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Lindsay A. Green  
*University of New Hampshire, Durham*

Arthur C. Mathieson  
*University of New Hampshire, Durham, Arthur.Mathieson@unh.edu*

Christopher D. Neefus  
*University of Rhode Island, chris.neefus@unh.edu*

Hannah M. Traggis  
*University of New Hampshire, Durham*

Clinton J. Dawes  
*University of New Hampshire, Durham*

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Short communication

Lindsay A. Green*, Arthur C. Mathieson, Christopher D. Neefus, Hannah M. Traggis and Clinton J. Dawes

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Abstract: Blackler first recorded Colpomenia peregrina in the Northwest Atlantic based on collections from Nova Scotia, Canada. Five decades later we found large quantities of C. peregrina in Maine, USA, even though it was absent during earlier floristic studies in this region. Thus, C. peregrina has undergone a rapid southern expansion along the Northwest Atlantic coast. While the causes of such an expansion are unknown, it could have a major effect on both shellfish cultivation and native seaweeds within New England because of competitive interactions and increased drag.

Keywords: brown alga; Colpomenia peregrina; introduction; Northwest Atlantic Ocean.

*Corresponding author: Lindsay A. Green, Department of Biological Sciences, University of New Hampshire, Durham, NH 03824, USA, e-mail: lindsay.green@unh.edu

Arthur C. Mathieson: Department of Biological Sciences, University of New Hampshire, Durham, NH 03824, USA; and Jackson Estuarine Laboratory, University of New Hampshire, Durham, NH 03824, USA

Christopher D. Neefus: Department of Biological Sciences, University of New Hampshire, Durham, NH 03824, USA

Hannah M. Traggis: Department of Biological Sciences, University of New Hampshire, Durham, NH 03824, USA

Clinton J. Dawes: Jackson Estuarine Laboratory, University of New Hampshire, Durham, NH 03824, USA; and Department of Biology, University of South Florida, Tampa, FL 33620, USA

The genus Colpomenia (Endlicher) Derbès et Solier is a globular, saccate brown alga found within temperate and tropical waters worldwide (Boo et al. 2011). Colpomenia peregrina Sauvageau is an epiphytic member of the Scytosiphonales that grows on a variety of hosts (Blackler 1967). It has a single plastid with one pyrenoid per cell, a heteromorphic life history alternating between an erect parenchymatous saccate gametophyte and a pseudoparenchymatous sporophytic crust, with only the gametophyte producing plurilocular gametangia (Clayton 1979, Cho et al. 2001). The gross morphological similarities between C. peregrina and Leathesia marina (Lyngbye) Decaisne may have previously caused confusion as to the presence of the former species (Blackler 1967).

Although C. peregrina is native to the Northwest Pacific, its type location is Brittany, France (Sauvageau 1927), where it was initially characterized during the early 20th century (Blackler 1967, Vandermeulen and DeWreede 1986). In France, C. peregrina earned the nickname “oyster thief” and was distinguished from C. sinuosa (Mertens ex Roth) Derbès et Solier by its smoother, thinner thalli and described as a new species. Blackler (1967) reported that it spread rapidly on the east coasts of England and Scotland, as well as Holland (1921), Denmark (1939), the Mediterranean (1939), and Norway (1949); its rapid establishment and extensive spreading in Europe have been well documented.

The initial presence of C. peregrina in the Northwest Atlantic was documented by Blackler (1964) based on a 1960 collection by A.R.A. Taylor from Atkins Point, Nova Scotia, Canada (44°54′26″ N, 62°18′32″ W; Figure 1). Subsequently, it was recorded from two southwestern Nova Scotia sites during the 1960s: Goodwin Island (43°31′44″ N, 65°47′05″ W, collected by C. MacFarlane and G.M. Miligan; Figure 1) and Polly Cove (44°29′19″ N, 63°53′33″ W, collected by D. Pace; Figure 1) followed by a fourth Nova Scotia site at Bon Portage Island (43°28′00″ N, 65°41′60″, collected by E. Fraser, D. Smith, and K. Lynch; Figure 1). Bird and Edelstein (1978) confirmed these early records and re-designated some specimens that were improperly identified as L. marina. In the late 1970s, C. peregrina was documented from another Nova Scotia site at Hospital Reef on Cape Sable Island (43°27′09″ N, 65°39′18″ W,
collected by J.S. Wilson; Figure 1) followed by a collection in the early 1980s from Grand Barachois Lagoon (46°48'20" N, 56°14'26" W, pers. comm., R.G. Hooper) located between Saint Pierre and Miquelon Islands, Newfoundland.

In the summer of 2011, we initially collected from extensive populations of *C. peregrina* at Fort Foster, Kittery, Maine (43°03’55” N, 70°40’54” W, Figure 1). Subsequently, re-evaluation of earlier photographic records from mid-coastal Maine confirmed that it had been misidentified as *L. marina* during the summer of 2010 on Great Chebeague Island in Casco Bay, Maine (43°43’57” N, 70°07’26” W, collected by A. and D. Swafford and J. Coyer; Figure 1) and in West Boothbay Harbor (43°50’53” N, 69°38’36” W, collected by P. Thayer; Figure 1). Earlier collections in the same geographic areas (Mathieson et al. 2007, 2008a,b, 2010) suggest that it was not present prior to 2007 and re-evaluations of herbarium vouchers of *L. marina* (n=121) collected from Newfoundland to Delaware during 1938–2005 confirmed the absence of *C. peregrina*. Given its rapid establishment and expansion in other geographic areas (Blackler 1964), we set out to determine the extent to which it had become established in the Gulf of Maine (i.e., Bay of Fundy to Cape Cod, Massachusetts) and to genetically verify that it was the same species previously documented in Nova Scotia (Bird and Edelstein 1978).

We collected from low intertidal and shallow subtidal zones from mid-coastal Maine to southern Massachusetts, including two offshore islands within the Isles of Shoals in New Hampshire/Maine (Figure 1). Herbarium sheet vouchers were prepared and deposited in the Albion R. Hodgdon Herbarium (NHA) at the University of New Hampshire. Biomass estimates of *C. peregrina* were made during the summer of 2011 by SCUBA diving along metered transects at three different sites: Fort Foster Kittery, Maine, Appledore Island, Maine (42°59’04” N, 70°37’06” W) and Star Island, New Hampshire (42°58’31” N, 70°36’58” W). Eight randomly selected destructive quadrats (0.25 × 0.25 m) were taken along a 100 m transect run perpendicular from the shore down through the intertidal and extending into the shallow subtidal. All biomass within each quadrat was scraped from the underlying substrata. Algal specimens were identified to species (Mathieson et al. 2008b for taxonomic references) and damp-dried weight and percent composition quantified. Measurements of thallus diameter and height were recorded for all *C. peregrina* samples (n=528), as well as epiphytic host specificity.

Representative specimens from Maine (Fort Foster and Appledore Island), New Hampshire (Fort Stark, Newcastle 43°03’22” N, 70°42’41” W and Star Island), and Massachusetts (Niles Beach, Gloucester 42°35’52” N, 70°39’23” W and Sandwich, Cape Cod 41°46’28” N, 70°29’30” W, collected by J.S. Wilson; Figure 1) followed by a collection in the early 1980s from Grand Barachois Lagoon (46°48’20” N, 56°14’26” W, pers. comm., R.G. Hooper) located between Saint Pierre and Miquelon Islands, Newfoundland.

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Material for molecular voucher specimens collected in the 1960s and 1970s from Nova Scotia, Canada (Polly Cove, Bon Portage Island, and Hospital Reef) were also analyzed. Total DNA was extracted from 2×2 mm samples using a NucleoSpin Plant II kit (Machery-Nagel, Düren, Germany), following the manufacturer’s instructions. The extracted DNA was stored at 4°C for use the same day or -20°C for later use in amplifying the cox3 gene.

Specific primer pairs for amplification and gene sequencing were newly designed as follows. F1: ATAACATGRTGRTGGWGAAT 5′ to 3′ and R1: GARCCATACATGACATA 5′ to 3′. An Eppendorf Mastercycler (Model 533, Hamburg, Germany) was used for all PCR amplifications. A TaKaRa ExTaq reaction kit (Takara Shuzo, Shiga, Japan) was used with a total reaction volume of 50 μl consisting of 5.0 μl 10× ExTaq buffer, 5.0 μl of 25 mM MgCl₂, 50 μM dNTP mixture, 0.4 μM of each primer, 1.25 U TaKaRa ExTaq, and 4.0 μl of DNA solution (containing between 0.5 and 1.0 μg DNA). PCR was performed with an initial denaturation step at 94°C for 4 mins, followed by 35 cycles of 30 s at 94°C, 30 s at 56°C, and 1 min at 72°C, and a final extension cycle at 72°C for 10 mins. The PCR products were purified using gel electrophoresis and sequenced at the HCGS DNA Sequencing Facility, University of New Hampshire.

Twenty-one cox3 sequences were used for the phylogenetic analysis: 20 previously published (Silberfeld et al. 2010, Boo et al. 2011) and one representative from the Northwest Atlantic (present study). Maximum-likelihood phylogenetic analysis was performed using MEGA 4.0 software (Tamura et al. 2007) with the Hasegawa-Kishino-Yano (HKY)+Γ model. Bootstrap values were calculated using 1000 replicates.

Representative samples from Maine, New Hampshire and Massachusetts, USA plus early voucher specimens from Nova Scotia, Canada all yielded identical cox3 sequences that were not an exact match to any previously
published sequence. Based on the phylogenetic analysis, our Northwest Atlantic samples fell within *C. peregrina*, which showed less variability than *C. claytoniae* (Figure 2). All novel sequences from this study have been submitted to GenBank (Accession numbers JX101661-JX101675).

Biomass surveys showed an abundance of *C. peregrina* within the low intertidal and shallow subtidal zones at the three study sites. Due to the nature of random sampling some quadrats lacked *C. peregrina*, resulting in a wide variation in biomass. The highest average biomass was recorded at Appledore Island (50.7±31.7 g m⁻², mean±SE) followed by Star Island (28.3±10.0 g m⁻²) and Fort Foster (6.9±4.3 g m⁻²). *Colpomenia peregrina* had a wide range of statures, with diameters ranging from 0.1 to 5.8 cm at Appledore Island, 0.1–9.0 cm at Star Island, and 0.1–7.0 cm at Fort Foster. Similarly, heights ranged from 0.1 to 4.8, 0.1 to 3.7, and 0.1 to 4.0 cm at Appledore Island, Star Island, and Fort Foster, respectively.


During the summer and fall of 2011, extensive populations of *C. peregrina* were recorded at 11 sites in mid-coastal to southern Maine, while it was also collected as far south as Cape Cod, Massachusetts during the spring of 2012 (Figure 1). Genetic analysis confirmed the identification of *C. peregrina* using representative samples from Maine to Massachusetts, with each being identical to original voucher specimens from Nova Scotia (Figure 2). Biomass surveys indicate the species is highly abundant (6.9–50.7 g m⁻²), morphologically variable, and grows on a wide range of hosts.

In the Northwest Pacific, fertile plants of *C. peregrina* can be seen year round. High temperatures and low salinities during the summer as well as low temperatures during winter can limit its growth (Vandermeulen and DeWreede 1986). We initially assumed that *C. peregrina* was a spring-summer annual; however, extensive monthly collections at Fort Foster have shown that it is present year round, particularly within the shallow subtidal zone. Our biomass estimates are slightly higher than previously recorded by Vandermeulen and DeWreede (1986) from the Northeast Pacific. They suggested that the saccate form of *C. peregrina* was opportunistic, even though both its photosynthetic rates and grazing by herbivores were low. Our observations are consistent with the occurrence of low herbivory rates, as no holes or cuts were observed.

Introduced species can cause major problems by altering natural communities and causing economic losses (Mathieson et al. 2007, 2008a,b). Currently there are approximately 120 introduced seaweed taxa known worldwide (Nyberg and Wallentinus 2005), and about 25 within the Northwest Atlantic (Mathieson et al. 2007, 2008a,b, Hofmann et al. 2010, Schneider 2010). The introduction of *C. peregrina* to France caused significant economic losses to the oyster industry (Blackler 1967). Although no deleterious impacts of *C. peregrina* on the New England shellfish industry have been reported, it has the potential to become a major pest considering its rapid expansion during the past 2 years. In New England, the eastern oyster industry was valued at US $117.6 million in 2010 (pers. comm., National Marine Fisheries Service). Since *C. peregrina* is an epiphyte on several native seaweeds within the low intertidal and shallow subtidal zones, it may also impact hosts by increasing drag and competing for space and limited resources.

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