

Medusae, siphonophores, and ctenophores as planktivorous predators in changing global ecosystems

Claudia E. Mills

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Medusae, siphonophores, and ctenophores are planktivorous predators operating at higher trophic levels in marine ecosystems of a wide range of productivity. It has been hypothesized that high-productivity ecosystems such as areas of upwelling tend towards food chains dominated by larger phytoplankton, large copepods, and ultimately many species of fish rather than gelatinous predators; ecosystems with lower productivity are characterized by small flagellate phytoplankton, small copepods, and ultimately numerous medusae and ctenophores. Evidence is provided that medusae, siphonophores, and ctenophores are actually important predators in both sorts of planktonic ecosystems, although uneven reporting in the literature may be cause for underestimates of the importance of these carnivores in some systems. As world fisheries begin to experience serious declines, it is relevant to recognize that the carnivorous “jellyfishes” are ubiquitous and are thus opportunistically positioned to utilize secondary production that is ordinarily consumed by fish.

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C. E. Mills: Friday Harbor Laboratories, University of Washington, 620 University Road, Friday Harbor, WA 98250, USA.

Introduction

Parsons and Greve (1977, 1979) suggested that high productivity or upwelling ecosystems tend towards food chains dominated by larger phytoplankton, accompanied by the production of large copepods, and many species of fish, whereas low productivity or convergent ecosystems, on the other hand, are characterized by small flagellate phytoplankton, with the production of small copepods, and eventually large numbers of medusae and ctenophores. Only the endpoints of this hypothesis, the progression through food chains dominated ultimately either by fishes or jellyfishes, will be discussed here. I hope to convince the reader that in fact, medusae, siphonophores, and ctenophores (referred to inclusively as jellyfishes) are present nearly to the same extent in upwelling systems as in regions of low productivity. By accidents of weather, ship locations, and home residence locations of scientists who study these animals, the numerical abundance and importance of the planktonic carnivorous jellyfishes has been overlooked on a global scale.

Observations from blue-water diving and manned submersibles

When the dual food chain hypothesis was originally proposed (Greve and Parsons, 1977), it was partially supported by the work of Greve (1971, 1972), who showed through field and laboratory observations that the tentaculate ctenophore *Pleurobrachia pileus* grew best in rather low concentrations of small species of copepods. Large copepods were shown to be detrimental in different ways to both adult and larval *P. pileus*, as large copepods can escape being eaten, often tearing the tentacles of larger ctenophores in the process, and they sometimes even prey on larval ctenophores. Parsons (1979), in further support of jellyfish being an endpoint in low productivity waters, cited the important role of gelatinous zooplankton in areas such as the Sargasso Sea as portrayed by direct observation in the new technology of blue-water diving.

A series of papers in the late 1970s reporting observations of the fragile gelatinous zooplankton by SCUBA divers (Hammer, 1975; Hammer *et al.*, 1975; Harbison *et al.*, 1977, 1978; and others) reminded many

Table 1. A comparison of the diversity of medusae, siphonophores, and ctenophores collected on submersible cruises using the "Johnson-Sea-Link I" and "II" in two locations. Studies were of approximately equal duration in water columns of nearly the same depth.

	High productivity	Low productivity
	North Atlantic off New England	Alboran Sea Mediterranean
	23 dives 800 m bottom	26 dives 700–800 m bottom
Hydromedusae	10	9
Siphonophores	13	13
Scyphomedusae	6	3
Ctenophores	9	11

oceanographers about the existence and putative importance of the gelatinous macrozooplankton. Most of the blue-water SCUBA diving in the 1970s and early 1980s occurred in the Caribbean and Sargasso Seas, and thus Parsons and perhaps others concluded that gelatinous zooplankton was most plentiful and most important in oligotrophic waters. In fact, the locations of these dives had more to do with the logistics of blue-water diving than with the expected presence or absence of gelatinous zooplankton. Blue-water diving at nearly any location in the world would yield the same sorts of fragile epipelagic planktonic jellyfish in substantial numbers (G. R. Harbison and L. P. Madin, pers. comm.). For example, during the past 15 years there has been an ongoing program of blue-water diving off Santa Barbara, California, by faculty and students at the University of California. These dives, often in upwelling conditions (in direct contrast to the oligotrophic and stable water masses of the Sargasso Sea) have revealed similar taxa, often with even the same cosmopolitan epipelagic species (S. H. D. Haddock, pers. comm.), as were described in the early blue-water diving papers from the Sargasso Sea and Caribbean.

Comparable submersible studies of the water column also reveal similar numbers of species of jellyfish in high and low productivity regions of the ocean. Table 1 summarizes findings from three submersible cruises, two in the NW Atlantic off Massachusetts (August–September 1986 and August 1987) and one in the Alboran Sea (Mediterranean) off the coast of Morocco (April 1991), in which the same taxonomic specialists working on medusae (R. J. Larson and C. E. Mills), siphonophores (P. R. Pugh), and ctenophores (G. R. Harbison) participated. In each location, specimens were collected on approximately the same number of dives in a water column of nearly the same depth. Species diversity of the three groups of gelatinous carnivores was remarkably similar between the two

locations. The total numbers of jellyfishes were much greater off New England than in the Alboran Sea, reflecting the greater numbers of all categories of zooplankton at that location. The key to discovering similarities in jellyfish diversity between these two sites was the inclusion of specialists interested in all three groups in both scientific parties. Although only a small number of jellyfish species are recognized by most oceanographers, a similarly diverse group of medusae, siphonophores, and ctenophores is present in most parts of the open ocean, but it is rare for oceanographic expeditions to have anyone on board interested in working with these animals.

Jellyfishes and radical ecological change

The most graphic example to date of a highly productive ecosystem, that has converted from supporting a number of valuable commercial fisheries to having few fishes and high numbers of jellyfishes, is the Black Sea and its adjacent Sea of Azov. The Black Sea has a history over the past several decades of increased pollution and eutrophication, coupled with a number of unfortunate introductions of alien marine invertebrates that have done much to change the ecology of both benthos and the water column (Zaitsev, 1992). There were 26 commercial fishes in the Black Sea in the 1960s; disappearance of most of the large predatory fishes led to an increase in small predatory fishes over the next decades, so that by the 1980s only five commercial fishes were left (Zaitsev, 1992), and nearly all of the zooplankton production was going into these planktivorous fishes (Studenikina *et al.*, 1991).

The ctenophore *Mnemiopsis leidyi*, accidentally introduced from the eastern seaboard of America to the Black Sea in 1982, and entering the Sea of Azov a few years later, has now succeeded in nearly completely replacing the planktivorous fishes in the Sea of Azov as the terminal zooplankton consumer, and is also very important in the ecosystem of the entire Black Sea basin (Shushkina and Musayeva, 1990; Studenikina *et al.*, 1991; Harbison and Volovik, 1995). Although a number of investigations in the USA have been devoted to the study of *Mnemiopsis leidyi* in its native environment, where it may be very abundant in bays in the summertime, it is unlikely that oceanographers really understood the importance of this lobate ctenophore, nor could anyone have predicted the general collapse of the fishing industries in the Sea of Azov, and to a lesser extent, the Black Sea, brought about by its arrival there.

Another potentially alarming situation is portended by large numbers of the siphonophore *Nanomia cara* observed in the Gulf of Maine in August 1992 (C. H. Greene, pers. comm.) and September 1993 (by the author participating on a cruise with C. H. Greene). While using the manned submersible "Johnson-Sea-

Link I" to search for copepods and euphausiids in 300 m deep Wilkinson Basin, we found instead large numbers of *N. cara*. These siphonophores, which varied in length from about 20 to 50 cm, were located only in the lower half of the water column by day, but migrated up and were throughout the entire water column at night. My estimates of distances between *N. cara* colonies, on the 18 September 1993 dusk submersible dive, when the animals were still in the process of vertically migrating, indicated fairly uniform densities of about 0.2 m^{-3} in the upper 50 m, densities varying patchily from 1 to 10 m^{-3} between 50 and 250 m, and densities up to $50\text{--}100 \text{ m}^{-3}$ individuals from 250 m to the bottom at 270 m. In addition to *N. cara*, three species of ctenophores (*Pleurobrachia pileus*, *Euplokamis dunlapae*, and *Bolinopsis infundibulum*) and a hydromedusa (*Aglantha digitale*) were also very numerous, and taken as a group, also spanned the entire water column. A similar aggregation of *N. cara* was reported by Rogers *et al.* (1978) in the same part of the Gulf of Maine, persisting at least from autumn 1975 through June 1976 and with estimated densities up to 7 m^{-3} as observed by manned submersible.

We do not know if the 1975–1976 and 1992–1993 *Nanomia* siphonophore blooms in the Gulf of Maine are regular cyclical events or if they are highly unusual, and we do not know the full duration of either episode. Other submersible observations in canyons south of Woods Hole, Massachusetts, have revealed *Nanomia cara* to be a normal and common member of the nearby NW Atlantic deep water community (Larson *et al.*, 1988). The extraordinary numbers of *N. cara* seen on submersible dives in 1992 and 1993 indicate that gelatinous predators must be a driving force in the water column ecology during such blooms.

With the general collapse of NW Atlantic fisheries from over-harvesting, it will be difficult to determine the effect of the recent *Nanomia cara* bloom on the pelagic Gulf of Maine ecosystem. A few interesting correlations can be made for the 1975–1976 *Nanomia* bloom, by looking back at records of fish recruitment in the area during those years. Myers *et al.* (1990) report that the 1975 and 1976 year classes of haddock and herring had average to good recruitment, but that yellowtail flounder, silver hake, redfish, and cod had unusually poor recruitment in those years. At present, we cannot determine whether the siphonophore was responsible for these poor fish recruitments. The diet of *N. cara* in the Gulf of Maine has yet to be investigated, but a similar species, *N. bijuga*, has been reported to feed primarily on copepods, chaetognaths, and shrimp in the Gulf of California (Purcell, 1981a) and primarily on *Euphausia pacifica* in Monterey Bay, California (B. H. Robison, pers. comm.) (Table 3).

Greve (1994) reported an unprecedented invasion of the siphonophore *Muggiaea atlantica* into the German

Table 2. Natural diets of some co-occurring hydromedusae in late spring 1987, Friday Harbor, Washington, USA, as determined from gut contents of field-collected animals. All species of medusae were collected from the same water masses. Since the same pool of potential prey was available to all species of hydromedusae, differences in diets reflect selectivity on the part of the medusae. Mechanisms for prey selection are discussed by Mills (1981b) and Purcell and Mills (1988). The number of medusae of each species examined is designated by "n". Prey consumed is presented as percentage of the total number of prey items in the guts. From unpublished data of the author.

Predator species	Prey consumed	
	%	Principal types
<i>Aequorea victoria</i> (n = 100)	90	Hydromedusae and ctenophores
<i>Catablema nodulosa</i> (n = 90)	78	Hydromedusae
	11	Invertebrate eggs
<i>Mitrocomella polydiademata</i> (n = 125)	45	Larvaceans and houses
	40	Invertebrate eggs
<i>Phialidium gregarium</i> (n = 125)	73	Invertebrate eggs
	11	Larvaceans and houses
<i>Bougainvillia principis</i> (n = 125)	85	Barnacle nauplii
<i>Aglantha digitale</i> (n = 36)	71	Copepods and copepod nauplii
	27	Invertebrate eggs, especially copepods

Bight in 1989. The population of this small, copepod-eating siphonophore reached 500 m^{-3} in summer 1989, temporarily reducing resident copepods to near zero. Although such a bloom has not yet been repeated, Greve warns of the potential for general disruption of this pelagic ecosystem if the normal dominant gelatinous predator *Pleurobrachia pileus* is replaced by *Muggiaea atlantica*, whose ecology is quite different, not being susceptible to the ctenophore-specific secondary carnivore *Beroe gracilis*.

Parsons (1979) questioned whether the removal of top predators by the world's fisheries might favour the establishment of increased numbers of jellyfish. *Nanomia* blooms in the Gulf of Maine might be an interesting example of such a phenomenon. In the Black Sea, predatory fishes were removed from the system, but the remaining small planktivorous fishes seem genuinely to have been outcompeted by newly arrived *Mnemiopsis* ctenophores. The result, however, in these two situations appears to be the same – domination of a previously highly productive ecosystem by jellyfishes following human-mediated events such as over-fishing, introductions of alien species to a new environment, and general

Table 3. Natural diets of some siphonophores as determined from gut contents of field-collected animals. The number of colonies of each species examined is designated by "n". Prey consumed is presented as percentage of the total number of prey items in the guts. Most siphonophores appear to be highly selective carnivores (discussed further by Purcell (1981a, 1984)).

Predator species	Location (season)	Prey consumed		Reference
		%	Principal types	
<i>Nanomia bijuga</i> (n = 53)	Gulf of California	25	Copepods	Purcell (1981a)
		20	Chaetognaths	
		16	Shrimp	
<i>Nanomia bijuga</i> (n = > 100)	Central California	Mostly Euphausiids		B. H. Robison (pers. comm.)
<i>Rosacea cymbiformis</i> (n = 47)	Gulf of California	58	Copepods	Purcell (1980, 1981c)
		25	Crab zoea larvae	
<i>Rhizophysa eysenhardti</i> (n = 646)	Gulf of California (summer)	100	Fish larvae	Purcell (1981b)
<i>Bassia bassensis</i> (n = 60)	Gulf of California	100	Copepods	Purcell (1981a)
<i>Muggiaea atlantica</i> (n = 117)	Washington State (Oct. 1980)	100	Copepods	Purcell (1982)

habitat degradation. Parsons (1993) pointed out that studying "the role of jellyfish, teleosts, and other predators in maintaining the natural ecology of the sea is practically a nonexistent research enterprise".

Jellyfishes in fjords

Medusae, siphonophores, and ctenophores form a major part of the macroplankton of temperate fjords, which are generally considered to be high productivity systems. Extensive year-round collections in the San Juan Archipelago, between British Columbia, Canada, and Washington State, USA, yielded nearly 65 different species of medusae, siphonophores, and ctenophores in the region (Mills, 1981a and unpublished). Mackie and Mills (1983) and Mackie (1985) reported an assemblage of about 6–8 hydromedusae, 3–5 siphonophores, and 4–5 ctenophores which were sufficiently abundant to be easily seen and counted from a manned submersible in fjords of British Columbia, Canada. Occurring in horizontal bands of varying thicknesses, some of these species reached localized densities of up to 10 animals m^{-3} . They clearly formed an important community of mesopelagic predators, about which little is known.

Norwegian fjords appear to have a very similar assemblage of gelatinous predators to those found in the fjords of British Columbia, but characterization of these jellyfishes is only now in progress (F. Pagès, pers. comm.). Fosså (1992) describes a Norwegian fjord whose ecology appears to have changed in response to

an unusual population increase of a normally resident jellyfish. While many Norwegian fjords are populated by the cosmopolitan and usually deep-water scyphomedusa *Periphylla periphylla*, only a small number of fjords contain large *P. periphylla* populations. This medusa has been present in Lurefjorden near Bergen since at least the 1940s, but in 1973 it suddenly experienced a population explosion, that, as measured by the nuisance factor in fishing nets, has continued to increase over the past 20 years. Presumably as a result of the increased numbers of medusae (which feed nearly exclusively on copepods (Table 4)), the pelagic fish community in Lurefjorden is now different from nearby fjords not dominated by jellyfish. The two species of mesopelagic fish, *Maurollicus muelleri* and *Benthoosema glaciale*, known to be important copepod predators in nearby fjords, are not present in Lurefjorden, where the whiting, *Merlangius merlangus*, often reported to be associated with medusae, is the dominant larger fish. It seems quite likely that the increased presence of *P. periphylla* has driven these changes.

Feeding by medusae, siphonophores, and ctenophores

A limited number of field studies have reported the natural diets of one or more jellyfishes in a particular ecosystem. A representative selection of this information, chosen for large sample sizes, is given in Tables 2–5. Whereas scyphomedusae and ctenophores appear to be fairly generalist predators, eating a wide variety of

Table 4. Natural diets of some scyphomedusae as determined from gut contents of field-collected animals. The number of medusae of each species examined is designated by "n". Prey consumed is presented as percentage of the total number of prey items in the guts (except Brewer data, where % is number of medusae containing that type of prey). Although most scyphomedusae that have been studied feed to some extent on all available zooplankton, nearly all authors below saw evidence of varying degrees of selectivity in their studies on scyphomedusae.

Predator species	Location (season)	Prey consumed		Reference
		%	Principal types	
<i>Aurelia aurita</i>	Kiel Bight		All zooplankton esp. mollusc larvae and copepods	Schneider and Behrends (1994)
<i>Chrysaora quinquecirrha</i> (n = 390)	Chesapeake Bay, USA (summers 1987–1990)	55–71	Copepods (also sometimes numerous rotifers, cladocerans, ctenophores, polychaetes, and bay anchovy eggs)	Purcell (1992)
<i>Cyanea capillata</i> (n = hundreds)	Connecticut (springs 1980–1986)	80–85 15–30 2–12 10	Copepods Decapods and mysids Amphipods Fish	Brewer (1989)
<i>Periphylla periphylla</i> (n = 39)	Norway (March 1992)	97	Copepods	Fosså (1992)
<i>Pseudorhiza haeckeli</i> (n ≈ 35)	S. Australia (summers 1984–1986)	41 33	Fish eggs and larvae Copepods	Fancett (1988)
<i>Stomolophus meleagris</i> (n = 165)	Gulf of Mexico (summers 1986–1987)	63 16 9 8	Bivalve veligers Copepods and nauplii Tintinnids Gastropod veligers	Larson (1991)

available zooplankton, individual species of hydromedusae and siphonophores tend to have highly specialized diets, but these latter animals usually occur in multi-species assemblages that may together have a more general effect. A bias exists in many feeding studies, because the diet of the gelatinous predator was examined near some harvestable resource. Studied in a different location, a predator might show different dietary preferences. An example of this is the NE Pacific hydromedusa *Aequorea victoria*, shown by Purcell (1989) to consume herring larvae in bays where herring are known to spawn, but never found by the present author to consume any kind of larval fishes in a nearby habitat without spawning herring. We do not yet have sufficient knowledge to distinguish between the effects of excessive predation by a bloom of one or the other of these groups of jellyfishes, if such exists.

Conclusions

Medusae, siphonophores, and ctenophores are present in all of the world's oceans, often in great numbers. It appears that these resident jellyfishes are in a position to take advantage of changes in local ecology, and that

newly introduced alien jellyfishes can also sometimes take great advantage of imbalances brought about by pollution, overfishing, or other major environmental change. Systems previously dominated by many species of fishes seem capable of switching over to domination by jellyfish with surprising ease and rapidity. We do not know whether these switches are reversible by human-mediated environmental change. Our knowledge of the specific biology and ecology of most medusae, siphonophores, and ctenophores can be described as non-existent to fragmentary, so that our understanding of how jellyfishes contribute to either healthy or radically altered ecosystems is still poor.

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Table 5. Natural diets of some ctenophores as determined from gut contents of field-collected animals. The number of each species examined is designated by "n". Prey consumed is presented as percentage of the total number of prey items in the guts. With the exception of *Beroe*, ctenophores that have been studied tend to be generalists and their diets appear to reflect prey availability.

Predator species	Location (season)	Prey consumed		
		%	Principal types	Reference
<i>Pleurobrachia bachei</i> (n = >1000)	California (year round 1971)	79	Copepods	Hirota (1974)
		12	Cladocera	
<i>Pleurobrachia pileus</i> (n = 1352)	Scotland (spring 1966)	56	Copepods	Fraser (1970)
		25	Cladocera	
		7	Barnacle larvae	
		6	Invertebrate eggs	
<i>Mnemiopsis leidyi</i> (n = 3300)	Virginia USA (year round 1965-1967)	55	Copepods	Burrell and Van Engel (1976)
		11	Barnacle nauplii	
		11	Mysids	
		9	Annelid larvae	
<i>Mnemiopsis leidyi</i> (n = 192)	Black Sea (May-June 1990)	46	Bivalve larvae	Tiskhon-Lukanina, <i>et al.</i> (1992)
		37	Cladocera	
		12	Copepods	
<i>Beroe</i> spp. (n = hundreds)	Worldwide (year round)	100	Ctenophores, salps	G. R. Harbison (pers. comm.)

References

- Brewer, R. H. 1989. The annual pattern of feeding, growth, and sexual reproduction in *Cyanea* (Cnidaria: Scyphozoa) in the Niantic River Estuary, Connecticut. *Biological Bulletin*, 176: 272-281.
- Burrell, V. G. Jr., and Van Engel, W. A. 1976. Predation by and distribution of a ctenophore, *Mnemiopsis leidyi* A. Agassiz, in the York River estuary. *Estuarine and Coastal Marine Science*, 4: 235-242.
- Fancett, M. S. 1988. Diet and prey selectivity of scyphomedusae from Port Phillip Bay, Australia. *Marine Biology*, 98: 503-509.
- Fosså, J. H. 1992. Mass occurrence of *Periphylla periphylla* (Scyphozoa, Coronatae) in a Norwegian fjord. *Sarsia*, 77: 237-251.
- Fraser, J. H. 1970. The ecology of the ctenophore *Pleurobrachia pileus* in Scottish waters. *Journal du Conseil International pour l'Exploration de la Mer*, 33: 141-168.
- Greve, W. 1971. Okologische Untersuchungen an *Pleurobrachia pileus*. 1. Freilanduntersuchungen. *Helgoländer Wissenschaftliche Meeresuntersuchungen*, 22: 303-325.
- Greve, W. 1972. Okologische Untersuchungen an *Pleurobrachia pileus*. 2. Laboratoriumsuntersuchungen. *Helgoländer Wissenschaftliche Meeresuntersuchungen*, 23: 141-164.
- Greve, W. 1994. The 1989 German Bight invasion of *Muggiaea atlantica*. *ICES Journal of Marine Science*, 51: 355-358.
- Greve, W., and Parsons, T. R. 1977. Photosynthesis and fish production: hypothetical effects of climatic change and pollution. *Helgoländer Wissenschaftliche Meeresuntersuchungen*, 30: 666-672.
- Hamner, W. M. 1975. Underwater observations of blue-water plankton: logistics, techniques, and safety procedures for divers at sea. *Limnology and Oceanography*, 20: 1045-1051.
- Hamner, W. M., Madin, L. P., Alldredge, A. L., Gilmer, R. W., and Hamner, P. P. 1975. Underwater observations of gelatinous zooplankton: sampling problems, feeding biology, and behavior. *Limnology and Oceanography*, 20: 907-917.
- Harbison, G. R., Biggs, D. C., and Madin, L. P. 1977. The associations of Amphipoda Hyperideae with gelatinous zooplankton - II. Associations with Cnidaria, Ctenophora and Radiolaria. *Deep-Sea Research*, 24: 465-488.
- Harbison, G. R., Madin, L. P., and Swanberg, N. R. 1978. On the natural history and distribution of oceanic ctenophores. *Deep-Sea Research*, 25: 233-256.
- Harbison, G. R., and Volovik, S. P. 1995. The ctenophore, *Mnemiopsis leidyi*, in the Black Sea: a holoplanktonic organism transported in the ballast water of ships. In *Non-indigenous and introduced marine species*. Ed. by D. Cottingham. NOAA Technical Report.
- Hirota, J. 1994. Quantitative natural history of *Pleurobrachia bachei* in La Jolla bight. *Fishery Bulletin*, 72: 295-335.
- Larson, R. J. 1991. Diet, prey selection and daily ration of *Stomolophus meleagris*, a filter-feeding scyphomedusa from the NE Gulf of Mexico. *Estuarine, Coastal and Shelf Science*, 32: 511-525.
- Larson, R. J., Harbison, G. R., Pugh, P. R., Janssen, J. A., Gibbs, R. H., Craddock, J. E., Mills, C. E., Miller, R. L., and Gilmer, R. W. 1988. Midwater community studies off New England using the *Johnson Sea-Link* submersibles. NOAA National Undersea Research Program Research Report, 88-4: 265-281.
- Mackie, G. O. 1985. Midwater macroplankton of British Columbia studied by submersible *PISCES IV*. *Journal of Plankton Research*, 7: 753-777.
- Mackie, G. O., and Mills, C. E. 1983. Use of the *Piscis IV* submersible for zooplankton studies in coastal waters of British Columbia. *Canadian Journal of Fisheries and Aquatic Science*, 40: 763-776.

- Mills, C. E. 1981a. Seasonal occurrence of planktonic medusae and ctenophores in the San Juan Archipelago (N.E. Pacific). *Wasmann Journal of Biology*, 39: 6–29.
- Mills, C. E., 1981b. Diversity of swimming behaviors in hydromedusae as related to feeding and utilization of space. *Marine Biology*, 64: 185–189.
- Myers, R. A., Blanchard, W., and Thompson, K. R. 1990. Summary of North Atlantic fish recruitment 1942–1987. Canadian Technical Report of Fisheries and Aquatic Sciences, 1743: 1–108.
- Parsons, T. R. 1979. Some ecological, experimental and evolutionary aspects of the upwelling ecosystem. *South African Journal of Science*, 75: 536–540.
- Parsons, T. R. 1993. The need for a holistic approach to ocean ecology. *Limnology and Oceanography*, 38: 1590–1592.
- Purcell, J. E. 1980. Influence of siphonophore behavior upon their natural diet: evidence for aggressive mimicry. *Science*, 209: 1045–1047.
- Purcell, J. E. 1981a. Dietary composition and diel feeding patterns of epipelagic siphonophores. *Marine Biology*, 65: 83–90.
- Purcell, J. E. 1981b. Feeding ecology of *Rhizophysa eysenhardti*, a siphonophore predator of fish larvae. *Limnology and Oceanography*, 26: 424–432.
- Purcell, J. E. 1981c. Selective predation and caloric consumption by the siphonophore *Rosacea cymbiformis* in nature. *Marine Biology*, 63: 283–294.
- Purcell, J. E. 1982. Feeding and growth of the siphonophore *Muggiaea atlantica* (Cunningham 1893). *Journal of Experimental Marine Biology and Ecology*, 62: 39–54.
- Purcell, J. E. 1984. The functions of nematocysts in prey capture by epipelagic siphonophores (Coelenterata, Hydrozoa). *Biological Bulletin*, 166: 310–327.
- Purcell, J. E. 1989. Predation on fish larvae and eggs by the hydromedusa *Aequorea victoria* at a herring spawning ground in British Columbia. *Canadian Journal of Fisheries and Aquatic Sciences*, 46: 1415–1427.
- Purcell, J. E. 1992. Effects of predation by the scyphomedusan *Chrysaora quinquecirrha* on zooplankton populations in Chesapeake Bay, USA. *Marine Ecology Progress Series*, 87: 65–76.
- Purcell, J. E., and Mills, C. E. 1988. The correlation between nematocyst types and diets in pelagic Hydrozoa. In *The biology of nematocysts*, pp. 463–485. Ed. by D. A. Hessinger and H. M. Lenhoff. Academic Press, Orlando.
- Rogers, C. A., Biggs, D. C., and Cooper, R. A. 1978. Aggregation of the siphonophore *Nanomia cara* in the Gulf of Maine: observations from a submersible. *Fishery Bulletin*, 76: 281–284.
- Schneider, G., and Behrends, G. 1994. Population dynamics and the trophic role of *Aurelia aurita* medusae in the Kiel Bight and western Baltic. *ICES Journal of Marine Science*, 51: 359–367.
- Shushkina, E. A., and Musayeva, E. I. 1990. Structure of planktic community of the Black Sea epipelagic zone and its variation caused by invasion of a new ctenophore species. *Oceanology*, 30: 225–228.
- Studenikina, Ye. I., Volovik, S. P., Mirzoyan, A., and Lutz, G. I. 1991. The ctenophore *Mnemiopsis leidyi* in the Sea of Azov. *Oceanology*, 31: 722–725.
- Tiskhon-Lukanina, Ye. A., Reznichenko, O. G., and Lukashcheva, T. A. 1992. Diet of the ctenophore *Mnemiopsis* in inshore waters of the Black Sea. *Oceanology*, 32: 496–500.
- Zaitsev, Yu. P. 1992. Recent changes in the trophic structure of the Black Sea. *Fisheries Oceanography*, 1: 180–189.