

A vegetation and soil survey method for surveillance monitoring of rangeland environments.

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22

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25 Abstract

26 Ecosystem surveillance monitoring is critical to managing natural resources and especially so under
27 changing environments. Despite this importance, the design and implementation of monitoring
28 programs across large temporal and spatial scales has been hampered by the lack of appropriately
29 standardised methods and data streams. To address this gap, we outline a surveillance monitoring
30 method based on permanent plots and voucher samples suited to rangeland environments around the
31 world that is repeatable, cost-effective, appropriate for large-scale comparisons and adaptable to
32 other global biomes.

33 The method provides comprehensive data on vegetation composition and structure along with soil
34 attributes relevant to plant growth, delivered as a combination of modules that can be targeted for
35 different purposes or available resources. Plots are located in a stratified design across vegetation
36 units, landforms and climates to enhance continental and global comparisons. Changes are
37 investigated through revisits. Vegetation is measured to inform on composition, cover and structure.
38 Samples of vegetation and soils are collected and tracked by barcode labels and stored long-term for
39 subsequent analysis. Technology is used to enhance the accuracy of field methods, including
40 differential GPS r plot locations, instrument based Leaf Area Index (LAI) measures, and three
41 dimensional photo-panoramas for advanced analysis. A key feature of the method is the use of
42 electronic field data collection to enhance data delivery into a publicly-accessible database.

43 Our method is pragmatic, whilst still providing consistent data, information and samples on key
44 vegetation and soil attributes. The method is operational and has been applied at more than 704
45 field locations across the Australian rangelands as part of the *Ecosystem Surveillance* program of
46 the Terrestrial Ecosystem Research Network (TERN). The methodology enables continental
47 analyses, and has been tested in communities broadly representative of rangelands globally, with
48 components being applicable to other biomes.

49 Here we also recommend the consultative process and guiding principles that drove the
50 development of this method as an approach for development of the method into other biomes. The
51 consistent, standardised and objective method enables continental, and potentially global analyses
52 than were not previously possible with disparate programs and datasets.

53 1 Introduction

54 Ecosystems support our social and economic well-being and require our vigilance as to their
55 condition and management intervention to ensure continued functionality (Magnusson et al.,
56 2013;Andersen et al., 2014). The diversity of ecosystems has contributed to a multitude of methods
57 used to sample their composition, structure and function. Despite acknowledgement of the need for
58 integrated measurements and evidence-based decision making (Likens, 2010;Eyre et al.,
59 2011;Likens and Lindenmayer, 2011), endorsing a single approach to ecosystem monitoring
60 remains difficult because managers, researchers, policy makers and funding agencies have diverse
61 applications for the data collected, and may have invested considerable effort and monitoring time
62 in existing methods. This causes an integration problem when bringing together monitoring data
63 across large areas, and is particularly problematic for programs involving extensive, multi-
64 jurisdictional, logistically challenging, and sparsely populated areas, such as rangelands (Bastin et
65 al., 2009;Herrick et al., 2010).

66 Underpinning these issues is a need to report on environmental change over decadal, or
67 longer, time periods (Allen-Diaz et al., 1996;Likens and Lindenmayer, 2011), and requires
68 monitoring methodologies that are well described and flexible to deliver on future, unanticipated
69 needs (Burton et al., 2014;Bayne et al., 2015). The challenge is to agree on a method without
70 complete knowledge of the requirements of future monitoring programs, the threats to ecosystems
71 or the opportunities that may emerge via innovation and technology (Spellerberg,
72 2005;Lindenmayer et al., 2014).

73 Rangelands occur on all inhabited continents (Figure 1a) with the predominant land-use being
74 low-intensity or nomadic livestock grazing on native pastures (Linstadter and Baumann, 2013). In
75 Australia, variable rainfall is perhaps the major ecological driver of spatial patterns (Stafford Smith
76 and McAllister, 2008), with the influence of variability particularly evident in arid areas (Van Etten,
77 2009; Dickman and Wardle, 2012). Rangelands represent 46% of terrestrial ecosystems globally and
78 81% in Australia (Figure 1b), but remain relatively poorly studied (Sparrow et al., 2014).
79 Understanding broad scale change in rangelands remains difficult due to a lack of monitoring and
80 decadal ecosystem dynamics (White et al., 2012b).

81 Here we present an overview and rationale of a cohesive and robust ecosystem surveillance
82 method that builds on previous techniques (Bastin et al., 2009; Herrick et al., 2010; Taylor et al.,
83 2014) for characterising and monitoring rangeland ecosystems. Specific protocols are described in
84 the AusPlots field manual included in the supplementary material (White et al., 2012a). The method
85 is operational and has been implemented at over 704 sites across Australia (see Box 1 in Appendix
86 S3), producing publicly available data for ecological studies of Australian rangelands (Guerin et al.,
87 2016; TERN, 2019). While the primary purpose of the program is to detect changes over large scales
88 of space and over long periods of time, it also provides readily available resources to answer
89 pressing current questions. For example, this method incorporates data collection (Tokmakoff et al.,
90 2016) to address key long-standing questions for rangeland ecosystems (Morton et al., 2011),
91 including understanding the role of soils and plant traits on productivity (Bastin et al.,
92 2017; Gallagher et al., 2019 Accepted with Revisions). The combination of vegetation data and
93 samples can be used to extract trait data or genomic data to anticipate responses to environmental
94 stressors across large gradients (Westoby et al., 2002; Wright et al., 2004; Guerin et al., 2012; Caddy-
95 Retalic et al., 2017). Similarly, the soil and plant samples, along with the vegetation data, provide
96 an open-access resource to assess emerging priorities such as understanding the multi-functionality
97 of ecosystems, and especially of drylands (Maestre et al., 2012).

98

99 **1.1 Challenges to broad scale monitoring approaches**

100 Surveillance monitoring is defined as broad in scope, involving measurements of many
101 attributes and species across a spatially and temporally wide-ranging network of field locations,
102 placing it between landscape and targeted field monitoring in detail and spatial extent (Eyre et al.,
103 2011). Challenges to the design and implementation of surveillance monitoring programs stem from
104 practical and scientific considerations (Lindenmayer et al., 2014), as well as from the imperative for
105 monitoring. Firstly, it is necessary to identify knowledge gaps, such as in the geographic extent or
106 type of data available (Sparrow et al., 2014), or the questions arising from environmental or societal
107 changes (Sutherland et al., 2015). The large and remote nature and the difficulty in accessing these
108 rangelands (Dickman et al., 2014; Sparrow et al., 2014) also provides a significant challenge for
109 these operational programs. Secondly, practical constraints must be factored into monitoring
110 protocols. Given the high cost of travel and data acquisition, surveillance monitoring methods need
111 to maximise benefits from each visit, necessitating an efficient and integrated workflow from data
112 collection to delivery of new knowledge (Tokmakoff et al., 2016).

113 The adoption of new monitoring programs, or integration of existing ones, can be hampered
114 by poor communication of goals. A well-designed method may fail because practitioners are
115 resistant to change or have divergent goals or cultural expectations (Ens et al., 2014). Widespread
116 consultation and involvement is therefore key to successful engagement (see Box 2 in Appendix
117 S4). A challenge for these programs is to prioritise time for adequate engagement in the design and
118 evaluation of methods in an environment with pressure to provide rapid results. New technologies
119 and innovations should be considered for inclusion in situations where they provide increase
120 accuracy or efficiencies over traditional techniques.

121 A key motivation for developing a new rangeland monitoring method in Australia was to
122 overcome the lack of compatibility between existing jurisdictional data collection methodologies
123 (see Box 11.5 in Foulkes et al., 2014). Global efforts to monitor terrestrial ecosystems (Bastin et al.,
124 2017) need to build upon regional and local data collection and ideally include a set of essential
125 environmental variables to provide a common modelling framework and scalable data to build a
126 cohesive global synthesis (Schmeller et al., 2015).

127

128 **2 Method overview and rationale**

129 **2.1 Pragmatic site selection**

130 A site selection protocol for surveillance monitoring of rangelands needs to be scientifically
131 robust but also practical. Consideration needs to be made for site access both at the time of initial
132 survey, but also for continued access for repeat measures. We implement a two stage stratification
133 procedure where we; 1) choose a bioregion to sample within (an Australian wide landscape
134 classification similar to ecoregions sensu Olson et al., 2001), and; 2) determine where within the
135 bioregion to establish plots.

136 Given the biophysical and anthropogenic context, we use three strategies to determine
137 bioregions within which to sample. 1) Stratified sampling to cover biophysical and disturbance
138 gradients. 2) Setting a minimum number of plots per representative bioregion or vegetation type. 3)
139 Additional sites identified via gap analysis. Scatterplots of relevant variables can reveal poorly
140 sampled regions in environmental space (Guerin et al., 2017), while ecologically scaled measures of
141 environmental uniqueness can be mapped over poorly sampled areas to identify priority habitats
142 (Arponen et al., 2008).

143 Once target bioregions have been identified, a more detailed process is undertaken at finer
144 spatial scales. Within stratified units, plot locations can be randomised where practical to maximise
145 representativeness and statistical rigour (Michalcová et al., 2011), whereas random sampling
146 without stratification across large areas results in under-represented habitats (Michalcová et al.,
147 2011). Plots can, and regularly are, co-located with those established by third parties (see Table 2)
148 or legacy projects to extract value and enhance temporal depth. Political information is often
149 relevant, including policy drivers influencing jurisdictions and opportunities for co-investment.
150 Some land managers see standardised surveillance monitoring as an opportunity to capture robust
151 information on the assets they manage and are receptive to co-investment.

152 Whether driven by stratification, gap-filling or policy needs, it is essential that site selection
153 accounts for logistical considerations such as access permissions and feasibility, to make the
154 program achievable and increase likelihood that sites will be re-sampled.

155

156 **2.2 Plot size and layout**

157 The choice of plot size was guided by the need to optimise the balance of survey resources
158 and scientific rigour. While representativeness and robustness to small-scale variation increase with
159 plot size (assuming vegetation within is homogenous), so does the expense of data collection,
160 equating to fewer plots for fixed resources. Large, single, 50 ha plots have become standard for the
161 study of demographic dynamics in rainforest biomes (Harms et al., 2001), 1 ha plots are used for
162 other woody ecosystems (Phillips et al., 2009; Miede et al., 2010), whereas grasslands are typically
163 surveyed in smaller 1 m² plots (Borer et al., 2014).

164 Given the vastness and heterogeneity of rangelands, there is a need for many plots and
165 therefore one-hectare plots were chosen for this method. Additional reasons for this choice

166 included: 1) the potential to capture species vital rates and vegetation processes (mortality,
167 recruitment, fire, grazing and drought responses) whilst maintaining a practical sampling size; 2)
168 the benefit of consistent results and reduced coefficients of variation in basal area, crown area and
169 vegetative structure between plots (Clark and Clark, 2000) – capturing small-scale patchiness whilst
170 providing representativeness overall; 3) enhanced integration with other activities that use 1 ha plots
171 (Phillips et al., 2009; Jurgens et al., 2012; Wood et al., 2015; Karan et al., 2016), and 4) to provide
172 information at an appropriate scale for validation of medium and high resolution remotely sensed
173 products (Congalton and Green, 2008).

174 The monitoring plots are established with Differential Global Positioning System (DGPS)
175 technology, to locate and record the coordinates of plot and transect vertices (Figure 2a) with sub-
176 meter accuracy and metal poles located in the corners and center to aid in relocation for repeated
177 monitoring. Each plot is located entirely within a relatively homogeneous (at the 1 ha scale) area of
178 a particular vegetation community, and is intended to be representative of that vegetation
179 community.

180 Plots are co-located with existing sites where possible. These sites have been established for a
181 variety of reasons (See table 2), and co-locating with these sites enables data from both programs to
182 be combined or correlated, enabling greater temporal depth, richer contextual information and often
183 co-investment in site establishment.

184

185 2.3 Floristics and vegetation

186 Plant cover and species composition are key essential variables (Pereira et al., 2013) for any
187 ecological surveillance monitoring program and are core modules in our method. Careful
188 consideration was given to ensuring adequate sampling effort, confirmation of species names by
189 taxonomists, and flexible methods to record both quantitative estimates of abundance for commonly
190 occurring and locally abundant species and occurrences of less abundant species within the plot. To
191 achieve this, several techniques are combined. All vascular plant species observed within the plot
192 are sampled and recorded, with identifications confirmed later by herbarium botanists. These
193 samples are then stored indefinitely. Vegetation is also characterised quantitatively by cover,
194 composition, growth-form and height. For this, a line point-intercept method is used across 10 x
195 100 m transects in a grid (Figure. 2a). This configuration ameliorates the skewing effects that site
196 heterogeneity may have on cover, which are difficult to avoid in rangelands (Vetter, 2005). Data
197 collected using this configuration are less sensitive to local heterogeneity or micro-patterning of the
198 vegetation.

199 Many authors recommend the collection of a minimum of 1000 intercepts in rangelands to
200 quantify cover per species (Lodge and Gleeson, 1976; Holm et al., 1984; Friedel and Shaw,
201 1987; Vittoz and Guisan, 2007). Following this research, our method utilises 1010 point intercepts
202 along transects to determine vegetation cover per species across the plot. Abundance and
203 presence/absence data are obtained by combining the identified sample data with the point-intercept
204 data, with rarer species not intercepted being scored as present but not assigned a cover score. Each
205 intercept is also attributed with information assessing plant height (used to reconstruct vegetation
206 structure and as a surrogate for recruitment of woody species) and whether the vegetation
207 intercepted is dead or senescent, to indicate mortality. Mass recruitment/mortality events are also
208 recorded. Height profiles can be created and their changes through time analysed to indicate
209 changes in vegetation community structure or to assign strata.

210 Accurate measures of vegetation cover are important for tracking environmental change, and
211 have many applications (Vittoz and Guisan, 2007). Cover can be summarised to family, genus,
212 species, growth-form levels or as fractional cover– the fraction of photosynthetically active

213 vegetation, dead vegetation and bare substrate. Relative cover-abundance can be used in
214 downstream analysis, or to classify vegetation, for example into structural classes such as forest or
215 shrubland based on height and cover of growth-forms. Change in vegetation structure (Fig 2c) or
216 composition can be quantified, for example to detect woody weed encroachment. Raw data can be
217 converted to common cover measures (e.g. opaque canopy cover or projected foliage cover), or
218 summarised by the highest intercepted plant at each point. The location of each point-intercept is
219 recorded (Fig. 2), allowing detailed spatial patterning to be investigated as an alternative to gross
220 plot-wide metrics.

221 Basal area is measured at each plot using a basal wedge sweep, to inform the amount of stored
222 above ground biomass based on allometric equations (e.g. Eamus et al., 2000). Basal area is
223 averaged across the plot, while raw tree stem counts are also recorded. A structural summary is also
224 collected, identifying the most dominant species in each of the Ground, Mid-layer and Upper strata
225 following the procedure described in Thackway et al. (2008). This enables the vegetation to be
226 described at the level of an 'Association', equivalent to a Level 5 structural description in the
227 Australian National Vegetation Information System (applicable to all vegetation types), with cover
228 and height information being calculated from the point intercept data.

229

230 **2.4 Photo-points**

231 Photo-points have long been used for monitoring (Watson and Novelty, 2004) and inventory
232 programs (e.g. Brandle et al., 2005). In keeping with this tradition, photo-points are created with a
233 new method in which panoramas (a continuous 360-degree sweep of static digital photographs with
234 at least 50% overlap between frames) are collected at three points (Fig. 3a,b,c). These photographs
235 are comparable with historical photographs, and can be analysed using computer vision techniques
236 to determine basal area (White et al., 2012a). It is anticipated that other structural metrics will be
237 able to be extracted from these photosets in future.

238

239 **2.5 Soils**

240 The soil protocol quantifies variability within and between plots and over time using a pre-
241 defined standard (National Committee on Soil and Terrain, 2009). This field protocol is undertaken
242 at the same time as the vegetation modules to enable vegetation analyses to consider
243 contemporaneous soil characteristics.

244 A plot description records erosion, micro-relief, landform pattern and element, drainage,
245 disturbance and soil surface condition. Four further modules are collected: soil pit; bulk density;
246 soil sub-sites; and metagenomic samples. A 1 m deep pit in the southwest corner of the plot (Fig.
247 2a) enables description and photographic recording of the upper soil profile and measurement of
248 pH, electric conductivity, texture, colour and structure (White et al., 2012a). Soils can then be
249 categorised using a standard such as the Australian Soil Classification (ASC) system (Isbell and
250 Terrain, 2016). Bulk density is measured at three depths of the pit to enable conversion of soil
251 properties to volumetric measures (Table 1). Soil sub-sites are collected at nine locations across the
252 plot, targeting variability in microhabitat, to collect the same information as at the soil pit to a depth
253 of 30 cm and analyse small-scale variability (Figure 2a). Soil samples taken specifically to enable
254 metagenomic analysis of environmental DNA (e.g. targeting soil biota in various phyla or traces of
255 above-ground flora and fauna) are collected from the surface at each sub-pit and stored on silica
256 granules. Soil samples are air dried and retained for further analysis and access by researchers
257 (Grundy et al., 2015).

258

259 **2.6 Samples for re-use**

260 Many monitoring methods that record species and taxonomic determination rely on botanists
261 who can identify specimens in the field and vouchers may only be collected for obtuse species or
262 records of interest (Hosking et al., 2000). Field identifications are prone to error (Scott and Hallam,
263 2003;Lacerda and Nimmo, 2010) and the requirement for taxonomic expertise can inhibit delivery
264 of plots. To address this issue, the method mandates the collection of herbarium vouchers for all
265 vascular plant species observed, which are tracked using barcode labels. In addition to ensuring
266 consistent identification, barcoded voucher specimens are a resource for ongoing research.
267 Vouchers can resolve taxonomic issues, including the discovery of new taxa, updating species
268 ranges (Hosking et al., 2000), and support studies of ecophysiology and occupancy across space and
269 time (Guerin et al., 2012).

270 Additional plant tissue is collected from each species and stored in synthetic gauze bags with
271 a barcode linked to the voucher specimen. These bags are used to avoid contamination from foreign
272 plants (e.g. cotton). The bags are rapidly dried on silica granules, ensuring they can be used for
273 genetic or isotopic analysis.

274 Soil samples (approximately 500 g) are taken from each 10 cm depth from the soil pit and
275 sub-sites, and these are barcode-labelled, air dried and archived in a dedicated facility.

276

277 **2.7 Validating remotely sensed products**

278 To enhance application of collected data to the validation of remotely sensed products, plots
279 are marked out with sub-metre DGPS for spatial accuracy and where possible aligned to a locally
280 accepted map grid (e.g. Map Grid of Australia). This enables the plot to be accurately matched to
281 pixels from remotely sensed imagery. Locating plots in homogeneous areas increases the likelihood
282 that the entire plot falls within a single remote sensing-derived mapping unit.

283 Cover information validates products from mid-resolution satellite imagery. Our point-
284 intercepts are able to be converted to either opaque canopy cover or foliage projected cover, making
285 the data useful for both ecological and imagery validation purposes. Because cover can be
286 summarised at different levels, from species to fractional cover (Scarth et al., 2015) , multiple
287 applications are possible, for example validation of tree cover interpretation from imagery (Bastin et
288 al., 2017).

289 The LAI2200 instrument (LiCor, Nebraska, USA) is used to collect and calculate Leaf Area
290 Index (LAI) data. This information can be used to validate international LAI products (Schaefer et
291 al., 2015), and to assist with the calibration between LAI and foliage projected cover derived from
292 remotely sensed products.

293 Structural information collected, including basal area determined using the basal wedge and
294 photo-points, along with growth-form and vegetation height data from point-intercepts, is useful for
295 validating satellite, airborne and terrestrial LIDAR systems.

296

297 **2.8 Data availability**

298 Data from the program are collected directly on an Android tablet and sent to a database when the
299 field officers have mobile phone coverage (Tokmakoff et al., 2016). Data are subsequently
300 combined with confirmed species identifications received after samples have been submitted to a
301 relevant herbarium, and the combined dataset is curated in preparation for publication. As sites are
302 finalised, they are identified as ready to publish and pushed to TERN's AEKOS data delivery
303 portal. During this process, the location of threatened or highly collectable species is de-natured

304 (Lowe et al., 2017). The data are then made freely available on the web portal for discovery,
305 download and re-use (Turner et al., 2017) using a Creative Commons (CC BY 4.0) by attribution
306 licence, or via the R package *ausplotsR*. (Guerin et al., 2019).

307

308 **3 Discussion of methodology and applications**

309 The standardised, quantitative surveillance monitoring method and innovative workflow we
310 outline can be employed across jurisdictional borders, allowing the measurement of diverse
311 environments at continental and global scales to answer questions would be difficult to address
312 using disparate datasets. Streamlined data collection and management ensure rapid delivery to end-
313 users and help minimise error (Box 1 in Appendix S3). The archiving of samples means that data
314 and results can be verified downstream, allowing resilience to nomenclatural change and innovative
315 future re-use of samples, for example bio-discovery (Lemetre et al., 2017).

316 Our approach is multi-disciplinary, collecting data relevant at multiple levels of ecological
317 analysis from population genetics to remote sensing. By collecting these measures at the same plot
318 using consistent methods, interactions among patch-level variables can be investigated.

319 The photo-points module is innovative in allowing traditional photo-point based change
320 analysis whilst enabling three-dimensional computer vision analysis. Technology is also embraced
321 in the collection of LAI data, using a DGPS to mark out plots and the electronic workflow from
322 data collection to publication (Tokmakoff et al., 2016).

323

324 **3.1 Infrastructure stimulating ecological research**

325 **3.1.1 Leaf samples**

326 Genomic sequencing technologies now provide cost-effective information on species
327 identification (DNA barcoding) and population genetic structure that allows rapid species
328 identification, the detection of cryptic species and identification of regions of high genetic diversity,
329 all of which are useful in a conservation context. The archiving of plant tissue samples ensures
330 material will be available even if the populations do not persist. Access to samples facilitates work
331 by independent researchers that may otherwise be impeded by the cost of sample collection from
332 remote locations. Leaf samples have been incorporated in a number of studies (e.g. Christmas et
333 al., 2017). Leaf samples are also available for isotope analysis and the study of leaf chemical
334 components such as the study of Dong et al. (2017) where these samples were used to demonstrate
335 that Leaf Mass per unit area increases with aridity.

336

337 **3.1.2 Soils**

338 Investigation of soils has typically focused on agrarian zones, meaning soil characteristics for
339 rangelands have largely been interpolated from sparse data, with this being particularly so in
340 Australia. In addition to basic characterisation, the method archives soils for future analysis (e.g.
341 DNA metabarcoding and chemical analyses). Soil surface samples are collected, from which
342 biological activity can be quantified and related to soil parameters. These samples facilitate research
343 on soil–vegetation interactions, typically conducted in local research projects. For example,
344 Lemetre et al. (2017) analysed these soil samples and reported that turnover in bacterial
345 biosynthetic composition followed a latitudinal pattern but did not appear to be driven by changes
346 in major vegetation type, a finding that directs approaches to future sampling of soils for natural
347 product discovery.

348

349 **3.1.3 Floristics and vegetation**

350 Vegetation data collection has been designed for multiple purposes. For example, standard
351 community ecology analytics such as ordination of vegetation and environmental variables can
352 provide insight into spatial patterning of species composition (Fig 2d) and its drivers. The data also
353 enable tracking of composition and cover dynamics with high reliability, enabling practical
354 outcomes like reporting on responses to disturbance or grazing impacts. The collection techniques
355 also provide a useful inventory, providing information on distribution and abundance of species,
356 with management applications such as providing information on the distribution and abundance of
357 problematic woody weed species.

358 The analysis opportunities for vegetation data from this program have been identified in more
359 detail in Guerin et al. (2017), including assessment of cover and species dominance analysis. The
360 future opportunities enabled by the multi-disciplinary method described here are also articulated.
361 Vegetation classification studies have also been conducted using the dataset (Baruch et al., 2018).

362

363 **3.1.4 Validating remote sensing products**

364 The method provides information useful for validating remotely sensed image products at
365 multiple scales, such as vegetation and soil products derived from mid-resolution satellite imagery.
366 By recording the shortest distance to another vegetation type, plots can represent a bigger spatial
367 footprint and be useful for validating lower spatial, but higher temporal, resolution imagery. The
368 data have further potential to validate high-resolution spatial and spectral image products, as well as
369 radar and LIDAR imagery.

370 Growth-form and cover data from this method were compared by Bastin et al. (2017) to
371 values obtained from visual estimates of very high resolution imagery over the same sites. This
372 information was then used to quantify observer estimate errors and errors between different
373 observers for this study that quantified the amount of forest occurring in dryland biomes globally.
374 Bastin et al. identified that previous estimates of dryland forest cover were between 40 and 47%
375 lower than their study indicates, leading to an increase of around 9% to estimates of forest cover
376 globally compared to previous knowledge.

377 **4 Conclusion**

378 We present a surveillance monitoring method for rangeland ecosystems developed in
379 Australia but applicable to global context. The method is now implemented as part of the
380 surveillance monitoring program of TERN, under the guiding principles of widespread consultation,
381 continual adaptation and coverage of variables/attributes relevant to multiple disciplines, to meet
382 the needs of a diverse ecosystem science community (Pereira et al., 2013). The method is
383 standardised, quantitative and modular, providing robust baselines and clear protocols for sample
384 and data management. The method embraces new technologies alongside more traditional
385 techniques. Sample archiving ensures continued utility of collected data and enables subsequent
386 analysis. A streamlined and accurate dataflow enables rapid open access data provision. The
387 method has had proven application to analyse rangelands systems globally (Bastin et al., 2017), and
388 many components are suitable for other environments. In future, we anticipate adding additional
389 variables on other environmental parameters including fauna sampling, for which protocols have
390 been prepared and are undergoing consultation, to be implemented as resources permit.

391

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398

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400 BS, JF, and AJL developed the concept, BS, JF, E JL, GW, SCR, GG and SVL contributed to field
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 402 AJL contributed to obtaining funding, BS, JF drafted the manuscript, all authors contributed to
 403 method critique, sections of text for the manuscript and contributed critically to the drafts and gave
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405

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Figures

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631 **Figure 1.** Extent of rangelands: (a) globally and (b) within Australia (White et al. 2012b).

632

633 **Figure 2.** Monitoring plot schemas and examples of information recorded in the point-intercept
634 module. (A) Plot layout and locations of soil sampling, basal wedge sweeps and point-intercept
635 transects; (B)–(D) point-intercept data on: (B) substrate; (C) growth-forms; (D) species (White et al.
636 2012a).

637

638 **Figure 3.** Photo-point panoramas. (A) Configuration: three photo-points are established in an
639 equilateral triangle with 2.5 m sides around the plot centre. Photographs are taken in a 360°
640 panorama; (B) Height setup: central dropper and mounted camera lens at 1.3 m; (C) Dealing with
641 topography (White et al. 2012a).

Tables

642 **Table 1.** Modules in the AusPlots rangelands monitoring method.

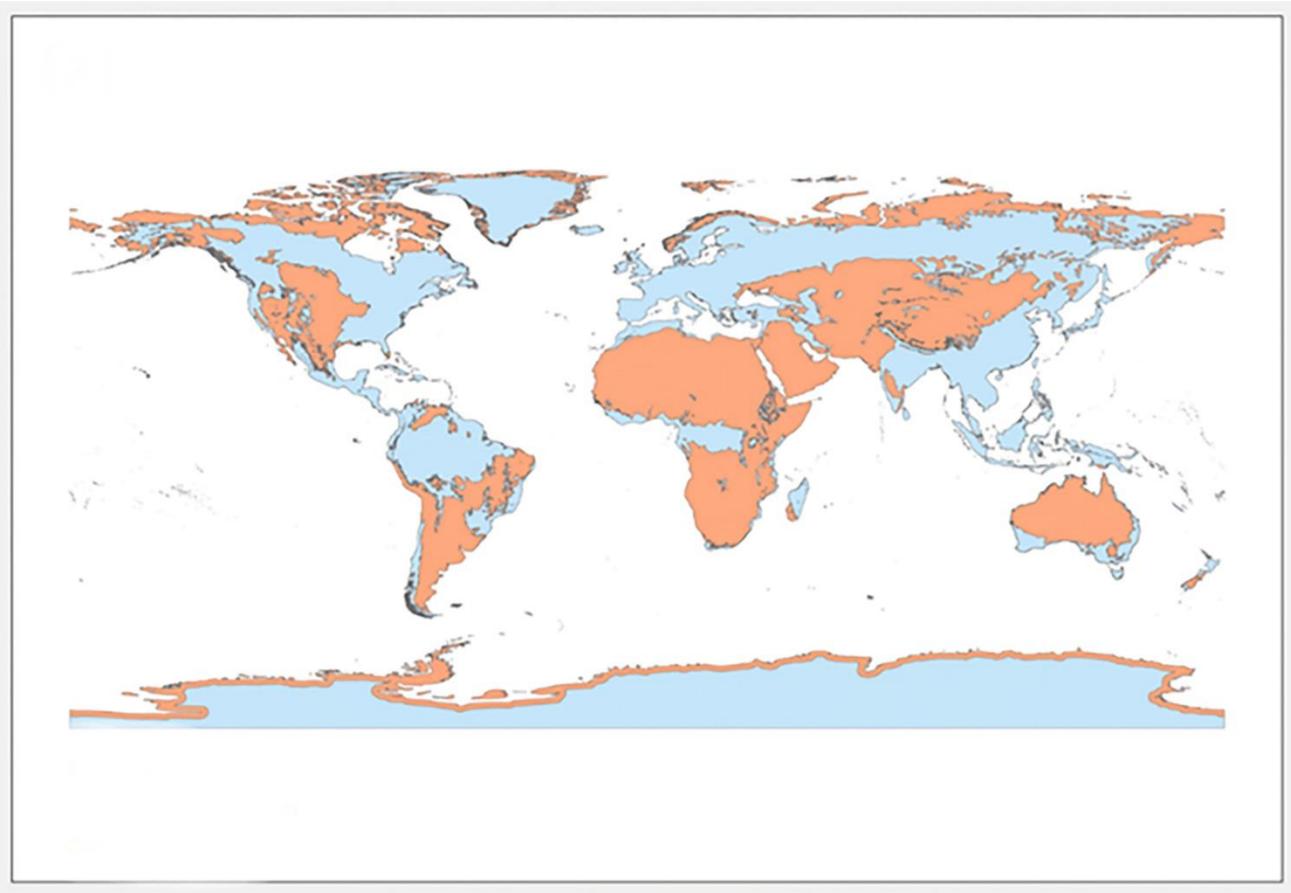
Module	Protocol	Time (mins)	Application
Plot layout	Accurate layout using DGPS; installation of permanent markers.	30	Accurate relocation; remote sensing validation
<u>Vegetation</u>			
Photo-panoramas	Collection of 360° photographs from three points	20	Computer vision analysis, point clouds and measures of basal area
Vouchering	Collection of vascular plant species	60-120	Taxonomy; spatial/temporal analysis of presence—absence
Tissue samples	Collection of single tissue samples from vascular plants (four from dominant species)	30-60	Genetic/isotopic analysis
Point-intercept	Collection of species, height, phenology, growth-form, senescence at 1010 points	180-360	Change in relative abundance, cover and structure; remote sensing validation
Basal area	Collection by species using basal wedge at nine points	20	Convertible to biomass
Structural summary	Recording of three dominant species in each of three strata (upper, mid ground)	5	Community descriptions
Leaf Area Index	Collection of at least 50 evenly spaced readings with the LiCor LAI 2200 LAI meter	20	Ecophysiological modelling; remote sensing validation
<u>Soils and Landscapes</u>			
Plot description	Record location, substrate, microtopography, erosion/disturbance	10	Assessment of characteristics/impact of disturbance
Soil pit characterisation	Collection of soil samples/data at 10 cm increments or identifiable horizons to 1 m	60-120	Characterisation and classification. Correlate with vegetation
Sub-site characterisation	Collection of nine samples in differing microhabitats at 0-10, 10-20 and 20-30 cm	60-90	Soil variability across plot
Bulk density	Collection of three measures at the soil pit at 0-10, 10-20 and 20-30 cm	60	Conversion to volumetric measures
Soil metagenomics	Collection of nine samples	30	Identify biota

644 **Table 2:** Co-location of sites with other projects

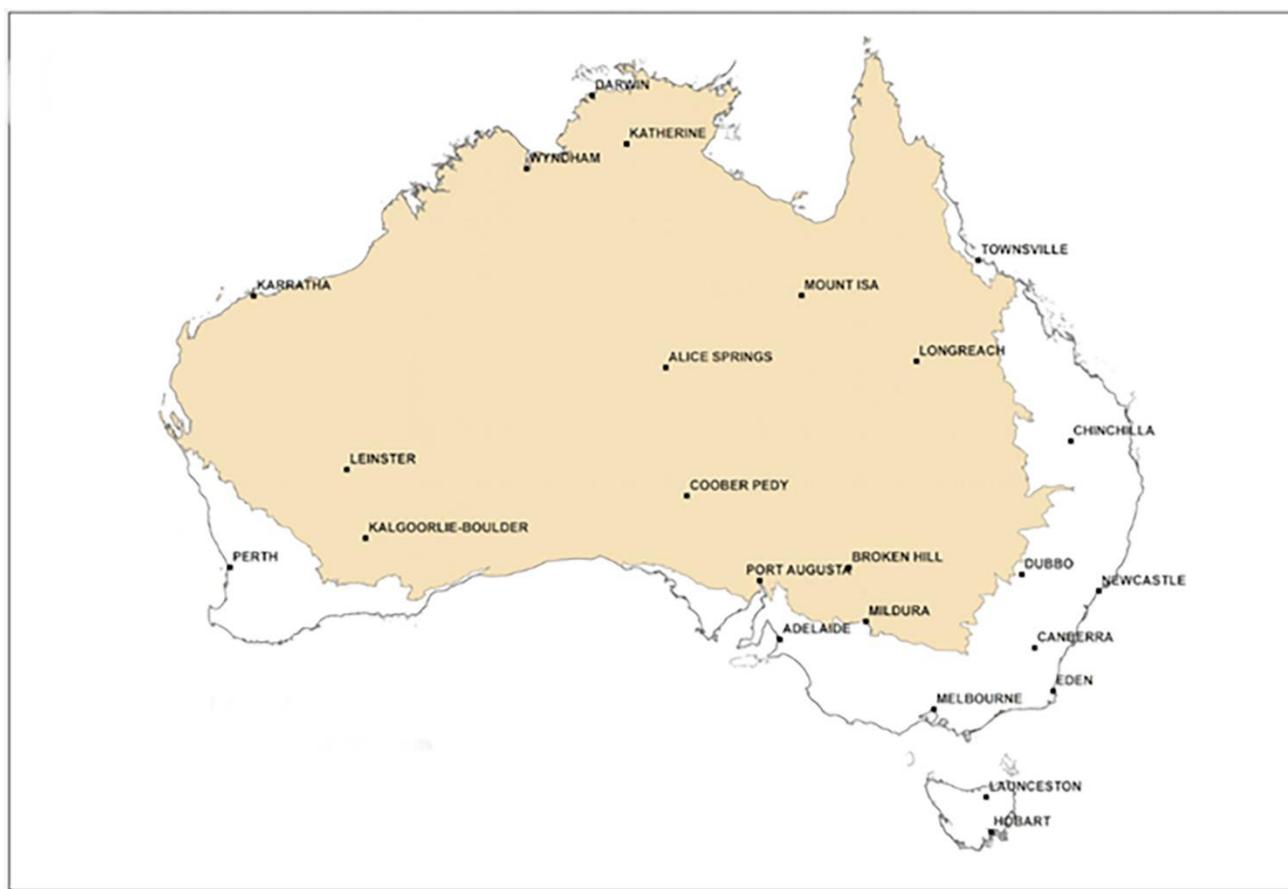
645

Project Type	Number of co-located projects	Number of co-located Plots
National Environmental Research Infrastructure - Process	7	40
National Environmental Research Infrastructure - Surveillance	4	43
Jurisdictional Process	1	9
Jurisdictional Surveillance	10	79
Non-Government Organisation	1	11
Totals	23	182

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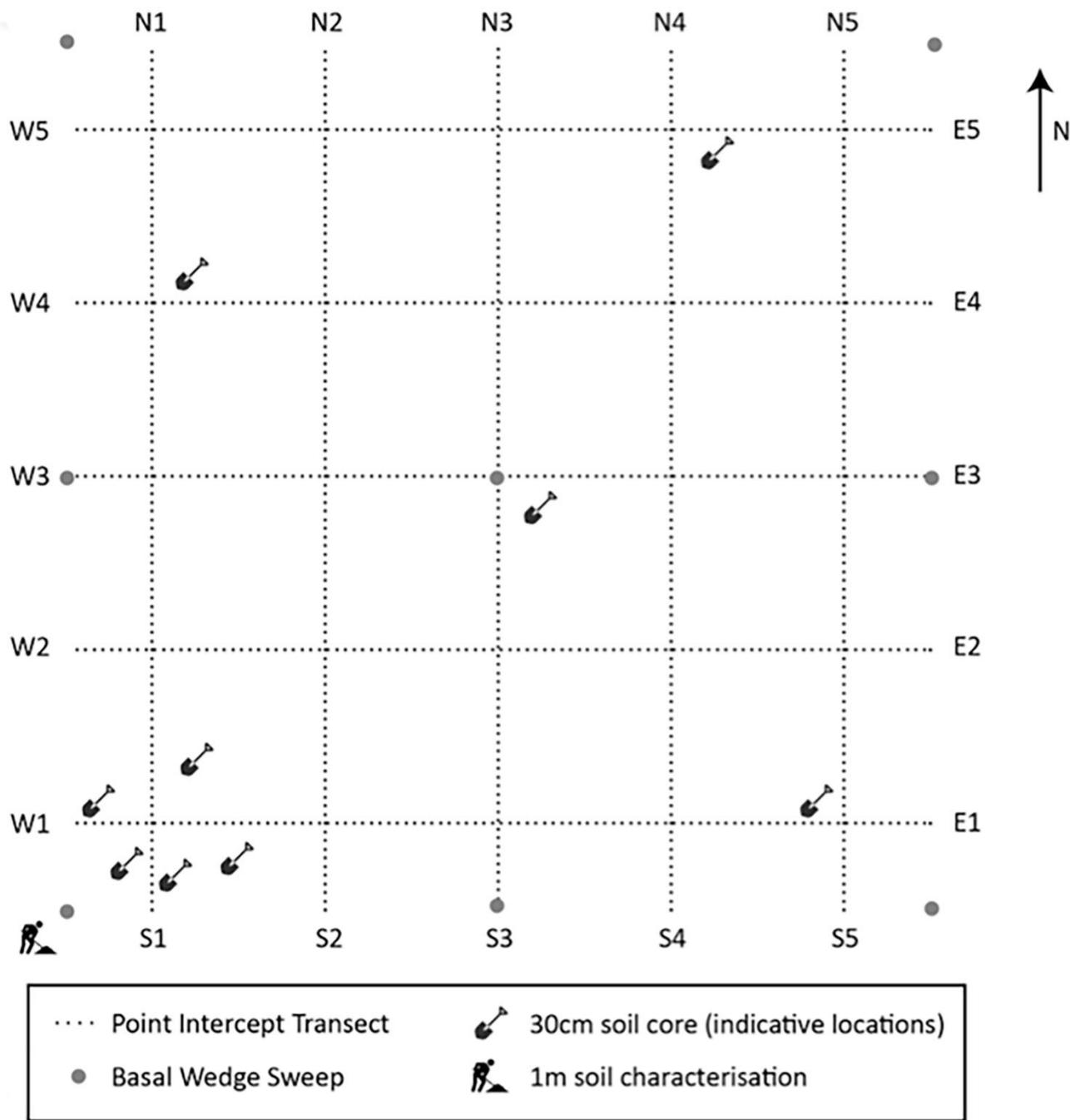
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652 **Figure 1.** Extent of rangelands: (a) globally and (b) within Australia (White et al. 2012b).

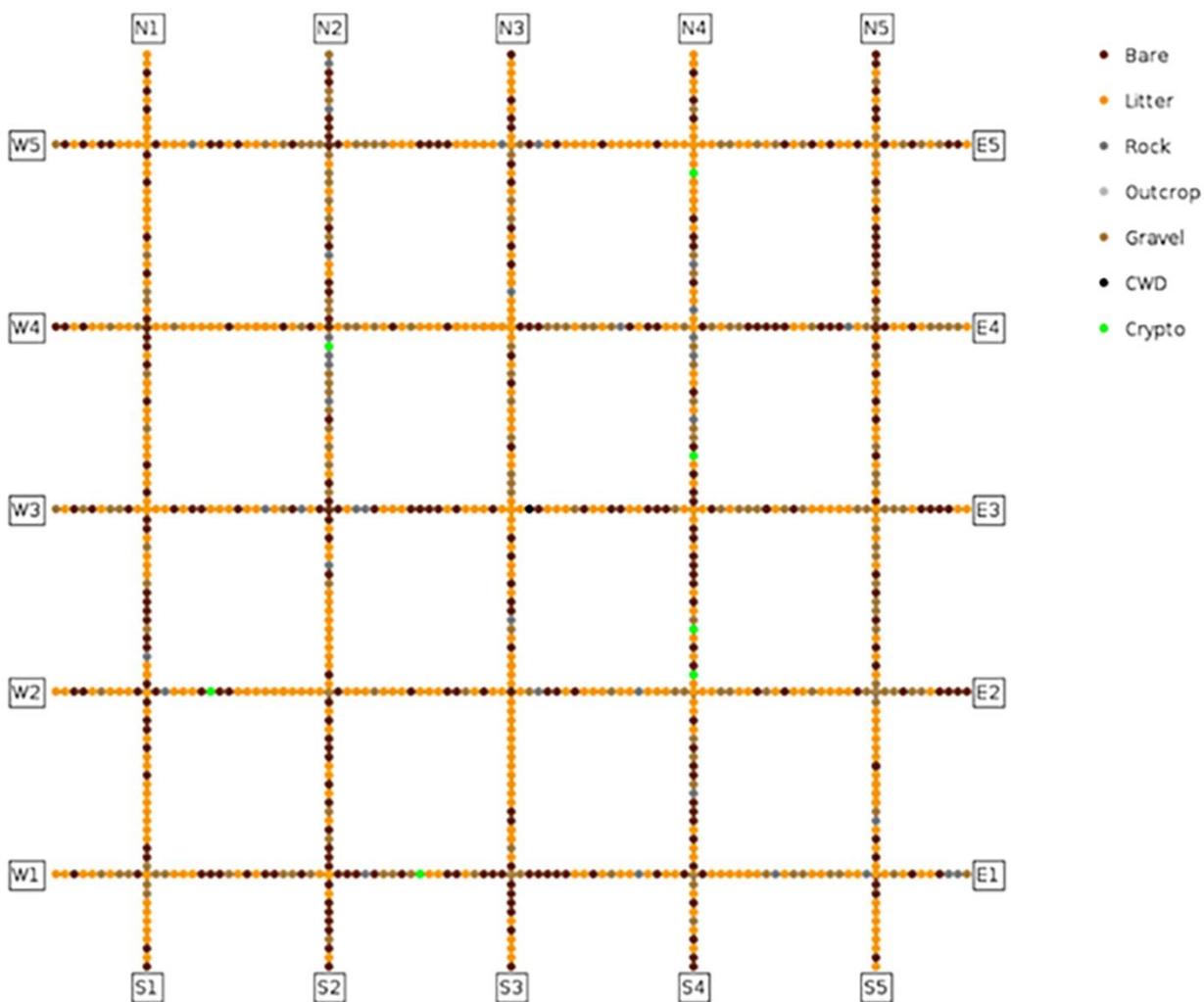
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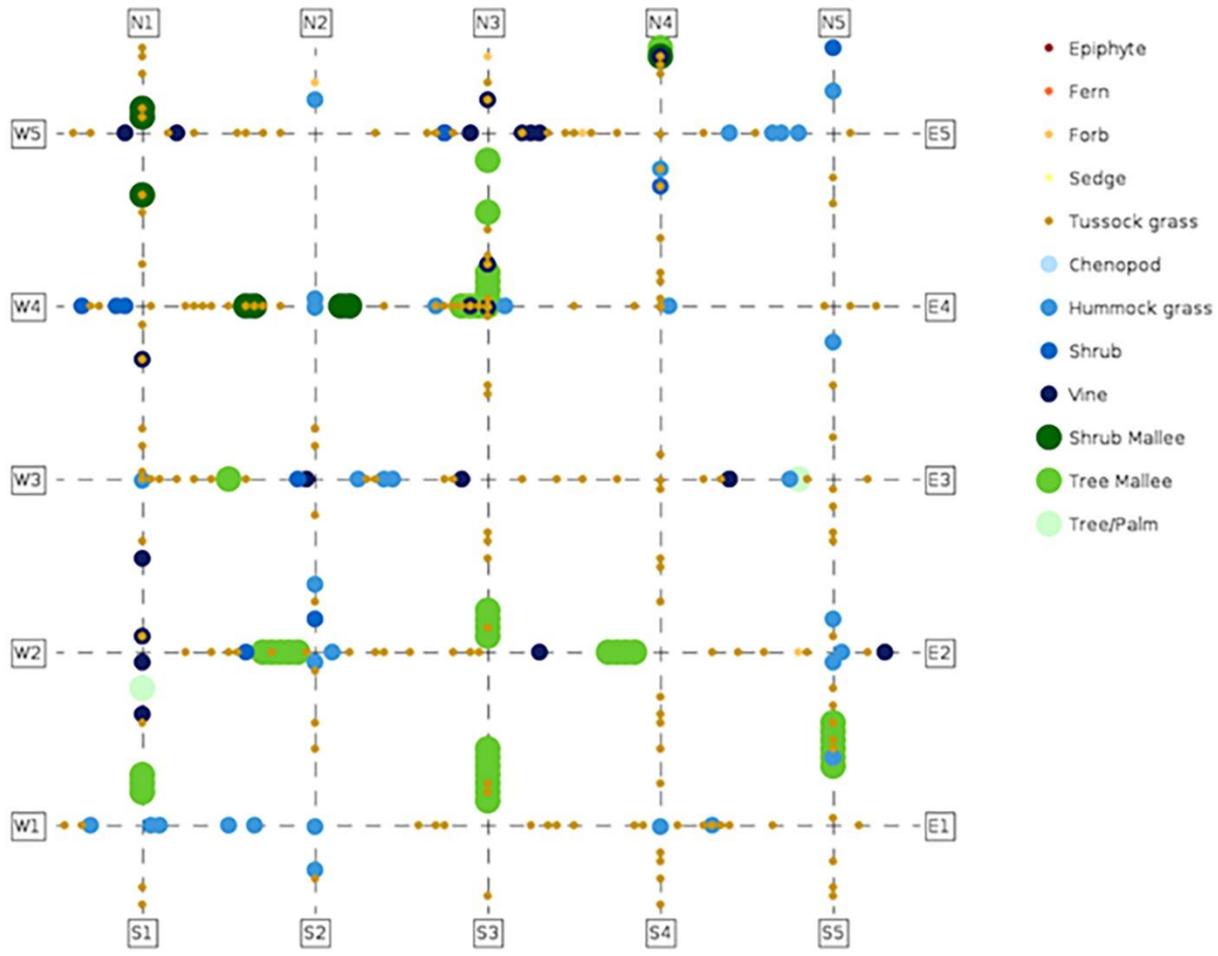


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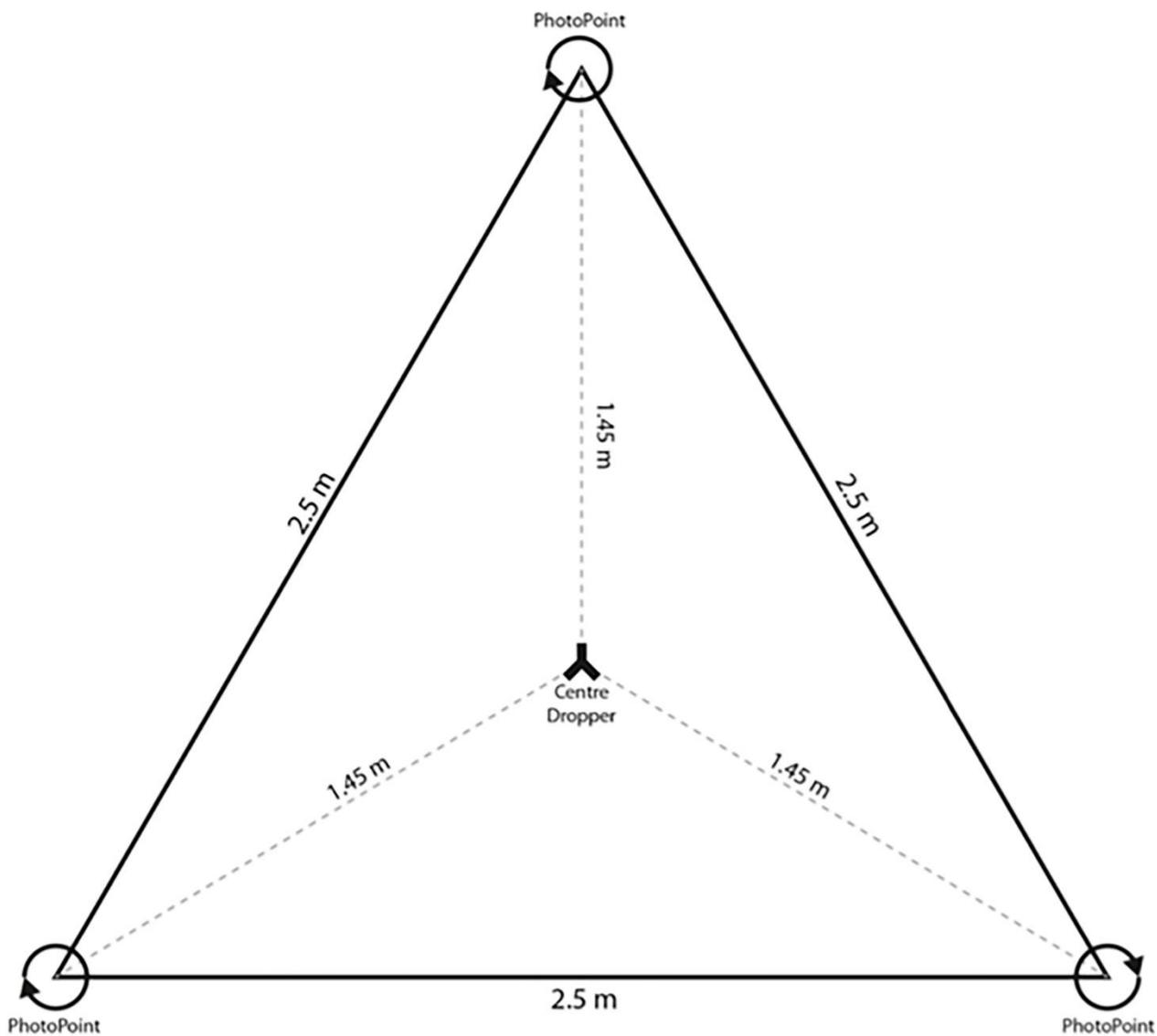


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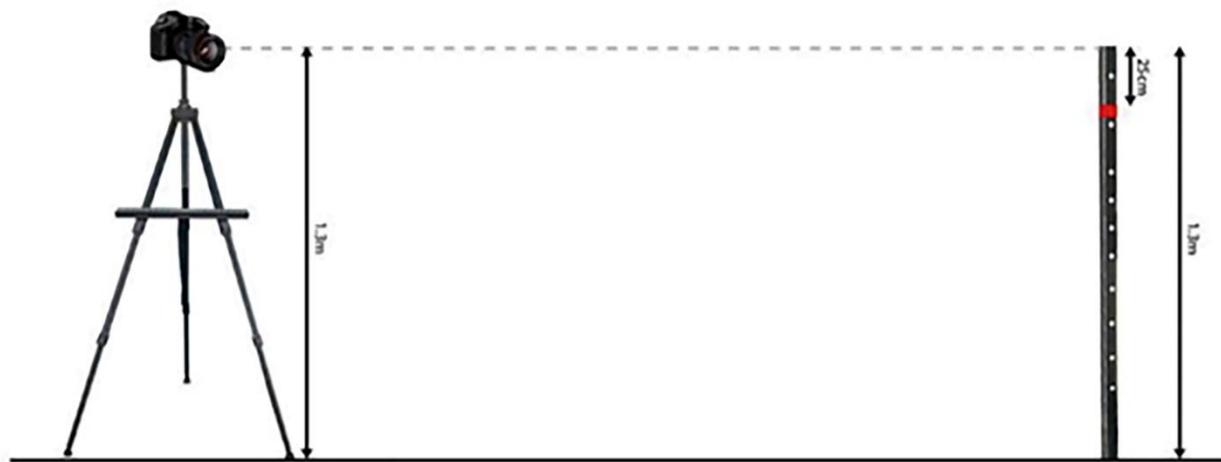
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665 **Figure 2.** Monitoring plot schemas and examples of information recorded in the point-intercept
 666 module. (A) Plot layout and locations of soil sampling, basal wedge sweeps and point-intercept
 667 transects; (B)–(D) point-intercept data on: (B) substrate; (C) growth-forms; (D) species (White et al.
 668 2012a).

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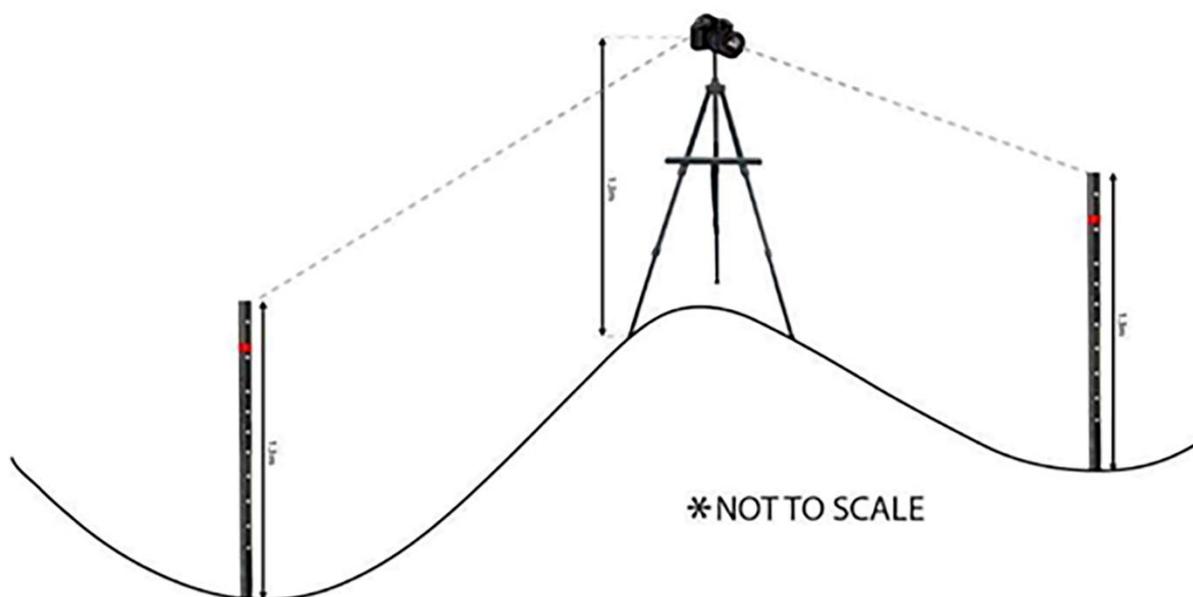
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678 **Figure 3.** Photo-point panoramas. (A) Configuration: three photo-points are established in an
679 equilateral triangle with 2.5 m sides around the plot centre. Photographs are taken in a 360°
680 panorama; (B) Height setup: central dropper and mounted camera lens at 1.3 m; (C) Dealing with
681 topography (White et al. 2012a).

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