Title: Patterns of pan-genome occupancy and gene co-expression under water-deficit in *Brachypodium distachyon*.

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ABSTRACT

Natural populations are characterized by abundant genetic diversity driven by a range of different types of mutation. The tractability of sequencing complete genomes has allowed new insights into the variable composition of genomes, summarized as a species pan-genome. These analyses demonstrate that many genes are absent from the first reference genomes, whose analysis dominated the initial years of the genomic era. Our field now turns towards understanding the functional consequence of these highly variable genomes. Here, we analyzed weighted gene co-expression networks from leaf transcriptome data for drought response in the purple false brome *Brachypodium distachyon* and the differential expression of genes putatively involved in adaptation to this stressor. We specifically asked whether genes with variable "occupancy" in the pan-genome – genes which are either present in all studied genotypes or missing in some genotypes – show different distributions among co-expression modules. Co-expression analysis united genes expressed in drought-stressed plants into nine modules covering 72 hub genes (87 hub isoforms), and genes expressed under controlled water conditions into 13 modules, covering 190 hub genes (251 hub isoforms). We find that low occupancy pan-genes are under-represented among several modules, while other modules are over-enriched for low-occupancy pan-genes. We also provide new insight into the regulation of drought response in *B. distachyon*, specifically identifying one module with an apparent role in primary metabolism that is strongly responsive to drought. Our work shows the power of integrating pan-genomic analysis with transcriptomic data using factorial experiments to understand the functional genomics of environmental response.

INTRODUCTION

Soil water availability is a critical factor determining plant growth, development, and reproduction (Bohnert, Nelson, & Jensen, 1995). Plants are able to cope with and acclimate to a range of soil water contents through the reprogramming of their physiology, growth, and development over time scales ranging from hours to seasons (Chaves, Maroco, & Pereira, 2003). Many of these acclimation strategies arise from altered transcriptional profiles (Fisher et al., 2016; Miao, Han et al., 2017). Drought-responsive gene regulatory pathways have been investigated extensively in model plant systems such as *Arabidopsis thaliana*, maize, and rice (Borah et al., 2017; Hayano-Kanashiro et al., 2009; Janiak, Kwa, & Szarejko, 2015; Nakashima, Ito, & Yamaguchi-Shinozaki, 2009; Nakashima, Yamaguchi-Shinozaki, & Shinozaki, 2014). A clear emerging theme, however, is that diverse species and varieties of plants exhibit diverse stress response mechanisms (Des Marais et al., 2012; Juenger, 2013; Pinheiro & Chaves, 2011), often controlled by complex regulatory networks. Understanding the genetic control of this phenotypic diversity is a priority for understanding the response of natural populations to climate change, and for designing resilient crop species (Benfey & Mitchell-Olds, 2008).

Recent studies have brought attention to the remarkable variation in gene content among plant populations (Alonge et al., 2020; Gao et al., 2019; Gordon et al., 2017; Haberer et al., 2020), reflected in a species' pan-genome. A pan-genome refers to the genomic content of a species as a whole, rather than the composition of a single, reference, individual genotype (Koonin & Wolf, 2008). In practice, pan-genomes are estimated by deeply resequencing the genomes of a diversity panel of genotypes, often using a reference genome to aid in final assembly and annotation (Lei et al., 2021). In the diploid model grass *Brachypodium distachyon*, genomic analysis of 56 inbred natural "accessions" revealed that the total pan-genome of the species comprised nearly twice the number of genes in any single accession (Gordon et al., 2017). Remarkably, only 73% of genes in a given accession are found in at least 95% of the other accessions (Gordon et al., 2017) – so-called "core genes" (universal or nearly universal genes; Koonin & Wolf, 2008) and "soft-core genes" (found in at least 95% of accessions; Kaas et al. 2012) – suggesting that a large number of genes are unique to subsets of accessions or even to individual accessions. The list of core genes in *B. distachyon* is enriched for annotations associated with essential cellular processes such as primary metabolism. Lower-occupancy genes, or "shell genes," are found in 5-94% of accessions and their annotations are enriched for many processes related to environmental response, including disease resistance. Similar patterns have been observed in the pan-genomes of *Arabidopsis thaliana*, barley, sunflower, and an ever-growing number of additional plant species (Bayer et al., 2020; Contreras-Moreira et al., 2017; Hübner et al., 2019). The DNA sequences of core genes bear the hallmark of strong purifying

73 selection and are typically expressed at a higher level and in more tissues as compared to shell genes.
74 Shell genes may be the result of gene duplications or deletions in the ancestor of a subset of studied
75 genotypes and, indeed, the vast majority of shell genes in *B. distachyon* appear to be functional, as
76 homologs are found in other species' genomes (Gordon et al., 2017).

77 The preceding observations raise the intriguing possibility that shell genes may represent segregating
78 variation that could be shaped by natural selection and thereby facilitate local adaptation or adaptive
79 responses to a variable environment. Multiple studies in *Arabidopsis thaliana* demonstrate the role of
80 segregating functional gene copies – effectively large-effect mutations – in shaping whole-plant response
81 to the abiotic environment (Monroe et al., 2016, 2018). The phenotypic effect size of a mutation can
82 determine the likelihood that the mutant will become fixed in a population, with large-effect mutations
83 more likely than not to confer deleterious phenotypes that may be removed from populations by natural
84 selection (Fisher, 1930). The observation that any two accessions of *B. distachyon* likely differ in the
85 presence or absence of hundreds of functional gene copies begs the question as to how potentially
86 function-changing gene deletions escape the purging effects of purifying selection. Pan-genomics requires
87 that we reconceptualize how we interpret "gene loss" as we move beyond a reference-genome view of
88 genome function. Genes identified as "missing" from a subset of accessions might represent deletions of
89 genes whose function was no longer constrained by purifying selection in some novel environment,
90 duplicated genes that originated in a subset of accessions and thus are absent in other accessions, or
91 paralogs that were both present in a common ancestor and then reciprocally lost in subsets of accessions.

92 The pleiotropic effect of a mutation can be affected by the number of genes with which it interacts
93 (Jeong et al., 2001); if a gene has relatively few interacting partners then its presence or absence in a
94 particular accession may have a small fitness effect and thus be maintained in populations. Similarly, the
95 efficacy of selection to purge deleterious alleles may be reduced if a gene is only expressed in a subset of
96 environments experienced by a species (Paaby & Rockman, 2014). In the context of pan-genomes, we
97 consider gene presence/absence polymorphisms as mutations, and we explore functional gene turnover by
98 testing the hypothesis that shell genes and core genes differ in their topological positions in
99 environmentally responsive gene co-expression networks.

100 Gene co-expression networks are widely used to interpret functional genomic data by assessing
101 patterns of correlation among genes via a threshold that assigns a connection weight to each gene pair
102 (Langfelder & Horvath, 2008; Zhang & Horvath, 2005). Sets of genes with similar expression profiles are
103 assigned to modules by applying graph clustering algorithms (Mao et al., 2009). Genes, or "nodes," in
104 such networks show considerable variation in the extent to which their expression co-varies with other

genes. As such, co-expression networks are generally considered "scale-free" in the sense that few nodes have many neighboring nodes while many nodes have few neighboring nodes (Guelzim et al., 2002). Modules are often comprised of genes with similar functions (Stuart et al., 2003; Wolfe, Kohane, & Butte, 2005). High connectivity "hub" genes that show a large number of interactions with other genes within a weighted co-expression network are candidates for key players in regulating cellular processes (Albert, Jeong, & Barabási, 2000; Carlson et al., 2006; Dong & Horvath, 2007). As such, hub genes might be expected to encode essential cellular functions and thus show pleiotropic effects when mutated or deleted. By contrast, genes with fewer close co-expression relationships are often situated on the periphery of networks and might, therefore, exhibit fewer pleiotropic effects when missing or mutated (Des Marais et al., 2017a; Masalia, Bewick, & Burke, 2017; Porth et al., 2014). In this context, we hypothesize that pan-genome core genes may be over-represented among co-expression network "hub-genes," as both appear to be involved in core cellular processes and may therefore show deleterious effects when deleted. Conversely, we hypothesize that pan-genome "shell genes" – whose patterns of expression and thereby phenotypic effects are more restricted and condition-specific – will be enriched among lowly connected (non-hub) genes in gene co-expression networks.

Here, we study the relationship between a plant's pan-genome and its gene co-expression networks using *Brachypodium distachyon. Brachypodium* is a small genus of the subfamily Pooideae (Poaceae) that contains ~20 species distributed worldwide (Catalán et al., 2016; Scholthof et al., 2018; Hasterok et al. 2022). The annual diploid species *B. distachyon* is a model for temperate cereals and biofuel grasses (Vogel et al. 2010; Mur et al. 2011; Catalán et al. 2014; Scholthof et al. 2018); a reference genome for one *B. distachyon* accession, Bd21 (IBI, 2010) is now complemented by 54 deeply resequenced natural accessions (Gordon et al., 2017). Recent studies demonstrate the utility of *B. distachyon* and its close congeners for elucidating the evolution and ecology of plant-abiotic interactions, focusing especially on responses to soil drying, aridity, and water use strategy (Des Marais & Juenger, 2016; Des Marais et al., 2017b; Monroe et al. 2021; Fisher et al., 2016; Handakumbura et al., 2019; Manzaneda et al., 2015, 2012; Martínez et al., 2018; Skalska et al., 2020; Verelst et al., 2013). In the present study, we first identify and characterize gene co-expression modules associated with response to soil drying. We then test the hypothesis that the occupancy of pan-genes – whether they are part of the shell or core gene sets of the pan-genome – is associated with their connectivity in the *B. distachyon* gene co-expression network. Our work demonstrates the dynamic nature of plant genomes and sets up future work on the functional consequences of diversity on the evolution of gene regulatory networks.

137 **MATERIALS AND METHODS**

138 **Plant material, experimental design, total-RNA extraction and 3' cDNA tag libraries preparation**

139 Sampling herein follows our earlier work documenting physiological and developmental response of
140 33 diploid natural accessions of *Brachypodium distachyon* (L.) P. Beauv. to soil drying (Figure 1; Table
141 S1) (Des Marais et al., 2017b). The sampled accessions were inbred for more than five generations (Filiz
142 et al., 2009; Vogel et al., 2006, 2009) and represent the geographic and ecological diversity of *B.*
143 *distachyon* across the Mediterranean region. Whole genome resequencing data is available for all studied
144 accessions (Gordon et al., 2017).

145 A total of 264 individual plants from the 33 accessions were grown under two greenhouse conditions,
146 restriction of water (drought, D) and well-watered (water, W). We sampled four biological replicates per
147 treatment and accession combinations [33 accessions x 4 replicates x 2 treatments (D and W)]. For a full
148 description of the growth and treatment conditions, please see Des Marais, Lasky, et al. (2017b). In short,
149 seeds were stratified at 6°C for 14d and then greenhouse-grown in 600 mL of Profile porous ceramic
150 rooting media (Profile Products) in Deepot D40H pots (650 mL; Stuewe & Sons). For the first 21d of
151 growth, all plants were watered to field capacity every other day. Daytime high temperatures ranged from
152 23°C to 28°C and night-time lows from 14°C to 18°C. On day 21 four treatment regimes were
153 implemented: Cool Wet, Cool Drought, Hot Wet, and Hot Drought, with Drought and Wet plants
154 spatially randomized within single blocks of Hot or Cool conditions. The hot treatment raised air
155 temperatures in the plant canopies by ~10°C. Because temperature was confounded with experimental
156 block in the design used for the current study, we did not include a temperature effect in any of the
157 statistical models used herein. Well-watered plants (hereafter "Water") were watered to field-capacity
158 every second day with fresh water, whereas drought plants (hereafter "Drought") were hand-watered daily
159 by pipette such that the soil water was reduced by 5% each day. The final soil water content of Drought
160 plants on day 11 of treatment – 32d post-germination – was 45% field capacity, which corresponds to a
161 decrease in water potential of 1.2 MPa as compared to field capacity in this growth media (Des Marais et
162 al., 2012).

163 For each plant, the two youngest, fully expanded leaves of the tallest tiller were excised with a razor
164 blade at the base of the lamina and flash-frozen on liquid nitrogen. Tissue was ground to a fine powder
165 under liquid nitrogen using a Mixer Mill MM 300 (Retsch GmbH). RNA was extracted using the Sigma
166 Spectrum Total Plant RNA kit, including on-column DNase treatment, following the manufacturer's
167 protocol, and quantified using a NanoDrop spectrophotometer (Thermo Scientific). We used a RNA-Seq

168 library protocol (3' cDNA tag libraries with fragment of 300-500 bp) for sequencing on the Illumina
169 HiSeq platform adapted from Meyer et al. (2011). This Tag-Seq method yields only one sequence per
170 expressed transcript in the RNA pool, allowing for higher sequencing coverage per gene as a function of
171 total sequencing effort (Tandonnet & Torres, 2017).

172 **Pre-processing of sequences, quantifying transcript abundances, normalizing procedures, and**
173 **controlling batch effects**

174 Sequencing was carried out using an Illumina HiSeq2500 platform (100 bp Single-End, SE,
175 sequencing). Quality control of SE reads was performed with FastQC software. Adapters and low quality
176 reads were removed and filtered with Trimmomatic-0.32 (Bolger, Lohse, & Usadel, 2014). Total numbers
177 of raw and filtered SE reads for each accession and treatment are shown in Table S2. Quantifying the
178 abundances of transcripts from RNA-Seq data was done with Kallisto v0.43.1 (Bray et al., 2016). To
179 accommodate the library preparation and sequencing protocols (3' tag from fragments of 300-500 bp),
180 pseudoalignments of RNA-Seq data were carried out using as references 500 bp from the 3' tails of the *B.*
181 *distachyon*_314_v3.1 transcriptome (IBI 2010;
182 https://phytozome.jgi.doe.gov/pz/portal.html#!info?alias=Org_Bdistachyon). We applied estimated
183 average fragment lengths of 100 bp and standard deviations of fragment length of 20. Resulting numbers
184 of transcripts per million (TPM) were recorded.

185 Exploratory analysis of the data set and the subsequent filtering and normalization of transcripts
186 abundance between samples, and the *in silico* technical replicate steps (bootstrap values computed with
187 Kallisto), were conducted with the Sleuth package (Pimentel et al., 2017). A total of 16,386 targets
188 (transcripts/isoforms) were recovered after the normalizing and filtering step using Sleuth. This program
189 was also used for batch-correction of data and of differentially expressed (DE) genes. To account for
190 library preparation batch effects, date of library preparation was included as a covariate with condition
191 variable in the full model (Table S2).

192 **Co-expression network analysis of normalized transcripts abundance**

193 Co-expression networks for the Drought and Water (control) data sets were carried out using the
194 transcripts per million (TPM) estimates and the R package WGCNA (Langfelder and Horvath 2008). We
195 analyzed 16,386 transcripts that were filtered and normalized for 127 Drought and 124 Water individual
196 samples (individual plants). After the removal of putative outliers, we retained 121 Drought and 108
197 Water samples that were used for network construction.

198 Identical parameters were used for the Drought and the Water data sets to construct their respective
199 co-expression networks. The *BlockwiseModules* function was used to perform automatic network
200 construction and module detection on the large expression data set of transcripts. Parameters for co-
201 expression network construction were fitted checking different values. We chose the Pearson correlation
202 and unsigned network type, soft thresholding power 6 (high scale free, R^2>0.85), a minimum module size
203 of 30, and a medium sensitivity (deepSplit = 2) for the cluster splitting. The topological overlap matrix
204 (TOM) was generated using the TOMtype unsigned approach. Module clustering was performed with
205 function *cutreeDynamic* and the Partitioning Around Medoids (PAM) option activated. Module merging
206 was conducted with mergeCutHeight set to 0.30. K_{diff} was calculated using WGCNA to estimate the
207 relationship between connectivity among genes within *vs* between co-expression modules.

208 Both isoform and gene counts were calculated. Isoform counts included all transcripts identified (e.g.,
209 Bradi1g1234.1; Bradi1g1234.2; Bradi1g1234.3) and gene counts only included different genes expressed,
210 thus different isoforms from the same gene were reduced to a single gene count in each case (e.g.,
211 Bradi1g12345.1 and Bradi1g12345.2 are two isoforms of one gene, Bradi1g12345).

212 **Detection of highly connected nodes (hub genes/ isoforms) within co-expression networks**

213 Three representative descriptors of modules, module eigengene (ME), intramodular connectivity
214 (k_{IM}), and eigengene-based connectivity (k_{ME}; or its equivalent module membership, MM) were
215 calculated using the WGCNA package. Briefly, ME is defined as the first principal component of a given
216 module and is often considered to represent the gene expression profiles within the module. k_{IM} measures
217 how connected, or co-expressed, a given gene is with respect to the genes of a particular module. Thus,
218 intra-modular connectivity is also the connectivity in the subnetwork defined by the module. MM is the
219 correlation of gene expression profile with the module eigengene (ME) of a given module. MM values
220 close to 1 or -1 indicate genes highly connected to the module. The sign of MM indicates a positive or a
221 negative relationship between a gene and the eigengene of the module (Langfelder & Horvath, 2010).
222 Genes with absolute MM value over 0.9 were considered "hub genes." Correlations between MM values
223 transformed by a power of $\beta = 6$ and k_{IM} values were also calculated.

224 **Pan-genome occupancy of clustered, hub, and differentially expressed genes across accessions**

225 Because the *B. distachyon* accessions studied herein comprise a subset of those included in the
226 original pan-genome (Gordon et al. 2017), we re-ran the clustering procedures used in our earlier analysis
227 with only the 33 accessions used here. We clustered CDS sequences from the annotated genomes of each

228 of the studied accessions to define core, soft-core, and shell genes with the software
229 GET_HOMOLOGUES-EST v03012018 (Contreras-Moreira et al., 2017) using the OMCL algorithm (-
230 M) and a stringent percent-sequence identity threshold (–S 98). The resulting pan-genome matrix was
231 interrogated to identify "core" genes observed in all 33 accessions, "soft-core" genes observed in 32 and
232 31, and "shell" genes observed in 30 or fewer accessions. Occupancy was defined as the number of
233 accessions that contain a particular gene model. We tested whether each module showed an excess or
234 deficit of shell genes as compared to genome averages of pan-gene occupancy using a Fisher's Exact
235 Test, as implemented in the R programming language (R Core Team, 2022).

236 **Enrichment and GO/KEGG annotation of clustered genes**

237 Gene ontology (GO) and the Kyoto Encyclopedia of Genes and Genomes (KEGG) annotations for
238 the *B. distachyon* 314 (Bd21 acccession) v.3.1 reference genome were retrieved
239 (http://phytozome.jgi.doe.gov/; IBI 2010). Gene lists were tested for functional enrichments with the
240 PANTHER (Protein ANalysis THrough Evolutionary Relationships) overrepresentation test
241 (http://www.pantherdb.org). The original *B. distachyon* 314 (Bd21 accession) v.3.1 gene ids were
242 converted to v.3.0 with help from Ensembl Plants (Howe et al., 2020) to match those in PANTHER16.0
243 (Mi et al., 2021). Tests were conducted on all genes and on both conditions -- Drought and Water --
244 applying the Fisher's Exact test with False Discovery Rate (FDR) multiple test correction. This analysis
245 was applied on different data sets: all genes, and pan-genome core, soft-core, and shell genes for each co-
246 expressed module.

247 **Drought versus Watered modular structure preservation and comparison between consensus and**
248 **set-specific modules**

249 Permutation tests were performed to check for preservation of the module topology in the Drought
250 (discovery data) and the Water (test data) networks using both the approach of Langfelder et al.
251 (Langfelder, Luo, Oldham, & Horvath, 2011) as well as the modulePreservation function of the NetRep
252 (Ritchie et al. 2016) R package with null="all" (include all nodes) option for RNA-Seq data. All NetRep
253 test statistics (Module coherence, Average node contribution, Concordance of node contributions, Density
254 of correlation structure, Concordance of correlation structure, Average edge weight, and Concordance of
255 weighted degree) were evaluated through permutation analysis; therefore, a module was considered
256 preserved if all the statistics had a permutation test P-value < 0.01. Searching for modules that could play
257 a role in drought response, we focused on Drought modules that were unpreserved in the Water network
258 (P-value > 0.01 in at least 1 of the seven statistics presented in Ritchie et al. (2016). The Consensus

259 modules (Cons) and the respective relationships between Consensus and Drought (D) and Water (W)
260 modules were performed as described in Langfelder and Horvath (2008).

261 **Annotations of upstream DNA motifs in the co-expression modules**

262 Genes assigned to modules in the Drought and Water networks were further analyzed with the
263 objective of discovering DNA motifs putatively involved in their expression in each module. Motif
264 analysis was carried out using a protocol based on RSAT::Plants (Ksouri et al., 2021). This approach
265 allowed us to discover DNA motifs enriched in the promoter regions of co-expressed genes and to match
266 them to curated signatures of experimentally described transcription factors. First, -500 bp (upstream) to
267 +200 bp (downstream) sequences around the transcription start sites of the genes detected in each module
268 and 50 random negative controls from other gene promoter regions of equal size were extracted from the
269 *B. distachyon* Bd21 v3.0 (Ensembl Plants version 46) reference genome. Then, the peak-motifs (Thomas-
270 Chollier et al., 2012) option was used to discover enriched motifs applying a 2nd order Markov genomic
271 model, and GO enrichment was computed for them. The analyses generated a report with links to similar
272 curated motifs in the database footprintDB as scored with normalized correlation (Ncor) (Sebastian and
273 Contreras-Moreira 2014). For each module a highly supported DNA motif was selected according to
274 Ncor, e-value, and the number of sites (i.e., putative cis-regulatory elements, CREs) used to compile the
275 motif. The matrix-scan tool (Turatsinze et al., 2008), with a weight threshold set to 70% of the motif
276 length (60% for the small module D8), was used to scan the discovered motifs and to identify individual
277 genes within each module harboring putative CREs.

278 All the protein sequences predicted for each co-expression module were analyzed using iTAK (Plant
279 Transcription factor & Protein Kinase Identifier and Classifier) online (v1.6) (Zheng et al., 2016) to
280 annotate their respective transcription factors, transcriptional regulators and related protein kinases
281 (classification system by Lehti-Shiu and Shiu (2012)).

282 **Differentially expressed (DE) isoforms/genes**

283 In order to determine how many isoforms and genes were differentially expressed between the two
284 treatments (D vs W), the two data sets were analyzed through the sleuth_result function (Pimentel et al.,
285 2017). This function computes likelihood ratio tests (LRT) for null (no treatment effect) and alternative
286 (treatment effect) hypotheses, attending to the full and reduced fitted models. A significant q-value ≤ 1E-
287 6 threshold was fixed to detect DE isoforms. The 50 most differentially expressed genes (25 up-regulated

288 and 25 down-regulated) were classified based on fold-change of the average TPM values between the
289 drought and water treatments.

290 **Data availability**

291 The *Brachypodium distachyon* filtered RNA-Seq data were deposited in the ENA (European
292 Nucleotide Archive; https://www.ebi.ac.uk/ena) with the consecutive accession numbers from
293 ERR6133302 to ERR6133575. All the R scripts, supplementary Excel files, TPM counts and adjacency
294 matrices are available in the Github repository identified by the following doi: xxxxxxxxxx created using
295 Zenodo.

296

297 **RESULTS**

298 Our RNA-Seq dataset comprised 121 Drought and 108 Water samples, obtained from 33 inbred
299 accessions of *B. distachyon* (Figure 1). After the filtering and quality control steps, we identified 16,386
300 transcripts in our analyses comprising 4,941 isoforms of 3,789 differentially expressed (DE) genes
301 between the water and drought treatments (Supplementary files S1; S2). Of these genes, 2,808 were
302 upregulated (3,591 isoforms) and 980 downregulated (1,350 isoforms) in D vs W conditions. One gene
303 (Bradi1g09950) showed both up-regulated (Bradi1g09950.2) and down-regulated (Bradi1g09950.1)
304 isoforms under drought conditions (Supplementary file S2).

305 **Modular distribution of genes in the gene co-expression networks**

306 The 33 genetically diverse accessions, along with random experimental variance, provided a wealth
307 of expression variance that was leveraged to estimate major axes of variation – co-expression modules –
308 from the RNA-Seq data. The modular distribution of genes in the Drought and Water gene co-expression
309 networks showed differences both in the number and the size of the modules (Figures S1a,b and S2a,b;
310 Tables 1a,b and S3a,b). The Drought co-expression network comprised 9 modules (D1-D9) containing a
311 total of 5,020 isoforms (min = 38, max = 2,477 isoforms per module), corresponding to 4,006 genes (min
312 = 27, max = 1,986 genes per module). The largest D1 module contained 15.1% of the isoforms (16.1% of
313 the genes) whereas two modules (D2, D3) clustered 4-6% of the isoforms and genes each and the
314 remaining modules clustered ≤ 2% of isoforms and genes each (Fig. S1a; Table 1a). 11,366 isoforms
315 (69.4%; 8,313 genes, 67.4%) were not clustered in any Drought module using our criteria (gray or "zero"
316 D0 module) (Figs. S1a and S2a; Table 1a). The Water co-expression network showed 13 modules (W1-

317 W13) containing a total of 6,711 isoforms (min = 48, max = 1,866 isoforms per module), corresponding
318 to 5,407 genes (min = 40, max = 1,439 genes per module). The largest W1 contained 11.4% of the
319 isoforms (11.6% of the genes) whereas six modules (W2-W7) clustered over >2-6% of the isoforms and
320 genes and the remaining six modules ≤ 2% each (Fig. S1b; Table 1b). 9,675 isoforms (59.0%; 6,934
321 genes, 56.0%) were not clustered in any Water module (gray or "zero" W0 module; Figs. S1b and S2b;
322 Table 1b). Hereafter, co-expression modules identified in the Drought RNA-Seq dataset will be labeled
323 with the prefix "D" while those modules identified in the Water (control) dataset will have the "W"
324 prefix.

325 While the genes comprising co-expression modules likely represent functionally related genes,
326 modules are not discrete entities and many genes within a module are likely co-expressed with genes in
327 other modules. The largest Drought modules showed a positive (D1, D2, D3, D4, and D5) or slightly
328 negative (D6 and D7) mean k_{diff}, while the smallest modules had more negative values (D8 and D9)
329 (Table 1a). Negative k_{diff} values for a node indicate that connectivity out of the module is higher than
330 intra-modular connectivity. Similarly, the Water modules showed positive mean k_{diff} values in the largest
331 modules (W1, W3, W5, W6, W7, and W9) or slightly negative (W4), with the exception of one large and
332 one intermediate modules (W2 and W8) both having negative values similar to those of the smallest
333 modules (Table 1b). High positive linear correlations between MM values and k_{IM} values were recovered
334 in both Drought (Fig. S3a) and Water (Fig. S3b) networks, thus validating the criterion of high MM
335 (>0.9) for selecting hub genes. Collectively, these results suggest that modules of both Drought and Water
336 networks were statistically well-supported but also that many of their genes likely share transcriptional
337 activity with genes included in other modules.

338

339 **Preservation and correspondence of Drought and Water networks**

340 We tested the hypothesis that some co-expression modules are only observed under one of our two
341 treatment conditions using two approaches. First, a permutation test was performed using NetRep (Ritchie
342 et al., 2016) to test for the preservation of module topology in the Drought versus the Water networks,
343 defined as discovery and test data sets, respectively. This test comprises seven topological statistics on
344 each module and condition (drought vs water), quantifying the replication of the structural relationship
345 between nodes composing each module under the null hypothesis that the module of interest is not
346 preserved. All Drought and Water modules were topologically preserved according to the seven NetRep
347 statistics (permutation test P-values < 0.01). We further tested for correspondence between Drought (D)

348 and Water (W) set-specific and Drought-Water Consensus (Cons) co-expression modules using WGCNA
349 (Fig. 2a,b). Consensus modules are those shared by two or more networks (Langfelder & Horvath, 2007).
350 We found that several Drought specific modules were comprised of no or very few isoforms included in
351 the Consensus modules (Fig. 2a). Thus, the Drought modules D5 (green), D7 (black) and D9 (magenta)
352 did not show a clear correspondence with consensus modules. The seemingly conflicting results between
353 NetRep – which detected module preservation between Drought and Water modules – and WGCNA –
354 which identified three modules that were not preserved – reflected the different sensitivities of these two
355 approaches to changing co-expression relationships within modules. This suggested that all modules were
356 preserved in a broad sense between treatments, but that nodes within modules D5, D7, and D9 could have
357 slightly different co-expression relationships than those observed in the corresponding modules in the
358 Water treatment. We also found Water specific modules (Fig. 2b), W1 (turquoise), W6 (red), W8 (pink),
359 and W12 (tan), that did not overlap with the Consensus modules or overlapped only with the non-co-
360 expressed gray module. W1 and W6 overlapped with the gray consensus module, whereas the smaller
361 modules W8 and W12 did not overlap with any consensus module.

362 Seven modules in the Drought network and eleven in the Water network showed a significant GO
363 term enrichment (Table 2a,b; Supplementary file S3). Both networks shared modules enriched for
364 different biological processes. Potentially equivalent modules between the D and W networks were
365 inferred attending to the common consensus module with which they overlapped (Figure 2a,b) and their
366 GO enrichments (Fig. 2c). Thus, the D1 and W2 modules matched Cons1 and were GO-enriched in
367 nitrogen, amide, and peptide metabolic and biosynthetic processes, D2 and W10 matched Cons4 and were
368 enriched in the photosynthesis, D3 and W3 matched Cons3 and were enriched in processes of transport
369 and locations of compounds, D4 and W9 were enriched in nucleic acid metabolic processes, and D8 and
370 W13 in nitrogen, amide, and peptide biosynthetic and metabolic processes. However, some modules
371 showed enrichment in biological process unique to one or the other co-expression network. For example,
372 D5 is enriched in genes involved in protein folding, response to heat, temperature, and abiotic stimulus,
373 and W6 in genes predicted to be involved in cell wall organization or biogenesis. Collectively, these co-
374 expression analyses suggest that the detected modules represent groups of functionally related genes, and
375 that many functional relationships among genes were conserved in the Water and Drought networks.

376 **Regulatory motifs of genes in the Drought and Water modules**

377 We detected statistically over-represented sequence motifs upstream of the gene sequences in several
378 modules. These motifs represent putative *cis*-regulatory elements (CREs) located in proximal promoter
379 regions of co-expressed genes in the Drought (Table 3a; Fig. S4a) and Water (Table 3b; Fig. S4b)

380 networks. A generally low but variable proportion of genes harboring putative CREs in their proximal
381 promoters was detected in each module. The drought (Table 3a) and water (Table 3b) modules showed
382 between 3.5-54.2% and 0.1-23% of genes with the predicted CREs, respectively. Genes in the same
383 module shared a conserved regulatory architecture. For example, calmodulin-binding CREs were
384 enriched in the D4 (associated with nucleobase-containing compound, heterocycle and nucleic acid
385 metabolic processes GO terms) and W9 modules (also enriched for nucleobase-containing compound GO
386 processes, among others). We also observed enriched motifs in treatment-specific modules, like the
387 proximal promoters of 27.8% of the genes in module D5 contained CREs similar to those bound by
388 transcription factor B-3 (Table 3a) (Scharf et al., 2012), known to regulate heat shock responses in
389 *Arabidopsis thaliana* (Bechtold et al., 2013; Guo et al., 2016; Nover et al., 2001).

390 Additionally, all the predicted encoded proteins in each module were analysed to annotate the
391 transcription factors (TF), transcriptional regulators (TR), and kinases (Table S4a,b). For example, TRs
392 annotated in the D5 module – identified above as putatively drought-specific -- are involved in responses
393 to abscisic acid, heat stress, water deprivation and defense, and in zinc, chromatin, and metal ion binding
394 (Table S4a).

395 **Hub nodes of the Drought and Water networks**

396 Hub nodes were detected in both the Drought (Table 1a) and the Water (Table 1b) networks. A total
397 of 87 (0.53%; 1.73% excluding grey module) hub node isoforms from 72 (0.58%; 1.8% excluding grey
398 module) hub genes were identified in the Drought network, and 251 (1.53%; 3.74% excluding grey
399 module) hub node isoforms from 190 (1.53%; 3.51% excluding grey module) hub genes in the Water
400 network. Roughly, more than twice per-module fraction of hubs was detected in the Water network
401 (1.53% hub nodes/ 1.53% hub genes; Table 1b) compared to the Drought network (0.53% hub nodes/
402 0.58% hub genes; Table 1a).

403 **Pan-genome analyses: occupancy of all clustered and hub genes**

404 The studied pan-genome subset contained 34,310 pan-genome clusters (hereafter "pan-genes") which
405 were classified by the number of accessions with a gene model represented in a given cluster to determine
406 their occupancy. We found 16,057 (46.8%) core gene clusters with at least one member in every
407 accession. We analyzed these core genes along with the 5,642 (16.4%) clusters from the soft-core pan-
408 genome (occupancy in 31 or 32 accessions) to account for gene annotation errors and uncertainty with
409 orthology assignments. In contrast, there were 12,611 (36.8%) shell genes (i.e., observed in fewer than 31

410 accessions). Of the 34,310 pan-genes, 15,848 were represented by sequenced RNA tags, after our filtering
411 and normalizing steps. The occupancy of the expressed pan-genes was 12,137 (76.6%) core, 1,869
412 (11.8%) soft-core and 1,842 (11.6%) shell genes. The discrepancy between the total number of pan-genes
413 and that of genes represented by RNA-Seq tags may have been caused by the filtering of lowly expressed
414 genes. Additionally, many genes in the genome or pan-genome were likely not expressed in leaf tissue at
415 the developmental stage assessed herein. However, our results were consistent with prior studies which
416 found that pan-genome core genes are frequently more highly expressed (Gao et al., 2019; Gordon et al.,
417 2017; Tao et al., 2021) and therefore more likely to be sampled in RNA-Sequencing libraries (Figure S5).

418 The distribution of pan-gene occupancy varied considerably among modules in both Drought and
419 Water networks (Table 4a,b; Fig. 3a,b). Most modules in both co-expression networks showed ratios of
420 shell genes consistent with the genome averages (Fisher's Test; Table 4a,b), with notable exceptions.
421 Among Drought modules, D8 showed a considerable excess of shell genes (63.2%). Conversely, D2
422 (10.0%) and D3 (11.2%) each show a deficit of shell genes. Water co-expression modules showed
423 considerably more variation in pan-gene occupancy. Most large modules in the water network showed
424 significant deficits of shell genes, though W4 (27.8%), W9 (21.8%), W11 (25.5%), W12 (27.5%), and
425 W13 (60.0%) had an excess of shell genes relative to the genome averages (note that W11, W12, and
426 W13 were very small modules with fewer than 60 genes each).

427 To investigate the proportion of putative hub genes in each module that were members of the shell
428 gene sets, greater departures from genome averages were required to reach statistical significance given
429 the comparatively small number of these genes (Table 4c,d; Fig. 3c,d). Among the Drought modules, D8
430 and D9 did not show hub genes whereas other modules showed a predominance (>80%) of core or soft-
431 core hub genes with the exception of D7, whose four hub genes were shell genes (Table 4c). Among the
432 Water modules, W9 had a high proportion of shell hub genes (33.3%) as well as W4 and W12 though
433 these were poorly enriched with hub genes, whereas W7 showed a low percentage of hub shell genes.
434 Only the shell genes of module W7 are significantly under-represented among the hub genes of the Water
435 modules, according to the Fisher test (Table 4d).

436 As expected, core and soft-core genes were mostly enriched in the same GO terms as those in their
437 respective complete modules (Table 2a,b). Similarly, shell genes corresponded to genes involved in the
438 same biological process as those included in their complete modules. However, the most significant GO
439 term of the shell genes of D1 was photosynthesis, which was not found to be significant among core
440 genes. Similarly, the shell genes of D8 were enriched in photosynthesis, nitrogen, amide, and peptide

441 biosynthetic and metabolic processes terms, while core gene sets did not show any significant
442 enrichments (Table 2a,b and supplementary file S3).

443 We analyzed the pan-genome occupancy of genes with putative CREs in the modules with at least 10
444 detected genes. These corresponded predominantly to core genes (>50%) in both networks, whereas 7-
445 22% of the genes with predicted motifs in the Drought network and 0-27.3% in the Water network
446 corresponded to shell genes (Table 3a,b).

447 **Enrichments of differentially expressed genes with a pan-genomic perspective**

448 72.2%, 17.8% and 10% of the differentially expressed genes with upregulated isoforms in the drought
449 condition were core, soft-core, and shell genes, respectively. On the other hand, 67.5%, 20.9% and 11.6%
450 of the genes with downregulated isoforms in the drought condition were core, soft-core, and shell genes,
451 respectively (Table 5; supplementary file S2). 59.3% of DE isoforms (58.7% of DE genes) and 47.4% DE
452 isoforms (44.6% DE genes) of Drought (Table S5a) and Water (Table S5b) networks, respectively, were
453 not assigned to any modules (i.e., they are members of the gray or zero module). Among the hub nodes
454 (isoforms), 64.4% (56/87) and 34.3% (86/251) of them correspond to DE isoforms of the Drought (Table
455 S6a) and Water (Table S6b) networks, respectively.

456 Five of the nine Drought co-expression modules had a predominance (>50%) of upregulated DE
457 isoforms except for the large modules D2 and D3 and the small modules D8 and D9 (Table S5a).
458 Similarly, among Water modules, only one large module (W4) and three small modules (W10, W11 and
459 W13) had a predominance of downregulated DE isoforms in the drought condition compared to the water
460 condition (Table S5b).

461 21 out of the 25 most strongly upregulated genes did not cluster with other genes in the drought co-
462 expression network (i.e., they were members of the grey D0 module), while the majority of these strongly
463 upregulated genes did cluster in a module in the water network (Table S7). The 25 most strongly
464 upregulated genes by drought showed a range of predicted functions which included two predicted
465 dehydrins (Decena et al. 2021), two ABA-associated proteins, and two lipid transfer proteins (LTPs)
466 (Table S7). LTPs were among the most highly induced transcripts in an *Arabidopsis thaliana* experiment
467 that imposed very similar soil drying conditions to those imposed in the present study (Des Marais et al.,
468 2012). The 25 most strongly downregulated genes showed markedly different patterns than the most
469 strongly upregulated genes. Most of the genes cluster in D2 and W10, and four of these genes (three
470 clustered in D2 and one in W10) were hub genes (Table S7). Both of these modules were enriched for

471 genes involved in photosynthesis (Table 2a,b) and the annotations for many of these genes suggested
472 associations with the light reactions of photosynthesis.

473

474 **DISCUSSION**

475 Large scale transcriptome data sets have been used to construct co-expression networks for gene and
476 gene regulation discovery in model plant systems and crops (Aoki et al, 2007, 2016; Masalia et al., 2017;
477 Miao et al., 2017). The co-expression network approach allows testing hypotheses on gene functions from
478 their putative regulatory interactions with other functionally known genes classified in the same modules
479 (Mochida et al., 2011), and on links between signalling pathways and phenotypic response to
480 environmental stress (Des Marais et al., 2012). However, the mechanisms of and consequences for
481 genetic diversity in environmentally responsive gene regulatory networks are less often considered (Sun
482 & Dinneny, 2018). Our system-level approach allowed us to construct a drought-responsive gene co-
483 expression network from leaf tissue transcriptome profiles of *B. distachyon* accessions and to identify
484 modules of putatively co-regulated genes within it. We integrated these network hypotheses with
485 information about gene presence/absence variation as represented in the *B. distachyon* pan-genome.

486 **Regulatory control of *Brachypodium* response to soil drying**

487 *B. distachyon* is an annual species native to seasonally dry environments in the Mediterranean, where
488 it has likely evolved mechanisms to tolerate short-term soil drying during the growing season as well as
489 unpredictably timed end-of-season drought (López-Álvarez et al., 2015). Several past studies have
490 identified mechanisms of response to soil drying comprising transcriptomic, metabolic, physiological, and
491 developmental plasticity (Bertolini et al., 2013; Chen et al., 2016; Decena et al., 2021; Des Marais, et al.,
492 2016, 2017b; Fisher et al., 2016; Gordon et al., 2014; Handakumbura et al., 2019; Luo et al., 2011;
493 Manzaneda et al., 2015; Priest et al., 2014; Ruíz et al., 2016; Verelst et al., 2013), as well as considerable
494 genetic diversity of response (GxE; Des Marais et al., 2017b; Handakumbura et al., 2019). Priest et al.
495 (2014) provided the first transcriptomic assessment of response to drying, exposing the Bd21 accession to
496 a simulated severe drying stress by removing plants from soil to desiccate on a lab benchtop. These
497 authors observed a strong transcriptional signature of down-regulated photosynthesis, cell division, and
498 cell growth. Subsequent work imposing a more gradual soil drying stress in Bd21 found the opposite
499 pattern, directly observing sustained cell division and transcriptomic patterns of altered primary
500 metabolism, rather than outright downregulation (Verelst et al., 2013). Indeed, studies imposing moderate

501 drying on diverse *B. distachyon* accessions revealed increased leaf mass per area and greater root biomass
502 in several accessions in response to drying (Des Marais et al., 2017b; Handakumbura et al., 2019), both of
503 which require considerable investment of carbohydrates. In the present study we do, however, observe
504 several strongly downregulated genes with annotated functions related to the light reactions of
505 photosynthesis as well as RuBisCO assembly and function (Table S7).

506 How can we reconcile these transcriptional signatures of reduced photosynthesis with the observation
507 that carbohydrate-intensive processes like root growth continue under drying? The effects of soil drying
508 on photosynthesis are complex, and the reduction of internal leaf CO_2 (c_i) caused by stomatal closure can
509 affect the redox status of cells (Pinheiro & Chaves, 2011). As the Calvin Cycle reduces available CO_2, it
510 is also a strong sink for energy captured by the photosystems. As this sink is lowered by decreased c_i,
511 continued high irradiance can lead to increased expression of photoprotective mechanisms and decreased
512 expression of photochemistry as cells try to protect themselves from excess energy (Demmig-Adams &
513 Adams, 1996). As a result, studies of soil drying responses often observe decreased activity of the
514 photosystems and increased expression of, for example, photorespiration or other energy sinks (Wingler
515 et al., 1999). While we have no direct measurements of photorespiration or the quantum yield of
516 photosystem II in the current study, our observation of decreased expression of transcripts associated with
517 photosystem proteins is consistent with these mechanisms.

518 In light of this past evidence for an important role of photosynthesis and primary metabolism in
519 drying response, we focus here on Drought module 5 (D5). D5 showed a low correlation with the
520 Consensus modules (Fig. 2a), consistent with the hypothesis that the genes in this module are involved in
521 regulating plant response to drought stress. Its co-expressed genes, both core and soft-core, are involved
522 in protein folding, response to heat, temperature, and abiotic stimulus (Table 2a). The four DE hub nodes
523 (three genes) of D5 were upregulated in the drought condition compared to the water condition (Table S8)
524 and each of these four hubs is a core gene. Molecular chaperones, especially heat shock proteins (HSPs),
525 were predominantly annotated in both co-expressed and upregulated DE genes (Table S8). Related to the
526 presence of chaperones, the annotated DNA motifs concurred with the GO enrichment of the protein
527 folding and the Heat stress transcription factor B-3 (Table 3a).

528 **Topological position of pan-genes**

529 One key conclusion with respect to the evolution of pan-genomes and gene regulation arise from our
530 analysis. First, pan-genes are non-randomly assorted among co-expression modules in *B. distachyon.* This
531 observation suggests that the functions conferred by some co-expression modules are likely under

532 stronger purifying selection than those conferred by other modules. These latter modules represent
533 possible regulatory variation on which natural selection may act. Such a prospect was anticipated by
534 Wagner and Altenberg when they argued that modularity allows for evolutionary tinkering (Wagner &
535 Altenberg, 1996). Our results point towards a role for segregating gene copies in generating this modular
536 variation. This conclusion extends earlier work indicated that low-occupancy genes tend to be enriched
537 for functional classes of genes putatively involved in local adaptation such as disease resistance and gene
538 regulation (Gordon et al., 2017).

539 Pleiotropy can have a strong effect on the rate of molecular evolution and on the roles that functional
540 gene variants might play in evolutionary change. Pleiotropy is often correlated with the position of a gene
541 or protein in biochemical and gene regulatory networks (Erwin & Davidson, 2009; Jeong et al., 2001),
542 which are now readily inferred from high dimensional datasets such as the genome-wide gene co-
543 expression networks studies herein. Here, we considered the case of potentially large-effect mutations –
544 segregating gene copies identified from a grass pan-genome – and ask whether such "pan-genes" are
545 unevenly distributed in gene co-expression networks. Focusing on the well-Watered (control)
546 environment, we find that shell genes – pan-genes found in fewer than 31 of our studied accessions -- are
547 statistically under-represented among the genes in five of the six largest (in terms of total number of
548 genes) co-expression modules (Table 4b). These large modules are enriched for GO terms comprising
549 essential processes such as protein synthesis, primary metabolism, various processes related to
550 phosphorus metabolism and signalling, and cell wall organization (Table 2b). Moreover, shell genes are
551 generally under-represented among module hub genes (diagnosed as those whose expression most highly
552 correlated with the module as a whole, and thus possibly the most topologically connected among genes
553 in a module) in these five Water modules (Table 4d). Collectively, these results support the hypothesis
554 that core pan-genome genes are centrally located in gene co-expression networks and involved in
555 biological processes likely to be under strong purifying selection.

556 Water module 4 (W4), comprising 590 genes, is the only large water module that is statistically
557 enriched for shell genes (Table 4b). Shell genes in W4 are associated with a range of GO terms including
558 processes related to photosynthesis (Table 2b). In general, the lists of shell genes in modules do not tend
559 to have strong GO enrichments, perhaps owing to the relatively small numbers of genes in these lists.
560 Interestingly, among the Drought modules, the only module for which shell genes do have an enrichment
561 (D8; Table 4a) includes GO terms associated with photosynthesis (Table 2a). Shell genes represent genes
562 found in some sampled accessions but missing in others, suggesting that *B. distachyon* may harbor
563 genetic diversity in molecular pathways related to photosynthesis. Whether these segregating variants

represent adaptive genetic diversity reflecting the broad geographical coverage of our sampling, or simply mildly deleterious copy number variants on their way to being lost will require a larger population sample. Previously, we demonstrated significant genetic variation among the same *Brachypodium* accessions used herein for leaf carbon content, leaf C:N ratios, and water use efficiency (WUE; Des Marais, et al.; 2017b). Among these, WUE was significantly associated with principal components summarizing climate diversity; it is possible that some of the segregating variation in photosynthesis gene presence/absence is involved in local adaptation to climate.

The co-expression network approach we employed here, in concert with a modern pan-genomic perspective on segregating genetic variation, allows us to identify subsets of genes involved in core metabolic processes such as photosynthesis, suggesting possible candidate genes regulating natural genetic variation in resource assimilation and growth. These genes make attractive targets for further hypothesis testing via genome editing. Collectively, our work demonstrates the importance of accounting for both gene copy number variation and regulatory interactions in studying genome function and evolution.

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DATA ACCESSIBILITY AND BENEFIT-SHARING

Data Accessibility

RNA Sequence data are available at the ENA (European Nucleotide Archive; https://www.ebi.ac.uk/ena) with the consecutive accession numbers from ERR6133302 to ERR6133575.

All the R scripts, supplementary Excel files, TPM counts and adjacency matrices are available in the Github repository identified by the following doi: xxxxxxxxxx created using Zenodo.

Experimental details and metadata for samples studied herein were reported previously (Des Marais et al. 2017b).

Benefit Sharing

All accessions used in the current study originated from collections generated by the *Brachypodium* research community over several decades. Benefits from this research accrue from the sharing of our data and results on public databases as described above.

AUTHOR CONTRIBUTION

D.L.D. and T.E.J. designed the research. D.L.D. performed the experimental manipulations and generated the data. D.L.D., R.S., and B.C.-M. performed the analyses. D.L.D. and R.S. wrote the primary draft of the manuscript, and all authors contributed to the final draft of the manuscript.

890 **TABLES**

891 **Table 1.** Statistics of the number and percentage of isoforms (all and hub isoforms) and genes (all and
892 hub genes), and mean k_{diff} (the difference between intra- and inter-modular connectivity) per module for
893 each Drought **(a)** and Water **(b)** co-expression networks. The quantity of total counts of genes in drought
894 and water networks is different because we counted the genes for each module independently. When
895 several isoforms of the same gene were clustered in different modules, the gene was counted multiple
896 times; however, it was counted only once if it clustered in the same module. For example, if D1 has three
897 isoforms (Bradi1g10000.1, Bradi1g10000.2 and Bradi1g10000.3) of the same gene, it was counted only
898 once (Bradi1g10000); when isoforms were found in different modules (e.g., Bradi1g10000.1 in D1,
899 Bradi1g10000.2 in D2, Bradi1g10000.3 in D3), the gene was counted three times.

900 **(a)**

Drought modules		isoforms (nodes) per module		genes per module		mean Kdiff
		all	hub (KME>0.9)	all	hub (KME>0.9)	
D0	Gray	11366 (69.4%)	nd	8313 (67.4%)	nd	nd
D1	Turquoise	2477 (15.1%)	4 (0.16%)	1986 (16.1%)	3 (0.15%)	7.9
D2	Blue	979 (6.0%)	20 (2.04%)	750 (6.1%)	13 (1.73%)	3.7
D3	Brown	750 (4.6%)	41 (5.47%)	627 (5.1%)	38 (6.06%)	10.1
D4	Yellow	318 (1.9%)	4 (1.26%)	258 (2.1%)	4 (1.55%)	3.7
D5	Green	214 (1.3%)	4 (1.87%)	169 (1.4%)	3 (1.78%)	0.8
D6	Red	111 (0.7%)	10 (9.01%)	95 (0.8%)	7 (7.37%)	-1.5
D7	Black	68 (0.4%)	4 (5.88%)	48 (0.4%)	4 (8.3%)	-0.2
D8	Pink	65 (0.4%)	0 (0%)	57 (0.5%)	0 (0%)	-5.9
D9	Magenta	38 (0.2%)	0 (0%)	27 (0.2%)	0 (0%)	-8.8
Total counts		**16386**	**87 (0.53%)**	**12330**	**72 (0.58%)**	
Total unique counts		**16386**	**87 (0.53%)**	**12137**	**72 (0.59%)**	
Total unique counts (excluding gray module)		**5020**	**87 (1.73%)**	**4006**	**72 (1.80%)**	

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(b)

Water modules		isoforms (nodes) per module		genes per module		mean Kdiff
		all	hub (KME>0.9)	all	hub (KME>0.9)	
W0	Gray	9675 (59.0%)	nd	6934 (56.0%)	nd	nd
W1	Turquoise	1866 (11.4%)	149 (7.98%)	1439 (11.6%)	115 (7.99%)	26.8
W2	Blue	870 (5.3%)	4 (0.46%)	727 (5.9%)	3 (0.41%)	-3.7
W3	Brown	827 (5.0%)	2 (0.24%)	696 (5.6%)	1 (0.14%)	0.3
W4	Yellow	701 (4.3%)	2 (0.29%)	590 (4.8%)	2 (0.34%)	-0.7
W5	Green	607 (3.7%)	12 (1.98%)	509 (4.1%)	10 (1.96%)	3.1
W6	Red	435 (2.7%)	9 (2.07%)	358 (2.9%)	8 (2.08%)	2.1
W7	Black	390 (2.4%)	26 (6.67%)	313 (2.5%)	23 (7.35%)	4.6
W8	Pink	288 (1.7%)	2 (0.69%)	245 (2.0%)	2 (0.82%)	-2.2
W9	Magenta	282 (1.7%)	22 (7.80)	239 (1.9%)	21 (8.79%)	5.8
W10	Purple	246 (1.5%)	5 (2.03%)	185 (1.5%)	2 (1.08%)	-1.4
W11	Greenyellow	89 (0.5%)	8 (8.99%)	55 (0.4%)	1 (1.81%)	-3.1
W12	Tan	62 (0.4%)	1 (1.61%)	51 (0.4%)	1 (1.96%)	-0.9
W13	Salmon	48 (0.3%)	9 (18.75%)	40 (0.3%)	1 (2.5%)	-1.6
Total counts		**16386**	**251 (1.53%)**	**12381**	**190 (1.53%)**	
Total unique counts		**16386**	**251 (1.53%)**	**12137**	**190 (1.57%)**	
Total unique counts (excluding gray module)		**6711**	**251 (3.74%)**	**5407**	**190 (3.51%)**	

Table 2. Summary of the enrichment analysis according to the statistically significant GO biological process for the genes (all, core, soft-core and shell genes) clustered in the Drought (D) (**a**) and Water (W) (**b**) modules applying the statistical overrepresentation test of Panther (http://pantherdb.org/) tool. The biological processes are summarized according to the lowest False Discovery Rate (FDR) values (see Supplementary file S3).

(a)

Modules		All genes	core genes	soft-core genes	Shell genes
D1	**turquoise**	nitrogen, amide and peptide metabolic and biosynthetic processes	nitrogen, amide and peptide metabolic and biosynthetic processes	nitrogen, amide and peptide metabolic and biosynthetic processes	photosynthesis, nitrogen, amide and peptide metabolic and biosynthetic processes
D2	**blue**	photosynthesis	photosynthesis	photosynthesis	*No statistically significant terms*
D3	**brown**	organonitrogen compound metabolic process, location/transport and phosphorylation	organonitrogen compound metabolic process, location/transport and phosphorylation	*No statistically significant terms*	*No statistically significant terms*
D4	**yellow**	nucleobase-containing compound, heterocycle and nucleic acid metabolic processes	*No statistically significant terms*	*No statistically significant terms*	*No statistically significant terms*
D5	**green**	protein folding, response to heat, temperature and abiotic stimulus	protein folding, response to heat, temperature and abiotic stimulus	response to heat, temperature, and protein folding	*No statistically significant terms*
D6	**red**	lipid and small molecule metabolic process	lipid and small molecule metabolic process	*No statistically significant terms*	*No statistically significant terms*
D7	**black**	*No statistically significant terms*	*No statistically significant terms*	*No statistically significant terms*	*No statistically significant terms*
D8	**pink**	photosynthesis, nitrogen, amide and peptide biosynthetic and metabolic processes	*No statistically significant terms*	*No statistically significant terms*	photosynthesis, nitrogen, amide and peptide biosynthetic and metabolic processes
D9	**magenta**	*No statistically significant terms*	*No statistically significant terms*	*No statistically significant terms*	*No statistically significant terms*

(b)

	Modules	All genes	core genes	soft-core genes	Shell genes
W1	**turquoise**	nitrogen compound, organic substance, primary metabolic processes	nitrogen compound, organic substance, primary metabolic processes	protein ubiquitination, protein modification by small protein conjugation	*No statistically significant terms*
W2	**blue**	peptide, amide and nitrogen biosynthetic and metabolic processes	peptide, amide and nitrogen biosynthetic and metabolic processes	peptide, amide and nitrogen biosynthetic and metabolic processes	peptide, amide and nitrogen biosynthetic and metabolic processes
W3	**brown**	protein, peptide, amide and ion transport/location	protein, peptide, amide and ion transport/location	protein and ion transport/location	*No statistically significant terms*
W4	**yellow**	photosynthesis, generation of precursor metabolites and energy, cellular nitrogen compound and organic substance biosynthetic processes, and electron transport chain	organic substance, primary, nitrogen compound, nucleobase-containing compound metabolic processes	*No statistically significant terms*	photosynthesis, generation of precursor metabolites and energy, cellular nitrogen compound and organic substance biosynthetic processes, and electron transport chain
W5	**green**	organonitrogen and phosphate-containing compounds, and phosphorus metabolic processes and protein phosphorylation	organonitrogen and phosphate-containing compounds, small molecule, protein and phosphorus metabolic processes and protein phosphorylation	*No statistically significant terms*	*No statistically significant terms*
W6	**red**	plant-type cell wall organization or biogenesis	plant-type cell wall organization or biogenesis	*No statistically significant terms*	*No statistically significant terms*
W7	**black**	glutathione, cellular amino acid, oxoacid, organic acid and sulfur compound metabolic processes	glutathione, cellular amino acid, oxoacid, organic acid and sulfur compound metabolic processes	*No statistically significant terms*	*No statistically significant terms*

W8	**pink**	small molecule, starch, proline and carbohydrate metabolic processes	small molecule, starch, proline and carbohydrate metabolic processes	*No statistically significant terms*	*No statistically significant terms*
W9	**magenta**	regulation of nucleic acid-templated transcription, and RNA, nucleobase-containing compound, macromolecule biosynthesis and metabolic processes	cellular protein modification process, protein dephosphorylation, protein modification process, macromolecule modification and dephosphorylation	regulation of nucleic acid-templated transcription, and RNA, nucleobase-containing compound, macromolecule biosynthesis and metabolic processes	*No statistically significant terms*
W10	**purple**	photosynthesis	photosynthesis	photosynthesis	*No statistically significant terms*
W11	**greenyellow**	*No statistically significant terms*	*No statistically significant terms*	*No statistically significant terms*	*No statistically significant terms*
W12	**tan**	*No statistically significant terms*	*No statistically significant terms*	*No statistically significant terms*	*No statistically significant terms*
W13	**salmon**	macromolecule, cellular nitrogen compound, peptide and amide biosynthetic processes	*No statistically significant terms*	*No statistically significant terms*	macromolecule, cellular nitrogen compound, peptide and amide biosynthetic processes

Table 3. DNA motifs and cis-regulatory elements (CREs) of co-expressed genes assigned to Drought (D) (**a**) and Water (W) (**b**) modules. Modules (numeric and color codes of the modules); Accession (footprintDB), Names (gene name); Consensus sequence of motif; Protein name (Swissprot); Ncor (normalized correlation score); e-value; sites (number of sites used to compile the DNA motif); Proportion of genes with CREs per module as detected by matrix-scan in -500 to +200 bp windows; Total genes with CREs [occupancy of core, soft-core and shell genes]. The dashes indicate that no results were retrieved.

(a)

Drought modules		Accession (footprintDB)	Names	consensus	Protein (Swissprot)	Ncor	evalue	sites	Genes with putative CREs in promoter	
									Proportion of genes with CREs per module	Total genes with CREs [occupancy of core, soft-core and shell genes]
D1	turquoise	MA1353.1 (JASPAR 2020), M0609 (AthalianaCistrome v4_May2016)	AT1G72740, AT1G72740. DAP, T04493	ssmaAAACCCTAGcy	Telomere repeat-binding factor 5 (MYB transcription factor)	0.741	1.40E-56	225	9.0%	178 [139 (78.1%); 26 (14.6%); 13 (7.3%)]
D2	blue	ABI4 (Athamap 20091028)	ABI4	scCACCaCCCrc	Ethylene-responsive transcription factor ABI4 (Protein ABSCISIC ACID INSENSITIVE 4)	0.697	5.80E-07	178	17.1%	128 [102 (79.7%); 17 (13.3%); 9 (7.0%)]

D3	**brown**	M1684_1.02 (CISBP 1.02)	T153999_1.02, WRKY60	gtCGGTCAACgk	Probable WRKY transcription factor 60 (WRKY DNA-binding protein 60)	0.877	1.30E-09	165	11.6%	73 [54 (74.0%); 13 (17.8%); 6 (8.2%)]
D4	**yellow**	MA1197.1 (JASPAR 2020), M0350 (AthalianaCistrome v4_May2016)	CAMTA1, CAMTA1.DAP, T27097	bgCGACGCGCTks	Calmodulin-binding transcription activator 1, AtCAMTA1 (Ethylene-induced calmodulin-binding protein b, EICBP.b) (Signal-responsive protein 2, AtSR2)	0.717	7.80E-18	69	19.4%	50 [26 (52.0%); 13 (26.0%); 11 (22.0%)]
D5	**green**	M0455 (AthalianaCistrome v4_May2016)	HSFB3.ampDAP, T08630	kkCTCTrGAAGgk	Heat stress transcription factor B-3, AtHsfB3 (AtHsf-05)	0.691	9.10E-13	89	27.8%	47 [36 (76.6%); 5 (10.6%); 6 (12.8%)]
D6	**red**	Not found	Not found	Not found	Not found	Not found	Not found	Not found	-	-
D7	**black**	AHL25_3 (Arabidopsis PBM 20140210)	AHL25	wcAAAATAAATak	AT-hook motif nuclear-localized protein 25 (AT-hook protein of GA feedback 1)	0.643	0.46	35	54.2%	26 [16 (61.6%); 5 (19.2%); 5 (19.2%)]

D8	**pink**	MA1274.1 (JASPAR 2020), M0256 (AthalianaCistrome v4_May2016)	OBP3, OBP3.DAP, T16682	mrArGAATArrrAARrdh	Probable plastid-lipid-associated protein 14, chloroplastic, AtPap14 (Fibrillin-11) (OBP3-responsive protein 1)	0.623	1.50E-12	14	3.5%	2
D9	**magenta**	Not found	Not found	Not found	Not found	Not found	Not found	Not found	-	-

(b)

	Water modules	Accession (footprintDB)	Names	consensus	Protein (Swissprot)	Ncor	evalue	sites	Genes with putative CREs in promoter	
									Proportion of genes with CREs per module	Total genes with CREs [occupancy of core, soft-core and shell genes]
W1	**turquoise**	M0587_1.02 (CISBP 1.02)	PK22848.1, T073801_1.02	wkwcTAAAATTTTAGwww	*No results*	0.59	6.80E-29	32	0.1%	2
W2	**blue**	At2g20350:M05132:TRANSFAC	At2g20350/M05132/TRANSFAC	mmAGGCCCATCwv	Ethylene-responsive transcription factor ERF120	0.59	1.90E-97	244	22.3%	162 [115 (71.0%); 25 (15.4%); 22 (13.6%)]

W3	**brown**	Not found	Not found	Not found	Not found	Not found	Not found	Not found	-	-
W4	**yellow**	MA1379.1 (JASPAR 2020), M0357 (AthalianaCistrome v4_May2016)	SOL1, SOL1.DAP, T15201	wwTTAAwwwddAAAwr	Carboxypeptidase SOL1, EC 3.4.17.- (Protein SUPPRESSOR OF LLP1 1)	0.662	1.10E-09	458	0.3%	2
W5	**green**	UP00581A_2 (UniPROBE 20160601)	vascular plant one zinc finger protein 2, VOZ2	sgAGTCAACGtcgv	Transcription factor VOZ2 (Protein VASCULAR PLANT ONE-ZINC FINGER 2, AtVOZ2)	0.88	1.90E-10	126	5.7%	29 [20 (69.0%); 4 (13.8%); 5 (17.2%)]
W6	**red**	M0523 (AthalianaCistrome v4_May2016)	MYB58.DAP, T03869	ytgCyACCAACCAvm	Transcription factor MYB58	0.837	1.10E-11	45	2.5%	9
W7	**black**	UP00582A_2 (UniPROBE 20160601)	ARABIDOPSIS THALIANA WRKY DNA-BINDING PROTEIN 18, At4g31800, AtWRKY18, F11C18.16, F28M20.10, WRKY transcription factor 1, WRKY transcription	mmsGGTCAAACGyr	WRKY transcription factor 18	0.698	7.40E-07	27	6.1%	19 [17 (89.5%); 2 (10.5%); 0 (0.0%)]

			factor 18, WRKY1							
W8	**pink**	Not found	Not found	Not found	Not found	Not found	Not found	Not found	-	-
W9	**magenta**	MA1197.1 (JASPAR 2020), M0350 (AthalianaCistrome v4_May2016)	CAMTA1, CAMTA1.DAP, T27097	ssaCCGCGTcss	Calmodulin-binding transcription activator 1, AtCAMTA1 (Ethylene-induced calmodulin-binding protein b, EICBP.b) (Signal-responsive protein 2, AtSR2)	0.755	4.50E-19	95	23.0%	55 [28 (50.9%); 12 (21.8%); 15 (27.3%)]
W10	**purple**	Not found	Not found	Not found	Not found	Not found	Not found	Not found	-	-
W11	**green-yellow**	Not found	Not found	Not found	Not found	Not found	Not found	Not found	-	-
W12	**tan**	Not found	Not found	Not found	Not found	Not found	Not found	Not found	-	-
W13	**salmon**	Not identified	Not identified	trAAATymwwwTTCAht	Not identified	0.504	0.00017	35	20.0%	8

Table 4. Occupancies [core (33 accessions); soft-core (31-32 accessions) and shell (≤30 accessions)] of the co-expressed genes **(a; b)** and hub genes **(c; d)** for each module of the Drought (D) and Water (W) co-expression networks. Asterisks indicate significant differences comparing shell and total genes between modules and network by Fisher test (*p ≤ 0.05; **p ≤ 0.01; ***p ≤ 0.001).

(a)

			Drought genes occupancy		
Drought modules		**Genes**	**Core (33)**	**Soft-core (32 or 31)**	**Shell (≤30)**
D0	**gray**	8313	5764 (69.3%)	1269 (15.3%)	1280 (15.4%)
D1	**turquoise**	1986	1422 (71.6%)	265 (13.3%)	299 (15.1%)
D2	**blue**	750	549 (73.2%)	126 (16.8%)	75 (10.0%)***
D3	**brown**	627	440 (70.2%)	117 (18.7%)	70 (11.2%)*
D4	**yellow**	258	160 (62.0%)	49 (19.0%)	49 (19.0%)
D5	**green**	169	123 (72.8%)	27 (16.0%)	19 (11.2%)
D6	**red**	95	62 (65.3%)	18 (19.0%)	15 (15.8%)
D7	**black**	48	28 (58.3%)	9 (18.8%)	11 (22.9%)
D8	**pink**	57	17 (29.8%)	4 (7.0%)	36 (63.2%)***
D9	**magenta**	27	20 (74.1%)	6 (22.2%)	1 (3.7%)
Total counts		**12330**	**8585 (69.6%)**	**1890 (15.3%)**	**1855 (15.0%)**
Total unique counts		**12137**	**8426 (69.4)**	**1869 (15.4%)**	**1842 (15.2%)**
Total unique counts (excluding gray module)		**4006**	**2813 (70.2%)**	**618 (15.4%)**	**575 (14.4%)**

(b)

			Water genes occupancy		
Water modules		**Genes**	**Core (33)**	**Soft-core (32 or 31)**	**Shell (≤30)**
W0	**gray**	6934	4788 (69.1%)	1006 (14.5%)	1140 (16.4%)*
W1	**turquoise**	1439	1094 (76.0%)	188 (13.1%)	157 (10.9%)***
W2	**blue**	727	544 (74.8%)	111 (15.3%)	72 (9.9%)***
W3	**brown**	696	500 (71.8%)	128 (18.4%)	68 (9.8%)***
W4	**yellow**	590	353 (59.8%)	73 (12.4%)	164 (27.8%)***
W5	**green**	509	375 (73.7%)	101 (19.8%)	33 (6.5%)***
W6	**red**	358	258 (72.1%)	67 (18.7%)	33 (9.2%)**
W7	**black**	313	216 (69.0%)	59 (18.8%)	38 (12.1%)
W8	**pink**	245	162 (66.1%)	50 (20.4%)	33 (13.5%)
W9	**magenta**	239	124 (51.9%)	63 (26.4%)	52 (21.8%)*
W10	**purple**	185	127 (68.6%)	36 (19.5%)	22 (11.9%)
W11	**greenyellow**	55	33 (60.0%)	8 (14.5%)	14 (25.5%)
W12	**tan**	51	31 (60.8%)	6 (11.8%)	14 (27.5%)
W13	**salmon**	40	13 (32.5%)	3 (7.5%)	24 (60.0%)***
Total counts		**12381**	**8618 (69.6%)**	**1899 (15.3%)**	**1864 (15.1%)**
Total unique counts		**12137**	**8426 (69.4)**	**1869 (15.4%)**	**1842 (15.2%)**
Total unique counts (excluding gray module)		**5407**	**3795 (70.2%)**	**890 (16.5%)**	**722 (13.3%)**

(c)

Drought modules		Hub Genes	Drought hub genes (KME>0.9) occupancy		
			Core (33)	Soft-core (32 or 31)	Shell (≤30)
D1	**turquoise**	3	3 (100%)	0 (0%)	0 (0%)
D2	**blue**	13	10 (76.9%)	1 (7.7%)	2 (15.4%)
D3	**brown**	38	20 (52.6%)	18 (47.4%)	0 (0%)
D4	**yellow**	4	3 (75%)	1 (25%)	0 (0%)
D5	**green**	3	3 (100%)	0 (0%)	0 (0%)
D6	**red**	7	4 (57.1%)	3 (42.9%)	0 (0%)
D7	**black**	4	0 (0%)	0 (0%)	4 (100%)
D8	**pink**	0	0 (0%)	0 (0%)	0 (0%)
D9	**magenta**	0	0 (0%)	0 (0%)	0 (0%)
Total counts		**72**	**43 (59.7%)**	**23 (31.9%)**	**6 (8.3%)**
Total unique counts		**72**	**43 (59.7%)**	**23 (31.9%)**	**6 (8.3%)**

(d)

Water modules		Hub Genes	Water hub genes (KME>0.9) occupancy		
			Core (33)	Soft-core (32 or 31)	Shell (≤30)
W1	**turquoise**	115	82 (71.3%)	18 (15.7%)	15 (13.0%)
W2	**blue**	3	2 (66.7%)	1 (33.3%)	0 (0%)
W3	**brown**	1	1 (100%)	0 (0%)	0 (0%)
W4	**yellow**	2	1 (50%)	0 (0%)	1 (50%)
W5	**green**	10	6 (60%)	3 (30%)	1 (10%)
W6	**red**	8	7 (87.5%)	1 (12.5%)	0 (0%)
W7	**black**	23	16 (69.6%)	7 (30.4%)	0 (0%)***
W8	**pink**	2	2 (100%)	0 (0%)	0 (0%)
W9	**magenta**	21	13 (61.9%)	1 (4.8%)	7 (33.3%)
W10	**purple**	2	2 (100%)	0 (0%)	0 (0%)
W11	**greenyellow**	1	1 (100%)	0 (0%)	0 (0%)
W12	**tan**	1	0 (0%)	0 (0%)	1 (100%)
W13	**salmon**	1	1 (100%)	0 (0%)	0 (0%)
Total counts		**190**	**134 (70.5%)**	**31 (16.3%)**	**25 (13.2%)**

Total unique counts	**190**	**134 (70.5%)**	**31 (16.3%)**	**25 (13.2%)**

Table 5. Pan-genome analysis of the differentially expressed (DE) isoforms according to their gene occupancy: core (33 accessions), soft-core (31-32 accessions) and shell (≤30 accessions).* One gene was represented by one up-regulated and one down-regulated isoform.

	Gene occupancy			
	core	**soft-core**	**shell**	**TOTAL**
upregulated	2027 (72.2%)	500 (17.8%)	281 (10%)	**2808**
downregulated	661 (67.5%)	205 (20.9%)	114 (11.6%)	**980**
up/down-regulated	1*	0	0	**1**
TOTAL	**2689**	**705**	**395**	**3789**

FIGURES

Figure 1. Summary of the experimental design and analyses performed in the 33 accessions of the model grass *Brachypodium distachyon* under drought (D) and water (control) (W) conditions.

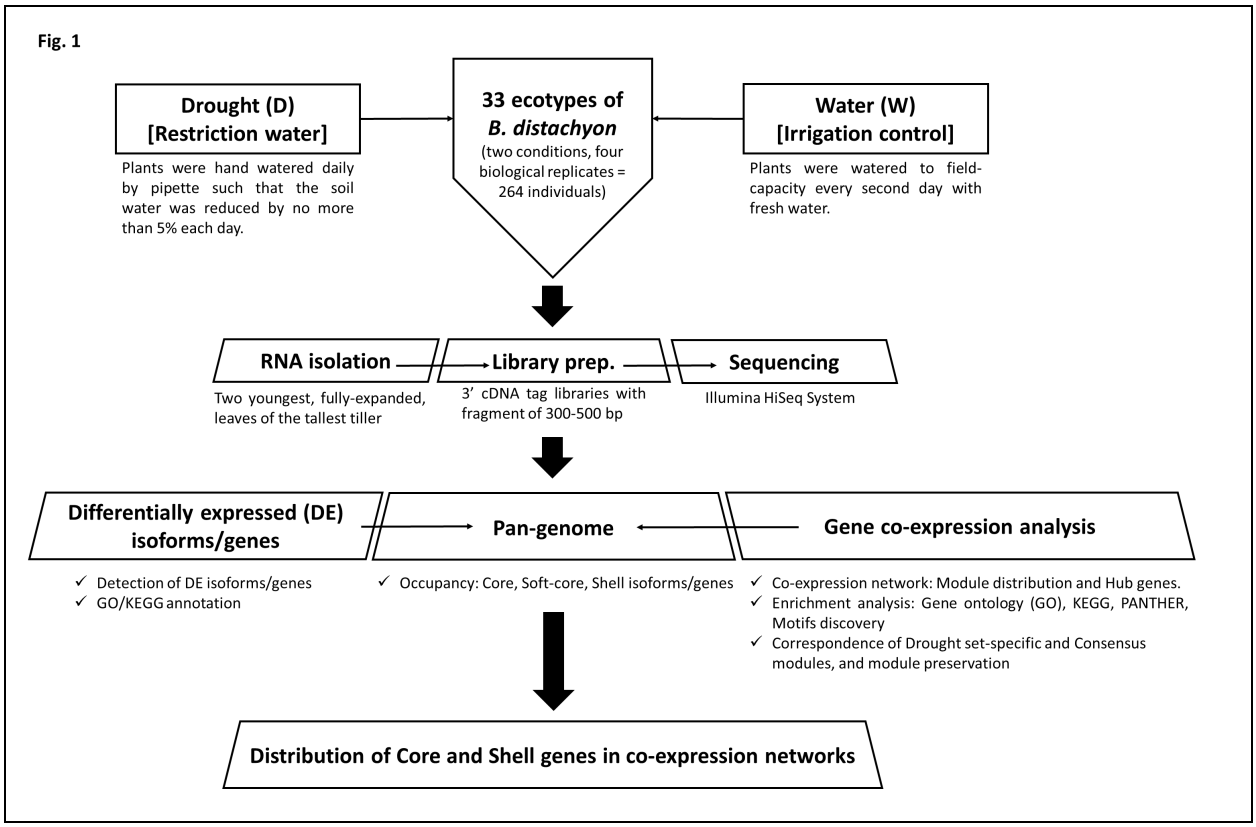

Figure 2 Correspondence (number of nodes) of Drought (D) (**a**) and Water (W) (**b**) set-specific and Drought-Water consensus (Cons) modules. Each row of the table corresponds to one Drought/Water set-specific module, and each column corresponds to one consensus module. Numbers in the table indicate node counts in the intersection of the corresponding modules. Coloring of the table encodes − log(p), with p being the Fisher's exact test p-value for the overlap of the two modules. The stronger the red color, the more significant the overlap is. (**c**) Comparison of the GO enrichments for each D and W module, according to the correlation with the common consensus module.

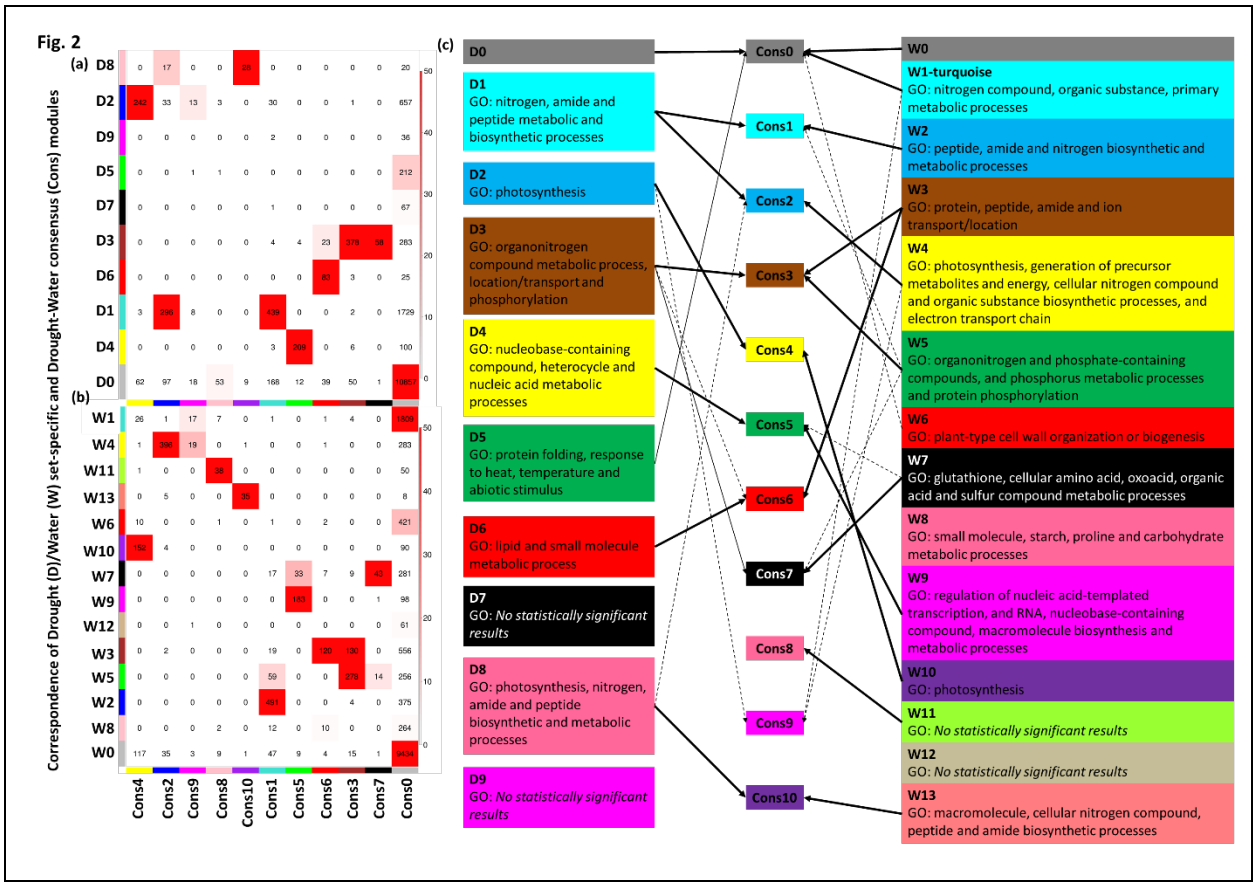
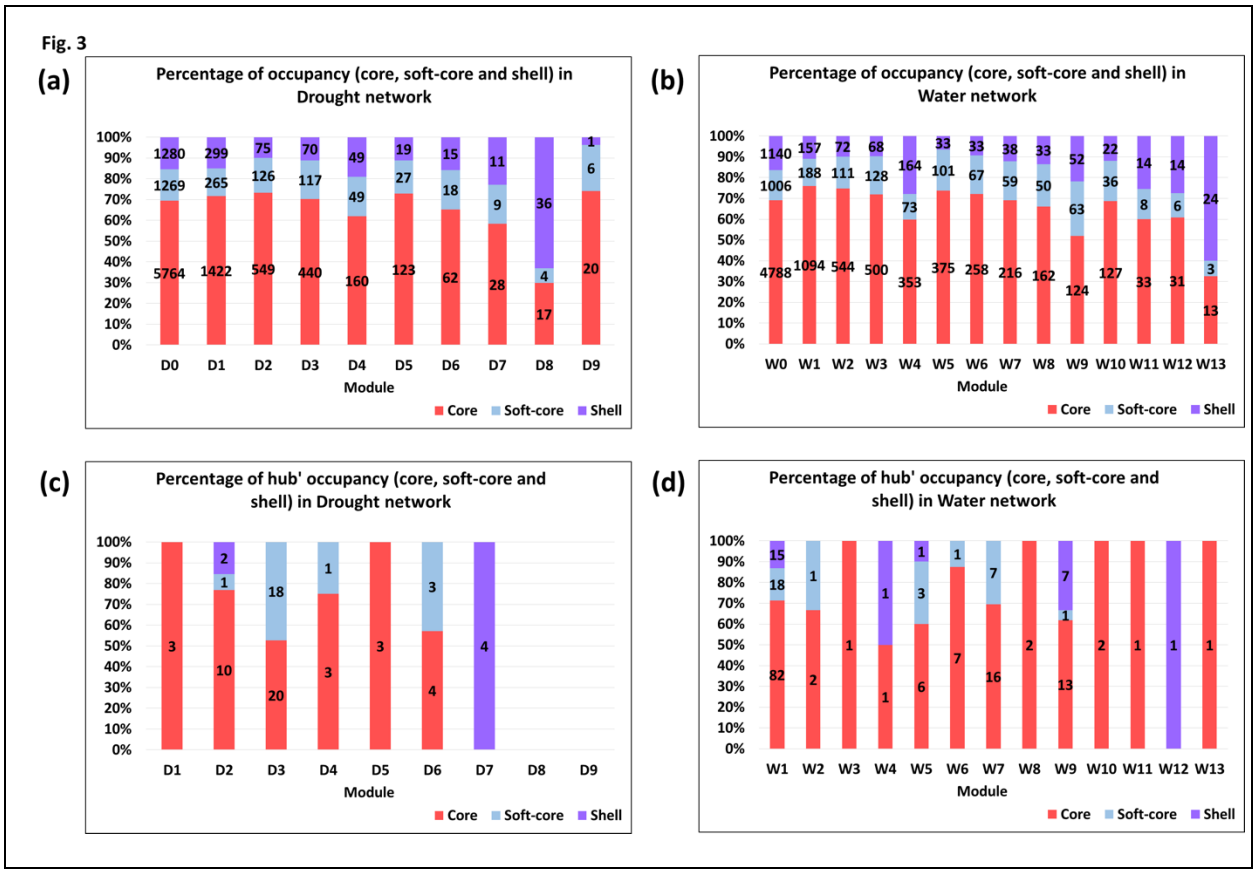
Figure 3. Proportion (%) of occupancies [core (33); soft-core (31-32) and shell (≤30)] of the co-expressed genes (**a; b**) and hub genes (**c; d**) for each Drought (D) and Water (W) network.