sleeves were occurring when placed on *Betula pendula* or *Pinus sylestris*  
344 tree-stems (Fig. 6). In those cases, the CH<sub>4</sub> concentrations developed in a disordered way with  
345 periods of increases immediately followed by sudden drops. These analytically monitored  
346 leakages were confirmed when checking the chamber fitting on the stems.

347 The determination of the coefficient of variation of the root-mean-square error CV(RMSE),  
348 often used to measure the relative differences between two populations of values, and which  
349 was calculated between the linear fitted slopes and the non-linear fitted slopes, was higher in  
350 the case of the manual sampling mode (0.69) as compared to the continuous flow mode (0.45).  
351 In other words, the difference between the linear and non-linear fittings was 53% higher in the  
352 manual mode as compared to the continuous mode. This went in parallel with the differences

353 between the average slope in the linear fitting and that from non-linear fitting which was 27%  
354 higher with the manual sampling mode as compared to 18% with the continuous mode.

355

## 356 **4 Discussion**

### 357 **4.1 Semi-rigid sleeve construction**

358 The semi-rigid sleeves are easy to assemble, lightweight, and can be locally sourced. The  
359 sleeves could easily be assembled on-site following transportation. This allows for minimal  
360 luggage or shipping space and low costs, a major asset in terms of logistics where remote  
361 fieldwork is concerned. The PET or PC sheets were precisely cut in advance whereas the  
362 framing with the Neoprene strips was done on-site. We made sure that all components were  
363 not emitting CH<sub>4</sub>, which might otherwise confound in-situ measurements. Nevertheless, the  
364 raw materials are commonly available internationally, could be found on-site and likewise  
365 tested. For small sleeves (stem diameters  $\leq 15$  cm) and middle-sized sleeves (stem diameters  
366  $\leq 25$  cm) the pre-shaped PET sheet can easily be constructed from soft drink PET bottles or  
367 PC water-fountain tanks. Larger sleeves (stem diameters  $> 25$  cm) can be built from flat PC  
368 sheets as the curvature and volume stability of the chamber becomes less compromised with  
369 larger stem diameters. Most important for the construction of the sleeves are the vertical  
370 wedges that keep the sheet equidistant from the stem along the radial periphery of the sleeve.  
371 The construction of a sleeve took about one hour and there was no requirement for specific  
372 machine tools and no adhesives were needed, as the Neoprene bands used were adhesive  
373 backed. For the production of large numbers of sleeve rectangular plastic sheets could be  
374 thermoformed using a specially designed mould (Throne, 1996).

375 The average CH<sub>4</sub> mass losses (2.2-3.3 %) from the sleeves after 20 minutes of deployment  
376 were two orders of magnitude greater as compared to the 0.038% mass loss after 30 minutes



377 of deployment reported by Hutchinson and Livingstone (2001) for a perfectly sealed chamber  
378 with a sole vent tube. Thereafter, our downscaled vent tube was proportioned to the CH<sub>4</sub>  
379 losses from the sleeves.

380

## 381 **4.2 Calibration of the semi-rigid sleeves**

382 All the chambers were reasonably precise (repeatable) in terms of total volume and the semi-  
383 rigid chambers (sleeves) performed equally compared to the rigid chambers. In terms of total  
384 volume inaccuracy, all chambers were below the threshold significance level of 5%.  
385 Moreover, the semi-rigid sleeves' total volume accuracy increased with increasing  $S_c/V_{tot}$ .  
386 Nevertheless, getting good accuracy is a matter of calibration as biases can be subtracted from  
387 the original readings.

388 The average 33 cm<sup>3</sup> greater  $V'_{tot}$  values as compared to  $V_{tot}$  for the large sleeve (Supplement  
389 S1) can be attributed to the volume of the wedges that were also undergoing a compaction  
390 when deployed as the interior periphery gets compressed. This tiny volume correction was not  
391 inserted in formula 4 for the sake of simplicity and because the difference with the calibration  
392 was still below 5%.

393 We added a known amount of CH<sub>4</sub> instantaneously to the chambers and followed its decline  
394 and associated chamber permeability. Thereafter, we can be aware of how well the chambers  
395 are doing in keeping the considered gas but not how well they do in minimizing the errors  
396 associated with the gas exchange processes between stems and the chamber. For those errors  
397 we referred to recommendations from other studies, such as: ensuring air-mixing, venting,  
398 reducing closure times, reducing chamber volume and considering non-linear fitting  
399 (Christiansen et al., 2011; Hutchinson and Livingstone, 2001; Juszczak, 2013; Pihlatie et al.,  
400 2013).

401

### 402 **4.3 Chamber permeability comparisons**

403 A reasonable mechanistic explanation to the fact that both semi-rigid sleeves were on average  
404 57% less permeable compared to the rigid chamber (Table 1) could come from the sleeve's  
405 smaller proportion of air contact lines between inside and outside the chambers thereby  
406 reducing opportunities for gas diffusion to occur. The difference in that proportion is similar  
407 in order of magnitude to the difference in permeability (Supplement S2). Moreover, it is  
408 possible that with an aging rigid chamber the permeability could increase faster than in the  
409 case of an aging semi-rigid sleeve as the proportion of fixed contact lines could be exposed to  
410 more cracks and unforeseen reduced air-tightness (Fig. 3, green lines).

411 This is also in line with the fact that for the same semi-rigid chamber design with the  
412 increasing  $S_c$ -to- $V_{tot}$  ratio, thus by increasing frame size, there is a concurrent decrease in the  
413 proportion of contact lines as well as a concurrent decrease in permeability. The rigid  
414 chamber had a much lower  $S_c$ -to- $V_{tot}$  ratio when compared to the sleeves and showed the  
415 greatest permeability. From our observations we can generalise the common trend found for  
416 all chamber types by saying that the larger the total volume of a stem chamber is, for a given  
417 gas exchange surface, the greater the expected permeability.

418 With the same logic and by considering the strong leverage effect of the concentration  
419 gradient ( $\Delta C$ ) between inside and outside the chamber, the advantage of the larger rigid  
420 chamber is that it keeps the concentration gradient more constant during the chamber  
421 deployment and therefore minimizes the non-steady-state gas saturation effect of the closed  
422 chamber system. However, this advantage loses its importance when semi-rigid sleeves are  
423 connected to precise gas analysers with analytical frequencies of up to 10 Hertz as the  
424 gradient effect can equally be minimized by reducing the closure times to a few minutes.

425 Additionally, by increasing the  $S_c$ -to- $V_{tot}$  ratio by 6 fold compared to rigid chambers and by  
426 mixing the enclosed gas through the continuous flow circulation, we also avoided the  
427 problems associated with large volume chambers (Hutchinson and Livingston, 2001, 1993).

428 Nevertheless, the only non-compressible time factor is the sleeve's equilibration period; a 90  
429 second period for the continuous air circulation to mix the entire headspace. This could be  
430 shortened by reducing the tube length, increasing the pump's flow-through or by installing a  
431 complementary fan if the sleeves were to be built much larger. In any case, the threshold time  
432 by which the sleeve headspace is mixed entirely can be monitored graphically while running  
433 every sample. Retrospectively, 90 seconds of equilibration, together with 3-minute closure  
434 time, conservatively characterised all replicates made for two different sleeve sizes (n=24).

435

#### 436 **4.4 Deployment in the field**

437 As expected, deployment of the semi-rigid sleeve was very straightforward and could be  
438 operated by a single person. The fact that the sleeves had a natural tendency to curve (pre-  
439 shaped) allowed them to stay in place when initially placed around the stem. This gave the  
440 researcher free hands to attach the straps subsequently. The whole setup takes two minutes to  
441 install and swapping the sleeves between different stem heights was also done much more  
442 efficiently in comparison to the rigid chamber deployment.

443 In theory all stem sizes could be fitted, the only limitation comes from the stem texture and  
444 this is valid for both semi-rigid sleeves as well as rigid chambers. In some situations, the tree  
445 bark had large crevices and it was necessary to prepare the stem prior to attachment of the  
446 sleeves or rigid chambers. The preparation was made by filling the crevices with mastic or  
447 play dough in the shape of a frame before the chamber or sleeve could be sealed to the stem.  
448 In some other situations it was enough to increase the thickness of the sleeves to reduce the

449 percentage of uncertainty in the chamber volume ( $V_c$ ). The impact of both crevices and  
450 bumps could be assessed with distance measurements made on photos taken on one side of  
451 the deployed sleeves.

452 Using five sleeve sizes it was possible to cover stem diameters ranging from 5 cm to 127 cm  
453 at breast height (DBH). Moreover, in terms of weight the two sleeves we tested were  
454 respectively 156 and 297 grams, compared to 3.3 kg for the rigid chamber. As a consequence,  
455 the whole collection of sleeves fitted in a single backpack and was light to carry.

456 Under changeable conditions such as varying sunlight intensities we recommend that the  
457 temperature inside and outside of the sleeve is measured, and to shade the sleeve. Otherwise,  
458 these varying conditions may alter the gas exchange processes between the stem and the  
459 atmosphere.

#### 460 **4.5 Sampling modes and regression fits**

461 In both cases, for manual sampling and continuous flow (Figs. 5 and 6), methane  
462 accumulation rates were better fitted with non-linear functions (quadratic or exponential).  
463 This confirms that the sleeve's closure system was sealing properly against the stems, as the  
464 headspace concentration change of a closed non-steady-state chamber (static chamber) will  
465 always remain non-linear and this is driven by the laws of diffusion (Hutchinson and  
466 Livingston, 2001). For the semi-rigid sleeve, the difference between both the  $R^2$  and the  
467 slopes between the linear fitted and the non-linear fitted concentration changes were roughly  
468 twice as small compared to those reported in the literature for soil chambers (Christiansen et  
469 al., 2011; Hutchinson and Livingston, 2001; Juszczak, 2013; Pihlatie et al., 2013).

470 Furthermore, the impact of the manual syringe sampling on the pressure fluctuation in the  
471 sleeve could be somewhat minimised by the fact that the chamber volume ( $V_c$ ), where the  
472 actual air mixing occurred, was increased by the additional dead volume added from the

473 analyser and tubing in continuous flow mode. Thus, the total volume ( $V_{\text{tot}}=V_c+V_{\text{dead}}$ ) was  
474 increased as much as 76% with the smaller sleeve. With rigid soil chambers this aspect is  
475 often not mentioned as in those cases the dead volume is negligible compared to the large  
476 chamber volume. In our case, for the manual sampling, over a 15 minutes period, we drew  
477 1.8% of the total volume from the larger sleeve (4 steps of 0.44%), which in terms of mass  
478 loss remains below the significance level of 5% and could be accounted for if more accuracy  
479 is needed. Although the repeated gas sampling minimises somewhat the pressure build up,  
480 recent studies have recommended avoiding manual sampling as much as possible because of  
481 associated pressure fluctuations (Christiansen et al., 2011; Juszczak, 2013).

482 The coefficient of variation of the root-mean-square error CV (RMSE) gave 53% higher  
483 coefficients for the manual sampling mode compared to the continuous flow mode thus  
484 indicating that the discrepancy between the linear fitting and the non-linear fitting is higher  
485 for the manual sampling mode. Moreover, as reported by some authors, fluxes calculated  
486 using linear fitting together with non-steady state chambers could be underestimated by as  
487 much as 40% (Christiansen et al., 2011; Pihlatie et al., 2013; Kutzbach et al., 2007). In our  
488 case, the underestimation was 27% for manual sampling mode and 18% for the continuous  
489 flow mode. As a consequence we would recommend using non-linear fitting (quadratic or  
490 exponential) together with manual sampling of the semi-rigid sleeves. In continuous flow  
491 mode, it is better to reduce the closure times as much as possible if planning to use linear  
492 fitting for greater simplicity. Both measures will contribute to improving line-fitting and  
493 estimating CH<sub>4</sub> accumulation rates.

494

## 495 **5 Conclusions**

496 Although all chamber types performed well, the semi-rigid design had numerous benefits  
497 including reduced gas permeability and an optimal  $S_c$ -to- $V_{tot}$  ratio. Furthermore, they can be  
498 easily constructed and transported in multiple sizes, are extremely light, cheap to build and  
499 fast to deploy. As an example, in three of our tropical campaigns it was possible to carry a  
500 complete collection of semi-rigid sleeves in a single backpack. The collection covered the  
501 sampling of all ecosystem stem-sizes. Alternatively, we could also build the chambers on-site  
502 after prior testing of the compounds for background emissions. The PET and PC sheets of the  
503 sleeves are sturdy and lasted the duration of the campaigns, while the closed-cell Neoprene  
504 strips could be used for several weeks in the field before they needed to be replaced.

505 Connecting the sleeves in continuous flow mode to fast and precise laser-spectroscopic gas  
506 analysers (CRD or OA-ICOS technologies) enables the combined analysis and air mixing of  
507 the sleeve's enclosed volume, as well as reducing the closure periods to no-more than three  
508 minutes, making linear fitting from initial rates less problematic. To ensure optimal accuracy  
509 of the concentration measurements, it is best to calibrate each individual sleeve's total volume  
510 by diluting a standard gas in the entire setup (chamber, connectors, tubes and analyser) prior  
511 to starting a measurement programme.

512 Finally, to make good estimates of the global importance of tree-stem  $CH_4$  emissions, it is  
513 essential to make measurements that cover all types of trees (species and morphotypes)  
514 present within the often remote ecosystems explored. This necessitates great adaptability in  
515 the chamber sizing and transport logistics. The semi-rigid sleeves meet these requirements  
516 without compromising the quality of the data collected.

517

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523

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603

#### 604 **Figure captions**

605 **Figure 1.** Smaller semi-rigid stem sleeve attached to a stem. The plastic PET sheet (a) has  
606 three imprinted circular rims (b) that ensured good stability and natural curvature of the sleeve.  
607 The circumference of the sheet was framed with a 1.5 cm thick and 3 cm wide expanded  
608 Neoprene strip (c) that sealed off the headspace located between the sheet and the stem. Inside  
609 this volume there were two vertical wedges (d) that kept the sheet at equidistance from the  
610 stem along the radial periphery of the sleeve. In its centre the sleeve was equipped with two

611 Snap-on rubbers with inserted three-way stopcocks (e) that were further connected to PVC  
612 tubes that went from the sleeve to the Ultraportable Greenhouse Gas Analyser. A coiled vent  
613 was placed in one corner of the sleeve (f) to regulate the pressure. The chamber was tightened  
614 to the stem with the help of two straps that perfectly aligned on top of the horizontal strips.

615 **Figure 2.** The three steps of the semi-rigid stem sleeve deployment. To ensure a good contact  
616 between the frame strips and the stem it was important to distribute the pressure of each strap  
617 all around the frames' periphery when tightening the sleeve. Close to the centre two Snap-on  
618 rubbers with inserted three-way stopcocks were pressed into the PET or PC plastic sheet.  
619 These stopcocks were connected to the two PVC tubes that circulated air in a continuous flow  
620 mode when connected to an Ultraportable Greenhouse Gas Analyser (UGGA).

621 **Figure 3.** Potential air contact path lines (chamber air versus ambient air) where gas diffusion  
622 can occur; a comparison between the acrylic rigid cylinder approach and the semi-rigid sleeve  
623 approach. The red lines represented the mobile contact lines that needed to be sealed properly  
624 every time the chambers were deployed and where most of the losses were likely to occur.  
625 The green lines represented the fixed contact lines which could have been leaking as a result  
626 of twisting forces on the joints leading to cracks.

627 **Figure 4.** 2-D Layout for the chamber volume ( $V_c$ ) calculation based on the stem diameter  
628 ( $D_{stem}$ ), the thickness of the chamber ( $T$ ), the sector covered by the chamber ( $K$ ) and the  
629 volume of the wedges ( $V_{wedge}$ ). Refer to the text for the volume calculations.

630 **Figure 5.** Contrasting methane concentration changes in the semi-rigid sleeve from enclosed  
631 gas samples measured in a manual mode (syringe) from tree-stems. In the first six runs (top  
632 quadrants) the concentration changes were regressed with a linear fit, while in the second set  
633 of runs they were regressed with a quadratic fit (non-linear). All runs 1-6 were measured on  
634 *Heisteria concinna* stems from a tropical lowland forest. The blue line corresponds to 95%

635 confidence intervals,  $RMSE$  = root-mean-square error,  $R^2$  = coefficient of determination,  $Y$  =  
636 methane concentration in ppmv.

637 **Figure 6.** Contrasting methane concentration changes in the semi-rigid sleeve from enclosed  
638 gas samples measured in continuous flow mode (UGGA) from tree-stems. In the first six runs  
639 (top quadrants) the concentration changes were regressed with a linear fit, while in the second  
640 set of runs (bottom quadrants) they were regressed with quadratic fit (non-linear). Runs 1, 2, 3,  
641 and 5 were made on *Betula pendula* stems, runs 4 and 6 were made on *Pinus sylvestris* stems,  
642 runs 3 and 6 show the concentration responses in situations where the sleeves were leaking.  
643 The blue line corresponds to 95% confidence intervals,  $RMSE$  = root-mean-square error,  $R^2$  =  
644 coefficient of determination,  $Y$  = methane concentration in ppmv.

**Table 1.** Chamber dimensions and mean permeabilities ( $P$ ) determined, for each replicated chamber ( $n = 3$ ), from the methane decline slope ( $Slope$ ), the total chamber volume ( $V_{tot}$ ), the initial concentration gradient between outside and inside ( $C_0 - C_{atm}$ ) and the gas exchange surface ( $S_c$ ).  $D$  = metallic cylinder diameter,  $L$  = peripheral length of the enclosure,  $H$  = height,  $T$  = thickness,  $C_0$  = initial enclosure concentration,  $C_{atm} = 1.8951$  ppmv,  $R^2$  = coefficient of determination of the decline regression,  $V_c$  = volume of the chamber,  $V_{tot} = V_c + V_{dead}$ ,  $V_{dead}$  = dead volume of the analyser plus the tubes =  $416 \text{ cm}^3$ . Values in brackets represent the standard error of the mean ( $\pm$ SEM).

Enclosure type	$D$ (cm)	$L$ (cm)	$H$ (cm)	$T$ (cm)	$S_c$ ( $\text{cm}^2$ )	$V_c$ ( $\text{cm}^3$ )	$V_{tot}$ ( $\text{cm}^3$ )	$C_0$ (ppmv)	$Slope$ ( $\text{mg m}^{-3} \text{ s}^{-1}$ ) $\times 10^{-4}$	$R^2$	$P$ ( $\text{m s}^{-1}$ ) $\times 10^{-7}$
Small sleeve	15	25	16	1.5	330	550	966 <sup>a</sup>	109.12 (2.00)	-21.40	0.930	8.30 (0.85) <sup>d</sup>
Large sleeve	15	30	24	1.5	594	990	1406 <sup>b</sup>	71.43 (1.14)	-9.86	0.922	4.77 (0.64) <sup>e</sup>
Rigid chamber	15	28	30	6.5	1413	13165	13581 <sup>c</sup>	9.58 (0.16)	-0.82	0.931	14.62 (1.86) <sup>f</sup>

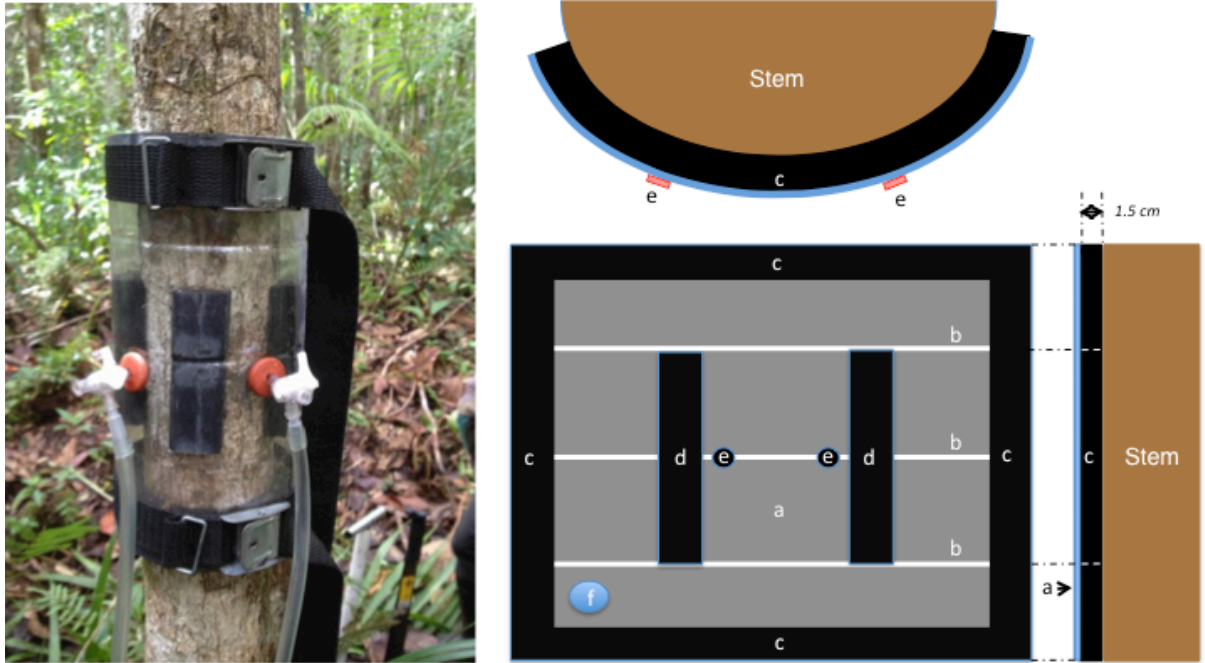
Volume inaccuracies: <sup>a</sup>  $\pm 3.4\%$ , <sup>b</sup>  $\pm 2.4\%$ , <sup>c</sup>  $\pm 4.1\%$ ; Permeability inaccuracies\*: <sup>d</sup>  $\pm 3.7\%$ , <sup>e</sup>  $\pm 2.8\%$ , <sup>f</sup>  $\pm 4.3\%$

\*Calculated from the error propagation formula:

$$\frac{dP}{|P|} \leq \sqrt{\left(\frac{dC}{|C|}\right)^2 + \left(\frac{dV}{|V|}\right)^2} + \sqrt{dC^2 + dC_{atm}^2} \cong \sqrt{\left(\frac{dV}{|V|}\right)^2} + 2$$



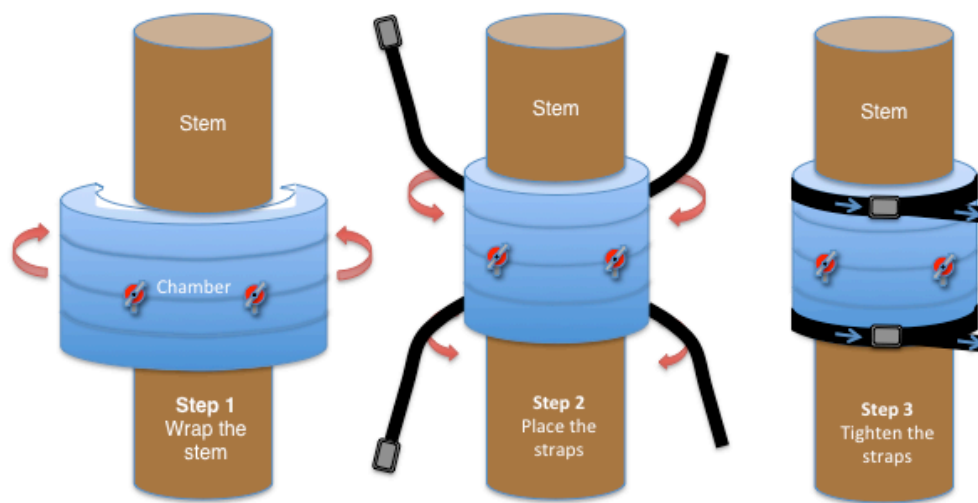
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651 Fig01

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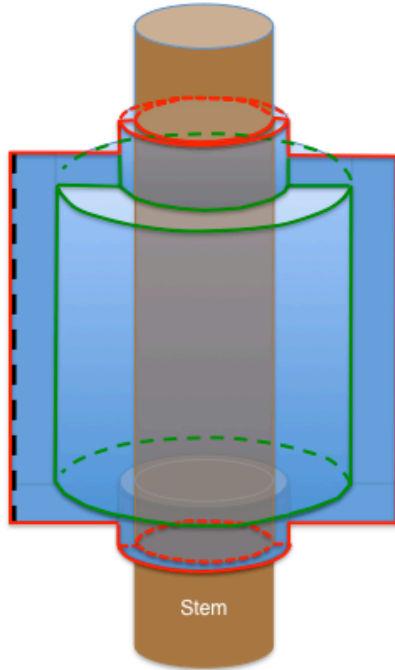
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654 Fig02

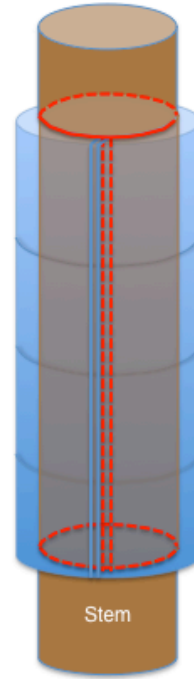
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Rigid stem chamber



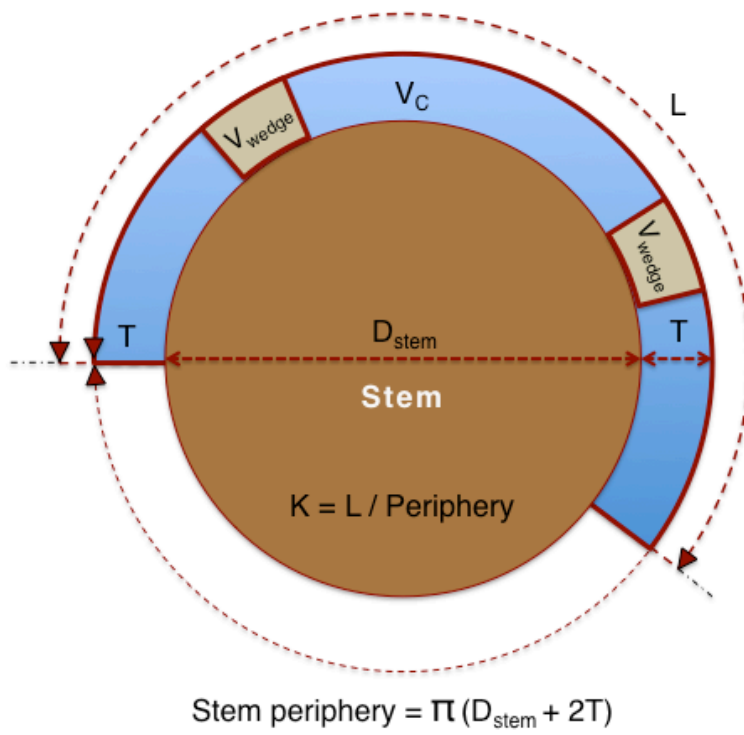
Semi-rigid stem sleeve



656

657 Fig03

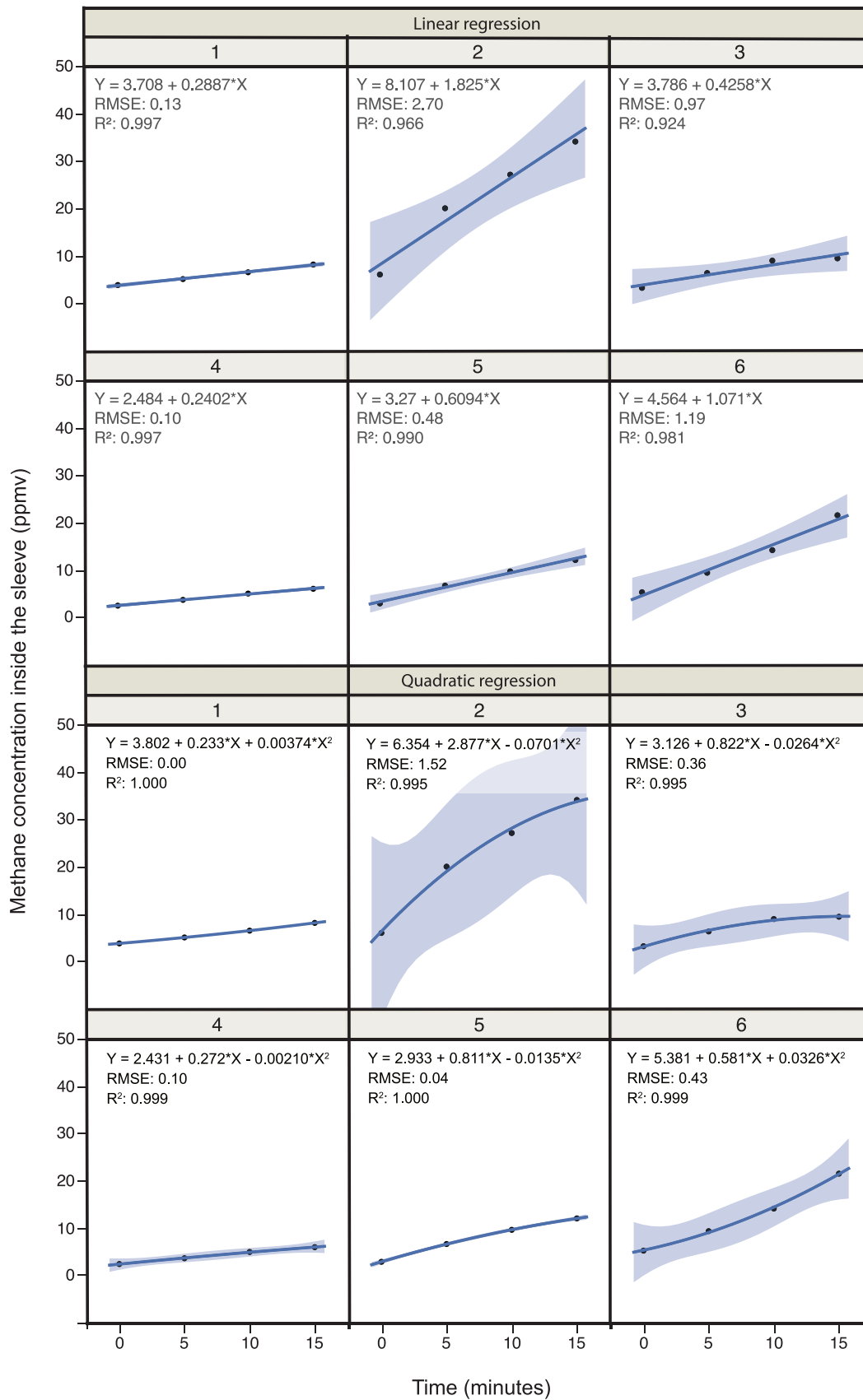
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660 Fig04

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663 Fig05

