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Species Profiles: Life Histories and Environmental Requirements of Coastal Fishes and Invertebrates (Pacific Northwest)

AMPHIPODS

by

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PREFACE

This species profile is one of a series on coastal aquatic organisms, principally fish, of sport, commercial, or ecological importance. The profiles are designed to provide coastal managers, engineers, and biologists with a brief comprehensive sketch of the biological characteristics and environmental requirements of the species and to describe how populations of the species may be expected to react to environmental changes caused by coastal development. Each profile has sections on taxonomy, life history, ecological role, environmental requirements, and economic importance, if applicable. A three-ring binder is used for this series so that new profiles can be added as they are prepared. This project is jointly planned and financed by the U.S. Army Corps of Engineers and the U.S. Fish and Wildlife Service.

Suggestions or questions regarding this report should be directed to one of the following addresses. Information Transfer Specialist National Wetlands Research Center

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CONVERSION TABLE

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Figure 1. A gammaridean amphipod (from Staude et al. 1977).

AMPHIPODS

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NOMENCLATURE/TAXONOMY/RANGE

Scientific name Amphipoda (Figure 1) Preferred common name . . Amphipods Class Crustacea Subclass Malacostraca Order Amphipoda Suborders . . Gammaridea, Hyperiidea, Caprellidea, Ingolfiellidea (Figure 2).

Geographic range: This report will focus largely on the suborders Gammaridea, Caprellidea, and Hyperiidea because of their importance in coastal areas of the

- ble marker half the

northeast Pacific Ocean (Figure 3). Gammaridea are the most abundant and diverse of the amphipods. Although primarily marine, they are also found in freshwater and certain moist terrestrial habitats (Reish and Barnard 1979). Marine Gammaridea are ubiquitously distributed. They are found in all regions, in all habitats, and at most depths. About 40% of the 80 gammaridean families are cosmopolitan in distribution; the remaining 60% are loosely associated with specific regions or zones (Barnard 1969; Bousefield 1978). Gammaridean distributions remain poorly known, but more recent studies (e.g., Barnard 1971) are



Figure 2. A, <u>Elasmopus</u> and B, <u>Eohaustorius</u>, both gammarid amphipods. C, <u>Caprella</u> <u>ferrea</u>, a caprellid amphipod. D, <u>Neocyamus</u> <u>physeteris</u> (female), a caprellid amphipod from sperm whale. E, <u>Phronima</u> <u>sedentaria</u>, a hyperiid amphipod that lives inside the tunic of urochordates. (A and B from Barnard 1975; C and D from McCain 1975; E from Barnes 1974. A-D reprinted with permission from the University of California Press; E reprinted with permission from Saunders College Publishing.)

finding more widespread distributions than were previously assumed. Off the Oregon Coast, 97 species of gammarids have been found from the surface to a depth of 2900 m (Barnard 1971), and 20 species divided among 11 families were in the upper 200 m (Pearcy 1972). About 200 gammarid species have been found in Washington waters (Staude et al. 1977). Some gammarid species dwell in subtidal or intertidal environments (Reish and Barnard 1979). The suborder Hyperiidea is entirely marine and pelagic; most members of the taxon live in the



Figure 3. Distribution of the ubiquitous amphipod suborders usemaridea and Hyperiidea in the coastal areas of the northeast Pacific Ocean.



bathyal zone, and some live in coastal waters (Bowman and Gruner 1973).

MORPHOLOGY/IDENTIFICATION AIDS

Animals of the order Amphipoda are distinguished by sessile, compound eyes, though some species are blind. A carapace is not present and the first, and sometimes the second, thoracic segments are fused with the "shrimplike" Α head. appearance results from lateral body compression. Gammarids and hyperiids have three pairs of pleopods (swimmerets); two or three pairs of uropods on the pleon (abdomen); at least eight pairs of thoracopods, counting the maxilliped; usually seven major leg pairs, called pereopods; and five or more pairs of gills. Males and females often can be distinguished morphologically. The head has five fused segments. two pairs each of antennae and maxillae, a heavily chitinized mandible, and a limblike maxilliped. There are seven freely articulated somites on the thorax (pereon). Coxal platelike lateral extensions of the thoracic pereon are developed from the first segment of each leg. Branchiae (gills) are fleshy and platelike and are attached medial to the coxae, 2-6 on each side. The abdominal region consists of three articulating segments on both anterior pleon and posterior urosome; the urosome has a terminal telson.

The following key (adapted from Barnes 1974; Kozloff 1974) is presented as an aid to separate the suborders of amphipods:

la. Pereon with seven apparent segments, all having well-developed appendages. Abdomen not vestigial. Body neither slender nor resembling that of a praying mantis. . . . 2

1b. Pereon with six apparent segments, some may have vestigial appendages; abdomen vestigial; head fused with second thoracic segment. Body slender and resembling that of a praying mantis (except for whale lice). Marine. Includes skeleton shrimp . . . Suborder Caprellidea.

2a. Eyes generally large, occupying most of head; coxae of pereopods small, often fused with the body, maxillipeds without palp; last two abdominal segments fused; body more or less transparent. Marine, and usually planktonic or associated with jellyfish or in tunics of dead salps Suborder Hyperiidea.

2b. Eyes usually present and conspicuous, but not large enough to cover most of the head; coxae of pereopods well developed, usually expanded. Marine, freshwater, and terrestrial... Suborder Gammaridea.

2c. Body elongate; coxae small; abdominal segments distinct; all but fourth and fifth pairs of abdominal appendages vestigial. Marine, interstitial. Rare. . . . Suborder Ingolfiellidea.

There currently exists no concise guide to amphipod species in the northwest Pacific. Publications of the National Museum of Canada, such as that by Conlan and Bousfield (1982), will eventually culminate in a comprehensive regional handbook on marine by gammarideans. Contributions Barnard (1975) and Staude et al. (1977) may be useful for identifying Gammaridea in restricted intertidal regions; the work of Bousfield (1978) freshwater described Gammaridea. Kozloff (1974) provided keys to the Caprellidea. Hyperiids can be identified to genus by using the descriptions published by Bowman and Gruner (1973).

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REASON FOR INCLUSION IN SERIES

Hyperiid amphipods are the third most abundant group of coastal marine crustacean zooplankton, following Copepoda and Euphausidea (Bowman and Gruner 1973). The benthic amphipods, especially Gammaridea, are an invaluable food source for many economically fish important and invertebrate limited Their mobility species. suggests that their distribution and abundance can be used as an indicator of environmental quality (Albright 1982). Omnivorous, opportunistic such as lysianassids (a feeders gammaridean family) recycle detritus and may help avert pollution by scavenging carcasses of larger animals following mass mortalities (Reish and Barnard 1979).

LIFE HISTORY

Female Amphipoda spawn via an amplexus (mating embrace) with males which lasts for hours or days. In swimming species the female swims with the male on top, or both swim on their Following ecdysis (molting) sides. and mating, eggs are laid through two ventral pores in the female's sixth thoracic sternite. Eggs can number from 1 to 200 or more. Thin, tube-dwelling gammarids have the fewest eggs, which tend to be large or contain large amounts of yolk. Because of the large size of the eggs, only one can be carried by some young females, while fully mature females carry three or four. Eggs hatch directly into juveniles resembling adults. In gammarids, one-quarter to one-half of the eggs may die before hatching. Juveniles are generally held in the brood pouch for a few hours to a few days after hatching before they are released (Barnard 1969: Reish and Barnard 1979).

Chang and Parsons (1975) found that the common inshore gammarid <u>Anisogammarus pugettensis</u> breeds year round in the Pacific Northwest, in contrast to beach and some intertidal amphipods of the cooler North Atlantic. Those species either have one brood per year or cease their reproductive activity during the coldest winter months. Females lay eggs during each of the last five or six molting stages, or at every other stage (Barnard 1969).

GROWTH CHARACTERISTICS

Growth is initially rapid in Gammaridea; molting initiates within several days of hatching and continues after maturity, slowing to every 20 to 30 days in the later stages of instar development. The average of development between (stage successive molts) lasts 15 days. Gammarids go through at least 12 instars; thus, the maximum lifespan estimates are a little more than 6 months, although some polar species are known to live 5 or 6 years (Reish and Barnard 1979).

Amphipod growth rates and lengths vary considerably. Adult amphipods range in size from less than 1 cm to about 28 cm, the largest being an undescribed lysianassid that was photographed in the abyssal Pacific Ocean (Schmidt 1968). Maximum growth rates of <u>A. pugettensis</u>, mentioned above, were 4.1% per week at 10 °C, increasing more than threefold to 14.3% per week at 20 °C (Figure 4), with higher efficiency at 20 °C. Growth relative to food intake in large (10 mg) individuals of this species was 47% to 72% when fed Enteromorpha (Chang and Parsons 1975).

POPULATION DYNAMICS AND IMPORTANCE TO FISHERIES

Amphipods are the main food item of many fish species, as well as other aquatic animals (Figure 5). Some pelagic species sometimes comprise the bulk of the diet of herring, mackerel, and Biscayan tunny (Schmitt 1968). Gammarids, on the basis of the Index of Relative Importance (IRI), were the most important food species for nearshore fishes in the Strait of Juan de Fuca (comprising more than half of



Figure 4. Growth of <u>Anisogammarus</u> <u>pugettensis</u> fed <u>Enteromorpha</u> <u>intes-</u> <u>tinalis</u> at 10 and 20 °C. (Reprinted with permission from the <u>Journal</u> of <u>the Fisheries Research Board</u> of <u>Canada</u>, from Chang and Parsons 1975).

the total IRI spectrum for 38% of the 55 fish species studied) and were the most important food item to tidepool fishes (Cross et al. 1978). For the most part, the gammarids were epibenthic rather than infaunal or pelagic. Cross et al. (1978) suggested that since hyperiid populations on which neritic fishes feed are naturally patchy, small localized perturbations are likely to create more patchiness. If adjacent areas remain unaffected, the neritic fish populations may not be adversely affected. However. sublittoral fishes, especially juvenile fishes, are more dependent on epibenthic prey, and thus more likely to be affected by perturbation. amphipod supply is although the often replenished by tidal action. Because of their relative isolation, tidepool fishes are most heavily affected by perturbation (Cross et al. 1978).

Mason (1974) hypothesized that delayed seaward migrations of juvenile

chum salmon (<u>Oncorhynchus keta</u>) and coho salmon (<u>Oncorhynchus kisutch</u>) may be attributable in part to the abundance of food organisms in rivers and estuaries. Abundant populations of gammarids in the upper estuary of Hyman Creek, British Columbia, constituted the main diet of the fry of these two salmon species. They also constituted the majority of the diet of chum fry at six nearby estuaries at low tide in the spring.

Corophium salmonis, а tubedwelling gammarid, is an abundant and preferred prey organism of chum salmon in the Skagit River salt marsh in Washington State (Congleton and Smith 1976). Though little is known of the seasonal abundances of C. salmonis, Albright (1982) found peak densities of the species in tide flats of Grays Harbor, Washington, in July and August. In the inner half of the bay they were the dominant organism on mud and sandy mud bottoms. Densities as high as 57,000/m² have been observed (Albright and Rammer 1976). From April through September production was 3.6-10.7 g/m², and turnover (production/mean biomass) was 7.2 to 8.6 g/m². In Grays Harbor, <u>C</u>. salmonis is an important prey item for dunlin (Calidris alpina), English sole (Parophyrys vetulus), and starry (Platichthys flounder stellatus) according to Smith and Mudd (1976) and for other fish species, as well as shrimp (Crangon spp.) and Dungeness crab (<u>Cancer</u> <u>magister</u>) (Albright 1982). Smith (1980) reported similarly high <u>C</u>. <u>salmonis</u> densities and predation on this amphipod by various species in other northwest estuaries.

Numerically, amphipods are the major component of the fauna of harbor pilings in California. Most are introduced species that have had little effect on indigenous amphipods in nearby areas (Barnard 1961; Reish 1964). Negligible economic loss due to fouling has resulted (Reish and Barnard 1979). In heavily polluted



Longfin Smelt Nemertean Worms

Figure 5. Fish and invertebrate predators of the amphipod <u>Corophium</u> <u>salmonis</u> (from Albright 1982).

sections of harbors, amphipods are reportedly absent both in the benthos and on pilings (Reish 1959).

Of the pelagic organisms in the upper 200 m off the Oregon coast, hyperiids comprise more than 10% of organisms by number; their abundance is known to vary seasonally (Van Arsdale 1967, cited by Pearcy 1972).

Two gammarid species have been examined for their potential as food in fish culture. Mass culture of <u>Anisogammarus pugettensis</u> was proposed by Chang and Parsons (1975) as an alternative to brine shrimp culture for young salmon; <u>A. pugettensis</u> can tolerate wide ranges of temperatures and salinities and thrives on a variety of plant and animal material.

It also scavenges dead fishes and uneaten fish food in ponds. However, its growth is slower than that of brine shrimp. Gammarus lacustris, found in shallow prairie lakes of the Hudson Bay drainage, meets dietary requirements for rainbow trout (Salmo gairdneri) 5 cm or greater, is easily captured, and can be harvested at a rate of 1,000 kg per ha per year. For most food ingredients it is comparable to or better than commercial feeds, and it improves body coloration and hence marketability of trout (Mathias et al. 1982).

ECOLOGICAL ROLE

Amphipods are considered the most efficient scavengers of sea bottoms

and shores, probably clearing up and recycling more organic shore debris than any other animal (Schmitt 1968). Griffiths and Stenton-Dozey (1981) described the importance of the gammarid Talorchestia capensis in consuming beached kelp in South This species and dipteran Africa. larvae consume some 60% to 80% of beached kelp within 2 weeks, and the gammarid is thought to make a significant contribution (through feces) to organic enrichment of the inshore marine system.

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Caine (1980), in an ecological comparison of two littoral species of caprellid amphipods in Washington State, indicated that each species has a different effect on its community. Deutella californica is a predator, but its removal did not alter community structure, even though it displays a preference for the of Obelia epibiotic community Caprella dichotoma. In contrast, laeviuscula is a periphyton scraper that has an enormous impact on the periphyton on <u>Zostera</u> <u>marina</u>, and thus increases available light for the seagrass, and permits its growth in areas where it would have otherwise Observations been excluded. on interspecific aggressive behavior indicate that <u>C. laeviuscula</u> dominant over other caprellids is dominant in protected habitats. Predatory caprellids did not appear to occur together where they would compete directly for food, while filterfeeding caprellids do compete to some extent for food (Caine 1977).

Reish and Barnard (1979) categorized gammarids and hyperiids by Nestlers include beachhabitat. hoppers of the gammarid family Talitridae, commonly found on sandy intertidal areas. High numbers occur under moisture-maintaining algal wrack, as discussed above. These must be transitory, species the authors speculated, because of frequent changes in tide and wrack accumulations. Species of six or

families seven gammarid (e.g., Ampelisca sp. and Photis sp.) construct tubes or cradles on soft or hard substrata, according to Barnard (1969). Corophium sp., common in estuaries where silting is heavy, forms masses of heavy tubes and creates currents with its abdominal appendages. The currents are strained by fringes of fine hairs on the appendages forward of the abdomen; then whatever is collected is scraped into the mouth (Kozloff 1973). Other species inhabit dwellings of other organisms. Many species are burrowers, especially in the gammarid families Haustoriidae, Oedicerotidae, Phoxcephalidae. and Elongated setae on the distal articles of the posterior pereopods are an adaptation for burrowing (Reish and Barnard 1979).

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A number of Gammaridea live on sedentary invertebrates such as corals, sponges, tunicates, anemones, and polychaetes. Their relationships with the hosts are not well understood (Reish and Barnard 1979).

Hyperiidea are primarily nek-They have well-developed tonic. swimming devices or buoyancy control, or are found in association with medusae or salps (Reish and Barnard 1979). sp., Phronima sometimes collected in plankton tows or along docks in the San Juan Islands, is found in empty salp tests (Kozloff 1974). Hyperiids may feed on the very organisms that host them, but may also use them as a base from which to forage, or they may feed on food captured by the host (Bowman et al. 1963). Their feeding habits are poorly understood. In one laboratory study of <u>Lestrigonus</u> SD. and Bougisia sp., food was shared with the host, Leptomedusa sp., when food supply was adequate, but when it was not, the amphipods fed on host tissue (Bowman and Gruner 1972). Parathemisto sp., a free-living hyperiid, preys on other plankters (Bowman 1960).









<u>Chelura</u> <u>terebrans</u>, a wood-borer found in California, is the best known amphipod pest. It enlarges holes in wood made by the isopod <u>Limnoria</u> sp. (Reish and Barnard 1979).

The swimming capability of epibenthic gammarids may reduce their susceptibility to predation. Feller and Kaczynski (1975) suggest that harpacticoid copepods were preferred over amphipods by juvenile chum salmon in Puget Sound during the spring because they are relatively easy to capture. Simenstad (1976) noted the same predatory habits for juvenile pink and chum salmon in Hood Canal, Washington, and found that juvenile salmon, in addition to preferring the less numerous and smaller harpactacoid copepods, also consumed gammarid eggs.

<u>conferv</u>iculus <u>Anisogammarus</u> is believed to defend or "buffer" its populations against predation by migrating fishes, such as juvenile chum salmon, by ecological adaptations. These adaptations decrease the foraging efficiency of the predator and include association with refuges in vegetation, clumping in refuges, association with structurally complex habitats and distributions related to riverflow and tides (Levings and Levy 1976). In Grays Harbor, Washington, however, mature male <u>C</u>. salmonis are subject to heavy predation beginning in April, when they wander over tideflats in search of females (Albright 1982).

Hyperiid swimming varies from feeble movement of appendages in <u>Cystisoma</u> sp. to rapid swimming in <u>Paraprone</u> sp. which has strong pleonal musculature (Bowman and Gruner 1973). Caprellids, attaching with posterior legs, feed by grasping food with their free anterior legs and antennae. Locomotion is accomplished with a loop-like movement in which the front legs attach while the rear ones release and reattach (Kozloff 1973).

In addition to being prey for many fish and invertebrate species, some pelagic amphipods comprise part of the crustacean diet of whales. Most of the grey whale diet on the west coast consists of six species of benthic amphipods (Matthews 1978). British gulls are also known to consume benthic amphipods (Schmitt 1968).

Locomotion in gammarids is largely by swimming; they are poorly balanced for walking. Even burrowers are strong swimmers. Small coupling hooks join pleopods, facilitating coordinated paddling motions. Some softbottom gammarids have elongated pereopods spread out like a spider's legs to prevent sinking into the mud. The body hangs upside down, lowering the center of gravity. Sediment burrowers possess strong and densely packed spines on their pereopods (Barnard 1969).

Caprellids, the suborder which includes skeleton shrimp, are largely intertidal and shallow subtidal. Their preference of substrate in the Pacific Northwest is not specific, but they do need to cling to something. Thus, they are found on algae, seagrasses, sponges, hydroids, and bryozoans, but not on bare sand or mud bottoms. Caprellids feed on diatoms, invertebrates, and possibly small detritus, and are prey for many fishes (including cod, blennies, and skates) and also for shrimp (McCain 1975). Whale parasites of the genus Cyamus are also in the caprellid suborder. This group includes about 18 hostspecific species. They lack a freeswimming stage; they leave the parental brood pouch and dig into the host with hooked dactyls (Schmitt 1968).



ENVIRONMENTAL REQUIREMENTS

Dissolved Oxygen

Pelagic gammarid and hyperiid amphipods have been collected from a scattering layer in deep, poorly oxygenated waters off southeastern Vancouver Island, British Columbia (Waldichuck and Bousfield 1962). Anisogammarus sp. and Allorchestes both common inshore gammarid SD., amphipod genera, were found in low dissolved oxygen environments (as low as 0.04 ppm at 12 °C) near sulfiterich paper pulp effluent. The former species was found in high numbers on the bottom (15 to 22 m) and the latter species, normally found in shallower waters, was observed near the surface. perhaps seeking more oxygenated water (Waldichuk and Bousfield 1962). Low oxvgen tolerance in either species remains to be determined, but Chang and Parsons (1975) observed that Anisogammarus pugettensis survived for several hours at 20% saturation levels. They also determined a Q_{10} of 1.6, lower than those of other crustaceans for which it is around 2 (Q_{10} is the factor by which the metabolic rate increases after a 10° increase in temperature). They suggest that this is an adaptation for coping with rapidly changing intertidal temperatures. Caprellids are known to leave eelgrass beds "in droves" at night when dissolved oxygen levels in the beds drop below 2 ppm.

Tolerances to low dissolved oxygen levels vary greatly among species; many are very sensitive to levels, especially low species restricted to areas where dissolved oxygen does not historically vary such as phoxogreatly. Groups cephalids (used as indicators of pollutant levels in sediment bioassays) appear much less tolerant to stressful conditions than many of the species discussed above (R. Albright, University of Washington; pers. comm.).

<u>Salinity</u>

Adult gammarids found in estuaries are fairly tolerant to a wide salinity range while many juveniles and embryos are not. Adult estuarine Corophium volutator survived in salinities of 2 to 59 ppt (McClusky 1967), but preferred a range of 10 to 30 ppt (McClusky 1970). Adult C. triaenonyx survived in a similarly wide range of salinities (Shyamasundari 1973). though juveniles could develop only at salinities of 7.5 to 37.5 ppt. For large numbers of individuals to survive and develop, 20 to 32.5 ppt were (Shyamasundari required 1976). Anisogammarus pugettensis, found naturally in 20 to 28 ppt salinities, cannot survive in freshwater but can survive at 11 ppt for at least 1 week (Chang and Parsons 1975). Some species, such as Phoxocephalid spp. or Ampeliscad spp., may have very narrow salinity tolerances. Other amphipods (e.g., <u>Gammarus</u> spp., <u>Hyalella</u> spp. and <u>Crangonyx</u> spp.) are found in freshwater.

Pollution and Dredging

Reish and Barnard (1979) observed that some amphipod species are more tolerant than others to organic pollution, but do not know what environmental factors cause the differences. It is known that some amphipods are sensitive to pollution in harbors. <u>Capitella</u> sp., a marine polychaete which is commonly used as a pollution indicator and which has a distribution that is often mutually exclusive to that of amphipods, is found in heavily polluted harbors. Capitella sp. is also found in unpolluted areas, such as deep sea bottoms off the coast of California, which are subject to freshwater inflow -- places where amphipods are notably absent (Reish and Barnard 1979).

The distribution of <u>Corophium</u> salmonis is influenced by sediment



type and depth (it prefers shallow, muddy sand substrates) more than by salinity. Other species of <u>Corophium</u> exhibit greater production near sewer outfalls -- an increase which is presumably attributable to organic enrichment (Birklund 1977).

Behavioral changes in amphipods exposed to sublethal quanities of oil have been noted and suggest a sensitivity to fresh oil. Beachhoppers are most likely to be affected by oil due to their occurrence in the high-tide wrack zone (Baker 1971), while species of <u>Ampelisca</u> show sensitivity in subtidal areas.

Dredging is likely to eliminate benthic amphipods, which live on or close to the substrate (Reish and Barnard 1979). However, McCaulley et al. (1977) suggest that in the event of dredging, adults are likely to move to nearby unaffected areas or juveniles may rapidly settle and repopulate the dredged areas (McCaulley et al. 1977). Crustaceans are generally very sensitive to pollution and, therefore, species dependent on them as food are indirectly affected by pollution.

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