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A reduced tillering trait shows small but important yield gains in dryland wheat production

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27 **Abstract**

28 Reducing the number of tillers per plant using a tiller inhibition (*tin*) gene has been considered as an
29 important trait for wheat production in dryland environments. We used a spatial analysis approach
30 with a daily time-step coupled radiation and transpiration efficiency model to simulate the impact of
31 the reduced-tillering trait on wheat yield under different climate change scenarios across Australia's
32 arable land. Our results show a small but consistent yield advantage of the reduced-tillering trait in
33 the most water-limited environments both under current and likely future conditions. Our climate
34 scenarios show that whilst elevated [CO₂] (e[CO₂]) alone might limit the area where the reduced-
35 tillering trait is advantageous, the most likely climate scenario of e[CO₂] combined with increased
36 temperature and reduced rainfall consistently increased the area where restricted tillering has an
37 advantage. Whilst long-term average yield advantages were small (ranged from 31 to 51 kg ha⁻¹ yr⁻¹),
38 across large dryland areas the value is large (potential cost-benefits ranged from AUD 23 to 60 MIL
39 yr⁻¹). It seems therefore worthwhile to further explore this reduced-tillering trait in relation to a
40 range of different environments and climates, because its benefits are likely to grow in future dry
41 environments where wheat is grown around the world.

42 **Keywords:** APSIM next generation, climate change, semi-arid environments, *Triticum aestivum*,
43 water use efficiency

44

45 **Introduction**

46 Wheat (*Triticum aestivum* L.) containing a reduced-tillering *tin* (tiller inhibition) gene (Atsmon and
47 Jacobs, 1977) has been proposed as an important breeding objective to increase grain yield in
48 dryland environments (Hendriks et al., 2015; Reynolds et al., 2009; Richards, 1988). The advantage
49 of the reduced-tillering trait is in its constraining of excessive tillering and therefore leaf area
50 development and water use during early- to mid-season when temperature and water availability
51 are favourable for tiller initiation (Richards, 1988; Sharma, 1995), shifting soil water availability from
52 pre- to post-anthesis crop growth when drought and temperature stress are frequent (Berry et al.,
53 2003; Richards, 1988; Stephens and Lyons, 1997).

54 Water availability has long been recognised as a key component and dominant variable affecting
55 crop production in dryland environments, where annual rainfall ranges from about 300 to 600 mm
56 and rainfall variability is substantial (Dawson, 1957). The importance of rainfall varies both
57 temporally and spatially, being a function of rainfall amount and distribution, as well as the ability of
58 respective soils to store water (Stephens and Lyons, 1997). The reduced-tillering trait can be
59 advantageous in some growing environments (i.e. growing seasons and locations) where post-

60 anthesis soil water is critically limiting (Duggan et al., 2005; Moeller and Rebetzke, 2017; Richards,
61 1988). Whereas in other environments with greater water availability, the free-tillering phenotypes
62 may be superior as tillering provides a greater potential to respond to increasing water availability.
63 The propensity to tiller allows for the development of a greater leaf area and the capacity to capture
64 resources (viz. water, CO₂, solar radiation, and mineral nutrients) needed for increasing biomass and
65 yield (Duggan et al., 2005; Mitchell et al., 2013; Sadras and Rebetzke, 2013). The greater growth and
66 water use, however, in turn may also reduce soil water availability later in the season, worsening
67 terminal drought so that only fewer number of tillers translate into fertile spike. Thus there is
68 uncertainty as to whether the reduced tillering-trait leads to increased or decreased yield when local
69 climate conditions vary. Despite the benefits reported in a number of experimental locations and
70 years (e.g. Mitchell et al. (2012) and Moeller and Rebetzke (2017)), there is a lack of systematic
71 quantification of long-term advantages of the reduced-tillering trait across a range of growing
72 environments, especially taking into account likely future conditions.

73 Global atmospheric [CO₂] is likely to reach 550 μmol mol⁻¹ by 2050 (Solomon et al., 2007), up from
74 just over 400 μmol mol⁻¹ at present (2020). Elevated [CO₂] (e[CO₂]) leads to a number of beneficial
75 growth and physiological responses, many of which are interpreted in the context of ameliorating
76 the negative impacts of drought (Leakey et al., 2009; Wullschleger et al., 2002). Elevated [CO₂]
77 decreases stomatal conductance (e.g. Ainsworth and Rogers (2007) and Houshmandfar et al. (2015)),
78 which, in turn, can translate into a proportional reduction in canopy level transpiration (e.g.
79 Houshmandfar et al. (2018) and Leakey et al. (2009)), depending on other determinates of canopy
80 level transpiration, e.g. leaf area index. The suitability of the reduced-tillering trait to a growing
81 environment may be different under e[CO₂], especially if combined with other likely climate change-
82 related events such as reduced rainfall and higher temperatures (Solomon et al., 2007).

83 Limited information is available about the reduced-tillering trait performance under e[CO₂]. Results
84 from field experiments under ambient [CO₂] (a[CO₂]) have also been highly variable from site to site
85 and from season to season (e.g. Mitchell et al. (2012) and Houshmandfar et al. (2019)). Experimental
86 approaches aimed at interpreting the interaction between genotype and environment are time-
87 consuming, especially where capturing the impact of long-term climate is needed (Asseng and
88 Turner, 2007; He et al., 2017). Crop simulation modelling is another approach allowing long-term
89 assessment under different growing conditions, and especially valuable where it is based on robust
90 data from rigorous field experiments (Christy et al., 2018; Zhao et al., 2019). In this current paper,
91 we used experimental data from two growing seasons at the Australian Grains Free-Air CO₂
92 Enrichment (AGFACE) facility together with other published data and crop simulation modelling to (i)
93 better understand the interaction between genetic and environmental components of the reduced-

94 tillering trait in wheat and (ii) extrapolate its potential long-term average benefits across Australia's
95 arable land (i.e. land capable of being used to grow crops). The potential yield advantages under the
96 present climate and likely future warmer and drier climate scenarios under e[CO₂] were considered.

97

98 **Materials and methods**

99 We analysed data from field experiments conducted in the AGFACE facility, and used published data
100 from multiple sites representing the main environment types in Australia's wheatbelt (Mitchell et al.,
101 2012), to validate the Agricultural Production Systems sIMulator (APSIM) (Brown et al., 2014;
102 Holzworth et al., 2018; Holzworth et al., 2014) to reproduce the reduced-tillering trait in wheat
103 under a range of growing environments and under e[CO₂]. The model was then used to extrapolate
104 potential long-term average benefits of the reduced-tillering trait across Australia's arable land. We
105 used APSIM because it has been widely used in wheat in climate change (e.g. Hochman et al. (2017)
106 and Wang et al. (2018)) and crop trait evaluation (e.g. Zhao et al. (2019)) studies, and has recently
107 been successfully tested against other AGFACE data (O'Leary et al., 2015). The model is capturing the
108 CO₂ effects on assimilation rates through modifiers of radiation use efficiency. Transpiration is a
109 function of daily dry matter increase multiplied by transpiration efficiency, which, in turn, depends
110 on vapour pressure deficit and [CO₂] (Holzworth et al., 2014; O'Leary et al., 2015). Actual
111 transpiration and assimilation rates are reduced if available soil water is inadequate to meet the
112 transpiration demand (Holzworth et al., 2014).

113 Simulations were set up according to the rules established for quantifying grain yield in previously
114 published studies under rainfed conditions, with non-limiting nutrients and well-controlled biotic
115 stresses. Under such conditions, wheat production is determined by the amount and distribution of
116 rainfall, solar radiation, temperature, atmospheric [CO₂], and fixed physical attributes of the soil
117 (Hochman et al., 2017). These settings were unchanged for the duration of the simulations (1962–
118 2018).

119

120 **The AGFACE experiments**

121 Two field experiments were conducted in 2011 and 2012 growing seasons at the AGFACE site
122 (Fitzgerald et al., 2016) located in Horsham, Victoria (Fig. 1 and Fig. 2). The site has a Mediterranean
123 type climate but with cooler and drier winters (Hutchinson et al., 2005). Long-term average annual
124 rainfall (1962–2018) of the area is 433.1 mm (standard deviation = 117.0 mm). Long-term average
125 maximum and minimum temperatures are 21.5 °C and 8.2 °C (Australian Bureau of Meteorology).

126 The soil type is a Vertosol clay with non-dispersive and pedal surface (Isbell, 2016), approximately
127 35% clay at the top increasing to 60% at 1.4 m depth. The experiment had eight plots (16 m in
128 diameter) of which four were ambient CO₂ (approximately 380 μmol mol⁻¹, average daytime [CO₂])
129 and four elevated CO₂ (centre concentration set at 550 μmol mol⁻¹). Each elevated CO₂ plot was
130 encircled by horizontal CO₂-release-tubes in an octagonal shape which were progressively raised as
131 the crop grew so that the CO₂ was injected about 15 cm above the canopy. A plot centre [CO₂] of 550
132 μmol mol⁻¹ was maintained for the elevated CO₂ treatment from sunrise to sunset starting from
133 germination. Average plot central [CO₂] were recorded every minute with an infrared gas analyser
134 (IRGA, SBA-4, PP Systems, Amesbury, MA, USA) located at the central part of each plot. The spatial
135 variations in the [CO₂] of the site were described by Mollah et al. (2009).

136 Two wheat (*Triticum aestivum* L.) genotypes contrasting in presence of the *tin* gene, “Silverstar” and
137 “Silverstar + *tin*”, were sown into two randomly allocated subplots (1.5 × 4 m, row spacing = 0.27 m),
138 one each in opposing halves of the ring under either rainfed or supplemental irrigation. Silverstar is
139 an early maturing cultivar initially bred for low rainfall environments but is also suitable for higher
140 yielding environments, as reported by Riffkin et al. (2003). Silverstar + *tin* was a BC₂F_{6.8} breeding line
141 SsrT65, derived by backcrossing a *tin* donor to the spring wheat variety Silverstar (Mitchell et al.,
142 2013). For the supplemental irrigation treatments, a total of 100- and 120-mm irrigation was applied
143 in five splits from 6 September to 18 October in 2011 and in four splits from 11 September to 29
144 October in 2012, respectively. This resulted in eight sets of environmental growing conditions: 2
145 years × 2 [CO₂] × 2 watering regimes. Sowing dates were 25 May in 2011 and 30 May in 2012. Annual
146 rainfall was 552.5 mm in 2011 and 301.8 mm in 2012.

147 Total aboveground biomass was measured at anthesis (DC65, Zadoks et al. (1974), DC: decimal code)
148 and physiological maturity (DC90, harvest). At each sampling date, plants were hand-harvested and
149 counted for number of plants and number of tillers m⁻² from 0.675 m² (5 rows × 0.27 m row spacing
150 × 0.5 m length, including edge rows) of each subplot. ‘Edge rows’ refers to the sub-plots arranged
151 within the closed canopy of the whole ring, so that any growth stimulation due to edge effects
152 between sub-plots would have been very small. Appreciable edge effects would lead to
153 overestimates of growth and yield, which would introduce significant positive deviations from the
154 1:1 lines in Fig. 3. The DC65 samples were dried for 72 h at 70°C. The DC90 samples were dried for
155 72 h at 40°C. The DC90 samples were further processed for grain yield. All parameters were
156 expressed on an area basis. Anthesis and physiological maturity dates were recorded for each
157 treatment year. A preliminary analysis of data from these experiments is given in Löw et al. (2015).

158

159 **The APSIM model parameterisation**

160 APSIM Next Generation Plant Modelling Framework (Holzworth et al., 2018) was tested against the
 161 AGFACE field data (Table 1), ensuring observed phenological stages of DC65 and DC90 were
 162 matched. Both Silverstar and Silverstar + *tin* were parameterised identically except for one
 163 parameter, potential branching rate (i.e. tiller initiation rate). The tiller inhibition gene restricts tiller
 164 number to a maximum of 4 tillers per plant (Duggan et al., 2005; Richards, 1988). Potential
 165 branching rate was therefore decreased from the APSIM default of 20 tillers plant⁻¹ (total tiller
 166 population including non-fertile tillers) in the cultivar Silverstar to 4 tillers plant⁻¹ in Silverstar + *tin*.
 167 The model reproduces the development of wheat leaves and tillers using a cohort approach based
 168 on the coordination of leaf and tiller initiation on main stem and tillers (Brown et al., 2014). Leaves
 169 and tillers that initiate at the same time belong to the same leaf or tiller cohorts and grow following
 170 the same pattern. Tillering (branching) is simulated with leaf number and a potential rate following
 171 the pattern of a Fibonacci series between germination and terminal spikelet. The actual branching
 172 rate is the potential branching rate reduced by water and nitrogen deficiencies, and further
 173 constrained by carbon assimilate supply (Evers et al., 2006). Tillering stops at terminal spikelet and
 174 tiller mortality occurs thereafter. Later initiating tillers with slower growth rate and the smallest
 175 tillers will die first. At the terminal spikelet, all tillers with less than four leaves stop growing new
 176 leaves (see Zhao et al., 2019).

177 For each season, the model was initialised to match measured sowing soil water content and
 178 available mineral nitrogen content through the soil profile (pooled across the experiment). Irrigation
 179 was applied by the model on the actual days of application. Due to variable seedling establishment
 180 across treatments, number of plants m⁻² were highly variable for different treatments and sampling
 181 dates (Table 1). Therefore, to have a meaningful test of the model, we used the actual (measured)
 182 number of plants m⁻² for each treatment simulation.

183

184 **TABLE 1** Summary of the observed number of plants, number of tillers, biomass at anthesis (DC65)
 185 and physiological maturity (DC90), as well as the harvested grain yield results from the AGFACE
 186 experiment. The values are averages from four replicates each (Löw et al., 2015). ± Standard error.

Sowing date	Rain (mm)	Irrigation (mm)	Measurement	Silverstar		Silverstar + <i>tin</i>	
				a[CO ₂]	e[CO ₂]	a[CO ₂]	e[CO ₂]
25/04/2011	552.5	0.0	Number of plants DC65 (m ⁻²)	110.8±6.0	138.0±32.6	169.2±35.8	109.7±18.9
			Number of tillers DC65 (m ⁻²)	539.6±49.0	581.0±66.2	237.2±52.8	303.3±85.5
			Biomass DC65 (kg ha ⁻¹)	7201±689	9648±461	7377±1624	8371±1223
			Number of plants DC90 (m ⁻²)	81.0±8.9	84.2±8.4	97.9±20.4	47.9±6.2
			Number of tillers DC90 (m ⁻²)	419.3±44.0	520.7±53.0	171.2±56.2	219.5±22.3

			Biomass DC90 (kg ha ⁻¹)	13252±908	18742±1323	8956±1753	11707±1388
			Grain yield (kg ha ⁻¹)	5305±635	7352±557	3889±833	4736±663
25/04/2011	552.5	100.0	Number of plants DC65 (m ⁻²)	115.0±12.5	153.3±7.3	122.6±22.1	111.3±21.0
			Number of tillers DC65 (m ⁻²)	503.8±66.1	670.2±22.2	388.9±110.4	256.2±35.9
			Biomass DC65 (kg ha ⁻¹)	8593±1046	11172±541	7721±1209	8976±886
			Number of plants DC90 (m ⁻²)	120.7±35.2	110.5±13.6	71.7±20.7	89.0±20.0
			Number of tillers DC90 (m ⁻²)	533.0±59.0	576.3±76.0	256.3±35.9	257.5±53.1
			Biomass DC90 (kg ha ⁻¹)	16033±595	20227±1578	9982±1081	13166±1501
			Grain yield (kg ha ⁻¹)	6503±612	8447±540	4183±461	5860±571
30/04/2012	301.8	0.0	Number of plants DC65 (m ⁻²)	121.9±5.1	88.5±11.8	107.0±14.0	100.7±8.8
			Number of tillers DC65 (m ⁻²)	506.6±47.1	463.7±58.6	168.7±12.0	235.1±6.9
			Biomass DC65 (kg ha ⁻¹)	7451±174	7290±700	5328±188	7417±116
			Number of plants DC90 (m ⁻²)	89.6±1.9	91.5±8.3	74.8±5.4	87.0±8.4
			Number of tillers DC90 (m ⁻²)	485.4±44.0	430.5±33.8	219.4±21.7	256.3±16.4
			Biomass DC90 (kg ha ⁻¹)	12487±1177	12678±841	9965±680	12365±1163
			Grain yield (kg ha ⁻¹)	5555±766	5741±225	4689±210	5635±596
30/04/2012	301.8	120.0	Number of plants DC65 (m ⁻²)	76.3±13.3	100.4±7.1	80.0±4.4	104.4±6.2
			Number of tillers DC65 (m ⁻²)	488.5±59.4	534.2±49.0	173.2±30.7	222.2±17.1
			Biomass DC65 (kg ha ⁻¹)	7493±732	9259±734	5532±712	7433±934
			Number of plants DC90 (m ⁻²)	85.2±9.3	84.4±8.1	87.0±8.4	89.6±12.4
			Number of tillers DC90 (m ⁻²)	436.8±23.1	530.2±81.5	198.3±30.3	236.9±3.7
			Biomass DC90 (kg ha ⁻¹)	13602±450	18557±1713	11312±1009	15321±715
			Grain yield (kg ha ⁻¹)	6078±253	8763±853	5568±559	7169±720

187

188 The model was further validated using published data in Mitchell et al. (2012). The paper reported
 189 grain yield in cultivar Silverstar and Silverstar + *tin* (BC₂F_{6,8} breeding lines SsrT02, SsrT14, and SsrT17)
 190 at five locations (Fig. 1), representative of the main environment types in Australia's wheatbelt. The
 191 locations were Gatton, Kingsthorpe, and Emerald in Queensland, Balaklava in South Australia, and
 192 Junee in New South Wales. The soils at Gatton, Kingsthorpe and Emerald were self-mulching, black-
 193 cracking clay Vertosols with high water-holding capacity. At Balaklava, the soil was a hard-setting,
 194 red-brown duplex soil with a sandy loam texture, and at Junee soil was a free-draining, red
 195 gradational loam (Mitchell et al., 2012). Long-term average annual rainfall (1962–2018) was 776.2
 196 mm in Gatton (standard deviation = 207.7 mm), 687.9 mm in Kingsthorpe (standard deviation =
 197 179.3 mm), 607.7 mm Emerald (standard deviation = 205.0 mm), 331.6 mm in Balaklava (standard
 198 deviation = 81.3 mm), and 519.4 mm in Junee (standard deviation = 147.9 mm).

199 The slopes of the relationships between simulated and observed values were compared using 95%
 200 confidence intervals calculated from the standard error (Lentner et al., 1982). Statistical analyses
 201 were performed and graphs were produced using R software (v 3.0.3) (R Core Team, 2000).
 202 Geospatial analyses were performed using ESRI ArcMap 10.6 (ESRI 2018. ArcGIS Desktop: Release
 203 10. Redlands, CA: Environmental Systems Research Institute). Water Use Efficiency (WUE) was
 204 defined as grain yield per unit water supply, calculated by dividing grain yield in kg ha⁻¹ by water
 205 supply in mm (water supply = seasonal rainfall + initial plant available water content in soil).

206

207 **The APSIM model long-term analyses at Horsham and other validation sites**

208 The APSIM model was used to investigate the productivity change resulting from the reduced-
209 tillering trait in wheat at the AGFACE (Horsham) and other validation sites (Mitchell et al., 2012),
210 using SILO gridded daily climate data (Jeffrey et al., 2001) from 1962 to 2018. The long-term
211 modelling conducted at ambient and elevated [CO₂] of 380 and 550 μmol mol⁻¹, respectively for the
212 historic 57-year sequence (“historic climate”), plus two additional climate sequences of “historic
213 climate + 2°C warmer”, created by increasing the daily average temperature by 2°C across the 57-
214 year period, and “historic climate + 2°C warmer + 20% less rainfall”, created by increasing the daily
215 average temperature by 2°C and decreasing daily rainfall by 20% over the 57-year period. These
216 changes were selected to approximate a warmer and drier climate expected by 2050 (Christy et al.,
217 2018). Silverstar and Silverstar + *tin* sown each year on the same day after the autumn-break,
218 defined as at least 10 mm rainfall in a 5-day period between 14 April and 30 June. In total, 684
219 simulations were conducted for each location (2 genotypes × 2 [CO₂] × 3 climate scenarios × 57
220 years).

221

222 **The APSIM model spatial analysis across Australia**

223 The long-term analyses at the validation sites were extended across all privately owned, arable
224 agricultural land in Australia, the spatial region identified in Fig. 1. The spatial area was divided into
225 0.05° × 0.05° grid cells for modelling. For each grid cell within this region, the APSIM model was run
226 for 2 genotypes × 2 [CO₂] × 3 climate scenarios × 57 years. Upscaling to a total evaluated area of
227 68,083,675 ha, the APSIM model simulated 15,394,788 site-years of wheat growth (1962–2018).
228 Daily climate data for each grid cell were sourced from the SILO gridded daily climate data available
229 for each 0.05° × 0.05° across Australia (Jeffrey et al., 2001). Soil data for each grid cell were sourced
230 from the Soil and Landscape Grid of Australia (Grundy et al., 2015). The average annual crop yield
231 potential over the 57-year simulation period at each site and sowing time was based on the total
232 yield for each genotype divided by the number of crops sown in 57 years. To have a realistic
233 comparative analysis across the landscape, all forms of post sowing crop failure were included in the
234 calculation of average annual crop yield (Christy et al., 2018).

235

236 **Results**

237 **Model performance against experimental data**

238 Testing of observed (Table 1 and Mitchell et al. 2012) vs. simulated values indicated that the model
239 was able to accurately reproduce the observed phenology (no difference between the two
240 genotypes were observed, data not shown), tillers number (Fig. 3c), biomass (Fig. 3b), and grain yield
241 (Fig. 3a) for Silverstar and Silverstar + *tin* under ambient and elevated [CO₂]. The slopes of the
242 simulated vs. observed responses were near unity with a calculated root mean square error (RMSE)
243 ranging from 34 to 75 tiller m⁻² for number of tillers at anthesis, from 23 to 53 tiller m⁻² for number
244 of tillers at maturity, from 412 to 672 kg ha⁻¹ for biomass at anthesis, from 827 to 1730 kg ha⁻¹ for
245 biomass at maturity, and from 127 to 710 kg ha⁻¹ for grain yield for the two genotypes under
246 ambient and elevated [CO₂]. The parameter estimates for the slope of the relationship between
247 observed and simulated values were not significantly different from each other at any of the
248 measurement dates, i.e. the model accuracy in reproducing the two genotypes and CO₂ conditions
249 were statistically similar.

250 The lack of a significant treatment effect of the + *tin* gene raises questions as to the usefulness of the
251 gene at the Horsham experimental site or the suitability of the site to express any measurable
252 benefit. The latter appears more likely because in the years that the experiments were conducted
253 the annual rainfall + irrigation ranged from about 300 to 650 mm (Table 1), which is not considered
254 very dry. The impact of drier seasons across Australia was therefore explored with simulation
255 modelling.

256

257 **Long-term responses at Horsham and other validation sites**

258 The trait difference between the two genotypes (Silverstar and Silverstar + *tin*) was simulated over
259 57 years of present and future climate scenarios at Horsham and other validation sites (Fig. 4 and
260 Fig. 5). Silverstar + *tin* produced a greater grain yield than Silverstar in 38% of the site-years across all
261 sites and climate scenarios (Fig. 5). These yield advantages however did not result in a greater long-
262 term average yield of Silverstar + *tin* over Silverstar at any of the simulated sites and climate
263 scenarios, except for Emerald where the long-term average yields were greater by 19 to 78 kg ha⁻¹
264 yr⁻¹.

265

266 **Spatial analysis across Australia's arable land**

267 The application of the model across all privately owned arable agricultural land in Australia showed
268 an average yield advantage of 36.1 kg ha⁻¹ yr⁻¹ in Silverstar + *tin* over Silverstar in 26% of the total
269 evaluated area under the present-day climate (averaged for 57 years, Fig. 7 and Fig. 8, Table 2).

270 Under e[CO₂] conditions, grain yield of Silverstar + *tin* was greater than that of Silverstar in 18% of
 271 the total evaluated area but the average yield advantage was greater (43.4 kg ha⁻¹ yr⁻¹) (Fig. 8 and
 272 Table 2). The size of the area where Silverstar + *tin* had a greater yield than Silverstar and the size of
 273 the effect were least under + 2°C warmer climate but greatest under + 2°C warmer + 20% less
 274 rainfall (Fig. 8 and Table 2). These yield advantages were related to water supply, defined as growing
 275 season rainfall plus initial plant available water in soil. On average, the benefit seemed to be greater
 276 at growing seasons with water supply of < 184-185 mm under the present-day climate (a[CO₂] (a)
 277 and e[CO₂] (b), Fig. 6), < 165-167 mm under the + 2°C warmer (a[CO₂] (c) and e[CO₂] (d), Fig. 6), <
 278 153-156 mm under the + 2°C warmer + 20% less rainfall (a[CO₂] (e) and e[CO₂] (f), Fig. 6), and
 279 completely disappeared at growing seasons with water supply of greater than ~1000 mm (Fig. 6).

280

281 **TABLE 2** Grain yield in the areas where long-term average yield of Silverstar + *tin* is greater than
 282 Silverstar (areas coloured in green in Fig. 8). (a) historic climate and a[CO₂] (380 μmol mol⁻¹), (b)
 283 historic climate and e[CO₂] (550 μmol mol⁻¹), (c) historic climate + 2°C warmer and a[CO₂], (d) historic
 284 climate + 2°C warmer and e[CO₂], (e) historic climate + 2°C warmer + 20% less rainfall and a[CO₂],
 285 and (f) historic climate + 2°C warmer + 20% less rainfall and e[CO₂]. Water Use Efficiency (WUE) was
 286 defined as grain yield per unit water supply, calculated by dividing grain yield in kg ha⁻¹ by plant
 287 available soil water (mm) at sowing + growing season rainfall from sowing to physiological maturity.
 288 ±: standard deviation. AUD differences are based on the average area sown to wheat in Australia (=
 289 12.97 M ha yr⁻¹) and average price of AUD 260 t⁻¹ (2011-2016).

	a	b	c	d	e	f
Area (M ha)	18.14	12.20	15.26	11.49	26.89	23.77
Average water supply (mm)	266±48	253±39	258±57	242±44	218±54	208±43
Average yield in Silverstar + <i>tin</i> (kg ha ⁻¹ yr ⁻¹)	3530±717	3661±695	4263±782	2860±808	3050±675	3483±888
Average yield advantage of Silverstar + <i>tin</i> (kg ha ⁻¹ yr ⁻¹)	36.1±26.3	43.4±34.7	31.6±23.1	41.4±30.8	39.9±28.6	51.0±35.2
Average WUE in Silverstar + <i>tin</i> (kg ha ⁻¹ mm ⁻¹ yr ⁻¹)	13.39±2.08	17.02±2.72	11.91±1.59	15.26±2.13	13.13±2.36	16.78±2.95
Average WUE advantage of Silverstar + <i>tin</i> (kg ha ⁻¹ mm ⁻¹ yr ⁻¹)	0.14±0.12	0.19±0.18	0.13±0.09	0.18±0.15	0.19±0.13	0.26±0.19
AUD differences attributed to + <i>tin</i> genetic coefficient (\$MIL)	32.5	26.2	23.9	23.6	53.2	60.1

290

291 Discussion

292 We used a spatial modelling approach to compare grain yields of two genotypes with the same
 293 genetic background but contrasting in the expression of the reduced-tillering trait under different
 294 climate change scenarios across Australia's arable land. Our study region represents an important
 295 proportion of global wheat production and is typical of many dryland cropping environments
 296 throughout the world (e.g., CIMMYT Mega environment 1, 2, 4 and 8, Braun et al. (1996))
 297 experiencing significant changes in climate (Christy et al., 2018). The seasonal variability in climatic

298 variables (e.g. water availability and temperature) and sowing times in the spatial analysis allowed
299 evaluating the reduced-tillering trait in a range of different water-limited and favourable
300 environments. Our results show that the contribution of the reduced-tillering trait to grain yield is
301 strongly influenced by environment. Due to the complex physiological processes and their
302 interactions during the growing period for yield formation, the reduced-tillering trait may not be
303 beneficial when all growing seasons are considered but becomes important in low-yielding, water-
304 limited seasons (Fig. 6). Accounting for the long-term variability of rainfall, the reduced-tillering trait
305 showed a greater yield in 26% of the total evaluated area of 68.01 M ha, under the present-day
306 climate (Fig. 8 and Table 2). Genetic gain in dry environments is less reliable than in wetter regions
307 or where irrigation is available (Richards et al., 2002), because the variable environmental conditions
308 allow the most favourable environment \times genotype interaction only in a fraction of growing season.
309 Data from previous studies suggest an average overall rate increase of 18 kg ha⁻¹ yr⁻¹ by Australian
310 wheat genotypes from 1958 to 2007 (Sadras and Lawson, 2013). Our data suggests that under the
311 present-day climate the reduced-tillering trait has the potential to improve yield by 36 kg ha⁻¹ on 3.4
312 M ha of the total 12.97 M ha sown to wheat in Australia each year (equals to a potential annual cost-
313 benefit of AUD 32 MIL, see Table 2). The grain yield improvements reported herein and in Sadras
314 and Lawson (2013) are for attainable yields achievable through skilful use of the best available
315 technology (Connor et al., 2011).

316 Growth under the most likely future conditions (e[CO₂] + 2°C warmer + 20% less rainfall) advantage
317 the reduced-tillering trait compared to the present in a larger area, but temperature increase and
318 e[CO₂] in general mitigates that advantage somewhat (Fig. 8 and Table 2). In our simulations we
319 assumed a uniform temperature increase throughout the growing season. Water-saving effects of
320 e[CO₂] on transpiration efficiency may compensate for the increased evaporative demand caused by
321 this temperature increase, but this compensation seemed insufficient to overcome an additional
322 20% decrease in rainfall. Our extrapolations confirm the trait value for particularly water-limited
323 conditions, as predicted in previous evaluations of the reduced-tillering trait (e.g. Houshmandfar et
324 al. (2019)). The interactions between e[CO₂] and transpiration efficiency are not straightforward, as
325 potential water saving effects are dependent on timing and extent of drought and rainfall events
326 (Houshmandfar et al., 2016; Tausz-Posch et al., 2019). Our modelled results capture these
327 interactions well where they depend on the relationship between biomass growth, water supply and
328 transpiration efficiency, and are robust in the extrapolation of the relative differences between the
329 cultivars (Zhao et al., 2019).

330 We used a number of different lines that were all derived by backcrossing a *tin* donor to the spring
331 wheat variety Silverstar (Mitchell et al., 2013), to validate the model employing the exact same

332 setup. Tiller production is dependent on genetics-by-environment-by-management interactions
333 (Innes et al., 1981). The presence of the *tin* gene provides a genetic control on the potential number
334 of tillers produced by a plant, but this varied from line to line depending on unknown modifier genes
335 controlling the level of *tin* expression to affect tiller numbers (Mitchell et al., 2013). Reduced-tillering
336 lines can be therefore classified as restricted or semi-restricted lines (Mitchell et al., 2013), based on
337 the maximum number of tillers produced per plant (Mitchell, 2010). The lines we employed to
338 validate the model (viz. SsrT65, SsrT02, SsrT14 and SsrT17) were all restricted tillering lines, each
339 producing a maximum of 4 tillers plant⁻¹ (Mitchell, 2010).

340 In conclusion, we suggest a small but consistent yield advantage conferred by the reduced-tillering
341 trait in the most water-limited environments both under current and likely future conditions. Our
342 climate scenarios show that whilst increasing [CO₂] alone might limit the area where the reduced-
343 tillering trait is advantageous, the most likely climate scenario with CO₂ combined with increased
344 temperature and reduced rainfall consistently increased the area where restricted tillering has an
345 advantage. Whilst yield advantages are small, it seems worthwhile to further explore this reduced-
346 tillering trait in relation to a range of different environments and climates, because its benefits are
347 likely to grow in future dry environments.

348

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362

363 **Data Availability Statement**

364 The data that support the findings of this study are available from the corresponding author upon
365 reasonable request.

366

367 **References**

368 Ainsworth, E. A. & Rogers, A. (2007). The response of photosynthesis and stomatal conductance to
369 rising [CO₂]: mechanisms and environmental interactions. *Plant, Cell & Environment*, 30(3),
370 258-270. <https://doi.org/10.1111/j.1365-3040.2007.01641.x>

371 Asseng, S. & Turner, N. (2007). Modelling genotype × environment × management interactions to
372 improve yield, water use efficiency and grain protein in wheat. *Frontis*, 91-102.
373 https://doi.org/10.1007/1-4020-5906-x_8

374 Atsmon, D. & Jacobs, E. (1977). A Newly Bred 'Gigas' Form of Bread Wheat (*Triticum aestivum*:
375 Morphological Features and Thermo-photoperiodic Responses 1. *Crop Science*, 17, 31-35.

376 Berry, P., Spink, J., Foulkes, M. & Wade, A. (2003). Quantifying the contributions and losses of dry
377 matter from non-surviving shoots in four cultivars of winter wheat. *Field Crops Research*,
378 80(2), 111-121. [https://doi.org/10.1016/s0378-4290\(02\)00174-0](https://doi.org/10.1016/s0378-4290(02)00174-0)

379 Braun, H. J., Rajaram, S. & van Ginkel, M. (1996). CIMMYT's approach to breeding for wide
380 adaptation. *Euphytica*, 92(1-2), 175-183. https://doi.org/10.1007/978-94-015-8806-5_25

381 Brown, H. E., Huth, N. I., Holzworth, D. P., Teixeira, E. I., Zyskowski, R. F., Hargreaves, J. N., & Moot,
382 D. J. (2014). Plant modelling framework: software for building and running crop models on
383 the APSIM platform. *Environmental Modelling & Software*, 62, 385-398.
384 <https://doi.org/10.1016/j.envsoft.2014.09.005>

385 Christy, B., Tausz-Posch, S., Tausz, M., Richards, R., Rebetzke, G., Condon, A., ... & O'Leary, G. (2018).
386 Benefits of increasing transpiration efficiency in wheat under elevated CO₂ for rainfed
387 regions. *Global Change Biology*, 24(5), 1965-1977. <https://doi.org/10.1111/gcb.14052>

388 Connor, D. J., Loomis, R. S., & Cassman, K. G. (2011). *Crop ecology: productivity and management in*
389 *agricultural systems*. Cambridge University Press.
390 <https://doi.org/10.1017/cbo9780511974199>

391 Dawson, J. A. (1957). The productivity of water in agriculture. *Journal of Farm Economics*, 39(5),
392 1244-1252. <https://doi.org/10.2307/1234683>

393 Duggan, B., Richards, R., Van Herwaarden, A. & Fettell, N. (2005). Agronomic evaluation of a tiller
394 inhibition gene (*tin*) in wheat. I. Effect on yield, yield components, and grain protein.
395 *Australian Journal of Agricultural Research*, 56(2), 169-178.
396 <https://doi.org/10.1071/ar04152>

- 397 Evers, J. B., Vos, J., Andrieu, B. & Struik, P. C. (2006). Cessation of tillering in spring wheat in relation
398 to light interception and red: far-red ratio. *Annals of Botany*, 97(4), 649-658.
399 <https://doi.org/10.1093/aob/mcl020>
- 400 Fitzgerald, G. J., Tausz, M., O'Leary, G., Mollah, M. R., Tausz-Posch, S., Seneweera, S., ... & Norton, R.
401 M. (2016). Elevated atmospheric [CO₂] can dramatically increase wheat yields in semi-arid
402 environments and buffer against heat waves. *Global Change Biology*, 22(6), 2269-2284.
403 <https://doi.org/10.1111/gcb.13263>
- 404 Grundy, M. J., Rossel, R. V., Searle, R. D., Wilson, P. L., Chen, C., & Gregory, L. J. (2015). Soil and
405 landscape grid of Australia. *Soil Research*, 53(8), 835-844. <https://doi.org/10.1071/sr15191>
- 406 He, D., Wang, E., Wang, J. & Lilley, J.M. (2017). Genotype × environment × management interactions
407 of canola across China: A simulation study. *Agricultural and Forest Meteorology*, 247, 424-
408 433. <https://doi.org/10.1016/j.agrformet.2017.08.027>
- 409 Hendriks, P. W., Kirkegaard, J. A., Lilley, J. M., Gregory, P. J. & Rebetzke, G.J. (2015). A tillering
410 inhibition gene influences root–shoot carbon partitioning and pattern of water use to
411 improve wheat productivity in rainfed environments. *Journal of Experimental Botany*,
412 67(1), 327-340. <https://doi.org/10.1093/jxb/erv457>
- 413 Hochman, Z., Gobbett, D.L. & Horan, H. (2017). Climate trends account for stalled wheat yields in
414 Australia since 1990. *Global Change Biology*, 23(5), 2071-2081.
415 <https://doi.org/10.1111/gcb.13604>
- 416 Holzworth, D., Huth, N. I., Fainges, J., Brown, H., Zurcher, E., Cichota, R., ... & Snow, V. (2018). APSIM
417 Next Generation: Overcoming challenges in modernising a farming systems model.
418 *Environmental Modelling & Software*, 103, 43-51.
419 <https://doi.org/10.1016/j.envsoft.2018.02.002>
- 420 Holzworth, D. P., Huth, N. I., deVoil, P. G., Zurcher, E. J., Herrmann, N. I., McLean, G., ... & Moore, A.
421 D. (2014). APSIM—evolution towards a new generation of agricultural systems
422 simulation. *Environmental Modelling & Software*, 62, 327-350.
423 <https://doi.org/10.1016/j.envsoft.2014.07.009>
- 424 Houshmandfar, A., Fitzgerald, G. J., Armstrong, R., Macabuhay, A. A. & Tausz, M. (2015). Modelling
425 stomatal conductance of wheat: An assessment of response relationships under elevated
426 CO₂. *Agricultural and Forest Meteorology*, 214, 117-123.
427 <https://doi.org/10.1016/j.agrformet.2015.08.249>
- 428 Houshmandfar, A., Fitzgerald, G. J., Macabuhay, A. A., Armstrong, R., Tausz-Posch, S., Löw, M., &
429 Tausz, M. (2016). Trade-offs between water-use related traits, yield components and

430 mineral nutrition of wheat under Free-Air CO₂ Enrichment (FACE). *European Journal of*
431 *Agronomy*, 76, 66-74. <https://doi.org/10.1016/j.eja.2016.01.018>

432 Houshmandfar, A., Rebetzke, G. J., Lawes, R. & Tausz, M. (2019). Grain yield responsiveness to water
433 supply in near-isogenic reduced-tillering wheat lines—An engineered crop trait near its
434 upper limit. *European Journal of Agronomy*, 102, 33-38.
435 <https://doi.org/10.1016/j.eja.2018.11.003>

436 Hutchinson, M. F., McIntyre, S., Hobbs, R. J., Stein, J. L., Garnett, S., & Kinloch, J. (2005). Integrating a
437 global agro-climatic classification with bioregional boundaries in Australia. *Global Ecology*
438 *and Biogeography*, 14(3), 197-212. <https://doi.org/10.1111/j.1466-822x.2005.00154.x>

439 Innes, P., Blackwell, R., Austin, R. & Ford, M. A. (1981). The effects of selection for number of ears on
440 the yield and water economy of winter wheat. *The Journal of Agricultural Science*, 97(3),
441 523-532. <https://doi.org/10.1017/s0021859600036844>

442 Isbell, R. (2016). *The Australian soil classification*. CSIRO publishing.
443 <https://doi.org/10.1071/9781486304646>

444 Jeffrey, S. J., Carter, J. O., Moodie, K. B. & Beswick, A. R. (2001). Using spatial interpolation to
445 construct a comprehensive archive of Australian climate data. *Environmental Modelling &*
446 *Software*, 16(4), 309-330. [https://doi.org/10.1016/s1364-8152\(01\)00008-1](https://doi.org/10.1016/s1364-8152(01)00008-1)

447 Leakey, A. D., Ainsworth, E. A., Bernacchi, C. J., Rogers, A., Long, S. P., & Ort, D. R. (2009). Elevated
448 CO₂ effects on plant carbon, nitrogen, and water relations: six important lessons from
449 FACE. *Journal of Experimental Botany*, 60(10), 2859-2876.
450 <https://doi.org/10.1093/jxb/erp096>

451 Lentner, C., Diem, K., & Seldrup, J. (1982). Introduction to statistics. Statistical tables. Mathematical
452 formulae. *Geigy Scientific Tables*, 2, 215-218.

453 Löw, M., Tausz-Posch, S., Rebetzke, G., Dreccer, M. F., Chapman, S. C., Seneweera, S., ... & Tausz, M.
454 (2015). Is a reduced-tillering trait (*tin*) beneficial under elevated CO₂ in four FACE
455 environments? In *Building Productive, Diverse and Sustainable Landscapes, 17th Australian*
456 *Agronomy Conference, 20-24 September 2015, Hobart, Australia. Conference*
457 *Proceedings* (pp. 733-736). Australian Society of Agronomy Inc.

458 Mitchell, J. (2010). *Evaluation of reduced-tillering (tin gene) wheat lines for water limiting*
459 *environments in Northern Australia*. PhD Thesis: The University of Queensland.

460 Mitchell, J., Chapman, S., Rebetzke, G., Bonnett, D. & Fukai, S. (2012). Evaluation of a reduced-
461 tillering (*tin*) gene in wheat lines grown across different production environments. *Crop*
462 *and Pasture Science*, 63(2), 128-141. <https://doi.org/10.1071/cp11260>

463 Mitchell, J., Rebetzke, G., Chapman, S. & Fukai, S. (2013). Evaluation of reduced-tillering (*tin*) wheat
464 lines in managed, terminal water deficit environments. *Journal of Experimental Botany*,
465 64(11), 3439-3451. <https://doi.org/10.1093/jxb/ert181>

466 Moeller, C. & Rebetzke, G. (2017). Performance of spring wheat lines near-isogenic for the reduced-
467 tillering '*tin*' trait across a wide range of water-stress environment-types. *Field Crops*
468 *Research*, 200, 98-113. <https://doi.org/10.1016/j.fcr.2016.10.010>

469 Mollah, M., Norton, R. & Huzzey, J. (2009). Australian grains free-air carbon dioxide enrichment
470 (AGFACE) facility: design and performance. *Crop and Pasture Science*, 60(8), 697-707.
471 <https://doi.org/10.1071/cp08354>

472 O'Leary, G. J., Christy, B., Nuttall, J., Huth, N., Cammarano, D., Stöckle, C., ... & Farre-Codina, I.
473 (2015). Response of wheat growth, grain yield and water use to elevated CO₂ under a
474 Free-Air CO₂ Enrichment (FACE) experiment and modelling in a semi-arid
475 environment. *Global Change Biology*, 21(7), 2670-2686. <https://doi.org/10.1111/gcb.12830>

476 Team, R. C. (2000). R language definition. *Vienna, Austria: R foundation for statistical computing*.

477 Reynolds, M., Foulkes, M. J., Slafer, G. A., Berry, P., Parry, M. A., Snape, J. W., & Angus, W. J. (2009).
478 Raising yield potential in wheat. *Journal of Experimental Botany*, 60(7), 1899-1918.
479 <https://doi.org/10.1093/jxb/erp016>

480 Richards, R. (1988). A tiller inhibitor gene in wheat and its effect on plant growth. *Australian Journal*
481 *of Agricultural Research*, 39(5), 749-757. <https://doi.org/10.1071/ar9880749>

482 Richards, R., Rebetzke, G., Condon, A. & Van Herwaarden, A. (2002). Breeding opportunities for
483 increasing the efficiency of water use and crop yield in temperate cereals. *Crop Science*,
484 42(1), 111-121. <https://doi.org/10.2135/cropsci2002.0111>

485 Riffkin, P. A., Evans, P. M., Chin, J. F. & Kearney, G. A. (2003). Early-maturing spring wheat
486 outperforms late-maturing winter wheat in the high rainfall environment of south-western
487 Victoria. *Australian Journal of Agricultural Research*, 54(2), 193-202.
488 <https://doi.org/10.1071/ar02081>

489 Sadras, V. & Lawson, C. (2013). Nitrogen and water-use efficiency of Australian wheat varieties
490 released between 1958 and 2007. *European Journal of Agronomy*, 46, 34-41.
491 <https://doi.org/10.1016/j.eja.2012.11.008>

492 Sadras, V. & Rebetzke, G. (2013). Plasticity of wheat grain yield is associated with plasticity of ear
493 number. *Crop and Pasture Science*, 64(3), 234-243. <https://doi.org/10.1071/cp13117>

494 Sharma, R. (1995). Tiller mortality and its relationship to grain yield in spring wheat. *Field Crops*
495 *Research*, 41(1), 55-60. [https://doi.org/10.1016/0378-4290\(94\)00109-p](https://doi.org/10.1016/0378-4290(94)00109-p)

- 496 Solomon, S., Manning, M., Marquis, M., & Qin, D. (2007). *Climate change 2007-the physical science*
497 *basis: Working group I contribution to the fourth assessment report of the IPCC* (Vol. 4).
498 Cambridge university press.
- 499 Stephens, D. J. & Lyons, T. J. (1997). Rainfall-yield relationships across the Australian wheatbelt.
500 *Australian Journal of Agricultural Research*, 49(2), 211-224.
501 <https://doi.org/10.1071/a96139>
- 502 Tausz-Posch, S., Tausz, M. & Bourgault, M. (2019). Elevated [CO₂] effects on crops: Advances in
503 understanding acclimation, nitrogen dynamics and interactions with drought and other
504 organisms. *Plant Biology*, 22, 38-51. <https://doi.org/10.1111/plb.12994>
- 505 Wang, B., Liu, D. L., O'Leary, G. J., Asseng, S., Macadam, I., Lines-Kelly, R., ... & Xing, H. (2018).
506 Australian wheat production expected to decrease by the late 21st century. *Global Change*
507 *Biology*, 24(6), 2403-2415. <https://doi.org/10.1111/gcb.14034>
- 508 Wullschleger, S., Tschaplinski, T. & Norby, R. (2002). Plant water relations at elevated CO₂–
509 implications for water-limited environments. *Plant, Cell & Environment*, 25(2), 319-331.
510 <https://doi.org/10.1046/j.1365-3040.2002.00796.x>
- 511 Zadoks, J. C., Chang, T. T. & Konzak, C. F. (1974). A decimal code for the growth stages of cereals.
512 *Weed Research*, 14(6), 415-421. <https://doi.org/10.1111/j.1365-3180.1974.tb01084.x>
- 513 Zhao, Z., Rebetzke, G. J., Zheng, B., Chapman, S. C. & Wang, E. (2019). Modelling impact of early
514 vigour on wheat yield in dryland regions. *Journal of Experimental Botany*, 70(9), 2535-2548.
515 <https://doi.org/10.1093/jxb/erz069>

516

517 **Figure legends**

518

519 **Figure 1** Long-term average (1962–2018) annual rainfall (mm) within Australia's arable land for the
520 two wheat genotypes and the location of the experimental sites used to validate the APSIM model.
521 Bar graphs show the long-term average monthly rainfall from January to December, respectively.

522

523 **Figure 2** Daily min (black dashed line) and max (black solid line) temperatures (°C) as well as rainfall
524 (grey solid line) (mm) in 2011 (a) and 2012 (b) at the AGFACE (Horsham).

525

526 **Figure 3** Simulated vs. observed values of Silverstar (●○●○) and Silverstar + *tin* (■□■□). (a) grain
527 yield data from the AGFACE (Horsham) site at a[CO₂] (□○) and e[CO₂] (■●), and from the other
528 validation sites at a[CO₂] only (□○) (Mitchell et al., 2013). (b) biomass at anthesis (■□●○) and at

529 physiological maturity (■□●○) from the AGFACE site. (c) number of tillers at anthesis (■□●○)
530 and at physiological maturity (■□●○) from the AGFACE site (c). Tables on top of panels show
531 fitting parameter statistics for the linear fits to data subsets as indicated in the row titles to the left,
532 and in the first row of each table. RMSE: root mean squared error. SE: standard error. Dashed line is
533 1:1 line.

534

535 **Figure 4** Grain yield (kg ha^{-1}) for (a) Silverstar + *tin* at Horsham, (b) Silverstar + *tin* at Balaklava, (c)
536 Silverstar + *tin* at Emerald, (d) Silverstar at Horsham, (e) Silverstar at Balaklava, (f) Silverstar at
537 Emerald, (g) Silverstar + *tin* at Gatton, (h) Silverstar + *tin* at Junee, (i) Silverstar + *tin* at Kingsthorpe,
538 (j) Silverstar at Gatton, (k) Silverstar at Junee, and (l) Silverstar at Kingsthorpe showing lower (25th
539 percentile – $1.5 \times (75^{\text{th}} \text{ quantile} - 25^{\text{th}} \text{ quantile})$) and upper ($75^{\text{th}} \text{ percentile} + 1.5 \times (75^{\text{th}} \text{ quantile} -$
540 $25^{\text{th}} \text{ quantile})$) limits (whiskers), 25th-75th percentile (box) and median (horizontal line) over 57 years
541 (1962-2018) under (H) historic climate, (W) historic climate + 2°C warmer, and (D) historic climate +
542 2°C warmer + 20% less rainfall at a[CO₂] (380 $\mu\text{mol mol}^{-1}$) and e[CO₂] (550 $\mu\text{mol mol}^{-1}$). Asterisks are
543 points that fall outside the limits of the whiskers.

544

545 **Figure 5** Grain yield advantage (kg ha^{-1}) of Silverstar + *tin* over Silverstar showing lower (25th
546 percentile – $1.5 \times (75^{\text{th}} \text{ quantile} - 25^{\text{th}} \text{ quantile})$) and upper ($75^{\text{th}} \text{ percentile} + 1.5 \times (75^{\text{th}} \text{ quantile} -$
547 $25^{\text{th}} \text{ quantile})$) limits (whiskers), 25th-75th percentile (box), median (horizontal line) and mean (dot)
548 over 57 years (1962-2018) at (a) Horsham (AGFACE), (b) Balaklava, (c) Emerald, (d) Gatton, (e) Junee,
549 and (f) Kingsthorpe) under (H) historic climate, (W) historic climate + 2°C warmer, and (D) historic
550 climate + 2°C warmer + 20% less rainfall at a[CO₂] (380 $\mu\text{mol mol}^{-1}$) and e[CO₂] (550 $\mu\text{mol mol}^{-1}$).
551 Asterisks are points that fall outside the limits of the whiskers.

552

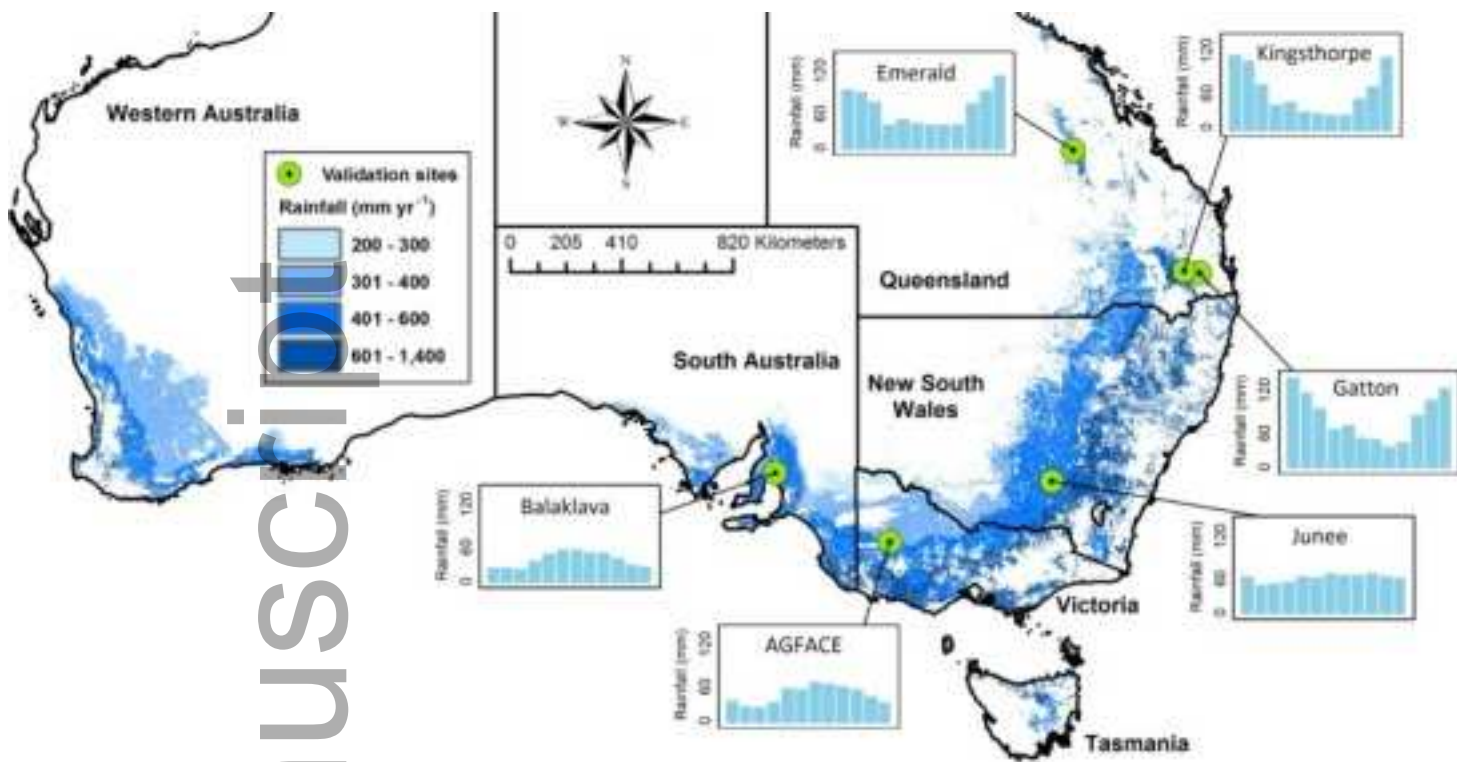
553 **Figure 6** Grain yield advantage (kg ha^{-1}) of Silverstar + *tin* over Silverstar in relation to water supply (<
554 3000 mm). Each point is coloured according to the percentage of results from simulations across
555 Australia's arable land (1962-2018) under (a) historic climate and a[CO₂] (380 $\mu\text{mol mol}^{-1}$), (b)
556 historic climate and e[CO₂] (550 $\mu\text{mol mol}^{-1}$), (c) historic climate + 2°C warmer and a[CO₂], (d)
557 historic climate + 2°C warmer and e[CO₂], (e) historic climate + 2°C warmer + 20% less rainfall and
558 a[CO₂], and (f) historic climate + 2°C warmer + 20% less rainfall and e[CO₂]. White points indicate
559 that there were no simulation results. Water supply is seasonal rainfall plus initial plant available
560 water content in soil.

561

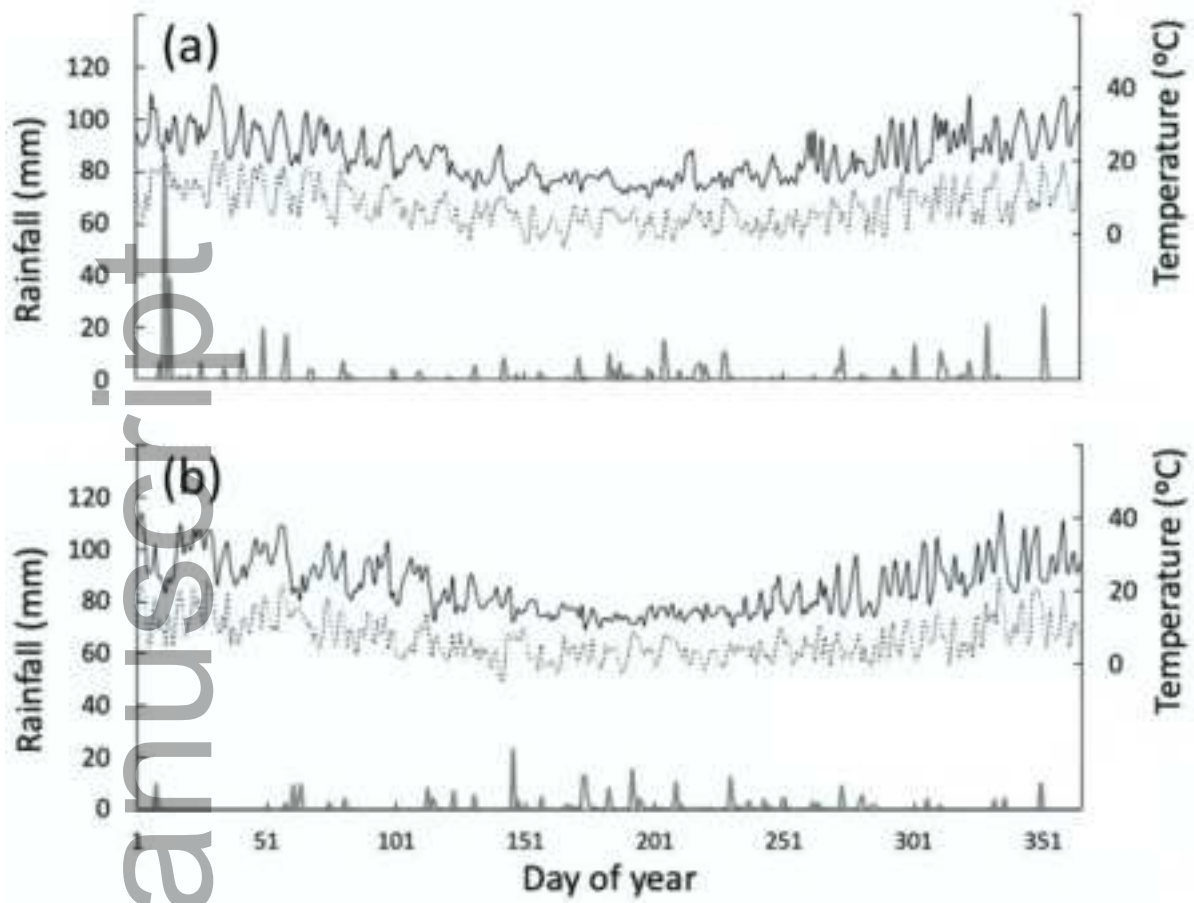
562 **Figure 7** Long-term average (1962-2018) grain yield ($\text{kg ha}^{-1} \text{ yr}^{-1}$) for (a) Silverstar + *tin* under historic
563 climate and $a[\text{CO}_2]$ ($380 \mu\text{mol mol}^{-1}$), (b) Silverstar + *tin* under historic climate and $e[\text{CO}_2]$ ($550 \mu\text{mol}$
564 mol^{-1}), (c) Silverstar under historic climate and $a[\text{CO}_2]$, (d) Silverstar under historic climate and
565 $e[\text{CO}_2]$, (e) Silverstar + *tin* under historic climate + 2°C warmer and $a[\text{CO}_2]$, (f) Silverstar + *tin* under
566 historic climate + 2°C warmer and $e[\text{CO}_2]$, (g) Silverstar under historic climate + 2°C warmer and
567 $a[\text{CO}_2]$, (h) Silverstar under historic climate + 2°C warmer and $e[\text{CO}_2]$, (i) Silverstar + *tin* under historic
568 climate + 2°C warmer + 20% less rainfall and $a[\text{CO}_2]$, (j) Silverstar + *tin* under historic climate + 2°C
569 warmer + 20% less rainfall and $e[\text{CO}_2]$, (k) Silverstar under historic climate + 2°C warmer + 20% less
570 rainfall and $a[\text{CO}_2]$, and (l) Silverstar under historic climate + 2°C warmer + 20% less rainfall and
571 $e[\text{CO}_2]$. Max and min are 0.01 and 0.99 percentiles of all long-term yields, respectively. Boxplots
572 represent frequently.

573

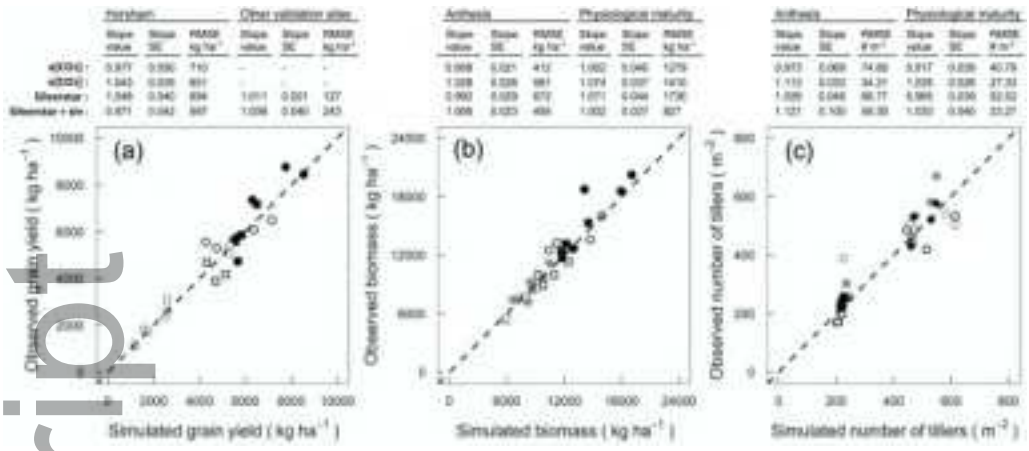
574 **Figure 8** Long-term average (1962-2018) yield advantage ($\text{kg ha}^{-1} \text{ yr}^{-1}$) of Silverstar + *tin* over
575 Silverstar under (a) historic climate and $a[\text{CO}_2]$ ($380 \mu\text{mol mol}^{-1}$), (b) historic climate and $e[\text{CO}_2]$ (550
576 $\mu\text{mol mol}^{-1}$), (c) historic climate + 2°C warmer and $a[\text{CO}_2]$, (d) historic climate + 2°C warmer and
577 $e[\text{CO}_2]$, (e) historic climate + 2°C warmer + 20% less rainfall and $a[\text{CO}_2]$, and (f) historic climate + 2°C
578 warmer + 20% less rainfall and $e[\text{CO}_2]$. Boxplots represent frequently.



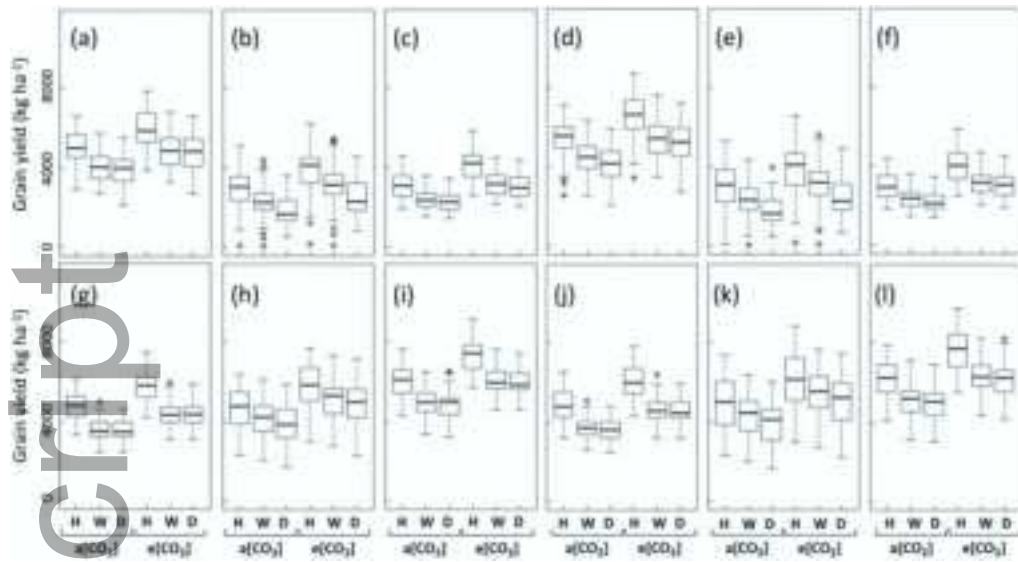
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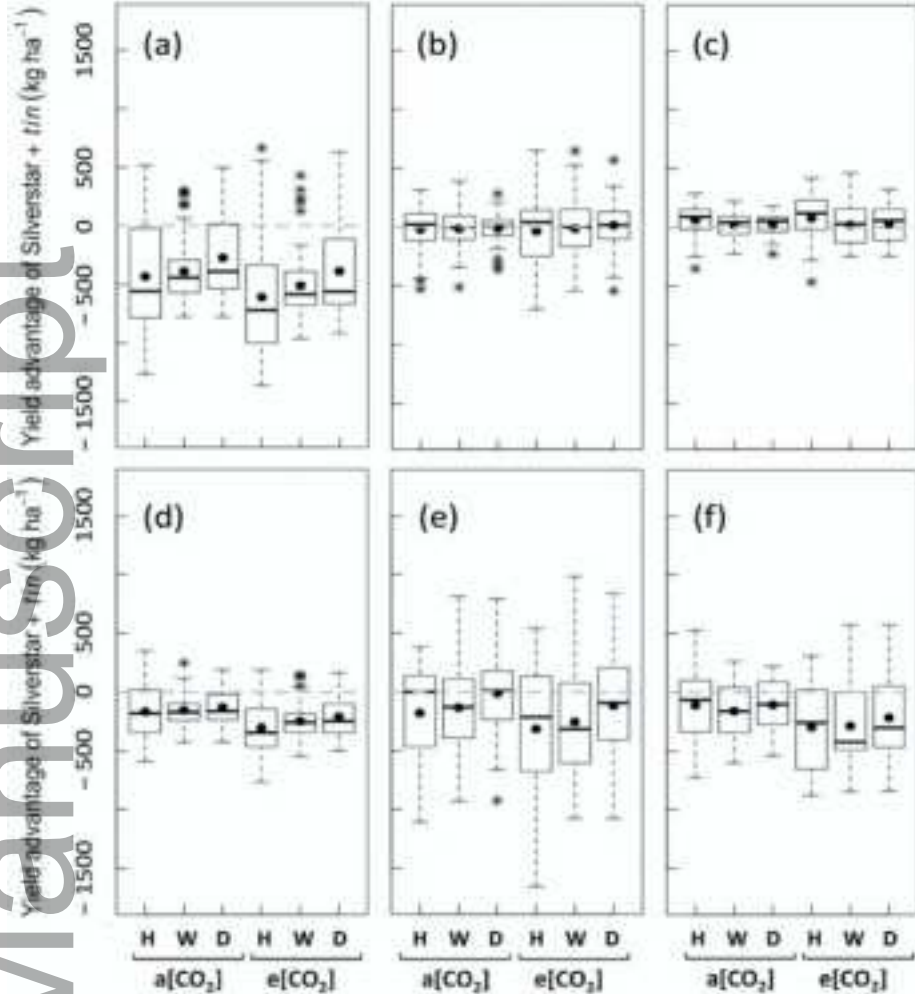
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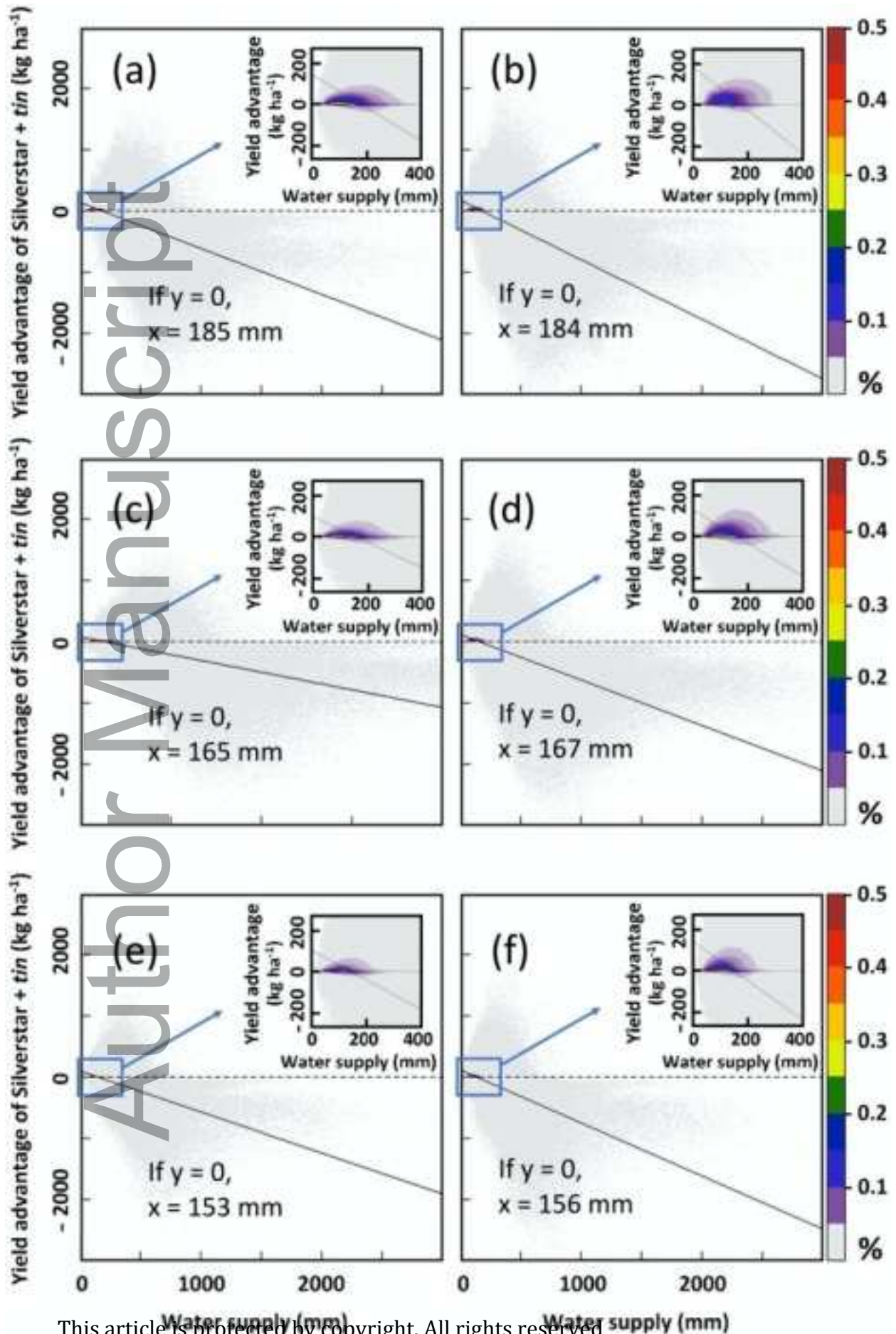
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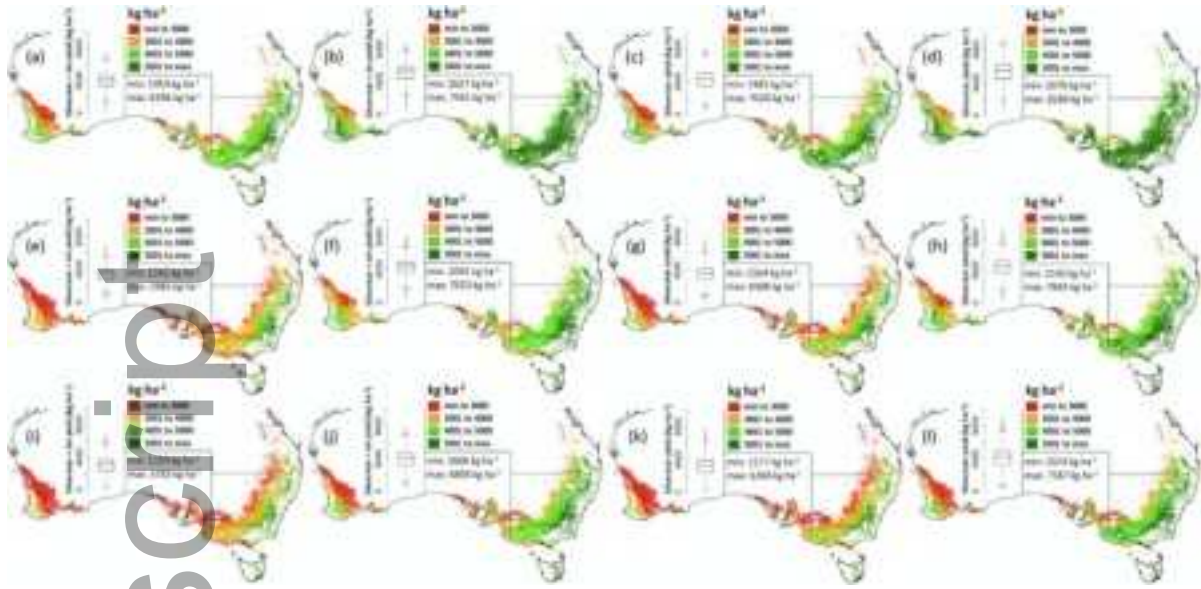
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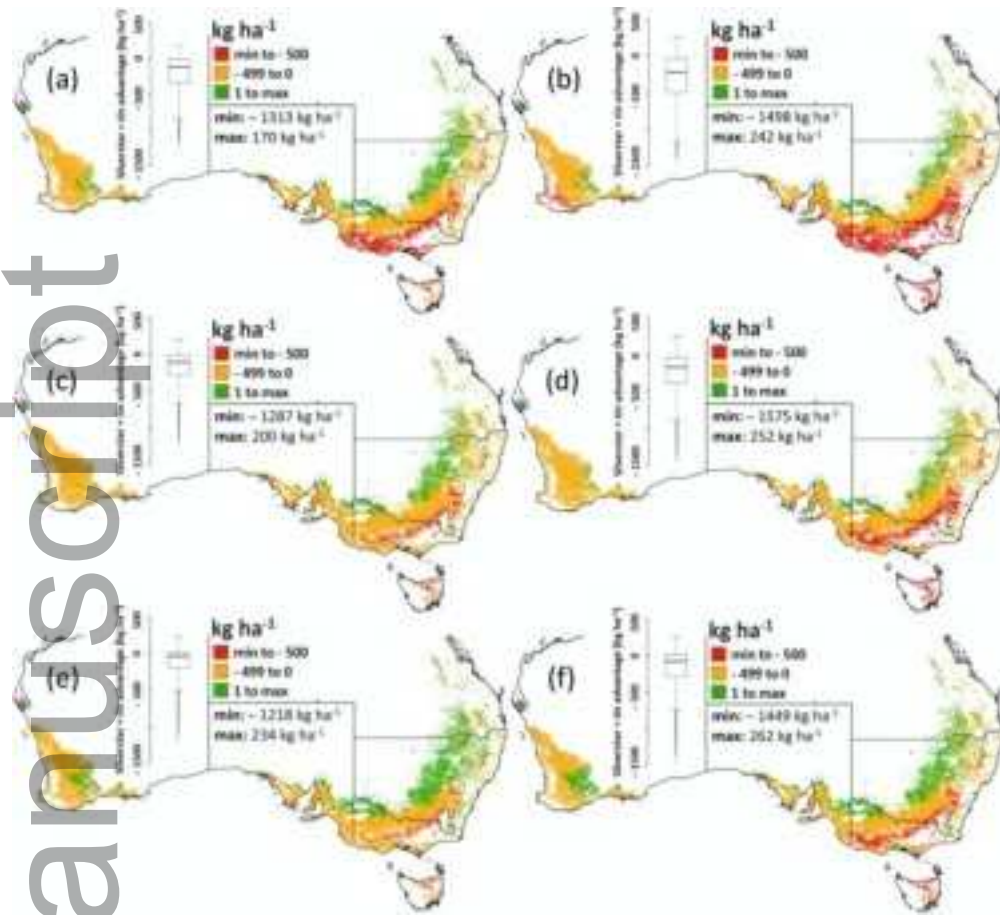
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