



Article (refereed) - postprint

Reiskind, Julia B.; Maberly, Stephen C. 2014. **A tribute to George Bowes: linking terrestrial and aquatic botany.**

Copyright © 2013 Elsevier B.V.

This version available <http://nora.nerc.ac.uk/508807/>

NERC has developed NORA to enable users to access research outputs wholly or partially funded by NERC. Copyright and other rights for material on this site are retained by the rights owners. Users should read the terms and conditions of use of this material at <http://nora.nerc.ac.uk/policies.html#access>

NOTICE: this is the author's version of a work that was accepted for publication in *Aquatic Botany*. Changes resulting from the publishing process, such as peer review, editing, corrections, structural formatting, and other quality control mechanisms may not be reflected in this document. Changes may have been made to this work since it was submitted for publication. A definitive version was subsequently published in *Aquatic Botany* (2014), 118. 1-3. [10.1016/j.aquabot.2013.11.008](https://doi.org/10.1016/j.aquabot.2013.11.008)

www.elsevier.com/

Contact CEH NORA team at
noraceh@ceh.ac.uk

1 **A tribute to George Bowes: linking terrestrial and aquatic botany**

2 Julia B. Reiskind¹ and Stephen C. Maberly²

3
4 1. Ecosystems Ecology, Department of Biology, University of Florida, Gainesville FL 32611
5 USA

6 2. Lake Ecosystems Group, Centre for Ecology & Hydrology, Lancaster Environment Centre,
7 Lancaster LA1 4AP UK

8
9
10
11 **Corresponding author:**

12 S.C. Maberly

13 Lake Ecosystems Group, Centre for Ecology & Hydrology, Lancaster Environment Centre,
14 Lancaster LA1 4AP UK.

15 Tel: +44 1524 595851.

16 E-mail: scm@ceh.ac.uk

17
18 Keywords: Aquatic, photosynthesis, Rubisco, terrestrial

20 George E Bowes was brought up in England and finished his PhD in 1967 at the University of
21 London. He then ‘crossed the pond’ to take a postdoctoral position at the University of Illinois
22 where he worked with Bill Ogren, followed by a year as a Carnegie Fellow at the Carnegie
23 Institute at Stanford University. In 1972 he moved to the Botany Department at the University of
24 Florida where he spent the rest of his career.

25 It was at Illinois that he and Bill Ogren undertook the defining work on Ribulose-bis-
26 phosphate carboxylase-oxygenase (rubisco), discovering the oxygenase reaction and its role in
27 photorespiration, and thus explaining the oxygen effect on photosynthesis in C₃ plants (Bowes et
28 al., 1971; Ogren and Bowes, 1971). They also found that, while rubisco is inhibited by oxygen,
29 phosphoenolcarboxylase (PEPC) is not, explaining the differential effect of oxygen on C₃ and C₄
30 plant photosynthesis (Bowes and Ogren, 1972). Bill Ogren (Ogren, 2003) provides some
31 fascinating information about the difficulties he and George faced in publishing this work and
32 persuading the photosynthesis community that it was correct. It is hard, now, to overestimate the
33 importance of this work. The name of the enzyme (then ribulose-diphosphate carboxylase) was
34 changed to recognize its oxygenase activity but more importantly this fundamental discovery
35 altered the course of photosynthesis research and influenced many fields including plant and
36 algal ecology and physiology, biochemistry and biogeochemistry, climate change and food
37 security.

38 Shortly after joining the University of Florida, George began to study aquatic plant
39 photosynthesis. His research trajectory was initiated by the discovery of 1) differing CO₂
40 compensation points among several aquatic species, and 2) the observation with Scott Holaday
41 of environmental impacts on intraspecific variation in CO₂ compensation points (Van et al.,
42 1976). His graduate students, Jocelyne Ascencio and Mike Salvucci, found that increased

43 temperature and photoperiod reduced the CO₂ compensation point and photorespiration in the
44 submerged macrophyte *Hydrilla verticillata* (Salvucci and Bowes, 1981; Ascencio and Bowes,
45 1983). The reduced compensation point was associated with increased activity of PEPC, and
46 several enzymes associated with C₄ photosynthesis, and decreased activity of photorespiratory
47 enzymes (Holaday and Bowes, 1980; Salvucci and Bowes, 1981). Finally, pulse-chase results
48 suggested that *Hydrilla* had a C₄-type photosynthetic system (Holaday and Bowes, 1980;
49 Salvucci and Bowes, 1983). An interesting area of complexity opened up when they found that
50 another submerged macrophyte, *Myriophyllum spicatum*, lacked the requisite enzyme activities,
51 despite showing the physiological characteristics of C₄ photosynthesis. Experiments with an
52 inhibitor of carbonic anhydrase suggested that this enzyme facilitates bicarbonate use as part of a
53 CO₂ concentrating mechanism (Salvucci and Bowes, 1982, 1983) underlining the diversity in the
54 way that freshwater macrophytes acquire inorganic carbon for photosynthesis. Subsequent
55 studies on *Hydrilla* indicated that Kranz anatomy, typical of many terrestrial C₄ plants, was
56 lacking. Immunogold labeling showed that there was an intracellular separation of Rubisco and
57 PEPC in leaves of C₄-type plants: rubisco was confined to the chloroplasts and PEPC to the
58 cytosol, i.e., there was an intracellular separation of the C₄ and Calvin cycles (Reiskind et al.,
59 1989). Later studies demonstrated that the chloroplast was the site where CO₂ was concentrated,
60 with CO₂ concentrations estimated to be 400 μM (Reiskind et al., 1997). *Hydrilla* was the first
61 known single-cell C₄ system, although this has been found subsequently in certain terrestrial C₄
62 plants (Voznesenskaya et al., 2001; Edwards et al., 2004).

63 In the mid-1990s George made a leap into studying the molecular details of the *Hydrilla*
64 C₄ system. Three key C₄-enzymes: PEPC, pyruvate Pi dikinase (PPdK) and NADP-malic
65 enzyme (NADP-ME) are up-regulated during induction of the C₄-photosynthetic state. Three

66 PEPC isoforms were identified in *Hydrilla* (Magnin et al., 1997; Rao et al., 2002). One of these
67 forms was up-regulated during induction, only expressed in C₄-type leaves and had the kinetic
68 characteristics of the C₄-isoform (Rao et al., 2008). Terrestrial C₄-PEPC isoforms possess a
69 serine moiety at the amino acid position 774 near the carboxy terminus, based on the *Flaveria*
70 sequence (Blasing et al., 2000). The *Hydrilla* C₄-PEPC isoform contains alanine at this position
71 like all C₃-isoforms, despite being similar kinetically to the C₄-isoform (Rao et al., 2008). Two
72 isoforms of PPdK and three isoforms of NADP-ME were also detected (Rao et al., 2006a; Rao et
73 al., 2006b; Estavillo et al., 2007). One NADP-ME isoform was from the chloroplast, up-
74 regulated in the light with kinetic characteristics intermediate between C₃ and C₄ NADP-ME
75 isoforms of terrestrial plants (Estavillo et al., 2007). Over many years, work by George Bowes
76 and his co-workers have established *Hydrilla* as a C₄-NADP-ME plant and it is now one of the
77 most completely studied C₄ plants on the planet.

78 Although continuing to work on freshwater aquatics, George also dived into the sea by
79 making annual collecting ‘cruises’ to the Bahamas and the Florida Keys. Two coenocytic green
80 macroalgae became the focus of this work: *Codium decorticatum* and *Udotea flabellum*.
81 Physiological studies showed that photorespiration in *Udotea* was low, as was PEPC activity
82 (Reiskind et al., 1988). However, the activities of phosphoenolpyruvate carboxykinase (PEPCK)
83 in both carboxylating and decarboxylating modes and the activities of the other requisite C₄-
84 cycle enzymes were sufficiently high to allow a C₄-like photosynthetic system to operate
85 (Reiskind et al., 1988). Treatment with a PEPCK inhibitor resulted in reduced photosynthetic
86 rates, increased O₂ sensitivity and reduced labeling of C₄-acids as initial products of
87 photosynthesis, suggesting that *Udotea* operated a C₄ system based on PEPCK with a spatial
88 separation of carboxylase (cytosol) and decarboxylase (chloroplast) activities (Reiskind and

89 Bowes, 1991). In an evolutionary sense, the C₄ system in *Udotea* appears to be the oldest known
90 of any photoautotroph. George continues to be active in marine research and has produced a
91 recent review with colleagues from around Florida on the impact of global climate change on
92 ocean acidification, and its effect on seagrasses and macroalgae (Koch et al., 2013). Meanwhile,
93 George's interest in rubisco continued in the terrestrial environment in collaboration with Mike
94 Salvucci and Gabriel Holbrook, and several agronomists at the University of Florida. Working
95 on *Nicotiana rustica*, Mike, Gabe and George reported a chloroplastic phosphatase that in the
96 light, and particularly in the presence of NADPH, degrades a naturally occurring inhibitor of
97 rubisco activity, Carboxy-arabitol-1 phosphate (Holbrook et al., 1989; Salvucci and Holbrook,
98 1989). This was another major discovery on the fundamental mechanisms of photosynthesis.

99 One key concern today that requires a knowledge of plant ecophysiology and
100 biochemistry, is the adequate production of food to support the growing human population
101 against the background of a changing climate. The current concentration of 400 ppm CO₂, up 75
102 ppm from the late 1960s, will tend to increase rates of photosynthesis as the oxygenase function
103 of rubisco will be suppressed, decreasing photorespiratory CO₂ loss and increasing ATP/NADPH
104 redirection to photosynthetic assimilation (Bowes, 1991; Long et al., 2004). Rising temperature,
105 however, may have a negative effect in some areas. A comparison of two rice cultivars exposed
106 to 350 and 700 ppm CO₂ under varying day/night temperature regimes showed that, while
107 photosystem II efficiency was largely unaffected, rubisco gene expression, protein content and
108 activity were adversely affected by elevated temperature (Gesch et al., 2003). Leaf
109 photosynthetic rates were negatively impacted at the higher CO₂ concentration and highest
110 temperature regime with some cultivar differences (Gesch et al., 2003). Agricultural productivity
111 could be increased further if the low photorespiratory rates and potentially higher productivity of

112 C₄ plants could be transferred into current important C₃ crops, such as rice. George has worked
113 actively in this area because *Hydrilla*, as a single-cell C₄ plant that lacks the structural
114 complexity of Kranz anatomy, is an excellent model for engineering a C₃ plant with C₄
115 characteristics (Bowes et al., 2002; Bowes, 2011).

116 In addition to George's scientific achievements, he was an excellent teacher of
117 undergraduate courses teaching not only the subject but also how to think scientifically and how
118 to ask and answer scientific questions. George's research lab was a mini United Nations with
119 representatives from around the world. This led to many collaborations, which are reflected in
120 the contributions in this issue. It was also a fertile ground for many jokes, especially between the
121 Yanks and the Brits, and a war fought some 235 years ago. Despite all of the above, George
122 found time to serve as Chair of the Botany Department (1998 – 2006). Faculty members describe
123 him as a fair and effective leader with the ability to maintain a balance among the different
124 disciplines.

125 Of course, this particular issue is also celebrating another aspect of George's academic
126 life, the editing of the scientific journal *Aquatic Botany*. George served on the editorial board
127 from 1982 before taking over as Editor in Chief in 1995 when J.M.A. Brown retired. He stood
128 down in 2013 after 19 years as editor. One feature of George's career is his contribution to
129 terrestrial, freshwater and marine science, which is proof that excellent science can cross
130 scientific disciplines. We have tried to represent some of this diversity in this Special Issue.

131

132 References

133 Ascencio, J., Bowes, G., 1983. Phosphoenolpyruvate carboxylase in *Hydrilla* plants with varying
134 CO₂ compensation points. *Photosynthesis Research* 4, 151-170.

135 Blasing, O.E., Westhoff, P., Svensson, P., 2000. Evolution of C₄ phosphoenolpyruvate
136 carboxylase in *Flaveria*, a conserved serine residue in the carboxyl-terminal part of the enzyme
137 is a major determinant for C₄-specific characteristics. *Journal of Biological Chemistry* 275,
138 27917-27923.

139 Bowes, G., 1991. Growth at elevated CO₂- photosynthetic responses mediated through rubisco.
140 *Plant Cell and Environment* 14, 795-806.

141 Bowes, G., 2011, Single-Cell C₄ Photosynthesis in Aquatic Plants. In: Raghavendra, A.S., Sage,
142 R.F. (Eds.), *C₄ Photosynthesis and Related CO₂ Concentrating Mechanisms*, pp. 63-80.

143 Bowes, G., Ogren, W.L., 1972. Oxygen inhibition and other properties of soybean ribulose 1,5-
144 diphosphate carboxylase. *Journal of Biological Chemistry* 247, 2171-&.

145 Bowes, G., Ogren, W.L., Hageman, R.H., 1971. Phosphoglycolate production catalyzed by
146 ribulose diphosphate carboxylase. *Biochemical and Biophysical Research Communications* 45,
147 716-&.

148 Bowes, G., Rao, S.K., Estavillo, G.M., Reiskind, J.B., 2002. C₄ mechanisms in aquatic
149 angiosperms: comparisons with terrestrial C₄ systems. *Functional Plant Biology* 29, 379-392.

150 Edwards, G.E., Franceschi, V.R., Voznesenskaya, E.V., 2004. Single-cell C₄ photosynthesis
151 versus the dual-cell (Kranz) paradigm. *Annual Review of Plant Biology* 55, 173-196.

152 Estavillo, G.M., Rao, S.K., Reiskind, J.B., Bowes, G., 2007. Characterization of the NADP malic
153 enzyme gene family in the facultative, single-cell C₄ monocot *Hydrilla verticillata*.
154 *Photosynthesis Research* 94, 43-57.

155 Gesch, R.W., Kang, I.H., Gallo-Meagher, M., Vu, J.C.V., Boote, K.J., Allen, L.H., Bowes, G.,
156 2003. Rubisco expression in rice leaves is related to genotypic variation of photosynthesis under
157 elevated growth CO₂ and temperature. *Plant Cell and Environment* 26, 1941-1950.

158 Holaday, A.S., Bowes, G., 1980. C₄ acid metabolism and dark CO₂ fixation in a submerged
159 aquatic macrophyte (*Hydrilla verticillata*) *Plant Physiology* 65, 331-335.

160 Holbrook, G.P., Bowes, G., Salvucci, M.E., 1989. Degradation of 2-carboxyarabinitol 1-
161 phosphate by a specific chloroplast phosphatase. *Plant Physiology* 90, 673-678.

162 Koch, M., Bowes, G., Ross, C., Zhang, X.-H., 2013. Climate change and ocean acidification
163 effects on seagrasses and marine macroalgae. *Global Change Biology* 19, 103-132.

164 Long, S.P., Ainsworth, E.A., Rogers, A., Ort, D.R., 2004. Rising atmospheric carbon dioxide:
165 Plants face the future. *Annual Review of Plant Biology* 55, 591-628.

166 Magnin, N.C., Cooley, B.A., Reiskind, J.B., Bowes, G., 1997. Regulation and localization of key
167 enzymes during the induction of Kranz-less, C₄-type photosynthesis in *Hydrilla verticillata*.
168 *Plant Physiology* 115, 1681-1689.

169 Ogren, W.L., 2003. Affixing the O to Rubisco: discovering the source of photorespiratory
170 glycolate and its regulation. *Photosynthesis Research* 76, 53-63.

171 Ogren, W.L., Bowes, G., 1971. Ribulose diphosphate carboxylase regulates soybean
172 photorespiration. *Nature-New Biology* 230, 159-&.

173 Rao, S., Reiskind, J., Bowes, G., 2006a. Light regulation of the photosynthetic
174 phosphoenolpyruvate carboxylase (PEPC) in *Hydrilla verticillata*. *Plant and Cell Physiology* 47,
175 1206-1216.

176 Rao, S.K., Fukayama, H., Reiskind, J.B., Miyao, M., Bowes, G., 2006b. Identification of C₄
177 responsive genes in the facultative C₄ plant *Hydrilla verticillata*. *Photosynthesis Research* 88,
178 173-183.

179 Rao, S.K., Magnin, N.C., Reiskind, J.B., Bowes, G., 2002. Photosynthetic and other
180 phosphoenolpyruvate carboxylase isoforms in the single-cell, facultative C₄ system of *Hydrilla*
181 *verticillata*. *Plant Physiology* 130, 876-886.

182 Rao, S.K., Reiskind, J.B., Bowes, G., 2008. Kinetic analyses of recombinant isoforms of
183 phosphoenolpyruvate carboxylase from *Hydrilla verticillata* leaves and the impact of substituting
184 a C₄-signature serine. *Plant Science* 174, 475-483.

185 Reiskind, J.B., Berg, R.H., Salvucci, M.E., Bowes, G., 1989. Immunogold localization of
186 primary carboxylases in leaves of aquatic and a C₃-C₄ intermediate species. *Plant Science* 61, 43-
187 52.

188 Reiskind, J.B., Bowes, G., 1991. The role of phosphoenolpyruvate carboxykinase in a marine
189 macroalga with C₄-like photosynthetic characteristics. *Proceedings of the National Academy of*
190 *Sciences of the United States of America* 88, 2883-2887.

191 Reiskind, J.B., Madsen, T.V., VanGinkel, L.C., Bowes, G., 1997. Evidence that inducible C₄-
192 type photosynthesis is a chloroplastic CO₂-concentrating mechanism in *Hydrilla*, a submersed
193 monocot. *Plant Cell and Environment* 20, 211-220.

194 Reiskind, J.B., Seamon, P.T., Bowes, G., 1988. Alternative methods of photosynthetic carbon
195 assimilation in marine macroalgae. *Plant Physiology* 87, 686-692.

196 Salvucci, M.E., Bowes, G., 1981. Induction of reduced photorespiratory activity in submersed
197 and amphibious aquatic macrophytes. *Plant Physiology* 67, 335-340.

198 Salvucci, M.E., Bowes, G., 1982. Photosynthetic and photorespiratory responses of the aerial and
199 submersed leaves of *Myriophyllum brasiliense*. *Aquatic Botany* 13, 147-164.

200 Salvucci, M.E., Bowes, G., 1983. Two photosynthetic mechanisms mediating the low
201 photorespiratory state in submerged aquatic angiosperms. *Plant Physiology* 73, 488-496.

202 Salvucci, M.E., Holbrook, G.P., 1989. Purification and properties of 2-carboxy-D-arabinitol 1-
203 phosphate. *Plant Physiology* 90, 679-685.

204 Van, T.K., Haller, W.T., Bowes, G., 1976. Comparison of photosynthetic characteristics of three
205 submerged aquatic plants. *Plant Physiology* 58, 761-768.

206 Voznesenskaya, E.V., Franceschi, V.R., Kiirats, O., Freitag, H., Edwards, G.E., 2001. Kranz
207 anatomy is not essential for terrestrial C₄ plant photosynthesis. *Nature* 414, 543-546.

208

209