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Ingvordsen, Cathrine Heinz; Gislum, René; Jørgensen, Johannes Ravn; Mikkelsen, Teis Nørgaard; Stockmarr, Anders; Bagger Jørgensen, Rikke

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Grain protein under future climate conditions of elevated temperature and carbon dioxide. A study of 108 spring barley accessions

Cathrine H. Ingvordsen

cahi@kt.dtu.dk

Centre for Ecosystems and Environmental Sustainability, Department of Chemical and Biochemical Engineering, Technical University of Denmark, Frederiksborgvej 399, DK-4000 Roskilde, Denmark

Corresponding author. Tel.: +45 23669751; email address: cahi@kt.dtu.dk (C. H. Ingvordsen).

René Gislum

Rene.Gislum@agrsci.dk

Crop Health, Department of Agroecology, Flakkebjerg, Aarhus University, Forsøgsvej 1, DK-4200 Slagelse, Denmark

Johannes R. Jørgensen

Johannes.Jorgensen@agrsci.dk

Crop Health, Department of Agroecology, Flakkebjerg, Aarhus University, Forsøgsvej 1, DK-4200 Slagelse, Denmark

Teis N. Mikkelsen

temi@kt.dtu.dk

Centre for Ecosystems and Environmental Sustainability, Department of Chemical and Biochemical Engineering, Technical University of Denmark, Frederiksborgvej 399, DK-4000 Roskilde, Denmark

Rikke B. Jørgensen

rijq@kt.dtu.dk

Centre for Ecosystems and Environmental Sustainability, Department of Chemical and Biochemical Engineering, Technical University of Denmark, Frederiksborgvej 399, DK-4000 Roskilde, Denmark

Key words: *climate change, grain protein harvested (GPH), Hordeum vulgare, near-infrared spectroscopy, tropospheric ozone*

Abbreviations: GPC: grain protein concentration; GPH: grain protein harvested; GWAS: Genome Wide Association Studies; MSC: multiplicative signal corrected; NIR: Near infrared radiation; PLSR: partial least squares regression; RERAF: Risø Environmental RiskAssessment Facility; RMSECV: root mean square error of cross validation; RMSEP: root mean square error of prediction; SEP: standard error of performance

1 **Abstract**

2 Climate change is predicted to decrease future grain yields and influence grain protein
3 concentration. In the present study a set of 108 spring barley accessions were cultivated under
4 predicted future levels of temperature, [CO₂] and [O₃] as single-factors and temperature and [CO₂]
5 in combination (IPCC SRES scenario A1FI). The found 8 % increase in grain protein concentration
6 under the combined treatment could not be depicted from the single factor treatments. Ozone as
7 single factor increased grain protein concentration with 6 %. In a future scenario with projected
8 lowered grain yield, harvesting as much protein as possible seems desirable. Grain protein
9 harvested only increased under elevated [CO₂] and was lowered 23 % in the future climate scenario
10 of elevated temperature and [CO₂]. Vast variation in the response of the 108 accessions was
11 identified. This variation should be further exploited to increase the grain protein harvested under
12 future climate change conditions.

14 **Introduction**

15 Climate change, with increased atmospheric concentration of the greenhouse gasses carbon dioxide
16 ([CO₂]) and ozone ([O₃]) together with rising temperature, is likely to decrease plant production in
17 the future and influence grain protein and quality (Danielsson *et al.*, 1999; Lobell and Field, 2007;
18 Wang and Frei, 2011; Collins *et al.*, 2013; IPCC, 2014a). According to the latest projections by
19 IPCC (Intergovernmental Panel of Climate Change) climatic conditions point to the worst-case
20 scenario (RCP8.5) unless actions are taken in the near future (IPCC, 2014b). In the worst-case
21 scenario temperature is predicted to rise 5 °C and [CO₂] to reach 1000 ppm compared to the 400
22 ppm of today. The increase of [O₃] is expected at 25 % reaching 40-77 ppb (Collins *et al.*, 2013;
23 Ellermann *et al.*, 2013). Numerous experimental studies have demonstrated the effect on cereal
24 grain yield by elevated temperature, [CO₂] and [O₃] as single- factors; increasing production by
25 [CO₂] (Wang *et al.*, 2013) and decreasing production by temperature (Luo, 2011) and [O₃] (Feng
26 and Kobayashi, 2009). Less studies have reported the effect on grain yield by the combination of
27 climatic factors (Mittler, 2006; Frenck *et al.*, 2011; Pleijel and Uddling, 2012; Dias de Oliveira *et al.*,
28 *et al.*, 2013). In studies of elevated temperature and [CO₂] combined, grain yield was found to
29 decrease by 14-53 % (Batts *et al.*, 1998; Clausen *et al.*, 2011; Ingvordsen *et al.*, 2014). The decrease
30 in cereal grain yield with a global temperature increase of > +3 °C is critical in the context of

31 maintaining a sufficient primary production, which can meet the needs of a growing world
32 population and improved living standards.

33 Grain protein concentration (GPC) has been reported to increase in response to abiotic stress
34 such as heat, drought and elevated [O₃] (Savin and Nicolas, 1996; Passarella *et al.*, 2008; Asare *et*
35 *al.*, 2011; Pleijel and Uddling, 2012), while GPC was decreased by elevated [CO₂] (Högy and
36 Fangmeier, 2008). Timing of the climate effect in plant development was further found to influence
37 the response in GPC (Rotundo and Westgate, 2009; Wang and Frei, 2011). Grain protein is decisive
38 for several end-uses, however, the effects on GPC by combined climatic factors is little studied
39 even though factors of climate change will appear concerted (Wang and Frei, 2011; Högy *et al.*,
40 2013).

41 The fourth major cereal of the world is barley (*Hordeum vulgare* L), which in temperate climates
42 is cultivated predominantly as a spring crop for feed to livestock and malt for use in brewing and
43 distilling industries. Barley has though, within the recent years, achieved increased attention for
44 human consumption due to its high nutritional value and potential health benefits (Baik and Ullrich,
45 2008). The diverse uses of barley grains cause different demands to the grain composition.
46 Generally, high protein content is preferable in barley for feed, whereas a low protein grain and
47 high starch content is preferred for malting purposes. Climate change alterations in protein content
48 can in the industrialized countries in the temperate zone also have substantial marked implications
49 with economic and social consequences.

50 In the context of climate change with projected decreased grain yields (IPCC, 2014a), the grain
51 protein harvested (GPH) is important for product quality and secured primary production. Few
52 studies have focused on the impact of climate change on cereal grain protein determining for the
53 quality and even fewer in the context of GPH in barley. The objective of this study was to examine
54 climate change effects to an array of accessions. Here we present the effect on grain yield and GPC
55 under the combination of elevated [CO₂] and temperature as under the single-factors elevated
56 [CO₂], temperature and [O₃] on 108 spring barley accessions.

57

58

59 **Material and methods**

60

61 *Plant material*

62 One hundred and eight 2- or 6-rowed primarily Nordic spring barley (*Hordeum vulgare* L.)
63 accessions were included in the study (Table 1). The set included 38 landrace accessions, 25 old
64 cultivars (before 1975), 41 modern cultivars (after 1975) and four breeder-lines. Accessions were
65 supplied by NordGen or Nordic barley breeding companies. For pedigree, breeder institute, and
66 provider please see S1.

67

68 *Experimental set up*

69 Five climate treatments were applied in the RERAF phytotron (Risø Environmental Risk
70 Assessment Facility) at the Technical University of Denmark, Roskilde
71 (http://www.eco.kt.dtu.dk/Research/Research_Facilitites/RERAF) to all 108 accessions throughout
72 their full lifecycle. The 108 accessions were a subset of the 138 accessions analysed by Ingvordsen
73 et al. (2014) for quantity of production. Within each of the five 24 m² chambers (height 3m) in the
74 phytotron, humidity, temperature and gasses were controlled as well as continuously monitored.
75 The five applied treatments can be seen in Table 2. They included (1) ambient (control) mimicking
76 present south Scandinavian summer of 19/12 °C (day/night), [CO₂] constantly at 385 ppm and no
77 O₃ added, (2) [CO₂] constantly at 700 ppm, (3) temperature elevated +5 °C (day and night), (4)
78 elevated temperature and [CO₂] combined at the level of the single-factor treatments and (5) [O₃]
79 constantly at 100-150 ppb (day and night). The climatic factors were mimicking levels predicted
80 ultimo 21st century, if greenhouse gasses are not substantially reduced (SCRES scenario A1FI,
81 IPCC, 2007). The CO₂ was supplied by Air Liquide A/S Denmark and O₃ by UV Pro 550A
82 generators (Crystal air products & services, Canada). Further details on RERAF are given by
83 Frenck *et al.* (2011) and Ingvordsen *et al.* (2014). Eight plants of each accession were grown in 11
84 L pots with 4 kg of sphagnum substrate (Pindstrup Substrate No. 6, Pindstrup Mosebrug A/S,
85 Denmark), where 10 g of NPK fertilizer (21-3-10, Yara) was applied at sowing. Water was applied
86 within one hour after the light was turned on by a surface dripping system delivering 4.4 L m⁻²
87 day⁻¹ in all treatments. To compensate for the drainage of the pot setup as well as root distribution
88 and water loss, water was applied above the average precipitation of Southern Scandinavia (236
89 mm; DMI, 2014). Watering was stepwise reduced from Zadoks growth stage (ZGS) 90 and ended at
90 ZGS 99 (Zadoks *et al.*, 1974). Light was supplied by 28 high-pressure mercury (1000 W or 400 W)
91 and 14 halogen (250 W) lamps in each chamber. The daily light cycle was 16/8 h (day/night) and
92 PAR (parabolic aluminized reflector) averaged at approximately 400 mol photons m⁻² s⁻¹ at canopy
93 height (ca. 1 m). To avoid possible chamber specific effects the treatments with its corresponding

94 batch of plants were rotated between the chambers on a weekly basis. In practice all plants were
95 exposed to ambient conditions for approximately 2 hours during the time of rotation and the time
96 necessary for the new chamber to reach the different treatment values.

97

98 *Grain yield*

99 Plants were harvested individually and after drying in constant temperature for a minimum of three
100 weeks, they were threshed and grain weight measured. After threshing grains were stored at 7 °C.
101 Number of grains was obtained by dividing with the weight of an enumerated sub-sample.

102

103 *Protein*

104 Total nitrogen (N)-analyses and following calculation of crude protein were performed by YARA
105 (Yara Analytical Services, Pocklington, England) on grain material via the Dumas Combustion
106 method on a LECO CNS TRUMAC. Crude protein was achieved on 17 accessions (stated in S1) in
107 each of the five treatments, and used to predict protein concentration in the remaining accessions.

108

109 *NIR measurements*

110 Spectral reflectance of whole grains from all accessions was obtained using a QFA-Flex 600F FT-
111 NIR instrument (Q-interline, Tølløse, Denmark). 1.5-7 g of the grains were placed in IR transparent
112 glass vials (height 6 cm, diameter 2.6 cm) and measured using a rotating sample device. The sample
113 was rotated at three rounds per minute. The measuring sample window at the rotating sample device
114 had a diameter of 6 mm, which provides an analysis surface of approximately 510 mm². Spectra
115 were collected at every 2 nm in the NIR region from 1100 to 2498 nm. One spectrum was obtained
116 for each sample as an average of 64 sub-scans. The spectra were reported as log (1/R).

117

118 *Statistics*

119 Principal component analysis (PCA) was performed on raw data as an explorative data analysis to
120 obtain a first overview of the data and to identify obvious outliers and delineate classes. Hotelling's
121 T-square versus residual plots was used to detect outliers. Partial least squares regression (PLSR)
122 models were developed on raw scatter corrected spectra, Savitsky-Golay first derivative (Savitzky
123 and Golay, 1964) averaging over 7 points and a second order polynomial spectra and multiplicative
124 signal corrected (MSC) (Geladi, 1985) spectra. Root mean square error of cross validation
125 (RMSECV) plotted against the number of PLSR latent variables for each pre-processing method

126 was used to select the optimum pre-processing method and the optimum number of latent variables
127 in the PLSR model. The optimum number of latent variables was chosen as the first local minimum
128 in the smooth declining RMSECV curve or the point, where this curve flattened. Random cross
129 validation with 10 segments and 10 iterations was used.

130 The performance of the PLSR model to predict protein were evaluated using the root mean
131 square error of prediction (RMSEP), standard error of performance (SEP) and bias. Initially the
132 obtained model was developed on 17 accessions per treatment and used to predict protein
133 concentration in the remaining accessions.

134 All analysis were carried out using MATLAB version 7.9.0 (R2009b) (The Mathworks, Inc.,
135 Natick, MA, USA) along with the PLS toolbox version 7.5.1 (Eigenvector Research, Inc., Manson,
136 WA, USA).

137 Following statistical analysis was carried out in R version 2.15.3(R core Team, 2013), and
138 SigmaPlot version 11.0, (Systat Software, Inc., San Jose California USA, www.sigmaplot.com) was
139 used for illustration.

140

141

142 **Results and discussion**

143 *Quality of applied treatments*

144 Atmospheric conditions of temperature and relative humidity were during cultivation in rather good
145 agreement with set points programmed in the RERAF phytotron. With regard to experimental levels
146 of [CO₂] the difference between the treatment of ambient and elevated [CO₂] was app. + 240 ppm
147 and on average 75 ppm lower than expected (Table 2). The increased [CO₂] in treatments with
148 ambient levels of [CO₂] is most probably due to that CO₂ cannot technically be removed, and the
149 large amount of plants seemed to have produced considerable quantities of CO₂ during respiration.

150

151 *Treatment effects on grain yield*

152 The effects of the single climatic factors on overall grain yield of the 108 accessions were reported
153 as a subset of 138 accessions accounted for in Ingvordsen et al. (2014). Grain yield was found in
154 agreement with previous studies, reporting increased grain yield at elevated [CO₂] (Ziska and
155 Bunce, 2007) and decreased grain yield at elevated [O₃] (Feng *et al.*, 2008) and elevated
156 temperature (Barnabás *et al.*, 2008) as under the two-factor treatment of elevated temperature and
157 [CO₂] combined (Clausen *et al.*, 2011).

158

159 *NIR and prediction of GPC*

160 A PLSR model based on NIR measurements and chemical measurement of N with subsequent
161 calculation of GPC was developed and used to predict GPC in the remaining accessions. Spectra for
162 all included accessions showed sufficient variance and clear peaks for further analysis (Fig. 1). A
163 good calibration model using 8 latent variables on MSC pre-processed NIR spectra showed an
164 $R^2=0.8$ with an RMSECV=1.34. Based on this calibration model the protein concentration was
165 predicted in the remaining accessions (Fig. 2).

166

167 *Treatment effects on GPC*

168 Elevated temperature as single-factor caused GPC to increase 29 % (Table 3). Several studies have
169 reported increased GPC from elevated temperature >35 °C (Savin and Nicolas, 1996; Majoul-
170 Haddad *et al.*, 2013) or around anthesis (Pettersson and Eckersten, 2007; Malik *et al.*, 2013). In the
171 present study, a constantly elevated temperature of +5 °C was also found to increase GPC. Högy *et*
172 *al.* (2013) found no change in GPC from a 2 °C increase in soil temperature, but decreased
173 concentrations of total non-structural carbohydrates, starch, fructose and raffinose. The increase in
174 GPC appears promising in terms of securing sufficient protein production under future climate
175 conditions, however, in a future climate [CO₂] is projected to increase concerted with temperature.

176 Under elevated [CO₂] the GPC decreased overall 5 % (Table 3). This was less than the 15 %
177 decrease found in a meta-analysis of barley with no significant difference between FACE, open-top
178 chambers and enclosure studies or if rooted in pots or field (Taub *et al.*, 2008). The less decrease
179 induced by elevated [CO₂] found in the present study might be due to the plant material tested, as
180 elevated levels of [CO₂] were in agreement (590-700 ppm). The material used in the meta-analysis
181 presumably covered four barley cultivars (Thule, Alexis, Jo1621 and Atem) (Kleemola *et al.*, 1994;
182 Thompson and Woodward, 1994; Sæbø and Mortensen, 1996; Fangmeier *et al.*, 2000), whereas the
183 present study included 108 accessions (Table 1 and S1).

184 Elevated [O₃] was found to increase overall GPC with 6 % (Table 3). Studies in wheat, which
185 has been reported more sensitive to [O₃] than barley (Mills *et al.*, 2007), have found GPC of wheat
186 to increase overall 7 % with averaged [O₃] of 58 ppb and exposure between 7-12 hours per day
187 (Feng *et al.*, 2008). The study by Feng *et al.* (2008) also reported 71 ppb [O₃] to cause further
188 increased protein concentration. In the present study [O₃] averaged 121 ppb on a 24 hours basis.
189 The similar increase in GPC from the double concentration and exposure-time to [O₃] may suggest

190 that barley is not very sensitive to O₃ or that barley has a different responds pattern to ozone than
191 wheat.

192 The 29 % increased GPC under elevated temperature was modified to 8 % under the
193 simultaneous exposure to elevated temperature and [CO₂] in the two-factor treatment (Table 3).
194 This result strongly points to the risk of misinterpretation of the combined effects, when deduced
195 from single-factor treatments. The combined effect of elevated temperature and elevated [CO₂] on
196 GPC was not found to be additive - an important point when considering the future effects of
197 climate change, where temperature, [CO₂] and [O₃] are predicted to increase concerted.

198

199 *Treatment effects on grain protein pr. grain*

200 Considering the quantity of grain protein in relation to the weight of a single grain (Table 3), the
201 picture changed from increase to decrease under elevated [O₃] and the two-factor treatment
202 compared to response in GPC. Under elevated [CO₂] the GPC and protein per grain decreased
203 similarly, 5 % (Table 3). The 29 % increase in GPC under elevated temperature was substantially
204 lowered to only 7 % when given on a pr. single grain weight basis. The decreases reflected the
205 diverse seed weights in the different treatments, where only the treatment with elevated [CO₂], had
206 more or less the same seed weight as found under ambient conditions (data not shown). However, a
207 suggested inhibition of the assimilation of nitrate into e.g. proteins under elevated [CO₂] (Bloom *et*
208 *al.*, 2010) could have engaged in the maintained and not increased grain weight by elevated [CO₂].
209 Further, the increase in GPH was found to be smaller than the increase in grain yield as previously
210 reported as indication on inhibition of the assimilation of nitrate under elevated [CO₂] (Pleijel and
211 Uddling, 2012).

212

213 *Treatment effects on GPH*

214 The increased GPC in the two-factor treatment that could potentially increase protein production
215 under future climate conditions vanished, when the treatment effect on actual harvested quantity
216 was considered (Table 3). Even though the GPC increased 8 % compared to ambient, the GPH was
217 found decreased by 23 % due to the decreased grain yield of 28 % in an atmosphere of elevated
218 temperature plus [CO₂]. The treatments effects on grain yield converted the potential increase in
219 GPH into an overall reduction, as was also seen in the single-factor treatments with elevated
220 temperature and [O₃]. In the single-factor treatment with elevated [CO₂] the opposite was observed,

221 as the decreased GPC was compensated for by the higher yield, and the resulting GPH was
222 increased compared to ambient (Table 3).

223 Since the findings of the present study are based on 108 accessions the overall effects reported
224 are considered robust with regard to barley, and the characteristics identified might be considered of
225 value in future breeding. Responsiveness to the elevated [CO₂] has been suggested as a breeding
226 target to increase grain yield under future climate conditions (Ziska and Bunce, 2007; Franzaring *et*
227 *al.*, 2013; Ingvordsen *et al.*, 2014). In the present study we found that the GPC under elevated
228 [CO₂] was decreased, though relatively little in comparison to the increased grain yield, suggesting
229 that a substantial increase in GPH could be envisaged from improved CO₂-responsiveness.
230 Harvested grain protein was found increased (13 %) under elevated [CO₂] - not from increased
231 protein pr. grain but from increased production of grains (Table 3) (Jablonski *et al.*, 2002;
232 Ingvordsen *et al.*, 2014). Application of additional nitrogen-fertilizer could potential ameliorate the
233 loss of protein in the grain under elevated [CO₂], however, Bloom *et al.*(2014) reported an
234 insignificant effect on GPC in wheat leaves under elevated [CO₂] due to inhibited nitrate
235 assimilation. The suggested inhibition of protein accumulation by elevated [CO₂] requests better
236 understanding of ammonium and nitrate use by crops under climate change conditions, an area that
237 has received little attention (Andrews *et al.*, 2013).

238

239 *Grain protein in the 108 accessions*

240 Among the 108 accessions, some differed to a greater or lesser extent from the overall responses to
241 the treatments, suggesting great diversity that could be exploited in breeding programs. No
242 significant difference in response to the climate treatments were observed between the group of
243 landrace and the group of cultivars (Fig. 3 and 4).

244 Considering the expected lower grain yield under future climate conditions harvesting as much
245 protein as possible is likely preferable. Under the two-factor treatment the 108 accessions decreased
246 in average 23 % in GPH relative to ambient, however, the individual accessions spanned from -60%
247 to 30 % GPH (Fig. 3). Two landraces (Kushteki and Moscou) and a 2-rowed Danish feed barley
248 cultivar (Jacinta) increased 30-33 % in GPH. All three accessions ranked in top ten for grain yield
249 of the 108 accessions, whereas only the feed barley ranked in the top (placed 2) in GPC in the two-
250 factor treatment. Another four accessions, two modern cultivars (Sebastian and Brage), a Finish
251 landrace (Luusua) and a breeder-line (Bor 05135) increased 13-16 % in GPH under elevated
252 temperature and [CO₂] in combination. Of these four accessions, only the landrace demonstrated

253 high rank (8) with regard to GPC under the two-factor treatment of all 108 accessions, and the
254 Danish cultivar (Brage) demonstrated high grain yield. The last two accessions, the Norwegian 6-
255 rowed cultivar and the Finish 2-rowed breeder-line demonstrated top-medium rank for GPC and
256 grain yield, where they ranked 52 and 21 in GPC and 22 and 14 in grain yield. When only
257 considering the performance in the two-factor treatment, all seven accessions rank in top 15 of the
258 108 accessions in GPH. That increased GPH was identified in landraces, cultivars and a breeder-
259 line as in 2- and 6-rowed suggest that beneficial genes for developing cultivars with high GPC and
260 grain yield are available from many sources.

261 Under elevated [CO₂], three accessions increased over 80 % in GPH. The accessions were two 2-
262 rowed old Swedish accessions (Arla and Pallas) and the 2-rowed Danish feed cultivar that also
263 demonstrated increased grain yield under the two-factor treatment (Jacinta). In the pedigrees of both
264 Jacinta and Arla the accession Bavaria (NGB6945) can be found (van Berloo and Hutten, 2005). In
265 addition both Jacinta and Arla showed high CO₂-responsiveness, both among the top five
266 accessions increasing most in grain yield under elevated [CO₂]. All three accessions had high grain
267 yields under elevated [CO₂], whereas under ambient conditions the old Swedish cultivars ranked
268 low (81 and 106) in GPC and grain yield (79 and 96).

269 Elevated temperature increased overall GPC the most and was only found decreased in eight
270 accessions being landraces and old cultivars. Overall grain protein harvested was decreased by 42
271 % but three accessions showed increased GPH under elevated temperature, all 2-rowed and
272 cultivars; an old Swedish (Mari), an old Danish (Odin) and a modern Danish (Sebastian).

273 One can speculate if the ability of Jacinta and Sebastian to produce high GPH in the two-factor
274 treatment was related to their suggested improved ability to secure high GPH in either of the single-
275 factor treatments of elevated [CO₂] or elevated temperature – or reverse; the performance in either
276 of the single-factor treatments contributed to the performance under the combined treatment.
277 However, the results from single-factor treatments were overall not found additive for the two-
278 factor treatment, and of the mentioned accessions only two were found in the top for GPH under
279 either of the single-factor treatments and the two-factor treatment. Considering more accessions
280 than the top three to five best ones, though revealed broader overlap of accessions producing high
281 GPH under the two-factor treatment and either of or both of the single-factors suggesting that high
282 performance under a single-factor treatment can be beneficial in the two-factor treatment.

283 Under elevated [O₃], several accessions decreased less than the averaged 11 %, and 14
284 accessions increased > 11 % with regard to GPH. Cultivar variation to [O₃] have previously been

285 reported in grain yield of soybean by Betzelberger et al. (2010) (with [O₃] applied eight hours a day
286 at 40-150 ppb) and for the set of present accessions in Ingvordsen *et al.* (2014). Two old cultivars
287 (Pallas and Juli) and an early modern cultivar (1978; Agneta) showed highest increased GPH under
288 elevated [O₃] and Agneta also ranked one with regard to GPH under the [O₃] treatment of all 108
289 accessions.

290

291 **Conclusions**

292 The massive variation in protein response to the applied climate treatments, emphasize that the
293 phenotypic differences should be exploited in breeding programs for abiotic stress tolerance. Likely,
294 a cascade of different genes encodes the different responses. Here, mining the genome with GWAS
295 (Genome Wide Association Studies) could help identifying some of the underlying genes, and the
296 link between these DNA markers and phenotypes could facilitate the breeding process.
297 Additionally, the identification of suitable genetic resources should be performed under treatments
298 of combined climatic factors, since the effects from the single-factors were found rarely to be
299 additive. The overall decreased GPH in the most realistic climate treatment, where [CO₂] and
300 temperature were elevated simultaneously, emphasizes the need to explore and exploit genotypes to
301 secure plant protein production under future climate conditions.

302

303

304 **Supplementary data**

305 Supplementary data are available at JXB online.

306 Supplementary Table S1. The 108 spring barley accessions included in present study with gene
307 bank number, accession type, row type, origin country/breeding country, collecting date/release
308 year, location/breeder institute, pedigree and marking of the 17 individual included in the chemical
309 analysis.

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Table 1. Overview of the accessions included. Old cultivars before 1975; modern cultivars after 1975.

	Landraces		Old cultivars		Modern cultivars ^a	
	2 rowed	6 rowed	2 rowed	6 rowed	2 rowed	6 rowed
Denmark	1	2	6	1	23	
Sweden	1	1	7	2	2	1
Finland ^b	2	5	1	4	2	6
Norway	1		1	3		4
Europe ^c	5	7			7	
non-Europe	1	4				
unknown	1	5				

^aincluding breeder-lines

^btwo landraces segregated either as 2 or 6 rowed and has not been included

^cnot including Scandinavian but Faroe Islands

Table 2. Experimental levels of manipulated climatic factors of applied treatments. Set points were; temperature (tmp): 19/12 °C (day/night) or 24/17 °C; [CO₂] (CO₂): 385 ppm or 700 ppm; [O₃] (O₃): 100-150 ppb; relative humidity 55/70 % day/night.

	<u>tmp</u> <u>day/night</u>	<u>[CO₂]</u> <u>(constant)</u>	<u>[O₃]</u> <u>(constant)</u>	<u>humidity</u> <u>day/night</u>
ambient	18.9±1.2/11.8±0.8	448.5±81.1	1.40±1.4	55.7±2.5/69.9±1.5
+CO ₂	19.0±1.2/12.5±2.1	684.7±41.1	0.98±1.7	55.3±5.1/69.4±5.9
+tmp	23.9±1.4/16.8±0.8	448.4±74.4	1.90±1.2	55.9±2.8/69.8±1.6
+tmp & CO ₂	23.8±1.3/16.9±0.9	688.3±38.2	1.50±1.4	56.0±2.9/69.8±1.8
+O ₃	18.9±1.2/11.9±1.0	443.1±67.5	121.1±32.8	55.7±2.4/69.8±1.7

Table 3. Overall averaged parameters for the 108 barley accessions cultivated under future levels of carbon dioxide (+CO₂), ozone (+O₃), temperature (+tmp) and under the two-factor treatment (+tmp & CO₂) as well as under ambient (amb). * specifies significant difference from the ambient treatment determined by t-test. Grain yield per plant and grain number per plant are from Ingvordsen *et al.*(Ingvordsen *et al.*, 2014)

	amb	+tmp & CO ₂	+ CO ₂	+O ₃	+tmp
Grain yield per plant (g)	6.85±1.29	4.92±1.18***	8.02±1.94***	5.82±1.38***	3.08±1.13***
<i>% different from ambient</i>		-28.12	17.10	-15.10	-54.98
Grain number per plant (#)	128.02±31.2	100.01±25.3***	149.93±17.1***	122.21±31.9	68.77±24.1***
<i>% different from ambient</i>		-21.88	17.11	-4.54	-46.54
Grain protein concentration (%)	13.97±1.82	15.06±1.97***	13.33±1.91*	14.76±1.96**	18.03±2.18***
<i>% different from ambient</i>		7.86	-4.85	5.68	29.11
Grain protein/grain (mg)	7.62±1.42	7.49±1.32	7.24±1.66	7.09±1.35**	8.14±1.62*
<i>% different from ambient</i>		-1.63	-4.87	-6.84	6.82
Grain protein harvested per plant (g)	0.95±0.20	0.74±0.19***	1.07±0.31**	0.85±0.18***	0.55±0.20***
<i>% different from ambient</i>		-22.53	12.46	-11.19	-42.26

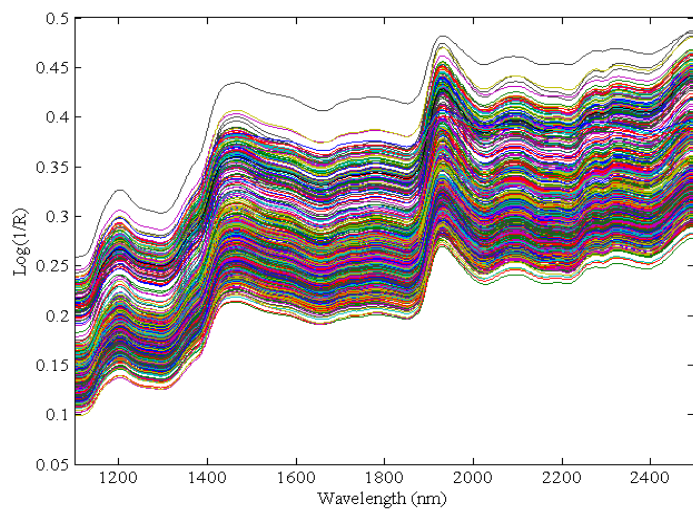


Fig. 1. Raw NIR spectra (1100 to 2498 nm) of all accessions.

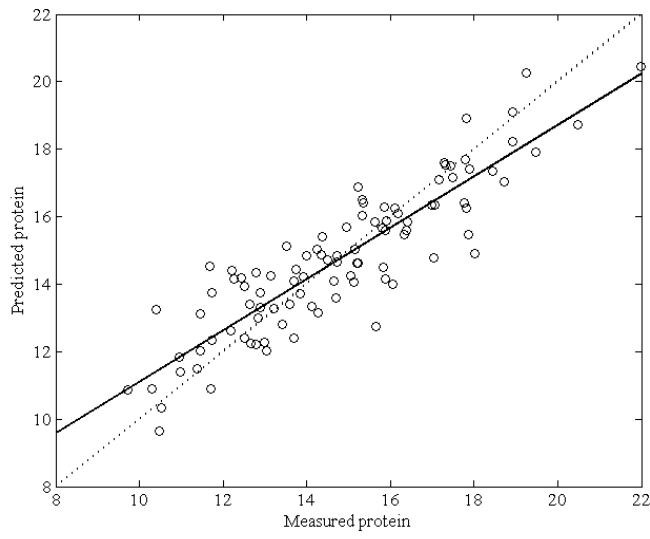


Fig. 2. Concentration of protein predicted using PLSR model vs. measured protein concentration. Full line indicate best fit with $R^2=0.8$ and $RMSECV=1.3392$. Dotted line has $R^2=1$

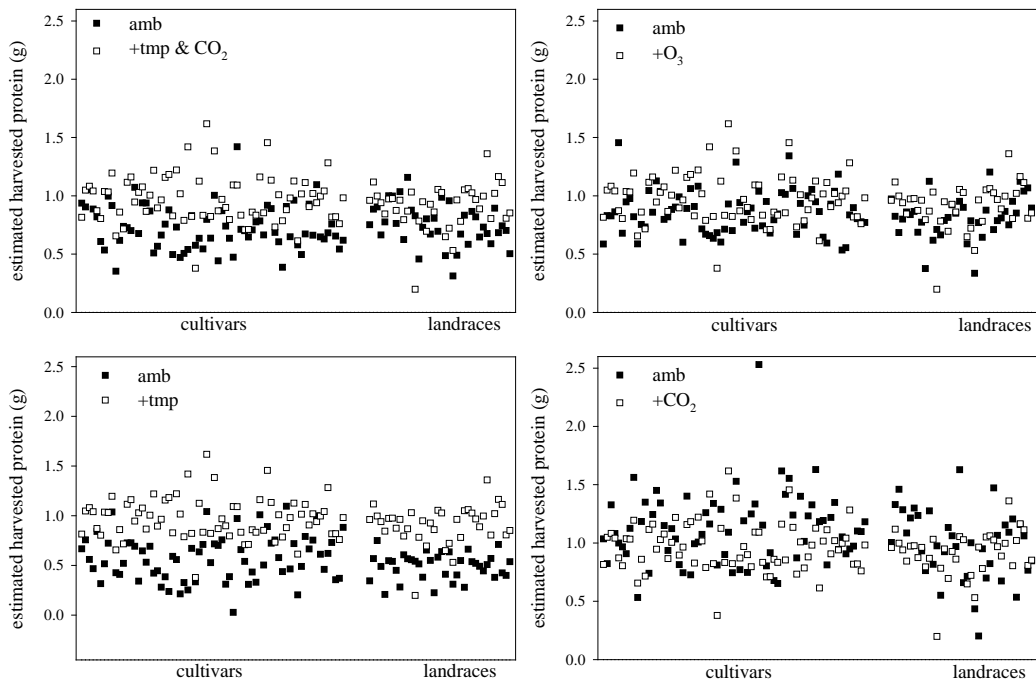


Fig. 3. Grain protein harvested (g/plant) of the 108 accessions in the five climate treatments (open) and ambient (closed). The order of accessions, left to right, is as in S1.

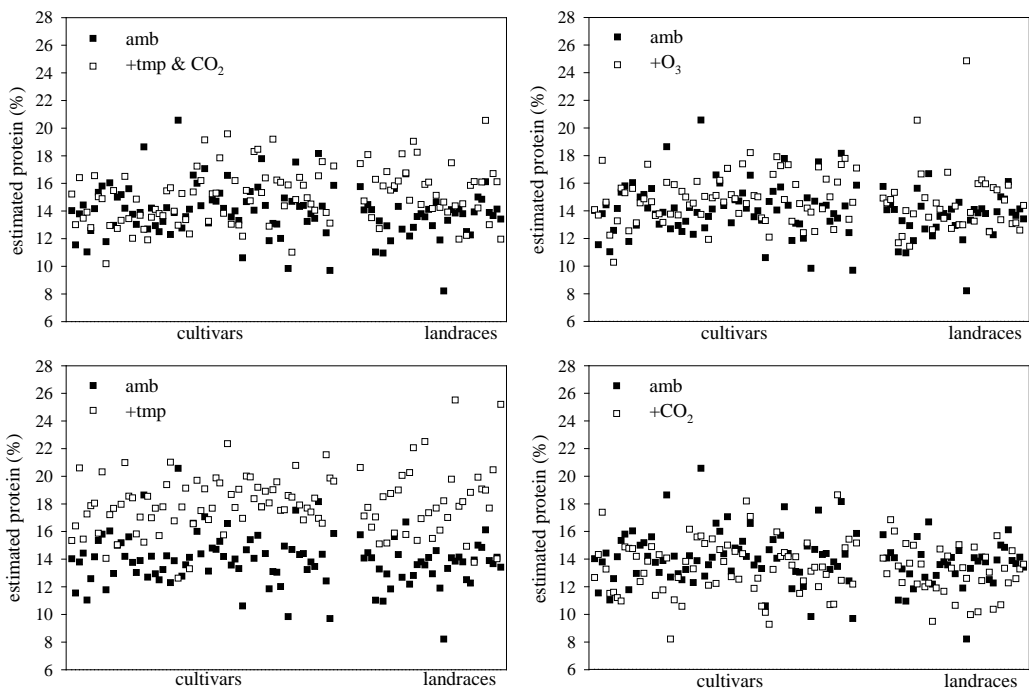


Figure 4. Grain protein concentration (%) of the 108 accessions in the five climate treatments (open) and ambient (closed). The order of accessions, left to right, is as in S1.