Productivity of kelp (*Laminaria* spp.) near the southern limit in the Northwestern Atlantic Ocean

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**ABSTRACT:** Annual primary production of kelp *Laminaria saccharina* was calculated from standing crop, density, tissue carbon content and growth rate measurements at 3 sites in Rhode Island, USA – 2 within Narragansett Bay and 1 in Rhode Island Sound. Results indicate similar seasonal growth patterns at all sites within Rhode Island. This pattern paralleled those in more northerly locations with rapid growth in late winter/early spring and minimal growth in summer. There was no reduction in production at the Rhode Island sites, although they are near the southern limit of this species. *L. digitata* was found at the most exposed sites where it comprised up to 23% of the total kelp density. The relative contribution of kelp to the total primary productivity varies according to season. The epiphytic community was most abundant in August and September. This study is the first report of 7 species of animals epiphytic on *Laminaria* spp. *Lacuna vincta* (Mollusca) was found to be an important grazer on the *Laminaria* blades especially in winter.

**INTRODUCTION**

Subtidal *Laminaria* zones are areas of intense primary productivity and large quantities of biomass in the northern hemisphere (see Kain, 1979 for review). Productivity of kelp has been measured in more northern areas, well within the latitudinal range of the species studied. Primary productivity of *Laminaria* spp. at the southern limit of its range is essentially unknown. Humm (1969) suggested that whereas some brown algae are limited at the southern end of their range by summer water temperatures when respiration exceeds photosynthesis, the same species may become adapted to warmer temperatures through natural variation. Hutchins (1947) proposed 2 relations that may control species distribution in the marine subtidal: (1) critical temperatures affecting individual survival; (2) critical temperatures needed to repopulate. The purpose of this study was to determine the extent to which *Laminaria* spp. were affected near their southernmost limit, Narragansett Bay and Rhode Island Sound (Taylor, 1972). The productivity of phytoplankton and zooplankton within Narragansett Bay has been measured (Smayda, 1957, 1973; Pratt, 1959, 1965; Furnas et al., 1976; Durbin and Durbin, 1981), but the productivity of macroalgae and vascular plants of Narragansett Bay have been regarded as insignificant (Kremer and Nixon, 1978). In this study, the density and standing crop of *L. saccharina* and *L. digitata* were measured, as were growth rate, mortality and productivity of *L. saccharina* in Narragansett Bay and Rhode Island Sound.

**MATERIAL AND METHODS**

**Site description**

Narragansett Bay (NB) (Fig. 1) opens to the south into Rhode Island Sound (RIS). This bay has a total area of 259 km\(^2\), a maximum length of 42 km and an average width of 6 to 8 km. Annual surface water temperatures range from -0.5 to 24°C. Mean depth of the bay is 9 m; maximum depth is 60 m. The common sediment type in the northern bay is a mixture of silt-clay, sand and gravel; sand is predominant in the south. Shale, granite, and metamorphic rock are common to the southern subtidal zone of the bay; while the northern subtidal zone substratum is a cobble-sand mixture. Twenty-six locations were surveyed using a combination of SCUBA and free diving to determine distribution, density and width of the subtidal *Laminaria* sac-
charina and L. digitata zone in Narragansett Bay and Rhode Island Sound. The locations were selected on the basis of substrata suitable for kelp attachment. Preliminary observations showed that the presence of kelp correlated with the following types of substrata: man-made, shale, granite, and offshore bedrock. A geological survey map of the substrata zones (Boothroyd and Al-Saud, 1978 in: Olsen et al., 1980) was used to determine the shoreline length within the kelp distribution range. The zones were measured with a map wheel (average of 5 measurements), and multiplied by a zone width derived from the distribution survey. The resulting value estimated the total area that could be occupied by kelp.

Sample station selection

Long-term sample stations (Fig. 1) were chosen from the distribution survey based on the following criteria: (a) existence of kelp beds of moderate to high density; (b) similar sample depths at all stations; (c) ease and safety of entering these stations throughout the year. The 2 stations in the southern portion of Narragansett Bay (NB) were: Fort Wetherill, Jamestown (41°28.5' N, 71°21.5' W) and Coaster's Harbor Island, Newport (41°29.0' N, 71°19.9' W). One station outside the Bay in Rhode Island Sound (RIS), Land's End, Newport (41°21.2' N, 71°18.6' W), was selected to compare with the 2 bay stations. The wave exposure at Coaster's Harbor Island, the northernmost study station, was more moderate than at the other stations, where it ranged from moderate to severe according to the season. Plants at Coaster's Harbor Island were harvested and tagged on a man-made stone wall, whereas at Ft. Wetherill and Land's End the substratum was cobble and bedrock.

Growth rate measurements

Meristematic growth rates of tagged Laminaria saccharina were determined using the standard hole punch procedure (Parke, 1948). Monthly measurements were recorded in situ using SCUBA at each of the 3 stations from April 1980 to June 1981. A hole 1.75 cm in diameter was punched 20 cm above the blade-stipe juncture to include the meristematic region of the blade (Parke, 1948; Mann, 1972b; Boden, 1979). Adult plants of uniform size were selected for growth measurements. Twenty individuals per station were tagged by placing a poultry wing tag at the base of the stipe. Holes were punched into the thallus after each sampling interval to measure the growth change during the next interval. These data (cm individual⁻¹ d⁻¹) were used in the subsequent calculations of primary productivity. The progress of the holes along the blades was recorded to determine the turnover of the blade length. Any plant lost during a sample interval was replaced by a newly tagged plant to maintain a sample size of 20 individuals at each station.

Biomass and density measurements

On the same days in which growth was measured, six 0.10 m² quadrats of Laminaria spp. were harvested at each sample station. During periods of severe storms the frequency of sampling was reduced. Before sampling, the depth of maximum Laminaria spp. density was estimated visually by divers swimming perpendicular to depth contours within the laminarian zone. Four sampling transects (A, B, C, D) within each study site were established, and quadrats were collected along these sampling transects. To avoid bias on the part of the divers, a random sampling technique was used to select the location of quadrats prior to the dive. For each quadrat to be collected, 3 location criteria were determined randomly: (1) a transect (A, B, C or D); (2) direction from maximum density (shallow or deep); (3) distance from maximum density. Once the samples were returned to the laboratory, the following features were measured for all harvested plants: plant density per quadrat, species composition per quadrat and individual blade length, wet weight and dry weight (dried to a constant weight in a plant dryer, 35°C). All individuals regardless of age class were counted to determine plant density.

The Kolmogorov-Smirnov test (Hollander and Wolfe, 1973) was used to determine whether the biomass and density data were normally distributed. The data were sorted and analysed by site (1) for the entire sample period and (2) at each sample date. A nonparametric 1-way analysis of variance (Kruskal-Wallis test; Hollander and Wolfe, 1973) tested for significant differences in biomass and density between sample stations.

Physical measurements

Subtidal water temperature and surface salinity were measured on each sampling date. Subtidal water temperature was measured by the divers at the depth of maximum kelp density (1.5 to 3 m below MLW). Incident solar radiation data were obtained from Eppley Laboratories, Inc. (Newport, R. I.). Light intensity was measured at various depths within the laminarian zone with a Li-Cor Quantum Meter (Li-185). Extinction coefficients of the 3 sample stations were
calculated from these light measurements. The mean annual extinction coefficients were $-0.150 \pm 0.030$ (SE) m$^{-1}$ at Coaster's Harbor Island (NB), $-0.211 \pm 0.033$ (SE) m$^{-1}$ at Pt. Wetherill (NB) and $-0.239 \pm 0.085$ m$^{-1}$ (SE) at Land's End (RIS).

### Carbon analysis

Six plants of *Laminaria saccharina* and 3 plants of *L. digitata* from the Pt. Wetherill sample station (collected March 1981) were dried to constant weight at 35 °C and ground separately using a Wiley mill (mesh size # 20). (Mann, 1972a found carbon to vary slightly in *L. longicruris* according to site or season.) The ground tissue was weighed (Perkin-Elmer Ultra Balance) and placed in preweighed aluminum boats. The samples were analysed for carbon content with a Horiba PIR-2000 Carbon Infrared Analyser while they were burned in a Lindberg Tube Furnace (compact style) at 950 °C. Combustion time depended on the return of the pen to the baseline. Oxygen was used as a carrier gas, at a flow rate of 1 l m$^{-1}$. The method was standardized using a solution of anhydrous D-glucose (Mallinkrodt Co.), distilled water and mercury (poison inhibited bacterial growth).

### Calculation of net annual primary productivity and P/B

Net annual primary productivity of the kelp for each station was calculated according to the following procedure:

1. Various lengths of the *Laminaria* blade from the meristematic region of adult plants were dried to constant weight. A step-wise polynomial regression (Snedecor and Cochran, 1967) of dry weight to blade length was used to determine the appropriate regression formula.

2. Data from growth rate measurements (mean blade length increase individual$^{-1}$ mo$^{-1}$) were substituted in the above regression for blade length to determine mean increase in dry weight individual$^{-1}$ mo$^{-1}$.

3. Mean increase in dry weight individual$^{-1}$ mo$^{-1}$ was multiplied by the ratio of carbon to dry weight to obtain the mean increase in carbon individual$^{-1}$ mo$^{-1}$. These monthly values were summed for the year to determine net primary productivity (g C individual$^{-1}$ yr$^{-1}$).

4. Net primary productivity (NPP) was multiplied by mean and median densities (individuals m$^{-2}$) of each station to obtain primary productivity (g C m$^{-2}$ yr$^{-1}$) on an area basis.

5. Production to biomass ratios (P/B) were computed for each sample station by multiplying the mean biomass produced per individual and the median density, then dividing the product by the median biomass of the harvested quadrats.

### Epiphyte assessment

The macroalgal epiphytes found on harvested *Laminaria* spp. were carefully removed from the blades, placed in a 10 % glutaraldehyde fixative with 0.1 M phosphate buffer, and stored at 10 °C until identified. Animals attached to, and associated with the *Laminaria* spp., were collected from the harvested individuals and stored at room temperature in a 70 % alcohol solution.

### RESULTS

**Distribution survey of *Laminaria* spp.**

*Laminaria saccharina* and *L. digitata* were limited to lower Narragansett Bay (NB) and Rhode Island Sound (RIS) (Fig. 1), where, in addition to suitable substrata, there was greater wave exposure and less turbidity than the upper Bay (Schenck and Davis, 1973). The length of shoreline that kelp could occupy was estimated as 53 km, and the mean kelp zone width as 6 m. Therefore, the surface area in Narragansett Bay that could support the laminarian species is 0.32 km$^2$. Not all the bedrock supports kelp, but kelp also occur sporadically on other substrata (e.g. cobble and sand). Thus the estimate of total area is reasonable. The kelp population was observed to have an extremely patchy distribution within the subtidal. However, *L. saccharina* grew sporadically in calmer waters 16 km...
north of the entrance to the Bay. Density was greatest on the southerly exposed shores within the Bay and Rhode Island Sound, and *L. digitata* could be found only at the stations with the greatest wave exposure.

**Seasonal growth of *Laminaria saccharina***

Mean monthly meristematic growth rates (cm d⁻¹) of individual plants showed a similar seasonal pattern at all 3 stations (Fig. 2). The maximum growth rate of 2 cm d⁻¹ occurred in May and June when the subtidal water temperature was 13°C and incident solar radiation was nearly 500 langleys d⁻¹ (Fig. 3). Little or no growth occurred in August and September when the subtidal water temperature was 17°C (Fig. 3) and the epiphyte community was observed to be most abundant. Salinity measurements at the 3 sample stations ranged from 30 to 34‰ through the year, and did not vary more than 1‰ from station to station. The highest mean annual growth rate was seen at Coaster’s Harbor Island (NB) whereas the lowest was at Land’s End (RIS) (Table 1). At the latter station, however, the greatest mean monthly growth rate was measured in spring 1981, but there was a slightly longer period of no growth during the summer of 1980 (Fig. 2).

Mean monthly mortality of the tagged individuals at all stations for the entire sample period (April 1980 to June 1981) was 29.3% (Fig. 2). No single plant survived the entire sample period. Maximum survival of a tagged plant was 9 mo. Maximum loss of tagged plants occurred in September at Coaster’s Harbor Island (NB) and in November/January at Ft. Wetherill (NB) and Land’s End (RIS) (Fig. 2).

**Seasonal biomass and density**

*Laminaria* at all 3 sample stations showed similar seasonal biomass fluctuations (Fig. 2). Standing crop
(g DW m\(^{-2}\)) was highest in early summer and lowest during winter. Many of the plants that survived the winter eroded to within 20 cm of the blade-stipe junction, thus reducing the biomass of the remaining population. This period was followed by a slow biomass increase in the spring. Land’s End (RIS) had the highest annual mean standing crop while Ft. Wetherill (NB) had the highest annual median standing crop (Table 1). The standing crop of Coaster’s Harbor Island (NB) was significantly lower than that of Ft. Wetherill (NB) (Kruskal-Wallis test, \(p < 0.002\)). However, the individuals at Coaster’s Harbor Island (NB) did not have a lower growth rate. The mean annual standing crops of \(L.\) saccharina at the sample stations within the Bay and Rhode Island Sound were 660 ± 54 g DW m\(^{-2}\) (median = 513 g DW m\(^{-2}\)) and 1000 ± 217 g DW m\(^{-2}\) (median = 536 g DW m\(^{-2}\)), respectively. The biomass lost from the experimental plants for the year sampled, based on the percentage of tagged plants lost and the mean individual dry weight of \(L.\) saccharina (15.1 g DW), was 699 g DW from Coaster’s Harbor Island (NB), 685 g DW from Ft. Wetherill (NB), and 1,125 g DW from Land’s End (RIS).

Blades of \(L.\) saccharina renewed their length between 3 and 4 times during the year. A turnover rate of 3.5 was multiplied by the mean standing crop of the Narragansett Bay stations to determine the mean annual biomass production of 2310 g DW m\(^{-2}\) yr\(^{-1}\). This value was multiplied by the calculated area of kelp cover (0.32 km\(^2\)) to obtain a total Narragansett Bay annual biomass of 7.4 \(\times\) 10\(^2\) metric tons (DW).

The density (individuals m\(^{-2}\)) of all age classes of \(Laminaria\) spp. showed a seasonal pattern similar to that of biomass (Fig. 2, Table 2) in that there was a rapid increase in density in the spring months and a decrease in density in the fall. The decrease in density was coincident with the greatest loss of tagged \(L.\) saccharina sporophytes (Fig. 2). \(L.\) digitata was found at 2

<table>
<thead>
<tr>
<th>Date</th>
<th>Coaster’s Harbor Island (NB)</th>
<th>Ft. Wetherill (NB)</th>
<th>Land’s End (RIS)</th>
</tr>
</thead>
<tbody>
<tr>
<td>May 1980</td>
<td>0</td>
<td>5 ± 2</td>
<td>2 ± 1</td>
</tr>
<tr>
<td>Jan 1981</td>
<td>0</td>
<td>10 ± 3</td>
<td>2 ± 1</td>
</tr>
<tr>
<td>Feb 1980</td>
<td>0</td>
<td>ND</td>
<td>ND</td>
</tr>
<tr>
<td>Mar 1980</td>
<td>0</td>
<td>5 ± 1</td>
<td>3 ± 2</td>
</tr>
<tr>
<td>Apr 1980</td>
<td>0</td>
<td>ND</td>
<td>ND</td>
</tr>
<tr>
<td>May 1980</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Jun 1981</td>
<td>0</td>
<td>2 ± 1</td>
<td>8 ± 5</td>
</tr>
<tr>
<td>Yearly mean</td>
<td>0</td>
<td>4</td>
<td>3</td>
</tr>
</tbody>
</table>

Table 1. \(Laminaria\) saccharina. Meristematic growth rate, standing crop, density, net productivity, and P/B in kelp beds in Narragansett Bay (NB) and Rhode Island Sound (RIS), 1980–1981. Median values in parentheses

<table>
<thead>
<tr>
<th>Stations</th>
<th>Mean meristematic growth rate (cm d(^{-1}) ± SE)</th>
<th>Mean and median standing crop (g DW m(^{-2}) ± SE)</th>
<th>Mean and median density (Ind. m(^{-2}) ± SE)</th>
<th>Percentage of (L.) saccharina of total density (%)</th>
<th>Annual net primary productivity (g C m(^{-2}) yr(^{-1}))</th>
<th>P/B</th>
</tr>
</thead>
<tbody>
<tr>
<td>Coaster’s Harbor Island (NB)</td>
<td>0.909 ± 0.214</td>
<td>606 ± 93 (367)</td>
<td>76 ± 11 (40)</td>
<td>100</td>
<td>1202 (633)</td>
<td>5.2</td>
</tr>
<tr>
<td>Ft. Wetherill (NB)</td>
<td>0.802 ± 0.152</td>
<td>715 ± 161 (632)</td>
<td>243 ± 95 (90)</td>
<td>77–100</td>
<td>3531 (1291)</td>
<td>4.5</td>
</tr>
<tr>
<td>Land’s End (RIS)</td>
<td>0.796 ± 0.286</td>
<td>1000 ± 217 (536)</td>
<td>95 ± 23 (50)</td>
<td>93–100</td>
<td>1099 (577)</td>
<td>3.6</td>
</tr>
</tbody>
</table>

Table 2. \(Laminaria\) digitata. Monthly density (Ind. m\(^{-2}\) ± SE) and standing crop (g DW m\(^{-2}\) ± SE) in kelp beds in Narragansett Bay (NB) and Rhode Island Sound (RIS), 1980–1981. ND = No data
of the 3 stations sampled (Ft. Whetherill and Land’s End) and it always had a lower density than L. saccharina (Table 1). Neither young nor reproductive L. digitata sporophytes were common in the quadrats. The recruitment of young L. saccharina sporophytes (< 50 cm in length) was most pronounced at Ft. Wetherill (NB) and Land’s End (RIS). It was these young sporophytes that had settled in dense patches on bare substrata (before the settlement of other macroalgae) which appeared to be the individuals that survived and later became reproductive. The large standard error of the spring density measurements gives an indication of the patchy kelp distribution (Fig. 2). The majority of individuals sampled in spring were first-year plants less than 80 cm. The greatest density measured in spring 1981 in 1 quadrat was 1990 individuals m⁻². Young sporophytes were found throughout most of the year. Sporophytes that settled during summer and fall, on the other hand, could attach only to other macroalgae already established (e.g. Chondrus crispus). The young laminarian plants were loosely attached to these algae and could easily be dislodged. Young sporophyte recruitment at Coaster’s Harbor Island (NB) was much lower than at the other stations. It is important to note that the population sampled at Coaster’s Harbor Island (NB) grew exclusively on vertical substrata (man-made wall) which may have affected the establishment of sporophytes. The greatest yearly mean and median density of kelp was measured at Ft. Wetherill (NB) (243 ± 95 and 90 individuals m⁻², respectively), where L. digitata was most abundant (Tables 1 and 2). The density at Ft. Wetherill (NB) was significantly higher than the density at Coaster’s Harbor Island (NB) (Kruskal-Wallis test, p < 0.005) and Land’s End (RIS) (p < 0.05).

Median values have been included for standing crop and density results because the results of the Kolmogorov-Smirnov test indicated that the data were not normally distributed over the entire sample period. This is a result of few occurrences of very high values which would inflate the mean values. Therefore, the median may be a more accurate measure of the kelp population. There were few instances when biomass and density data were normally distributed at a particular station and sample date.

### Epiphyte community

The epiphyte community was observed to be most abundant during August and September when kelp growth was negligible. At this time, the kelp were moribund and could be torn by wave action. Further observations indicated that epiphytes were at a minimum in spring during maximal growth of kelp. Table 3 is a species list of the plant and animal epiphytes found within the kelp beds. Some genera (e.g. *Spirorhiz, Ulva*) have been previously identified attached to or associated with *Laminaria* spp. (Tokida, 1960; Smith, 1964; Kain, 1979; Edwards, 1980). This is the first report of 7 species of animal epiphytes (e.g. *Craterolophus convolvus, Bugula turrita*). The predominant epiphytes on the blades were the invertebrates *Bugula turrita* (Bryozoa) and *Obelia* spp. (Coelenterata), while macroalgal epiphytes were more ephemeral, perhaps because of earlier colonization and dominance of animal epiphytes. Only 1 epiphytic animal, *Lacuna vincta*, grazed heavily on the blades and blade tips. Its activity was greatest during winter when *L. digitata* blades were eroded to within 20 cm of the blade-stipe juncture. Epidermal cells of the kelp were clearly damaged and sometimes absent in areas where *L. vincta* was present. *L. vincta* was found on both species of kelp but was observed to be most abundant on *L. saccharina* throughout the year. When a kelp stand was adjacent to a bed of *Mytilus edulis* at the upper Bay site, byssal threads from the mussels were found attached to the kelp, reducing the movement of the blade. During extreme wave exposure, the blades would tear if anchored by the byssal threads.

<table>
<thead>
<tr>
<th>Species Phylum/Division</th>
</tr>
</thead>
<tbody>
<tr>
<td>Animals</td>
</tr>
<tr>
<td><em>Obelia</em> sp.¹</td>
</tr>
<tr>
<td><em>Craterolophus convolvus</em></td>
</tr>
<tr>
<td><em>Mitrella lanata</em>¹</td>
</tr>
<tr>
<td><em>Lacuna vincta</em>²</td>
</tr>
<tr>
<td><em>Littorina littorea</em></td>
</tr>
<tr>
<td><em>Mytilus edulis</em>²</td>
</tr>
<tr>
<td><em>Asterias forbesii²</em></td>
</tr>
<tr>
<td><em>Spirorhiz spirillum¹</em></td>
</tr>
<tr>
<td><em>Didemnum candidum</em></td>
</tr>
<tr>
<td><em>Cryptosula palliaria²</em></td>
</tr>
<tr>
<td><em>Bugula turrita</em></td>
</tr>
<tr>
<td><em>Balanus sp.</em></td>
</tr>
<tr>
<td><em>Anoplopactylum lenus</em></td>
</tr>
<tr>
<td>Plants</td>
</tr>
<tr>
<td><em>Ulva lactuca³</em></td>
</tr>
<tr>
<td><em>Palmeria palmata¹</em></td>
</tr>
<tr>
<td>*Ceramium spp.*³</td>
</tr>
<tr>
<td>*Ectocarpus spp.*³</td>
</tr>
</tbody>
</table>

Previously identified as an epiphyte by:

1 Kain, J. M. (1979), reported on *Laminaria* spp.
2 Edwards, A. (1980), reported on *Laminaria hyperborea*
3 Tokida, J. (1960), reported on *Laminaria* spp.
4 Smith, R. (1964), reported on *Laminaria* spp.

### Table 3. Epiphytes commonly associated with Laminaria spp. in Narragansett Bay and Rhode Island Sound, 1980-1981.

Asterisks: species observed > 9 mo of sample period.
A linear regression of blade length and dry weight is the best fit for adult *Laminaria saccharina* (Fig. 4). This regression formula is:

\[
\text{Dry weight (g DW)} = (0.15615) \text{ blade length (cm)} + 1.6
\]

\[r^2 = 0.80\]

![Graph showing the relationship between blade length and dry weight for adult *Laminaria saccharina*.](image)

with a correlation coefficient of 0.89 (range = 0.84 to 0.94 \(\alpha = 0.05\)). *L. saccharina* had a higher percent of tissue carbon (31.5 ± 3.4 %) than *L. digitata* (26.9 ± 6.0 %), but this difference was statistically not significant.

**Net annual primary productivity**

The kelp stand at Ft. Wetherill (NB) had a higher primary productivity on an area basis than Coaster's Harbor Island (NB) and Land's End (RIS) (Table 1). The mean and median primary productivity for the 2 stations within Narragansett Bay (2366 and 962 g C m\(^{-2}\) yr\(^{-1}\), respectively) were calculated by pooling data from the Narragansett Bay sample stations. The productivity when using median density was approximately half the productivity using the mean density. It can be argued that these productivity values are an overestimate because no distinction was made between juvenile and adults in the density measurements. The maximum productivity of *Laminaria saccharina* individuals (50 to 60 % of the total) occurred in late winter and spring (Table 4), and the lowest was in August and September (0.4 to 2.2 % of the total). The mean productivity per individual at the 3 stations for the year was 15.43 ± 0.99 g C individual\(^{-1}\) yr\(^{-1}\).

**DISCUSSION**

The vegetative growth pattern of *Laminaria saccharina* at the 2 sample stations in Narragansett Bay was similar to that in Rhode Island Sound. The pattern agrees with results reported for *L. longicruris* in St. Margaret's Bay, Nova Scotia (Mann, 1972b; Chapman and Craigie, 1977; Gagne et al., 1982) (*L. saccharina* is thought to be conspecific with *L. longicruris* [Chapman, 1974]). This pattern has been regarded as a typical growth cycle of *Laminaria* spp., with rapid growth from late winter to late spring and minimal growth in summer, independent of environmental conditions (Kain, 1979). In some *Laminaria* populations, growth

<table>
<thead>
<tr>
<th>Date</th>
<th>Coaster's Harbor Island (NB)</th>
<th>Ft. Wetherill (NB)</th>
<th>Land's End (RIS)</th>
</tr>
</thead>
<tbody>
<tr>
<td>May 1980</td>
<td>2.03</td>
<td>2.20</td>
<td>2.79</td>
</tr>
<tr>
<td>Jun</td>
<td>1.76</td>
<td>1.53</td>
<td>1.28</td>
</tr>
<tr>
<td>Jul</td>
<td>0.42</td>
<td>1.45</td>
<td>0.68</td>
</tr>
<tr>
<td>Aug</td>
<td>0.07</td>
<td>0.32</td>
<td>0.14</td>
</tr>
<tr>
<td>Sep</td>
<td>0.19</td>
<td>0.21</td>
<td>0.12</td>
</tr>
<tr>
<td>Oct–Nov</td>
<td>1.88</td>
<td>–</td>
<td>0.22</td>
</tr>
<tr>
<td>Dec–Jan 1981</td>
<td>2.82</td>
<td>–</td>
<td>4.14</td>
</tr>
<tr>
<td>Feb–Mar</td>
<td>3.53</td>
<td>2.54</td>
<td>–</td>
</tr>
<tr>
<td>Apr</td>
<td>3.12</td>
<td>1.81</td>
<td>–</td>
</tr>
<tr>
<td>Annual total</td>
<td>15.82</td>
<td>14.31</td>
<td>16.16</td>
</tr>
</tbody>
</table>

*Projected values based on Coaster's Harbor Island data.*
differs from the typical pattern (Reynolds, 1974; Gerard and Mann, 1979; Gagne et al., 1982). This difference has been attributed to the concentration and duration of ambient nitrogen supplies; when the nitrogen supply is abundant, growth is correlated with the light cycle (Chapman and Craigie, 1977; Gagne et al., 1928). The growth pattern of Rhode Island plants resembled *L. longicruris* individuals in St. Margaret's Bay which were adapted to short periods of ambient nitrogen availability (Gagne et al., 1982). The ambient nitrogen and inorganic tissue nitrate cycles of plants in Narragansett Bay (Fig. 3; Asare and Harlin, 1983) are similar to those of St. Margaret's Bay (Gagne et al., 1982). This similarity adds support to the hypothesis that *Laminaria* growth is adapted to a combination of local environmental factors.

The mean annual biomass of *Laminaria saccharina* and *L. digitata* at the 2 sample stations in Narragansett Bay (0.779 kg DW m⁻² [median = 0.513 kg DW m⁻²]) and the 1 in Rhode Island Sound (1.092 kg DW m⁻² [median = 0.536 kg DW m⁻²]) are lower than those measured for comparable *Laminaria* spp. in St. Margaret's Bay, Nova Scotia (summer) (Mann, 1972a) and Helgoland, F. R. Germany (late summer/early fall) (Lining, 1969) (Table 5). The 2 previous studies had a harvesting time frame of 3 to 4 mo, when standing crop might have been at its maximum, as demonstrated by the results of this study (Fig. 2).

There was a 30% difference between the blade-length turnover rate calculated for *Laminaria longicruris* in Nova Scotia (3.5 times yr⁻¹) (Mann, 1972b) and the rate calculated for *L. saccharina* in Narragansett Bay (3.3 times yr⁻¹). However, Mann (1972b) has found increases in biomass to vary according to size classes of plants, and calculated a biomass turnover rate ranging from 4 to 10 times yr⁻¹. The biomass turnover rate was not calculated for the present study. However, if Mann's (1972b) biomass turnover rate (rather than blade-length turnover rate) was multiplied by the Narragansett Bay standing crop, the mean annual kelp biomass for Narragansett Bay sample stations (2310 g DW m⁻² yr⁻¹) would double.

For the entire Bay, the relative productivity of phytoplankton is 2 orders of magnitude higher than the kelp production, when the area of occurrence of kelp (0.32 km²) is considered. Within the kelp beds, the productivity of kelp (based on either mean or median density) is 2 to 10 times greater than the productivity of phytoplankton (308 g C m⁻² yr⁻¹) (Furnas et al., 1976). Therefore, kelp within the narrow range of its distribution provides greater primary productivity in the lower Bay than previously realized.

There was a narrow temporal overlap in the distribution of primary production of phytoplankton and kelp in Narragansett Bay. Approximately 60% of the kelp production occurred from January to June (Table 4) when 17% of the phytoplankton production occurred (Furnas et al., 1976). Sixty% of the total kelp production equals 4.5 x 10⁸ g C yr⁻¹ while 17% of the total phytoplankton production is 4.6 x 10⁸ g C yr⁻¹ in the lower Bay. The relative contribution of kelp to total primary production varies with the season: in winter, the carbon production of the kelp is ~ 10% that of the phytoplankton; in summer, this value is much less.

Narragansett Bay kelp productivity exceeds the kelp productivity in Nova Scotia calculated by Hatcher et al. (1977), due to a difference in kelp density at the 2 study areas (Table 5). The primary productivity of individuals (g C individual⁻¹ yr⁻¹) in Nova Scotia was double (28 g C individual⁻¹ yr⁻¹) (Hatcher et al., 1977) that of individuals in Narragansett Bay (15.43 ± 0.99 g C individual⁻¹ yr⁻¹) (Table 4), suggesting greater productivity on an individual basis (rather than an areal basis) in northern areas. Table 5 compares productivity of other kelp communities. The Narragansett Bay kelp productivity was comparable to that in Spain (Lapointe et al., 1981), and it was greater than the productivity of *Laminaria saccharina* in Scotland (Johnston et al., 1977) and of *L. longicruris* in Nova Scotia (Hatcher et al., 1977). This was noteworthy considering that Narragansett Bay is near the southern limit for the species (Taylor, 1972).

Production to biomass (P/B) ratios provide information useful in describing the relative stage of a biological community (Odum, 1968). The P/B ratios in this study ranged from 5.2 at Coaster's Harbor Island to 3.6 at Land's End. The P/B decreased from sheltered to exposed areas (Table 1), consistent with the results from St. Margaret's Bay (Mann, 1972b), even though P/B ratios from Nova Scotia were double those of Narragansett Bay. A similarity of temporal P/B ratios was found between those calculated by Lapointe et al. (1981) (where P/B decreased from 0.03 in January to 0.01 in July), and those of this study (where P/B decreased from 0.02 to 0.005 for the same months). This decrease in P/B is thought to be the result of succession from an immature to a mature community (Odum, 1968). The temporal pattern of P/B may also reflect the seasonal changes of the subtidal community where frequent abiotic disturbances cause a decline of the relative production of the community.

*Laminaria* spp. density in Rhode Island was variable and the annual mean and median densities in the subtidal were greater than those in Europe (Kain, 1979). The high density and variability measured in spring was the result of the patchy distribution of first year plants, as had been noted previously in European *Laminaria* populations (Kain, 1979). *L. japonica* var. *ochotensis* (in Japan) is the only other species that has a density comparable to that of *L. saccharina* (in Rhode
Table 5. Representative standing crop and net primary production of *Laminaria* spp. communities

<table>
<thead>
<tr>
<th>Location</th>
<th>Reference</th>
<th>Standing crop</th>
<th>Net production</th>
<th>Period sampled</th>
<th>Species</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>WW kg m⁻²</td>
<td>DW kg m⁻²</td>
<td>(g C m⁻² yr⁻¹)</td>
<td></td>
</tr>
<tr>
<td>Helgoland, F.R. Germany</td>
<td>Lüning (1969)</td>
<td>12.71</td>
<td>2.67*</td>
<td>–</td>
<td>Aug–Nov</td>
</tr>
<tr>
<td>Nova Scotia, Canada</td>
<td>Mann (1972a)</td>
<td>16.00</td>
<td>3.36*</td>
<td>–</td>
<td>Jun–Aug</td>
</tr>
<tr>
<td>Nova Scotia, Canada</td>
<td>Hatcher et al. (1977)</td>
<td>–</td>
<td>–</td>
<td>143–428</td>
<td>May–May</td>
</tr>
<tr>
<td>Loch Creran, Scotland</td>
<td>Johnston et al. (1977)</td>
<td>–</td>
<td>–</td>
<td>120</td>
<td>Jan–Dec</td>
</tr>
<tr>
<td>Lagoon Pt., Alaska USA</td>
<td>Calvin and Ellis (1978)</td>
<td>14.50</td>
<td>3.05*</td>
<td>–</td>
<td>May</td>
</tr>
<tr>
<td>Narragansett Bay and R.I. Sound USA</td>
<td>This study</td>
<td>–</td>
<td>0.2–6.0</td>
<td>2366 (mean) 962 (median)</td>
<td>Apr 80–Jun 81</td>
</tr>
</tbody>
</table>

* Estimated values from a 21% conversion of wet weight to dry weight
** Estimated from daily measurements

Island) (Kaneko and Nihara, 1977). *L. saccharina* remains the dominant species over *L. digitata* at any depth within the subtidal zone in Rhode Island.

Herbivore grazing of *Laminaria* spp. in Rhode Island was primarily due to *Lacuna vincta*, a prosobranch gastropod. *L. vincta* caused extensive damage on *Laminaria* populations in New Hampshire (Fralick et al., 1974). The green sea urchin, *Strongylocentrotus droebachiensis*, grazes on kelp in Nova Scotia (Mann, 1973). However, *S. droebachiensis* is limited to deeper waters in Rhode Island and is not found within the kelp beds; therefore, sea urchin grazing was not evident (own obs.).

This study supports earlier estimates that Narragansett Bay is a phytoplankton-based system (Kremer and Nixon, 1978). However, the data show that the macroalgae are more productive than previously credited, especially the Laminariales (which represent only a portion of the total macroalgae). Our values for the primary productivity of kelp may be an underestimate because factors such as spore release, biomass turnover (rather than blade-length turnover), and exudation, have not been included. This study and others (Mann, 1973; Johnston et al., 1977; Lapointe et al., 1981) have shown that kelp provide substrata, habitat and food for other subtidal inhabitants. Thus, the laminarians may contribute even more to the Narragansett Bay ecosystem than the scope of this study has examined.

In conclusion, *Laminaria saccharina* and *L. digitata* are as productive near their southern range limit as observed in studies well within the distributional range. Net primary productivity, growth rates, standing crop and tissue carbon were within the range and/or similar to results of other kelp studies (Kain, 1979). However, there were dissimilar results in the following: (a) population density was extremely variable but on the average the density was higher than in Nova Scotia for *L. longicruris* (Hatcher et al., 1977) and in Great Britain for *L. hyperborea* (Kain, 1979); (b) the rate of blade-length turnover was less than that measured in Nova Scotia (Mann, 1972b); (c) productivity of individual plants was ~50% less than for *L. longicruris* in Nova Scotia (Hatcher et al., 1977); and (d) P/B ratios for the Rhode Island population of *L. saccharina* were approximately half those of the *L. longicruris* population in Nova Scotia.

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