1	Application of stable carbon isotopes for reconstructing salt-marsh floral zones and
2	relative sea level, New Jersey, USA
3	
4	Andrew C. Kemp <sup>ab*</sup> , Christopher H. Vane <sup>c</sup> , Benjamin P. Horton <sup>d</sup> , Simon E. Engelhart <sup>d</sup> ,
5	and Daria Nikitina <sup>e</sup>
6	
7	<sup>a</sup> School of Forestry and Environmental Studies, Yale University, New Haven, CT 06510, USA.
8	<sup>b</sup> Yale Climate and Energy Institute, Yale University, New Haven, CT 06510, USA.
9	<sup>c</sup> British Geological Survey, Kingsley Dunham Centre, Keyworth, Nottingham NG12 5GG, UK.
10	<sup>d</sup> Sea Level Research, Department of Earth and Environmental Science, University of Pennsylvania,
11	Philadelphia, PA 19104, USA.
12	<sup>e</sup> Department of Geology and Astronomy, West Chester University, West Chester, PA 19380, USA.
13	* Corresponding author, andrew.kemp@yale.edu; tel 203 436 3978

#### 14 Abstract

We investigated use of  $\delta^{13}$ C values in bulk organic sediment to reconstruct the botanical 15 16 origin of samples preserved in coastal sedimentary archives as a proxy for relative sea 17 level in New Jersey, USA. Modern transects at three sites demonstrated that low and 18 high salt-marsh floral zones dominated by C<sub>4</sub> species (Spartina alterniflora and Spartina *patens*) were associated with sediment  $\delta^{13}$ C values between -18.9‰ and -15.8‰ and 19 20 occurred at elevations from mean tide level (MTL) to mean higher high water (MHHW). 21 Brackish transitional settings vegetated by Phragmites australis with Iva fructescens and 22 Typha sp ( $C_3$  species) and freshwater upland samples ( $C_3$  species) were characterized by bulk sediment  $\delta^{13}$ C values of -27.0% to -22.0% and existed above MHHW. Parallel 23 24 transects at one site suggested that intra-site variability was not discernible. The utility of  $\delta^{13}$ C values for reconstructing relative sea level in New Jersey is limited by an inability to 25 26 differentiate between brackish sediments related to sea level and freshwater upland 27 samples. To facilitate this distinction in a 4.4 m core, we used presence or absence of 28 agglutinated for a precision of the second s -18.9‰ was derived from a vegetated salt marsh and formed between MTL and MHHW. 29 30 Sediment more depleted than -22.0‰ and containing agglutinated foraminifera formed in 31 a brackish transitional zone between MHHW and HAT. Sediment more depleted than 32 -22.0‰ and lacking foraminifera formed above MHHW and maybe unrelated to former 33 sea level. Samples with intermediate values (-22.0% to -18.9%) formed between MTL and HAT. Radiocarbon dates suggest that a transition from brackish to salt marsh  $\delta^{13}C$ 34 35 values recorded in the core took approximately 350 years (1807-1452 years BP). 36 Keywords

37 Stable carbon isotope, salt marsh, New Jersey, sea level, Spartina

### 38 1 Introduction

39 Stable carbon isotopes have been used to determine the botanical and environmental 40 origin of organic material preserved in coastal sedimentary archives (Chmura and 41 Aharon, 1995; Wilson et al., 2005a; Lamb et al., 2006; González and Törnqvist, 2009). 42 In temperate regions, the transition between freshwater, salt marsh and marine settings 43 presents a strong environmental and elevational gradient, which is reflected in the stable 44 carbon isotopic signature of plants and bulk-organic sediments (Chmura et al., 1987; Matson and Brinson, 1990; Goñi and Thomas, 2000).  $\delta^{13}$ C values are the  ${}^{13}$ C: ${}^{12}$ C ratio 45 46 measured in samples and expressed in parts per mil (‰) compared to a standard reference 47 sample (Pee Dee Belemnite, PDB). During photosynthesis, land plants preferentially concentrate the <sup>12</sup>C isotope to varying degrees; this fractionation of atmospheric carbon is 48 49 recorded in plant tissues. Species using  $C_3$  (Calvin-Benson) and  $C_4$  (Hatch-Slack) photosynthetic pathways are associated with  $\delta^{13}$ C values of -34‰ to -23‰ and -17‰ to 50 51 -9‰ respectively (Chmura and Aharon, 1995; Lamb *et al.*, 2006). Along the temperate 52 northeast and mid-Atlantic coasts of the USA, salt marshes are predominantly vegetated 53 by grasses (e.g. Spartina spp.) utilizing the C<sub>4</sub> pathway (Middleburg *et al.*, 1997; Johnson 54 et al., 2007; Tanner et al., 2007). In contrast, freshwater uplands from elevations above 55 the uppermost limit of tidal inundation are associated with plants using the C<sub>3</sub> pathway (Middleburg et al., 1997; Lamb et al., 2006). As the dominant input to salt-marsh 56 sediments is likely derived from vascular vegetation (Malamud-Roam and Ingram, 2001; 57 Lamb *et al.*, 2006), bulk sediment  $\delta^{13}$ C values can be used as a proxy for describing the 58 59 dominant vegetation at the time of deposition (Malamud-Roam and Ingram, 2004). However, measurements of  $\delta^{13}$ C in bulk sediment also include allochthonous material 60

61	that may be derived from fresh, brackish or marine environments as either dissolved or
62	particulate matter (Lamb et al., 2006; Gebrehiwet et al., 2008). Further difficulties arise
63	from the presence of C <sub>3</sub> plants that are tolerant of tidal submergence (e.g. Juncus
64	roemerianus), which limits use of this technique on the southeastern and Gulf coasts of
65	the USA where C <sub>3</sub> species dominate both freshwater upland and salt-marsh environments
66	making them indistinguishable from one another on the basis of bulk-sediment $\delta^{13}C$
67	values alone (Chmura and Aharon, 1995; Kemp et al., 2010).
68	

.

•.1 1• 1

1 1 1 1 0 0 1 1 1 1

<u>\_1</u>

69 The ability to distinguish between sediments derived from salt-marsh and freshwater upland environments presents a means to use  $\delta^{13}$ C values as a sea-level indicator by 70 recognition of these floral environments in organic sedimentary sequences (Wilson et al., 71 2005b; Lamb *et al.*, 2007). This approach is applicable in study areas where  $C_3$  and  $C_4$ 72 73 plants have existed (with different distributions) over the period under consideration (Wilson et al., 2005a) and requires an understanding of the influence of post-depositional 74 diagenesis on measured  $\delta^{13}$ C values (DeLaune, 1986; Ember *et al.*, 1987; Fogel *et al.*, 75 76 1989). The precision of this approach may be increased if salt marshes can be further divided using  $\delta^{13}$ C values into floral zones (high and low salt marsh) characterized by 77 78 varying proportions of  $C_3$  and  $C_4$  inputs (Edwards, 2007). To be used as a sea-level indicator, it is necessary to quantify the relationship between bulk sediment  $\delta^{13}$ C values 79 80 (or the vegetation types they represent) and elevation in the tidal frame (Shennan, 1986; 81 van de Plassche, 1986). This relationship is formalized by the indicative meaning, which 82 is the elevational range occupied by a sea-level indicator (indicative range) in relation to 83 a contemporaneous tide level (reference water level).

85	In this study, we investigate the use of $\delta^{13}$ C values from bulk-organic sediments to
86	identify salt-marsh floral zones and be used as sea-level indicators in southern New
87	Jersey, USA. The $\delta^{13}$ C values of modern (surface) bulk sediments and plants were
88	measured in samples collected along salt-marsh transects reflecting an environmental and
89	elevational gradient at three study sites (Figure 1). Replicate transects at one site (Leeds
90	Point) facilitates investigation of intra-site variability. We use this modern dataset in
91	tandem with foraminiferal data (Kemp et al.) to interpret $\delta^{13}$ C values measured in a 4.4 m
92	core of organic sediment as changes in floral composition. This application provides a
93	means to consider the strengths and limitations of bulk sediment $\delta^{13}C$ values as sea-level
94	indicators in southern New Jersey salt marshes and similar regions. Preliminary results
95	from radiocarbon dating estimate the timing and duration of a change in dominant plant
96	community preserved in the core (1807 to 1452 years BP).

## 98 2 Study Area

99 The central and southern Atlantic coast of New Jersey is characterized by a chain of 100 barrier islands, separating a back-barrier lagoon system from the open ocean. The coast 101 between Great Bay to the north and Cape May to the south (Figure 1) includes nine inlets 102 between barrier islands that typically decrease in size from north to south along the coast 103 (Ferland, 1990). These inlets facilitate exchange of water between the Atlantic Ocean 104 and lagoons. Large areas of formerly open-water lagoon have been infilled by vertical 105 accretion of salt-marsh sediment (Daddario, 1961; Meyerson, 1972; Thorbjarnarson *et* 

106	al., 1985; Psuty, 1986; Ferland, 1990). The resulting sequences of sediment provid
107	archives of Holocene sea-level and environmental change.

109 Modern salt marshes in this region form extensive platforms. Tidal flats are rare as the 110 coast is experiencing ongoing erosion (Dolan et al., 1979; Fitzgerald et al., 2008). A 111 low-marsh floral zone of *Spartina alterniflora* (tall form) is frequently present, while 112 high-marsh floral zones are characterized by Spartina patens, Spartina alterniflora (short 113 form) and Distichlis spicata (Daddario, 1961). The border between salt marshes and 114 freshwater upland is vegetated by Phragmites australis and Iva fructescens, with less 115 frequent occurrences of *Typha* sp. and *Scirpus* sp. This zone is typically narrow and 116 represents brackish conditions (Daddario, 1961; Stuckey and Gould, 2000). 117 118 The region has a semidiurnal tidal cycle and is microtidal. Tidal ranges (MLLW to 119 MHHW) are slightly larger on the ocean side of the barrier islands (1.4 m at Atlantic 120 City; Figure 1) than in the lagoons. Tidal ranges at the study sites around Great Bay were 121 estimated by VDatum (Yang et al., 2008) to be 1.1 m at Leeds point and Bass River and 122 1.3 m at Brigantine Barrier. 123

The three sites described in this study were also the focus of an investigation into the modern distribution of salt-marsh foraminifera (Kemp et al.). Leeds Point is situated on the west side of Great Bay (Figure 1). Salt-marshes in this area frequently exceed 1 km in width (Ferland, 1990). We sampled two transects (A-A' and B-B'; Figures 1a and 2a) that extended from freshwater upland, through a narrow (10-20 m wide) brackish zone

129	vegetated by <i>Phragmites australis</i> and <i>Typha</i> sp., a wide (up to 100 m) high-marsh floral
130	zone dominated by Spartina patens associated with Spartina alterniflora (short form) and
131	a narrow (less than 10 m) low-marsh floral zone bordering a tidal channel characterized
132	by low-density stands of Spartina alterniflora (tall form) and unvegetated muddy
133	sediment. Core EF10 was collected at the Leeds Point site in a high salt marsh
134	environment (Figure 1).
135	
136	We established a 50 m long transect (C-C') at the confluence of Bass River with Great
137	Bay (Figure 1). The transect ran from a brackish, transitional (salt-marsh to upland) zone
138	defined by Phragmites australis, through a high-marsh floral zone dominated by Spartina
139	patens and Spartina alterniflora (short form) and into a narrow (less than 10 m wide)
140	low-marsh floral zone of tall-form Spartina alterniflora (Figures 1b and 3a).
141	
142	The site at Brigantine Barrier is a back-barrier salt marsh (Figure 1). A 120 m transect
143	(D-D') encompassed the brackish transition from freshwater upland to salt marsh
144	dominated by Phragmites australis and Iva fructescens (less than 10 m wide), a high-

145 marsh floral zone defined by *Spartina patens* and *Spartina alterniflora* (short form) and a

- 146 low-marsh floral zone of patchy *Spartina alterniflora* (tall form) and exposed muddy
- 147 sediment (Figures 1c and 4a).

148

# 149 **3 Methods**

150 *3.1 Sampling Regime* 

151 At the three sites we established transects across the modern salt marsh, which were 152 positioned to include the full range of physiographic environments at each site (Figure 1). 153 Sampling stations reflected changes in elevation and vascular vegetation. At each station 154 we collected bulk surface (0-1 cm) sediment for analysis. Two parallel transects at Leeds 155 Point were used to consider the influence of intra-site variability. Sample elevations were 156 established using Real Time Kinematic (RTK) satellite navigation with a minimum of 157 2000 base station observations (Leica GPS 1200+). Individual samples were leveled to 158 base stations using a total station and VDatum was used to convert altitudes from 159 orthometric to tidal datums. We collected examples (leaf and stem) of living salt-marsh 160 plants (Spartina alterniflora, Spartina patens and Phragmites australis) from Leeds Point for comparison with measured bulk sediment  $\delta^{13}C$  values at stations where these species 161 162 were the dominant type of vegetation. 163

A core (EF10) was selected for analysis from the Leeds Point site following stratigraphic
investigation. The core was recovered in 50 cm sections using a Russian-type hand core.
It was sampled at a resolution of 5 cm in the laboratory to ensure that all stratigraphic
units were adequately represented. Each core sample consisted of a 1 cm thick section of
sediment.

169

170 *3.2 Stable carbon isotopes preparation and measurement* 

171 Modern and core bulk sediment samples were prepared for measurement of  $\delta^{13}$ C, C:N

and Total Organic Carbon (TOC) by treatment with 5% HCL for 18 hours. They were

then washed with deionized water. Plant samples were washed with deionized water to

174	remove sediment particles. All sample types were dried overnight in an oven at 40°C and
175	milled to a fine powder using a pestle and mortar. <sup>13</sup> C: <sup>12</sup> C and TOC analyses were
176	performed by combustion in a Costech Elemental Analyzer coupled on-line to a Optima
177	dual-inlet mass spectrometer, with $\delta^{13}C$ values calculated to the Vienna Pee Dee
178	Belemnite (VPDB) scale using a within-run laboratory standard (cellulose, Sigma
179	Chemical prod. no. C-6413) calibrated against NBS-19 and NBS-22. C:N ratios were
180	analyzed on the same instrument and ratios were calibrated through an acetanilide
181	standard. C:N results are presented on a weight to weight basis. Replicate analysis of
182	well-mixed samples indicated a precision of $\pm <0.1\%$ for $\delta^{13}C(1\sigma)$ . For measurements
183	of nitrogen, the precision was $\pm 0.16 (1\sigma)$ .

#### 185 *3.3 Radiocarbon ages*

186 Radiocarbon ages were obtained for five samples from core EF10. The selected samples 187 were identifiable macrofossils of common salt-marsh species that were determined to be 188 in growth position and had a known relationship to the former marsh surface. In addition 189 a woody fragment lying horizontally in the core was interpreted as having been deposited 190 on a former marsh surface. Each sample was cleaned under a microscope to remove 191 contaminating material such as adhered sediment or invasive younger roots and dried at 192 <50°C. Radiocarbon ages were calibrated individually using Calib 6.0.2 (Stuiver and 193 Reimer, 1993) and the IntCal09 calibration curve. We report original radiocarbon ages 194 and calibrated dates (with  $2\sigma$  calibrated uncertainty), expressed by convention as years 195 before present (BP) where zero is AD 1950 (Stuiver and Polach, 1977).

196

### **4 Results**

198 4.1 Characterization of modern salt-marsh sediments and plants

- 199 The  $\delta^{13}$ C, C:N and TOC composition of modern salt-marsh sediments was measured in
- 200 61 surface (0-1 cm) samples collected at the three study sites. Along Leeds Point transect
- 201 A (Figure 2, left panels, A-D), samples between 0 and 22 m (stations 1 to 5) had  $\delta^{13}$ C
- 202 values from -27.0‰ to -22.7‰, TOC of 4% to 32% and C:N ratios of 12.4 to 14.4.
- 203 These samples were associated with freshwater upland and *Phragmites australis* with *Iva*
- 204 *fructescens* environments. Samples from 30-108 m along the transect in areas vegetated
- by Spartina patens and Spartina alterniflora recorded  $\delta^{13}$ C values from -17.5% to
- 206 -15.8‰, TOC between 7.5 and 23.6% and measured C:N ratios from 12.1 to 16.6 (Figure

207 2).

208

- 209 A similar pattern was observed on the second transect from Leeds Point (Figure 2, right
- 210 panels, E-H). Samples between 0 and 25 m (stations 1 to 8) had  $\delta^{13}$ C values from

211 -26.7‰ to -22.9‰, TOC of 3-32% and C:N ratios of 12.9 to 15.3. These samples were

- associated with freshwater upland and *Phragmites australis* with *Iva fructescens*
- 213 environments. Between 34 and 81 m (stations 9 to 18),  $\delta^{13}$ C values measured in 10
- samples from Spartina patens, Spartina alterniflora and unvegetated muddy zones varied

215 from -18.7‰ to -16.2‰ with TOC values of 7-24% and C:N ratios of 12.6-17.3.

- 217 At Bass River (Figure 3), a single sample (station 1) situated in a stand of *Phragmites*
- 218 *australis* had a  $\delta^{13}$ C value of -25.0‰, TOC of 2.6% and C:N ratio of 14.6. Samples
- 219 collected from Spartina patens, Spartina alterniflora and muddy unvegetated sediment

- 220 zones had measured  $\delta^{13}$ C values of -20.5‰ to -15.4‰, TOC values of 0-33% and C:N 221 ratios of 8.8 to 22.4 (Figure 3).
- 222

223	The transect at Brigantine Barrier included a single sample (station 1) situated in a stand
224	of mixed stand of <i>Phragmites australis</i> and <i>Iva fructescens</i> which had a $\delta^{13}$ C value of
225	-22.0‰, TOC of 26.2% and C:N ratio of 13.8 (Figure 4). Stations 2-12 were situated in a
226	mixed zone of Spartina patens, Spartina alterniflora and Salicornia spp. These samples
227	had $\delta^{13}$ C values of -18.9‰ to -16.1‰, measured TOC of 1% to 38% and C:N ratios
228	between 10.5 and 15.5. Three samples collected from unvegetated muddy sediment at
229	115-117 m (stations 13-15) had $\delta^{13}C$ values of -18.5‰ to -16.5‰, measured TOC of 5%
230	to 10% and C:N ratios between 9.5 and 14.1 (Figure 4).
231	
232	We measured $\delta^{13}$ C, TOC and C:N in stems and leaves from single examples of salt-marsh

233 plants collected at Leeds Point (Figure 1). The Spartina alterniflora (C4 photosynthetic

pathway) specimen had  $\delta^{13}$ C values of -12.4‰ and -13.0‰ for its stem and leaf

respectively, with TOC of 38.7% and 43.1% and C:N ratios of 92.9 and 35.5. *Spartina* 

236 *patens* (C<sub>4</sub> photosynthetic pathway) stem material had a  $\delta^{13}$ C value of -13.8‰, TOC of

43.7% and C:N of 52.9. A leaf from the same plant yielded a  $\delta^{13}$ C value of -14.0‰, TOC

of 29.9% and C:N of 36.6. An example of *Phragmites australis* (C<sub>3</sub> photosynthetic

pathway) recorded stem and leaf  $\delta^{13}$ C values of -25.2‰ and -24.6‰ respectively. TOC

240 measured from stem material was 47.5% compared with 41.9% in the leaf. C:N ratios

were 135.7 in the stem and 24.8 in the leaf.

# 243 *4.2 Characteristics of bulk sediments in core EF10*

244	We measured $\delta^{13}$ C, TOC and C:N in 91 samples (1 cm thick) of sediment recovered from
245	the upper 4.2 m of core EF10 which represents the sedimentary units above the basal
246	sand (Figure 6). $\delta^{13}$ C values from the lowermost section of the core (4.20 m to 3.35 m
247	depth) varied from -26.8‰ to -22.2‰. Between 3.35 m and 2.80 m there was a trend
248	toward less depleted $\delta^{13}$ C values (Figure 6), ten samples in this interval varied from
249	-24.8‰ to -19.1‰. There was relatively little variability in $\delta^{13}$ C from the upper 2.80 m
250	of the core, where measured values varied from -16.2‰ to -13.1‰. Measured TOC
251	between 4.20 m and 3.40 m in the core increased from 3% to 39.5% (Figure 6). A
252	reversal of this trend was observed between 3.40 m and 2.40 m with measured TOC
253	values that decreased from 39.5% to 8.6%. In the upper part of the core (top 2.40 m),
254	TOC increased to a peak of 32.8% at 1.40 m (Figure 6) and averaged 18.3%. There was
255	no clear trend in measured C:N ratios from core EF10 (Figure 6), which varied from 13.8
256	to 32.8 (average 20.9), with the exception of two anomalous data points at depths of 0.80
257	m (C:N of 45.7) and 0.05 m (C:N of 46.6).

258

#### 259 4.3 Radiocarbon ages

Four plant macrofossils from depths of 3.14 m, 2.82 m, 2.68 m and 2.45m in core EF10

261 were radiocarbon dated (Table 1). Three of the samples were identified as *Spartina* 

262 *patens* and yielded  $\delta^{13}$ C values consistent with this interpretation (Chmura *et al.*, 1987).

263 One macrofossil was a rhizome and of *Scirpus* sp. We also dated a horizontal fragment

of wood at 3.27 m. After calibration these dates spanned the interval from 1806 to 1378

265 years BP with  $2\sigma$  uncertainty from  $\pm 25$  to  $\pm 85$  years.

### 267 **5 Discussion**

#### 268 5.1 Modern distribution of salt-marsh plants

269 Salt-marsh plant communities form elevation-dependent floral zones because of their 270 differing tolerances to frequency and duration of inundation by saline water (Chapman, 271 1960; Redfield, 1972; Niering and Warren, 1980). This distinctive pattern provides a 272 means to reconstruct relative sea level by recognition of these floral zones in coastal sedimentary archives where organic material has accumulated, such as salt marshes, 273 274 infilled lagoons and estuaries (Shennan, 1986). To do so, requires that the elevational 275 range of each floral zone can be robustly estimated from modern salt marshes. In 276 southern New Jersey, we recognized three salt-marsh floral zones which are present at 277 each of the study sites. These zones were assigned a conservative (broad elevational 278 range to capture ecologically rare occurrences) indicative meaning (Table 2), similar to 279 those established by van de Plassche (1991) in Connecticut, USA. The narrow zone of 280 *Phragmites australis* (often with *Typha* sp. and *Iva fructescens*) between freshwater 281 upland and salt-marsh environments was given an indicative meaning of having formed 282 above mean higher high water (MHHW). No upper limit was established for this plant 283 community because *Phragmites australis* also occurs throughout the study region at sites 284 without marine influence such as freshwater marshes and the periphery of lakes and 285 ponds. High-marsh floral zones dominated by Spartina patens, Distichlis spicata and 286 stunted Spartina alterniflora were associated with elevations between MHW and MHHW 287 (Table 2). Measurements of the modern boundary between high-marsh and brackish transitional floral zones at the Leeds Point site (n=88) located it within 2 cm of MHHW. 288

289	Low-marsh floral zones characterized by Spartina alterniflora (tall form) were assigned
290	an indicative meaning of mean tide level (MTL) to MHW. At Leeds Point, 70
291	measurements of the modern boundary between Spartina alterniflora and the high-marsh
292	floral zone confirmed that it occurred within 1 cm of MHW with a $\pm 1\sigma$ confidence
293	interval of 6 cm.

295 The modern distribution of salt-marsh foraminifera was described at the same three study 296 sites by Kemp et al. (in review), who recognized five distinct assemblages. In freshwater 297 upland environments, foraminifera were absent in surface sediments. Foraminifera 298 require brackish, saline or marine conditions and are not found in sediments from 299 freshwater upland environments along the landward edges of salt marshes (Scott and 300 Medioli, 1978; Gehrels, 1994; Edwards et al., 2004). The brackish zone of Phragmites 301 australis, Iva fructescens and Typha sp. was inhabited by Haplophragmoides manilaensis 302 , or alternatively Jadammina macrescens with Trochammina inflata. High-marsh floral 303 environments were dominated by Arenoparrella mexicana and Tiphotrocha comprimata. 304 The low-marsh floral zone was uniformly dominated by Miliammina fusca. 305

306 5.2 Salt-marsh plant  $\delta^{13}C$  values

307 The physiological contrast between  $C_3$  and  $C_4$  photosynthetic pathways is reflected in

308  $\delta^{13}$ C values measured in living salt-marsh plants (Chmura and Aharon, 1995). Isotopic

309 differences arise from discrimination against  ${}^{13}$ CO<sub>2</sub> by the Rubisco enzyme in favor of

310 <sup>12</sup>CO<sub>2</sub>. The net effect of this process is smaller in plant species producing 4-carbon

311 sugars (C<sub>4</sub>) resulting in less depleted  $\delta^{13}$ C values (modal value -12‰) than in those

312	producing 3-carbon sugars (C <sub>3</sub> ), which have an average $\delta^{13}$ C value of -27‰
313	(Schlesinger, 1997; Choi et al., 2001; Fry, 2006). In this study, an example of
314	<i>Phragmites australis</i> (C <sub>3</sub> plant) yielded $\delta^{13}$ C values of -25.2‰ and -24.6‰. This result is
315	directly to comparable to other investigations, which have reported similar $\delta^{13}C$ values
316	(-29.4‰ to -24.6‰) for this species (Chmura and Aharon, 1995; Cloern et al., 2002).
317	The $\delta^{13}$ C values from <i>Spartina patens</i> (C <sub>4</sub> plant) in southern New Jersey (-14.0‰ and
318	-13.8‰) are within the range documented for this species in published studies along the
319	mid-Atlantic (Emery et al., 1967) and north east coasts of the USA (Middleburg et al.,
320	1997). Likewise, measured $\delta^{13}$ C values from an example of <i>Spartina alterniflora</i> (C <sub>4</sub>
321	plant) in this study (-13.1‰ and -12.4‰) fell within reported ranges for the species
322	(Ember et al., 1987; Chmura and Aharon, 1995; Goñi and Thomas, 2000; Gebrehiwet et
323	al., 2008).

# 325 5.3 $\delta^{13}C$ values in bulk surface sediments

In New Jersey, bulk surface sediments from low and high-marsh floral zones dominated 326 by Spartina alterniflora (tall form) and Spartina patens respectively, yielded  $\delta^{13}$ C values 327 of -18.9% to -15.4% (Figures 2 to 4). These values are comparable to those found along 328 329 the Gulf and Atlantic coasts of the USA for Spartina spp.-derived sediments. In Louisiana, average  $\delta^{13}$ C values of -16.5% to -16.2% were reported for low marsh 330 Spartina alterniflora (DeLaune, 1986; Chmura et al., 1987; Chmura and Aharon, 1995). 331 In North Carolina, sediments under *Spartina alterniflora* have been associated with  $\delta^{13}$ C 332 values of between -18.6‰ and -14.0‰ (Craft et al., 1988; Currin et al., 1995; Kemp et 333 *al.*, 2010). Similar sediments in South Carolina had  $\delta^{13}$ C values of -20.1‰ to -15.4‰ 334

335	(Ember <i>et al.</i> , 1987; Goñi and Thomas, 2000). In Georgia, values of -17.5‰ to -15.0‰
336	were measured (Fogel et al., 1989). Middleburg et al. (1997) showed that Spartina
337	sediments had $\delta^{13}$ C values of -19.5‰ to -14.1‰ in Massachusetts.
338	
339	Bulk-sediment $\delta^{13}$ C values from the brackish transition zone (dominated by <i>Phragmites</i>
340	australis, Iva Fructescens and Typha sp.) in New Jersey were between -27.0‰ and
341	-22.0‰ (Figures 2 to 4). Middleburg et al. (1997) reported a $\delta^{13}$ C value of -24.5‰ for
342	bulk sediment at the upland border of salt marshes vegetated by Phragmites australis,
343	Typha sp. and Scirpus sp. in Massachusetts. Whilst these species of vegetation are less
344	common in other regions, bulk sediment $\delta^{13}C$ values from the transition between
345	freshwater upland and salt marsh had an average value of -22.1‰ in Louisiana (Chmura

*et al.*, 1987).

347

Four bulk sediment samples from the freshwater upland at Leeds Point had  $\delta^{13}$ C values of 348 349 -26.5‰ to -25.1‰ (Figure 2). These values are similar to those reported for freshwater upland sediments in other studies that varied from -28.1% to -23.3%. In Louisiana, an 350 average  $\delta^{13}$ C value of -27.8‰ was provided for freshwater marshes (DeLaune, 1986; 351 352 Chmura et al., 1987), whilst forest sediments close to salt marshes in South Carolina had  $\delta^{13}$ C values of -28.8‰ to -27.5‰ (Goñi and Thomas, 2000). Bulk sediments above the 353 influence of astronomical tides were associated with  $\delta^{13}$ C values of -28.1% to -26.8% in 354 North Carolina (Kemp et al., 2010). In the San Francisco Bay estuarine system, 355 freshwater sediments were shown to have  $\delta^{13}$ C values of -27.2‰ to -23.3‰ (Cloern *et* 356

*al.*, 2002). Upland border sediments in Massachusetts were reported as having a  $\delta^{13}$ C 358 value of -24.5‰ (Middleburg *et al.*, 1997).

360	The range of $\delta^{13}$ C values reported for bulk surface sediments from <i>Spartina alterniflora</i>
361	and Spartina patens floral zones in New Jersey (-18.9‰ to -15.4‰) was more depleted
362	than living plant tissue (stems and leaves) from the same species (-14.0% to -12.4%). A
363	consistent depletion of bulk sediment $\delta^{13}C$ compared to <i>Spartina</i> spp. tissues has been
364	widely recognized (Haines, 1976; Ember et al., 1987; Benner et al., 1991). In North
365	Carolina, Spartina sediments up to 6.4‰ more depleted than corresponding plants have
366	been reported (Craft et al., 1988; Kemp et al., 2010). Similarly, Goni and Thomas (2000)
367	showed a difference of 4.0% to 6.8% in South Carolina. Bulk sediments up to 5.5%
368	more depleted than Spartina alterniflora tissue were recorded in Georgia (Fogel et al.,
369	1989; Benner <i>et al.</i> , 1991). Differences between $\delta^{13}$ C values from <i>Spartina</i> tissue and
370	sediment are a consequence of fractionation of carbon within plant tissues causing
371	cellulose and lignin from the same living plant to have different $\delta^{13}$ C values (Lamb <i>et al.</i> ,
372	2007). During early diagenesis following the plant's death, cellulose is decomposed by
373	bacterial and fungal communities at a rate several times faster than lignin (Benner et al.,
374	1987; Benner et al., 1991; Buchan et al., 2003), resulting in sediments 4‰ to 7‰ more
375	depleted than living Spartina spp. tissue (Ember et al., 1987; Fogel et al., 1989; Haddad
376	et al., 1992; Opsahl and Benner, 1995; Goñi and Thomas, 2000; Buchan et al., 2003). In
377	addition, bulk sediments include allochthonous material that can enhance or dampen
378	diagenetic differences to living plant material depending on its source (Lamb et al.,
379	2006).

381 In contrast to *Spartina* spp., further fractionation of carbon during early diagenesis was 382 not discernible (within measured ranges) between *Phragmites australis* plant tissue 383 (-25.2‰ to -24.6‰) and bulk sediment (-27.0‰ to -22.0‰). Field and laboratory experiments on *Phragmites australis* have shown that  $\delta^{13}$ C values from *Phragmites* 384 385 australis tissue underwent a change of less than 2‰ during early decomposition (Katalin 386 et al., 2006), which is less than the range reported for living examples of this species. 387 Beyond the period of initial decomposition, several investigations have shown that bulk sediment  $\delta^{13}$ C values are incorporated into coastal sedimentary archives in a manner 388 389 allowing reliable identification of floral zones after more than 3000 years (Byrne et al., 390 2001; Malamud-Roam and Ingram, 2004; Lamb et al., 2007). These studies suggested that fractionation of bulk sediment  $\delta^{13}$ C is most pronounced during the short period 391 392 following deposition of dead plant material and that bulk sediment underwent little further change. In core EF10, consistency of  $\delta^{13}$ C values between depths of 0.05 m and 393 394 2.80 m (14.8‰  $\pm$  0.8; 1 $\sigma$ ) suggests that no systematic, post-depositional shift can be 395 discerned in bulk sediments with a floral origin dominated by C<sub>4</sub> plants, corresponding to 396 approximately 1450 years (Figure 6). 397

398 At Leeds Point two parallel transects were established (Figures 1 and 2) to investigate 399 intra site variability in measured sediment  $\delta^{13}$ C values. Most previous studies used single 400 transects and sought to describe variability among sites (Wilson *et al.*, 2005a, 2005b; 401 Kemp *et al.*, 2010). Implicit in paleoenvironmental interpretations based upon single 402 transects is an assumption that small-scale (within a floral zone at a single marsh)

403	variability is not significant. On salt marshes, variability of this kind may occur due to
404	spatial changes in allochthonous inputs (Gebrehiwet et al., 2008) or differing
405	decomposition in pockets of aerobic rather than anoxic sediment. Transects at Leeds
406	Point showed a consistent pattern of measured $\delta^{13}$ C values (Figure 2b). Sediments from
407	floral zones dominated by C <sub>3</sub> plants on transect A-A' had an average $\delta^{13}C$ value of
408	-25.8‰ (-27.0‰ to -22.7‰) compared to -25.8‰ (-26.7‰ to -22.9‰) on transect B-B'.
409	Samples from high and low salt-marsh floral zones on transect A-A' yielded an average
410	$\delta^{13}$ C value of -16.7‰ (-17.6‰ to -15.8‰). Equivalent samples from transect B-B' had
411	an average $\delta^{13}$ C value of -16.8‰ (-17.6‰ to -16.2‰). Similarity between bulk sediment
412	$\delta^{13}C$ values along the two transects at Leeds Point suggests that small-scale spatial
413	variability was not significant.

. . ....

0.1 \* 1 \* 1

414

#### 415 5.4 Reconstructing Holocene relative sea-level changes

416 Establishing the botanical origin of bulk organic coastal sediments offers a means to

417 reconstruct sea level by estimating the elevation at which a sediment sample formed and

418 was deposited (Tornqvist et al., 2004; Tornqvist et al., 2006; Johnson et al., 2007).

Whilst there is no direct correlation between elevation and measured  $\delta^{13}$ C values (Kemp 419

420 et al., 2010) (Figure 5), recognition of floral zones in an appropriate stratigraphical

421 context allows relative sea level to be reconstructed. To consider the use of bulk

sediment  $\delta^{13}$ C values as a sea-level indicator in southern New Jersey we estimated the 422

- 423 indicative meaning of 91 samples from a core (EF10) at Leeds Point (Figures 1 and 5).
- 424 These estimates and interpretations are reliant upon the underlying assumption that plant

species have maintained their ecological preferences and physiography throughout theperiod under consideration, including the present.

427

We recognized four indicative meanings that could be assigned to samples in core EF 10(Table 2).

430

431 1)  $\delta^{13}C$  values less depleted than -18.9% formed between MTL and MHHW.

432 Measured  $\delta^{13}$ C values in modern bulk sediment did not distinguish between low and 433 high-marsh floral zones as both were dominated by C<sub>4</sub> plants (Figure 5). The range of 434 measured  $\delta^{13}$ C sediment values from these environments in New Jersey was -18.9‰ to 435 -15.4‰. As such, we recognized a salt-marsh environment as having  $\delta^{13}$ C values less 436 depleted than -18.9‰ and occupying an elevational range from MTL to MHHW, which 437 are the lower and upper tidal limits of vegetated modern salt-marshes in the study region 438 (Table 2). Under current tidal conditions at Leeds Point, the range from MTL to MHHW

439 is 0.59 m.

440

The sample at 1.8 m provides a unique example (in this core) of how  $\delta^{13}$ C values can be applied in relative sea level reconstructions. No foraminifera were present in the sample, which was unusual given the nature of nearby samples; it had a  $\delta^{13}$ C value of -15.2‰ (Figure 6). A bulk sediment  $\delta^{13}$ C measurement less depleted than -18.9‰ allowed this sample to be classified as having a salt-marsh origin typical of modern sites in New Jersey in light of its stratigraphic context and organic-rich nature. This example demonstrates how  $\delta^{13}$ C can be used to reconstruct relative sea level in some instances

where foraminifera (or other sea-level indicators) are not preserved or cannot be used forother reasons.

450

451 2)  $\delta^{13}C$  values more depleted than -22.0‰ and lacking agglutinated foraminifera formed 452 above MHHW.

453 Brackish (-27.0% to -22.0%) and freshwater upland (-26.5% to -25.1%) environments

454 in New Jersey could not be separated using  $\delta^{13}$ C values because both were dominated by

455 C<sub>3</sub> plants (Figure 5). As such, core samples having  $\delta^{13}$ C values associated with C<sub>3</sub> plants

456 (more depleted than -22.0‰) could only be said to have formed at an elevation above

457 MHHW. In studies seeking to reconstruct relative sea level, such samples should be

458 restricted to establishing freshwater limiting points, which constrain only the upper

459 altitude of former sea level (Shennan and Horton, 2002; Engelhart *et al.*, In Press).

460 Foraminifera were absent in all samples below 3.95 m and in a sample at 1.80 m (Kemp461 et al.; Figure 6).

462

463 3)  $\delta^{13}C$  values more depleted than -22.0‰ with presence of agglutinated for a minifera

464 *formed between MHHW and HAT.* 

465 In southern New Jersey, foraminifera are absent in modern freshwater upland sediments,

466 whilst modern brackish sediments included agglutinated taxa such as Jadammina

467 macrescens and Haplophragmoides manilaensis (Kemp et al. in review). Therefore

- 468 samples with  $\delta^{13}$ C values typical of C<sub>3</sub> plants and presence of foraminifera were
- 469 associated with brackish conditions and given a PME of MHHW to HAT (Table 2).

470 Under current tidal conditions at Leeds Point the elevational range between MHHW and471 HAT is 0.51m.

472

473 4) Samples with intermediate  $\delta^{13}C$  values (-22.0% to -18.9%) and presence of

474 *agglutinated foraminifera formed between MTL and HAT* 

475 The floral origin of samples with intermediate  $\delta^{13}$ C values is unclear, although presence

476 of agglutinated foraminifera indicates an intertidal origin. Samples of this nature were

477 given a PME of MTL to HAT to reflect this uncertainty, which corresponds to a 0.90 m

- 478 range at Leeds Point today (Figure 6).
- 479

480 Samples from core EF10 at depths below 3.95 m had  $\delta^{13}$ C values of -26.8‰ to -25.4‰

481 (Figure 6). Foraminifera were absent in all of these samples (Figure 6), therefore we

482 assigned a PME of above MHHW. Between 3.95 m and 2.80 m, 29 samples had

483 measured  $\delta^{13}$ C values from -27.0% to -16.2%. Agglutinated foraminifera were present

484 in all of these samples. Of this group, 24 samples were more depleted than -22.0‰ and

485 coupled with the presence of agglutinated foraminifera were considered to represent

486 brackish conditions and assigned a PME of MHHW to HAT. Five samples had

487 transitional  $\delta^{13}$ C values (between the ranges of C<sub>3</sub> and C<sub>4</sub> plants) and were assigned a

488 conservative PME of MTL to HAT. Measured  $\delta^{13}$ C values in 58 samples in the upper

489 2.80 m of core EF10 ranged from -18.4‰ to -13.1‰. These values are within the range

490 of modern sediments from vegetated salt marshes in southern New Jersey and we

491 assigned these samples a PME of MTL to MHHW (Figure 6).

492

Bulk sediment  $\delta^{13}$ C values that are synonymous with a dominance of C<sub>4</sub> plants may be 493 494 used to reconstruct relative sea level in instances where the sedimentary context of a 495 sample and modern transects supports its interpretation of having a salt-marsh origin. 496 Relative sea level is reconstructed by subtracting estimated PME from measured altitude 497 (with respect to the same tidal datum) for each sample. This scenario is applicable to the 498 northeast and mid-Atlantic coasts of the USA where C<sub>4</sub> plants are (and have been) the 499 dominant plant species on vegetated salt marshes (van de Plassche, 1991; Gehrels, 1994; 500 Middleburg et al., 1997). The absolute elevational range corresponding to MTL-MHHW 501 varies among and within these regions due to differences in tidal range, making the 502 potential precision of this approach geographically variable. The threshold used for 503 distinguishing such samples would vary slightly depending on the modern data used, but 504 was less than 2‰ among the modern sites we documented. Appropriate modern datasets 505 describing the distribution of plants with respect to tidal datums and of sufficient scope to 506 include salt marsh, brackish and upland floral zones are necessary to calibrate paleoenvironmental interpretations of  $\delta^{13}$ C values. 507

508

509 Understanding changes in plant community and subsequently relative sea level using core 510 samples with  $\delta^{13}$ C values typical of C<sub>3</sub> plants is made difficult by the inability to 511 distinguish freshwater environments that are not restricted to tidal limits from brackish 512 floral environments in the uppermost part of the tidal frame. Caution dictates that these 513 samples be used as freshwater limiting points in instances where only  $\delta^{13}$ C values are 514 available for interpretation. However, presence of agglutinated foraminifera in such 515 samples allows them to estimate PME, because their distribution is restricted to intertidal

516	environments. Indeed, the combination of a $C_3 \delta^{13}C$ value and presence of agglutinated
517	foraminifera in New Jersey (and similar regions) restricts estimated PME to the interval
518	between MHHW and HAT, which is a more precise interpretation than is possible from
519	$C_4  \delta^{13} C$ values either in isolation or with the presence of agglutinated foraminifera. One
520	reason for investigating the use of stable carbon isotopes to reconstruct sea level was to
521	provide an instrumental means to measure sea-level indicators rather than relying on time
522	consuming and specialist counting of microfossils such as foraminifera. However,
523	determining presence or absence of agglutinated foraminifera can be done quickly,
524	cheaply and with minimal consideration of taxonomy.
525	
526	In contrast to New Jersey and similar regions, high salt-marsh floral zones along the
527	southeastern Atlantic and Gulf of Mexico coasts are often dominated by the $C_3$ plant
528	Juncus roemerianus (Eleuterius, 1976). Although C <sub>4</sub> plants such as Distichlis spicata do
529	exist as patches in the high marsh, their dominance is frequently restricted to low marsh
530	floral zones (Chmura et al., 1987; Kemp et al., 2010). The difficulties of
531	paleoenvironmental interpretation in this region were recognized by (Chmura and
532	Aharon, 1995) and described with specific reference to relative sea-level reconstruction
533	by (Kemp et al., 2010). In such settings it is challenging to distinguish among fresh,
534	brackish and high salt-marsh floral zones using $\delta^{13}C$ values. Therefore the indicative
535	meanings of $\delta^{13}$ C values described from New Jersey are not applicable in regions with
536	different salt marsh biomes or to buried sediments that formed in these circumstances.
537	The modern geographic division between these salt-marsh ecological regions on the
538	Atlantic coast of the USA is shown by marked contrasts in the distribution of Juncus

539	roemerianus, which covers 49-77% of salt marsh area in North Carolina, less than 10% in
540	Virginia and Maryland and less than 0.1% in Delaware and states further north
541	(Eleuterius, 1976).

543 5.5 Implications for understanding salt-marsh evolution

544 Core EF10 provides some insight into how salt marshes were established at Leeds point 545 and the relative usefulness of stable carbon isotopes and foraminifera for establishing the 546 floral and environmental origin of sediments beneath salt marshes. The lowermost 547 section of core EF10 consists of sand and gravel that is likely a glacial outwash deposit 548 (Figure 6). This unit is overlain by unstructured, organic-rich, sediment (up to 40.8%) 549 TOC) from 4.2 m to 3.2 m, with plant macrofossils above 4.16 m determined to be 550 Phragmites australis. Agglutinated foraminifera were present above 3.95m (Figure 6). 551 We recognize this unit as a brackish salt marsh to upland transition at depths above 3.95 552 m and assigned indicative meaning reflecting this interpretation. Understanding the most basal section is difficult because  $\delta^{13}$ C values suggest a brackish or freshwater origin 553 554 which cannot be distinguished in the absence of foraminifera. The presence of 555 *Phragmites australis* plant macrofossils indicates a brackish origin, but could represent 556 downward growth into freshwater sediment. A salt-marsh peat between 2.8 m and 0.7 m and was recognized by  $\delta^{13}$ C values less depleted than -18.9% with agglutinated 557 558 foraminifera and abundant plant macrofossils typical of modern New Jersey salt marshes. 559 The uppermost 0.7 m of the core was composed of organic silt and likely reflects 560 anthropogenic alteration of the salt marsh (ditching).

561

562	The interval between 3.2 m and 2.8 m in core EF10 spans the environmental change from
563	a brackish transitional zone to salt-marsh floral community. Measured $\delta^{13}C$ values
564	displayed clear variability (-24.5‰ to -16.4‰) and were frequently transitional between
565	values associated with $C_3$ and $C_4$ floral zones (-22.0% to -18.9%; Figure 6). We propose
566	that this period of variability represents encroachment of the salt-marsh floral
567	environment that persists until today on a brackish transitional zone. Radiocarbon dates
568	show that this environmental change took approximately 350 years (1806 at 3.27 m to
569	1452 years BP at 2.82 m; Figure 6) to be manifest in measured $\delta^{13}$ C values, although
570	uncertainty in measurement and calibration can accommodate a period of between 207
571	and 501 years. A study focused on evolution of salt marsh conditions in North Carolina
572	concluded that it would take more than 200 years for bulk sediment to develop the
573	characteristics of a Spartina patens high salt marsh along the border of a freshwater
574	upland, even though the plant community is able to establish itself in three to five years
575	(Craft et al., 2002). However, the trajectories of such developments are likely to be non-
576	linear and effected by other changes during that time such as climate variability (Craft et
577	<i>al.</i> , 2002). Radiocarbon dates and $\delta^{13}$ C values from core EF10 suggest that bulk
578	sediment may have taken 350 years to reflect a change in salt-marsh floral community,
579	although it is not possible to distinguish between time taken for the dominant plant
580	species to change and time taken for bulk sediment to subsequently reflect this botanical
581	change.
582	

583 6 Conclusions

584	We investigated the use of $\delta^{13}$ C values measured in bulk organic sediment to establish the
585	botanical origin of samples from coastal sedimentary archives in New Jersey, USA as a
586	means to reconstruct relative sea level. Modern transects established at 3 sites with a
587	total of 61 samples showed that sediment derived from C <sub>4</sub> plants had $\delta^{13}C$ values from
588	-18.9‰ to -15.8‰ and included both a low salt-marsh zone vegetated by Spartina
589	alterniflora (tall form) and a high salt-marsh floral zone dominated by Spartina patens
590	and Spartina alterniflora (short form). In contrast, bulk sediment associated with C3
591	plants was characterized by $\delta^{13}$ C values of -27.0% to -22.0%. These environments
592	included brackish transitional zones vegetated by Phragmites australis with Iva
593	Fructescens and freshwater upland. A replicate modern transect at the Leeds Point site
594	demonstrated that there was no discernible intra-site variability between samples of the
595	same floral origin. Comparison of sediment $\delta^{13}C$ values with examples of living plants
596	from the study sites showed that Spartina spp. underwent diagentic change shortly after
597	deposition (up to 6.5‰), but then likely remained unchanged for 1500 years. Changes to
598	<i>Phragmites australis</i> were less than 2‰. We used 91 samples from a core collected at
599	the Leeds Point salt marsh to investigate the use of $\delta^{13}C$ values for establishing the
600	botanical origin of sediments. Four classifications of samples were proposed;
601	1) Those with $\delta^{13}$ C values less depleted than -18.9‰ were conservatively interpreted as
602	having formed on a vegetated salt marsh between mean tide level (MTL) and highest
603	astronomical tide (HAT). Such an interpretation remains valid in the absence of
604	foraminifera as shown by one example in core EF10.
605	2) Sediment more depleted than -22.0‰ and containing agglutinated foraminifera formed
606	in a brackish transitional zone between MHHW and HAT. This classification had the

greatest degree of vertical precision. Documenting presence or absence of agglutinated
foraminifera can be achieved quickly and with minimal taxonomic training, making this
combination a useful sea-level indicator.

610 3) Sediment more depleted than -22.0‰ and lacking foraminifera formed above MHHW

and maybe unrelated to former sea level. It was therefore is restricted establishing

612 limiting data. The inability of  $\delta^{13}$ C values to distinguish brackish and freshwater-derived

613 sediments is its primary limitation in New Jersey and similar regions.

614 4) Caution dictates that samples with intermediate  $\delta^{13}$ C values (-22.0% to -18.9%) be

615 interpreted as having formed between MTL and HAT, reflecting uncertainty in

616 determining floral origin.

617

618 Core EF10 records the change from a brackish transitional environment to a salt marsh

619 that persists until the present. A 0.4 m thick section spans the change between these two

620 environments and is typified by intermediate or transitional  $\delta^{13}$ C values. Radiocarbon

dating suggests that this change took place between 1807 and 1452 years before present.

622 This period is broadly similar to empirical predictions of the time needed for bulk organic

623 sediment to assume the characteristics of a new dominant vegetation, and specifically the

624 change from a  $C_3$  to  $C_4$  dominated zone.

625

#### 626 Acknowledgements

627 Funding for this study was provided by NICRR grant DE-FC02-06ER64298 and National

628 Science Foundation award EAR-0951686. C.H. Vane publishes with permission of the

629 Executive Director of British Geological Survey. We thank C. Kendrick and M. Leng for

630	stable carbon isotope measurements (National Isotope Geosciences Laboratory, UK).
631	Kemp thanks a graduate internship at the National Ocean Sciences Accelerator Mass
632	Spectrometry laboratory (Woods Hole) and Mark Roberts for radiocarbon support.
633	Engelhart was supported by the USDA Forest Service Global Change Research Program
634	of the Northern Research Station. This research was supported by Earthwatch Institute
635	Student Challenge Award Programs and we thank the students who participated so
636	enthusiastically in fieldwork. The Edwin Forsythe National Wildlife Refuge (U.S. Fish
637	and Wildlife Service) is acknowledged for their co-operation and for providing access to
638	study sites. Jerry Mead (Academy of Natural Science, Philadelphia) kindly provided us
639	with use of the RTK. This paper is a contribution to IGCP project 588 "Preparing for
640	coastal change" and PALSEA.

641 Figure Captions

642 **Figure 1:** Location of study sites in southern New Jersey (USA) at (A) Leeds Point, (B)

643 Bass River and (C) Brigantine Barrier. Surface (0-1 cm) sediment samples were

- 644 collected for analysis of stable carbon isotopes, C:N and total organic carbon along
- 645 transects at each sites. A core (EF10) was recovered from Leeds Point for analysis (A).
- 646

647 Figure 2: Stable carbon isotopes along two transects (A-A' and B-B') from Leeds Point.

648 Left panels show results from transect A (A-A'); right panels show results from transect

649 B (B-B'). (A, E) Elevation profiles of transects including zonation of vascular

650 vegetation. (B, F) Measured  $\delta^{13}$ C values from bulk surface sediment samples; (C, G)

total organic carbon (TOC) and (D, H) C:N ratios measured in bulk surface sediment

652 samples along the transects. Black and white circles represent samples with  $\delta^{13}$ C values

associated with  $C_3$  and  $C_4$  photosynthetic pathways respectively. In each panel, the error

- associated with each measurement is smaller than the symbol used.
- 655

```
Figure 3: Stable carbon isotopes along a transect (C-C') at Bass River. (A) Elevation
```

657 profile of the transect including zonation of vascular vegetation. (B) Measured  $\delta^{13}$ C

values from bulk surface sediment sample; (C) total organic carbon (TOC) and (D) C:N

ratios measured in bulk surface sediment samples along the transects. Black and white

660 circles represent samples with  $\delta^{13}$ C values associated with C<sub>3</sub> and C<sub>4</sub> photosynthetic

661 pathways respectively. In each panel, the error associated with each measurement is

smaller than the symbol used.

**Figure 4:** Stable carbon isotopes along a transect (D-D') at Brigantine Barrier. (A)

665 Elevation profile of the transect including zonation of vascular vegetation. (B) Measured

 $\delta^{13}$ C values from bulk surface sediment samples; (C) total organic carbon (TOC) and (D)

667 C:N ratios measured in bulk surface sediment samples along the transects. Black and

668 white circles represent samples with  $\delta^{13}$ C values associated with C<sub>3</sub> and C<sub>4</sub>

669 photosynthetic pathways respectively. In each panel, the error associated with each

670 measurement is smaller than the symbol used.

671

**Figure 5:** Relationship between elevation and measured  $\delta^{13}$ C values in bulk organic 672 673 sediment from vegetated sampling stations at three modern salt marshes. Elevations are 674 expressed as a standardized water level index (SWLI) to allow comparison among sites 675 with different tidal ranges. Tidal datums are shown for reference. Symbols represent the 676 floral environment from which samples were collected and symbol shading denotes site. Grey regions show elevation and  $\delta^{13}$ C thresholds used for defining environmental origin. 677 678 HAT = highest astronomical tide; MHHW = mean higher high water, MHW = mean high 679 water, MTL = mean tide level. Value used for HAT is from the Atlantic City tide gauge. 680 **Figure 6:** Measured values of  $\delta^{13}$ C, total organic carbon (TOC) and C:N ratios in 91 681

samples from core EF10. Measurement errors (<0.1‰) are smaller than the symbols used. Vertical dashed lines differentiating C<sub>3</sub>, C<sub>4</sub> and transitional values are limits established from the four modern transects. Filled circles show position of radiocarbon dates with mid-point ages. Downcore presence of agglutinated foraminifera typical of salt marshes (SMF) is shown by filled bars, while open bars show samples in which no

- 687 foraminifera were present (from Kemp et al., in review). Paleomarsh elevation (PME,
- right panel) was estimated for samples with  $\delta^{13}$ C values typical of C<sub>4</sub> salt-marsh plants as
- 689 mean tide level (MTL) to mean higher high water (MHHW). Samples with  $\delta^{13}$ C values
- 690 associated with C<sub>3</sub> plants and the presence of agglutinated foraminifera were assigned a
- 691 PME from MHHW to highest astronomical tide (HAT). Samples with  $\delta^{13}$ C values
- 692 associated with C<sub>3</sub> plants and no salt-marsh foraminifera were assumed to have formed
- above MHHW (indicated by the arrow). MLW = mean low water.

# 694 **Table 1: Radiocarbon ages**

Depth (m)	<sup>14</sup> C Age	δ <sup>13</sup> C	Macrofossil	Max BP	Min BP	Lab Code
3.27	$1880\pm30$	-12.69	Horizontal woody fragment	1728	1884	OS-87528
3.14	$1750\pm30$	-26.47	Scirpus sp.	1562	1731	OS-79178
2.82	$1550 \pm 25$	-14.4	Spartina patens	1383	1521	OS-66514
2.68	$1541 \pm 14$	-14.57	Spartina patens	1379	1517	OS-70445
2.45	$1502\pm14$	-13.24	Spartina patens	1349	1407	OS-70443

Radiocarbon ages on from core EF10. Ages at 2.68 m and 2.45 m were derived from extended AMS counting to reduce analytical uncertainty and are not reported following rounding conventions. Maximum and minimum are calibrated ages (using Calib 6.0.2 with IntCal09) before present (BP).  $\delta^{13}$ C was measured in a CO<sub>2</sub> aliquot collected during sample combustion and represents a value for the dated macrofossil and not the bulk sediment from which it was recovered. Radiocarbon ages were corrected for the effect of  $\delta^{13}$ C fractionation by the reporting laboratory.

# **Table 2**

Floral Zone	Dominant Vegetation	<b>Elevational Range</b>	$\delta^{13}C$ (‰)
Low salt marsh	Spartina alterniflora (tall form)	MSL to MHW	>-18.9
High salt marsh	Spartina patens Spartina alterniflora (short form)	MHW to MHHW	>-18.9
Brackish transition	Phragmites australis Typha sp. Iva fructescens	MHHW to HAT <sup>1</sup> Above MHHW <sup>2</sup>	<-22.0

705	Indicative meanings assigned to salt-marsh floral zones. These values provided estimates
706	of paleomarsh elevation for samples in core EF10. MSL = mean sea level, MHW = mean
707	high water, MHHW = mean higher high water, HAT = highest astronomical tide. For the
708	brackish transition zone, we used two different ranges depending on the presence $(^{1})$ , or
709	absence ( <sup>2</sup> ), of agglutinated salt-marsh foraminifera that are not present in modern
710	freshwater upland environments (above HAT).

## 711 **References**

712 Benner R, Fogel, ML, Sprague, EK. 1991. Diagenesis of belowground biomass of 713 Spartina alterniflora in salt-marsh sediments. Limnology and Oceanography 36: 1358-714 1374. 715 Benner R, Fogel, ML, Sprague, EK, Hodson, RE. 1987. Depletion of  $^{13}C$  in lignin and its 716 717 implications for stable isotope studies. Nature 329: 708-710. 718 719 Buchan A, Newell, SY, Butler, M, Biers, EJ, Hollibaugh, JT, Moran, MA. 2003. Dynamics 720 of bacterial and fungal communities on decaying salt marsh grass. Applied and 721 Environmental Microbiology 69: 6676-6687. 722 723 Byrne R, Ingram, BL, Starratt, S, Malamud-Roam, F, Collins, JN, Conrad, ME. 2001. 724 Carbon-Isotope, Diatom, and Pollen Evidence for Late Holocene Salinity Change in a 725 Brackish Marsh in the San Francisco Estuary. Quaternary Research 55: 66-76. 726 727 Chapman VJ. 1960. Salt Marshes and Salt Deserts of the World. Interscience Publishers, 728 New York. 729 730 Chmura GL, Aharon, P. 1995. Stable carbon isotope signatures of sedimentary carbon in 731 coastal wetlands as indicators of salinity regime. Journal of Coastal Research 11: 124-732 135. 733 Chmura GL, Aharon, P, Socki, RA, Abernethy, R. 1987. An inventory of <sup>13</sup>C abundances 734 735 in coastal wetlands of Louisiana, USA: vegetation and sediments. Oecologia 74: 264-736 271. 737 738 Choi Y, Wanng, Y, Hsieh, YP, Robinson, L. 2001. Vegetation succession and carbon 739 sequestration in a coastal wetland in northwest Florida: evidence from carbon isotopes. 740 Global Biogeochemical Cycles 15: 311-319. 741 742 Cloern JE, Canuel, EA, Harris, D. 2002. Stable Carbon and Nitrogen Isotope 743 Composition of Aquatic and Terrestrial Plants of the San Francisco Bay Estuarine 744 System. Limnology and Oceanography 47: 713. 745 746 Craft C, Broome, S, Campbell, C. 2002. Fifteen years of vegetation and soil development 747 after brackish-water marsh creation. Restoration Ecology 10: 248-258. 748 749 Craft CB, Broome, SW, Seneca, ED, Showers, WJ, 1988. Estimating sources of soil 750 organic matter in natural and transplanted estuarine marshes using stable isotopes of 751 carbon and nitrogen. Estuarine Coastal and Shelf Science 26: 633-641. 752 753 Currin CA, Newell, SY, Paerl, HW. 1995. The role of standing dead Spartina alterniflora 754 and benthic microalgae in salt marsh food webs: considerations based on multiple stable 755 isotope analysis. Marine Ecology Progress Series 121: 99-116.

756	
757	Daddario JJ. 1961. A lagoon deposit profile near Atlantic City, New Jersey. Bulletin of
758	the New Jersey Academy of Science 6: 7-14.
759	
760	DeLaune RD. 1986. The use of $\delta^{13}C$ signature of C-3 and C-4 plants in determining past
761	depositional environments in rapidly accreting marshes of the Mississinni River deltaic
762	nlain Louisiana USA Chemical Geology: Isotone Geoscience section 59: 315-320
763	
764	Dolan R Hayden B Rea C Heywood J 1979 Shoreline erosion rates along the middle
765	Atlantic coast of the United States Geology 7: 602-606
766	manie coust of the onlice states. Geology F. 662 666.
767	Edwards RI 2007 Sea Level Studies: Low energy coasts sedimentary indicators. In
768	Encyclonedia of Quaternary Science Flias SA editor Flsevier: Amsterdam: 2004-3005
769	Encyclopeana of Qualernary Science, Enas on eanor. Ensevier, ministeriaam, 2994 5005.
770	Edwards RI Wright AI van de Plassche O 2004 Surface distributions of salt-marsh
771	for aminifera from Connecticut USA: modern analogues for high-resolution sea level
772	studias Marina Micronaleontology 51: 1-21
773	siudes. Murine Micropaleoniology <b>51</b> , 1-21.
774	Flautarius I 1976 The distribution of Juncus roemerianus in the salt marshes of North
775	America Chesaneake Science 17: 280
776	America. Chesupeare Science 17. 207.
770	Ember I.M. Williams DE Morris, IT 1987 Processes that influence carbon isotone
778	variations in salt marsh sadiments Marine Ecology Progress Series 36: 33 12
770	variations in sait marsh seatments. Marthe Ecology 1 rogress Series <b>50</b> . 55-42.
780	Emery KO Wigley RI Bartlett AS Rubin M Barghoorn ES 1967 Freshwater neat on
781	the continental shelf Science 158: 1301-1307
782	ine continental shelf. Science 150. 1501-1507.
782	Engelhart SF Peltier WR Horton RP In Press Holocene relative sea-level changes
784	and algorial isostatic adjustment of the US Atlantic coast Geology
785	and glacial isosialle adjusiment of the 0.5. Manne coust. Geology
786	Ferland MA 1990 Holocene depositional history of the southern New Jersey harrier and
787	hack harrier regions IIS Army Corps of Engineers n 75
788	buck burner regions. OS miny corps of Engineers, p. 75.
789	Fitzgerald DM Fenster MS Argow RA Ruynevich IV 2008 Coastal Impacts Due to
790	Sea-Level Rise Annual Review of Earth and Planetary Sciences 36: 601-647
791	Sea Dever Rise. Initiaal Review of Darin and Flaneary Sciences 50. 001 017.
792	Fogel MI. Kent Sprague F. Gize AP Frey RW 1989 Diagenesis of organic matter in
793	Georgia salt marshes Estuarine Coastal and Shelf Science 28: 211-230
794	Georgia suit marsnes. Estaarine, Coastat and Sheij Selence 20. 211 250.
795	Fry B 2006 Stable Isotone Ecology Springer New York
796	TTY D. 2000. Suble Isolope Leology. Springer, New Tork.
797	Gebrehiwet T. Koretsky, CM. Krishnamurthy, RV 2008. Influence of Sparting and Juncus
798	on saltmarsh sediments III: organic geochemistry Chemical Geology 255. 114-110
799	on summarsh seaments 111, of same seconemistry. Chemical Ocology 255, 114-119.
800	Gehrels WR 1994 Determining relative sea-level change from salt-marsh foraminifera
801	and plant zones on the coast of Maine USA Journal of Coastal Research 10. 990-1009

- 802
- 803 Goñi M, Thomas, K. 2000. Sources and transformations of organic matter in surface 804 soils and sediments from a tidal estuary (North Inlet, South Carolina, USA). Estuaries 805 and Coasts 23: 548-564. 806 807 González JL, Törnqvist, TE. 2009. A new Late Holocene sea-level record from the 808 Mississippi Delta: evidence for a climate/sea level connection? Quaternary Science 809 Reviews 28: 1737-1749. 810 811 Haddad RI, Newell, SY, Martens, CS, Fallon, RD. 1992. Early diagenesis of lignin-812 associated phenolics in the salt marsh grass Spartina alterniflora. Geochimica et 813 Cosmochimica Acta 56: 3751-3764. 814
- 815 Haines EB. 1976. Stable carbon isotope ratios in the biota, soils and tidal water of a 816 Georgia salt marsh. Estuarine Coastal Marine Science 4: 609-616.
- 817 818 Johnson BJ, Moore, KA, Lehmann, C, Bohlen, C, Brown, TA. 2007. Middle to late 819 Holocene fluctuations of  $C_3$  and  $C_4$  vegetation in a Northern New England Salt Marsh,
- 820 Sprague Marsh, Phippsburg Maine. Organic Geochemistry 38: 394-403.
- 821
- 822 Katalin VB, Mátyás, P, Lajos, V, Noémi, T. 2006. A study of the decomposition of reed 823 (Phragmites australis) as a possible source of aquatichumic substances by measuring the 824 natural abundance of stable carbon isotopes. International Review of Hydrobiology 91: 825 15-28.
- 826
- 827 Kemp AC, Vane, CH, Horton, BP, Culver, SJ. 2010. Stable carbon isotopes as potential 828 sea-level indicators in salt marshes, North Carolina, USA. The Holocene 20: 623-636. 829
- 830 Lamb AL, Vane, CH, Wilson, GP, Rees, JG, Moss-Hayes, VL. 2007. Assessing  $\delta^{I3}C$  and
- 831 C/N ratios from organic material in archived cores as Holocene sea level and
- 832 palaeoenvironmental indicators in the Humber Estuary, UK. Marine Geology 244: 109-833 128.
- 834 835 Lamb AL, Wilson, GP, Leng, MJ. 2006. A review of coastal palaeoclimate and relative sea-level reconstructions using  $\delta^{13}C$  and C/N ratios in organic material. Earth-Science 836 837 Reviews 75: 29-57.
- 838
- 839 Malamud-Roam F, Ingram, BL. 2001. Carbon Isotopic Compositions of Plants and 840 Sediments of Tide Marshes in the San Francisco Estuary. Journal of Coastal Research *17*: *17-29*.
- 841
- 842
- Malamud-Roam F, Ingram, BL. 2004. Late Holocene  $\delta^{13}C$  and pollen records of 843
- 844 paleosalinity from tidal marshes in the San Francisco Bay estuary, California.
- 845 Quaternary Research 62: 134-145.
- 846

847 848	Matson EA, Brinson, MM. 1990. Stable carbon isotopes and the C:N ratio in the estuaries of the Pamlico and Neuse Rivers, North Carolina. Limnology and
849	Oceanography 35: 1290-1300.
850	
851	Meyerson AL. 1972. Pollen and paleosalinity analyses from a Holocene tidal marsh
852	sequence, Cape May County, New Jersey. Marine Geology 12: 335-357.
853	
854	Middleburg JJ, Nieuwenhuize, J, Lubberts, RK, van de Plassche, O. 1997. Organic
833	carbon isotope systematics of coastal marsnes. Estuarine Coastal and Shelf Science 45:
830 857	081-087.
837 858	Nieving WA Warren BS 1080 Vegetation Patterns and Processes in New England Salt
030 950	Marshop PioScience 20: 201 207
0 <i>39</i> 060	Marshes. Dioscience 50. 501-507.
000 061	Ongohl S. Bounou B. 1005. Early diagonagia of nagonalay plant tigging. Liquin and outin
801 862	Opsant S, Benner, R. 1995. Early alagenesis of vascular plant lissues. Lighth and culture decomposition and biogeochemical implications. Coochimica et Cosmochimica Acta <b>50</b> .
802 862	<i>decomposition and diogeochemical implications. Geochimica et Cosmochimica Acta</i> <b>59</b> .
803 864	4009-4904.
804 865	Deute ND 1096 Hologene and Invelie New Jergen Divisional Conservation 7, 156 167
803 866	Psuly NP. 1980. Holocene sea level in New Jersey. Physical Geography 7: 130-107.
800 967	Redfield AC 1072 Development of a New Evaluated soft marsh Ecological Managarha
807 868	AC. 1972. Development of a New England salt marsh. Ecological Monographs
000 860	<b>42</b> . 201-237.
809 870	Schlasinger WIL 1007 Clobal Discoschemistry An Anglusia of Clobal Change
870 971	Schlesinger WH. 1997. Global Biogeochemistry: An Analysis of Global Change.
0/1 072	Acquemic Press.
872 872	Spott DR Madiali ES 1078 Vartical zonations of marsh for aminifora as accurate
873	indiactors of formar sea loyals. Natura 272: 528 531
074 975	indiculors of former sed levels. Nature 272. 526-551.
876	Shannan I 1086 Flandrian soa layal changes in the Fanland II: Tondencies of soa layal
870	movement altitudinal changes and local and regional factors. Journal of Quaternamy
878	Science 1: 155-170
870	Science 1. 155-179.
880	Shennan I. Horton, R. 2002. Holocene land-and sea-level changes in Great Britain
881	Journal of Quaternary Science 17: 511-526
882	Journal of Qualernary Science 17: 511-520.
883	Stuckey IH Gould 11 2000 Coastal plants from Cape Cod to Cape Canaveral
884	University of North Carolina Press Chanel Hill
885	
886	Stuiver M Polach H4 1977 Reporting of $^{14}C$ data Radiocarbon 19: 355-363
887	Starver M, I blach, III. 1977. Reporting of C auta. Radiocarbon 19. 555 565.
888	Stuiver M. Reimer, P.I. 1993. Extended $^{14}C$ database and revised Calib 3.0 $^{14}C$ age
889	calibration program, Radiocarbon 35: 215-230
890	

891	Tanner BR, Uhle, ME, Kelley, JT, Mora, CI. 2007. $C_3/C_4$ variations in salt-marsh
892	sediments: An application of compound specific isotopic analysis of lipid biomarkers to
893	late Holocene paleoenvironmental research. Organic Geochemistry <b>38</b> : 474-484.
894	
895	Thorbjarnarson KW, Nittrouer, CA, DeMaster, DJ, McKinney, RB. 1985. Sediment
896	accumulation in a back-barrier lagoon, Great Sound, New Jersey. Journal of
897	Sedimentary Research 55: 856-863.
898	
899	Tornqvist TE, Bick, SJ, van der Borg, K, de Jong, AFM. 2006. How stable is the
900	Mississippi Delta? Geology <b>34</b> : 697-700.
901	
902	Tornqvist TE, Gonzalez, JL, Newsom, LA, van der Borg, K, de Jong, AFM, Kurnik, CW.
903	2004. Deciphering Holocene sea-level history on the US Gulf Coast: a high-resolution
904	record from the Mississippi Delta. Geological Society of America Bulletin 116: 1026-
905	1039.
906	
907	van de Plassche O. 1986. Sea-level research: a manual for the collection and evaluation
908	of data, In: van de Plassche O (Ed.). Geobooks, Norwich, p. 618.
909	
910	van de Plassche O. 1991. Late Holocene sea-level fluctuations on the shore of
911	Connecticut inferred from transgressive and regressive overlap boundaries in salt-marsh
912	deposits. Journal of Coastal Research 11: 159-179.
913	1 0
914	Wilson GP Lamb AL Leng MJ Gonzalez S Huddart D 2005a $\delta^{13}$ C and C/N as
915	potential coastal palaeoenvironmental indicators in the Mersey Estuary UK Quaternary
916	Science Reviews 24: 2015-2029
917	
918	Wilson GP Lamb AL Leng M.I. Gonzalez S. Huddart D. 2005b Variability of organic
919	$\delta^{13}C$ and $C/N$ in the Mersey Estuary II K and its implications for sea-level
020	reconstruction studies. Estuaring Coastal and Shelf Science 64: 685-608
921	reconstruction studies. Estudrine, Coustal and Shelf Science <b>04</b> . 005-070.
921	Vana 7 Myers FP Wona 4 White S 2008 Vdatum for Chesaneake Ray Delaware
023	Ray and Adjacent Coastal Water Areas: Tidal Datums and Sea surface Topography
024	NOAA Tachnical Momorandum NOS CS 15 U.S. Denartment of Commerce National
92 <del>4</del> 925	Oceanic and Atmospheric Administration Silver Spring Maryland p 110
925 076	occume una Annospherie Auministration, suver spring, marytana, p. 110.
027	
028	
140	











