EFFECTS OF LONGLINE OYSTER AQUACULTURE ON BENTHIC INVERTEBRATE COMMUNITIES IN HUMBOLDT BAY, CALIFORNIA

By

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ABSTRACT

EFFECTS OF LONGLINE OYSTER AQUACULTURE ON BENTHIC INVERTEBRATE COMMUNITIES IN HUMBOLDT BAY, CALIFORNIA

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Oyster aquaculture has had a commercial presence in Humboldt Bay for nearly 60 years and has experienced changes in scope and methodology as the industry has grown. The traditional method of bottom-culture oyster beds has been phased out, with longline oyster aquaculture becoming the common replacement. However, this transition has preceded much of the research regarding potential impacts to the broader ecosystem. The benthic invertebrate community of Humboldt Bay is a vital food source for many commercially important fishes, as well as for the many shorebirds that utilize Humboldt Bay. The importance of the invertebrate community to the ecosystem highlights the need to investigate how off-bottom culture affects invertebrate community composition. During the summer of 2017 and the winter of 2017/18, I collected benthic and epibenthic invertebrate samples from Humboldt Bay's North Bay. I then used multivariate analyses to compare the invertebrate community composition between eelgrass and mudflat habitats with and without aquaculture. I found that invertebrate communities responded most to the presence of structure and were not

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significantly different between aquaculture and eelgrass habitats. Transects conducted to measure eelgrass cover revealed significantly lower eelgrass coverage and shoot count when aquaculture was present. Eelgrass beds are important refuge areas for many juvenile fish species, as well as a vital food source for many migrating waterbirds. This study found that the benthic invertebrate communities were comparable between aquaculture and eelgrass habitats but that eelgrass densities were reduced in aquaculture habitats, which should be considered when managing current and future oyster aquaculture in Humboldt Bay.

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Appendix I. Plots of trends in compositional change of invertebrate communities within the Bird Island, East Bay, and Mad River regions of the North Bay. The x-axis indicates

INTRODUCTION

Benthic invertebrates (i.e. polychaetes, amphipods, isopods, molluscans, crustacea, etc.) are a vital part of estuarine food webs, and are essential to a sustainable aquatic ecosystem (Beaumont et al. 2007). Invertebrates are characterized by a diversity of feeding methods, from suspension and deposit feeders to active predators, and thereby forge connections that create the base of a healthy foodweb for most ecosystems (Herman et al. 1999). The response of the benthic invertebrate community can be used to gauge the health of an ecosystem, and tracking invertebrate communities through time can disclose whether a system is improving or worsening in its overall health (Pearson and Rosenburg 1978). In Humboldt Bay, an estuary located in northern California, the benthic invertebrate community also serves as a vital food source for many commercially important fishes, including several species of juvenile rockfish and an assortment of clupeids, surfperches, and hexagrammids (Simenstad and Fresh 1995; Pinnix et al. 2005). Also, of the more than thirty species of shorebirds that utilize Humboldt Bay, benthic invertebrates are necessary to the health and sustainability of many (Colwell 1994; Danufsky 2000). Within Humboldt Bay, intertidal habitats are naturally composed of eelgrass and/or open mudflat environments. However, in portions of the Bay, these habitats are modified by the addition of longline oyster aquaculture. The effects of these aquaculture beds, particularly to the benthic invertebrate community, are somewhat uncertain. To ensure a sustainable ecosystem for the fishes and birds of Humboldt Bay, it is essential to understand how benthic invertebrate communities differ between these

various habitats, and thus how any potential changes to the ecosystem (i.e., expansion of aquaculture) may impact the rest of the ecosystem.

The most diverse invertebrate communities have historically been associated with eelgrass beds, and Humboldt Bay contains over 30% of all remaining eelgrass in the state of California (Trianni 1996; Gilkerson and Merkel 2014). Eelgrass beds provide habitat complexity to the otherwise homogenous mud substrate which naturally characterizes much of Humboldt Bay (Skeesick 1963). When compared to bare sand/mud substrate, eelgrass beds have been shown to foster higher abundance and species density of benthic macroinvertebrates (Orth 1973; Stoner 1980; Orth et al. 1984; Bostrom and Bonsdorff 1997). Eelgrass beds increase habitat complexity and food availability, decrease flow rate, and create refuge from predation (Summerson and Peterson 1984; Simenstad and Fresh 1995).

Similar to eelgrass beds, oysters introduced via aquaculture provide areas of refuge and attachment surfaces for invertebrates. The oysters themselves are autogenic ecosystem engineers- the physical structures they create change the environment around them, impacting the biotic and abiotic resources available to the surrounding aquatic community (Jones et al. 1994; Gutierrez et al. 2003). Through increased habitat complexity they also support a population of prey species which can have impacts on the food web of the entire system (Ruesink et al. 2005). In a study by van der Zee et al. (2015), it was found that the addition of an ecosystem engineering mussel resulted in a shift in the species composition of benthic infauna, as well as an increase in the number and diversity of functional feeding groups represented in the intertidal ecosystem.

Although small changes in diversity are sometimes disregarded, the impacts to functional diversity can influence how that ecosystem operates (Tilman 2001; Micheli and Halpern 2005).

Aquaculture-produced shellfish is a growing market worldwide (Ababouch et al. 2016), but production in the United States is hindered by concerns about the uncertain environmental impacts (Costa-Pierce 2002; Dumbauld et al. 2009). Although many predict that aquaculture-produced protein will be vital to feeding the growing world's population, there is also a concern about the terrestrial crops and wild fish food inputs required for aquaculture production of fish species (Troell et al. 2014). Oysters produced within an open, ocean-connected system however, do not require such external inputs and do not add strain to terrestrial crops or wild fish populations. Humboldt Bay, with its connection to the Pacific Ocean, is one such fishery. As the largest producer of aquaculture oysters in the state of California, Humboldt Bay produces over 70% of California's oysters (Coast Seafoods Company 2016) and the oyster aquaculture industry has an economic impact of almost \$20 million to the region's economy (Richmond et al. 2018). The oysters produced in Humboldt Bay, Pacific and Kumamoto (*Crassostrea*)

gigas and *Crassostrea sikamea*, respectively), are cultivated using the cultch-on-longline aquaculture method, which requires settling oyster spat onto empty oyster shells and braiding them into the longline rope (Figure 1; Cote et al. 2017). Longline oyster culture in Humboldt Bay has replaced the traditional method of



Figure 1. Cultch-on-longline bed in Humboldt Bay, CA.

on-bottom oyster beds harvested via mechanical dredge (Chew 2001). On-bottom culture methods had a substantial direct impact on eelgrass, and the change to off-bottom methods was motivated by a desire to lessen the impacts to the benthic habitat (Gilkerson and Merkel 2014). However, the transition to longline aquaculture has preceded the scientific research to inform best management practices for this culture method; the majority of the oyster aquaculture research that exists today has examined the impacts of on-bottom oyster beds to the ecosystem, resulting in gaps of knowledge regarding the relationship between longline oyster culture and invertebrate communities (Dumbauld et al. 2005).

The introduction of oysters on longline can change the physical environment, with the potential to impact the associated benthic community. The physical structure of longline oyster culture can change the flow regime, while the oysters themselves eject biodeposits (feces and pseudofeces), the products of active suspension feeding (Newell and Landgon 1996; Newell and Koch 2004; Ruesink et al. 2005). These aggregated biodeposits have a faster rate of sinking than do non-aggregated particles in the water column, causing them to be more quickly incorporated into the bottom sediment (Kautsky and Evans 1987). However, the spatial extent of biodeposition from oysters depends upon how quickly those deposits settle to the substrate beneath the lines. Accumulation of feces and pseudofeces can result in over-enrichment and anoxia; but sufficient mixing, driven by hydrodynamic forces, could potentially spread biodeposits throughout the system. A study by Forrest and Creese (2006) found enhanced deposition and slowing of water flow within off-bottom oyster culture beds, and found these areas to have macrofaunal communities with composition patterns reflecting impact from disturbance. However, other studies have found oyster aquaculture habitats to foster similar benthic invertebrate communities as eelgrass beds (Hosack 2003; Rumrill and Poulton 2004; Hosack et al. 2006; Ferraro and Cole 2007). Due to the potential for changes to the physical habitat as a result of aquaculture, it is important to understand the differences in benthic invertebrate assemblages between eelgrass and mudflat habitats within Humboldt Bay, both with and without longline oyster aquaculture present.

In recent years, the aquaculture industry in Humboldt Bay has sought to expand the footprint of longline beds within the Bay. However, these proposals have been rejected due to the uncertain impact to the ecosystem (Weiner et al. 2017). Understanding how benthic invertebrate communities differ between habitats with and without longline oyster aquaculture will provide insight into how potential expansion of aquaculture practices may influence the ecosystem of Humboldt Bay and similar estuaries. Thus, the primary objectives of my thesis were to:

1. understand how benthic and epibenthic communities are affected by the presence of longline oyster aquaculture in Humboldt Bay.

compare seasonal (summer and winter) macroinvertebrate assemblages
between four habitat types: 1) eelgrass with aquaculture, 2) eelgrass without aquaculture,
mudflat with aquaculture, and 4) mudflat without aquaculture.

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METHODS

Study Site

Humboldt Bay, located in northern California, is the second largest enclosed bay in the state (Schlosser and Eicher 2012). Oyster aquaculture in Humboldt Bay primarily occurs in the North Bay, the region of the Bay where this study was focused. The North Bay is a shallow area characterized by extensive mudflats and drained by three channel systems converging to flow into the Central Bay and ultimately into the Pacific Ocean. The North Bay has a mean high water (MHW) area of approximately 14.2 square miles and a mean lower low water (MLLW) area of only 6.9 square miles (Skeesick 1963). For this study, aquaculture sites in the North Bay were delineated into three regions- Bird Island, East Bay, and Mad River (Figure 2**Error! Reference source not found.**). These t hree regions experience differences in flow regimes and turnover rates; physical parameters which can influence other abiotic factors as well as the biological characteristics of a region. To address this potential variability between regions, sampling sites were distributed equally within each region.



Figure 2. Humboldt Bay is located in northern California. This study focused on the North Bay subsection of Humboldt Bay, which was divided into three sampling regions: Bird Island, Mad River, and East Bay, where samples were collected from four habitat types: 1) eelgrass with aquaculture, 2) eelgrass without aquaculture, 3) mudflat with aquaculture, and 4) mudflat without aquaculture.

Sampling Methods

Within each of the three regions of the North Bay, I used ArcMaps's random sample tool (ArcMap 10.4.1) to locate sampling sites within each of four habitat types: 1) eelgrass with aquaculture, 2) eelgrass without aquaculture, 3) mudflat with aquaculture, and 4) mudflat without aquaculture. Based on previous studies, I selected at least five sites per habitat type per region to sample each season; this was found to be a sufficient number of samples to detect benthic macrofaunal community differences due to habitat type (Ferraro and Cole 2004). I conducted separate random sample site draws for the summer and winter sampling seasons (Appendix A).

The primary study goal was to quantify potential differences between benthic invertebrate communities in habitats with and without longline oyster aquaculture. To do this, a 2.5cm by 10cm tall core was collected using a PVC pipe with a vacuum seal when the mudflats were exposed during low tides (Figure 3). The diameter of the cores was selected based on a study by Ferraro and Cole (2004) which showed this volume to be sufficient to capture the diversity and abundance of representative benthic fauna. In addition to the cores, epibenthic organisms were sampled using an epibenthic pump similar to the one used by Toft et al. (2013). With the pump base resting on the sediment surface, the epibenthic pump pulled approximately five liters of water through the 500micron mesh bag attached to the output pipe of a hand bilge pump (Figure 3). For the early morning tides of the summer sampling season, epibenthic pump samples were collected on the incoming tide, following the collection of low tide core samples. During the winter season, low tides occurred in the evenings, so we conducted epibenthic pumps on the outgoing tide, prior to core collection, to avoid high water sampling after dark. For both seasons, samples were collected when the water was between 25-90cm deep.

Several environmental habitat variables were measured at all sampling sites. First, five 0.5m² quadrats (Figure 3) were evenly spaced along a 50m transect to measure eelgrass percent cover and eelgrass shoot count. Eelgrass percent cover was estimated by eye while the shoots were individually counted (Tallis et al. 2009). Cores were collected for analysis of carbon content and particle size of the sediment, as these factors have also been correlated with benthic community composition (Bott and Diebel 1982). The elevation of each sampling site was recorded from the ArcMap sampling map.



Figure 3. Left: Collection of 10cm deep sediment cores for benthic invertebrates and analysis for sediment characteristics. Center: Measuring eelgrass percent cover and shoot count using five 0.5m2 quadrats along a 50m transect. Right: Collection of epibenthic organisms using a manual bilge pump with a 500micron net attachment.

These samples were collected during both the summer and winter seasons to quantify community differences between the highly productive summer months and the less productive winter months. Summer samples were collected between June 22-28, 2017 and July 21-27, 2017 and winter samples were collected between December 2-7,

2017, January 2-5, 2018, and January 28 – February 2, 2018. Sampling dates were chosen to correspond with the lowest set of low tides during those months. During summer sampling, tides ranged from -0.17m to -0.55m, while the tidal range during the winter months was -0.09m to -0.48m.

Sample Processing

All samples were stored on ice in the field; following transport to Humboldt State University, sediment cores for carbon content and particle size analysis were stored at -80°C and -18°C, respectively. Samples collected for carbon content were analyzed using the loss on ignition protocol (Gavlak et al. 2005) and the equipment of the College of Natural Resources and Sciences Core Research Facility at Humboldt State University. Particle size analysis was performed using the sieve and hydrometer method for percent sand, silt, and clay (Day 1965). All samples collected for invertebrates were stored at 4°C to await sieving to remove invertebrates from the sample. Infauna sediment samples were washed with seawater through a series of stacked sieves (4mm to 2mm to 0.5mm) (Lewis and Stoner 1981), and the organic material left on each sieve was fixed in buffered 10% formalin and stained with Rose Bengal. The benthic pump samples were washed on the 0.5mm sieve and similarly fixed. The fixed samples were examined under a dissecting microscope, invertebrates were removed and placed in 70% ethanol for storage, and individual organisms were later identified to the taxonomic level indicated in Table 1. Forrest and Creese (2006) found similar taxonomic levels to be sufficient to detect spatial differences in soft-bottom invertebrate communities. Copepoda are generally considered

to be meiofauna (organisms which will pass through a 0.5mm mesh) (Watzin 1983), and as such were not counted in any sample, as those encountered were a result of entanglement in detritus within samples and were not targeted in this sampling scheme. As the aquatic invertebrate community was the target of this study, incidental terrestrial organisms were not identified.

Table 1. Breakdown of main	macrofaunal phyla into the taxonomic groups used for
classification. Superscript in	licates functional feeding group for that taxa (D= deposit,
Su= suspension, Sc= scaveng	er, P= predator, H= herbivore, O= omnivore, M= mixed.
Macrofaunal phylum	General groups used

Macrofaunal phylum	General groups used					
Arthropoda	Class level: Ostracoda ^{Su} ,					
	Order level: Amphipoda ^{Sc} , Cumacea ^{Sc} , Isopoda ^{Sc} , Tanaidacea ^{Sc}					
	Infraorder: Brachyura ^{Sc} , Caridea ^{Sc}					
	Family: Caprellidae ^M , Chironomidae ^D					
Mollusca	Class level: Bivalvia ^{Su} , Gastropoda ^H					
Annelida	Subclass: Oligochaeta ^D					
	Family level: Polychaeta:					
	Ampharetidae ^D					
	Capetellidae ^D					
	Cirratulidae ^D					
	Cossuridae ^D					
	Dorevilleidae ^P					
	Eunicidae ^{sc}					
	Glyceridae ^P					
	Lumbrineridae ^P					
	Maldanidae ^D					
	Nephytidae					
	Nereididae					
	Opheliidae					
	Orbiniidae					
	Ueniidae ²					
	Pholoidae ²					
	Phyllodicidae ⁻					
	Spionidae ²²					
Echinodermata	Phylum level: Echinodermata ^P					
Other phyla	Phylum level: Nemertea ^P , Cnidaria ^P ,					
	Platyhelminthes ^P , Sipuncula ^D					

Statistical Analyses

Eelgrass and sediment metrics

For the eelgrass measurements of percent cover and shoot count, differences between seasons, regions, and habitat types were evaluated. Because the mudflat habitats were chosen for their lack of eelgrass, only the two eelgrass habitat types- with and without aquaculture, were used for this analysis. Using R version 3.4.0, two-way analysis of variance (ANOVA) was conducted with factors of aquaculture (presence or absence), season (summer or winter), region (Bird Island, East Bay, and Mad River), and interaction terms. For all ANOVA comparisons, the assumptions of homogeneity of variance and normality were evaluated.

All site characteristics- elevation, sediment metrics (percent carbon, sand, silt, and clay), as well as eelgrass shoot count and percent cover, were summarized as an average per habitat per season.

Benthic invertebrates

Comparisons of benthic invertebrate communities in habitats with and without aquaculture were conducted to evaluate potential differences in community composition due to the addition of oyster longlines. For all analyses, the invertebrate counts for benthic cores and epibenthic pumps were combined for each site sampled. Taxa accumulation curves were generated for each habitat within each season. These plots show the rate of accumulation of new species with increasing numbers of samples (Ugland et al. 2003), and can be used to be determine if adequate sampling has occurred. No new taxa would be expected with increasing samples if the accumulation curve achieves an asymptote. If, however, the curve does not reach an asymptote, additional samples would likely continue to result in increasing numbers of taxa being encountered, indicating that an insufficient number of samples were collected. Seasonal comparison of the number of taxa encountered within each habitat type (1. eelgrass with aquaculture, 2. eelgrass without aquaculture, 3. mudflat with aquaculture, and 4. mudflat without aquaculture) was conducted using a two-way ANOVA with factors habitat type and season. Within each season, the number of invertebrates within the two eelgrass habitat types (with and without aquaculture) were also compared using one-way ANOVA with factor aquaculture presence or absence.

Patterns in invertebrate community structure within the North Bay were examined using non-metric multidimensional scaling (NMDS) and permutational multivariate analysis of variance (PERMANOVA). Due to their spatial locations within the North Bay, the three regions (Bird Island, East Bay, and Mad River) experience differences in flow regime and turnover rates- factors which can influence other abiotic factors. Due to their ability to influence the biotic community, seasons and regions were treated separately for all multivariate analyses. Because sites were placed a priori into habitat groups (eelgrass and mudflat with or without aquaculture), PERMANOVA can be used to determine if there are statistical differences between the habitat types. PERMANOVA partitions the variation in the space of the dissimilarity measure chosen when conducting the NMDS. In this case, the Bray-Curtis dissimilarity measure was used to create a dissimilarity matrix comparing sites based on taxa composition (Bray and Curtis 1957). The Bray-Curtis distance is commonly used for community composition datasets (Peterson and McCune 2001; McCune et al. 2002). PERMANOVA is suitable for multivariate community data because it does not make distributional assumptions of either the original data or the calculated dissimilarity matrix (Anderson 2017). To address homogeneity of spread, a dispersion test can be conducted to evaluate differences in variability between groups. The null hypothesis of the dispersion test is no difference in spread of the groups being compared (Anderson 2005, 2017). If a significant PERMANOVA result is obtained, a post-hoc test can be used to determine between which groups the differences occur.

Following statistical comparison of the habitat groups, non-metric multidimensional scaling (NMDS) can be used to visualize patterns in community composition. NMDS has several benefits for biotic community data; there is no assumption of multivariate normality and it is accepting of a large number of zero values, which is likely in taxa comparisons (Field et al. 1982; MacNally 1990; Clarke 1993; McCune et al. 2002; Ferraro and Cole 2007). For ecological community data, NMDS uses a multivariate dissimilarity matrix based on the taxa composition dataset to condense the data into a reduced number of dimensions, in which sites are plotted based upon their taxa similarities. For this multivariate analysis, sites were grouped by season and region. To prepare the taxa data, I first used the Hellinger transformation, which is commonly used for ordinations of taxa abundance data, and which gives low weight to rare taxa (Legendre and Gallagher 2001). With these transformed values, I again used the Bray-Curtis dissimilarity measure to create the dissimilarity matrix, as this measure has been proven to be among the most robust and effective ordination distances for community data when using non-metric multidimensional scaling (Faith et al. 1987).

The number of axes used to plot community similarities is a balance between maintaining the relationship between the similarity of sites in the original data and the similarity of sites in the synthesized and condensed data (McCune et al. 2002). Too many dimensions can result in the information being spread over too many axes, lessening the ability to discern covariation between the taxa composition of sampling sites. Generally, a stress value less than 0.2 would represent a useful ordination with low likelihood of misinterpretation, with lower stress levels indicating that the ordination better represents the actual data. A stress value greater than 0.3 represents an NDMS solution that should not be used, as it is uninformative and little better than a random placement of sampling sites (Field et al. 1982; Clarke and Warwick 2001). The correct number of dimensions was determined by plotting the stress level against different numbers of dimensions and determining at which dimension there was an asymptote (or elbow) in the stress level. This indicates that additional dimensions would not result in a large reduction in stress, and the dimension at which the elbow occurs should be the number of dimensions used for ordination plotting.

NMDS ordination can be used to visualize similarities between groupings of sampling sites. When sample units are grouped a priori to ordination mapping (i.e. by habitat type), an ellipse can be drawn about the centroid of a group, to represent the standard deviation of that grouping of sites (Oksanen et al. 2017). These ellipses can indicate the similarity between groups- ellipses which have no overlap would represent

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groups which have dissimilar taxa composition. In addition to the results of PERMANOVA analysis, which statistically analyzes similarities of groups, ordinations can be used to observe trends in group similarities.

To evaluate whether any habitat types were characterized by particular taxa, indicator taxa analysis was performed using the "indicspecies" package in R (De Caceres and Maintainer 2016). Indicator analysis assesses the statistical significance of taxa associations to specified groupings of sampling sites, providing information about the fidelity of taxa to certain habitats. By comparing taxa occurrence within the different habitats, taxa are assigned a strength of association which is compared to a permuted association level likely to occur by chance. If the observed association level is not found to be likely to occur by chance, then that taxa is determined to be significantly associated with a particular habitat (De Caceres and Maintainer 2016). Additionally, invertebrates were categorized into functional feeding groups (predator, herbivore, omnivore, suspension, deposit, and suspension/deposit feeder) (Ferraro and Cole 2007; Macdonald et al. 2010; Barnes and Hamylton 2015; van der Zee et al. 2015) to better understand how community structure is linked to broader ecological function (Fauchald and Jumars 1979; Jumars et al. 2015).

The correlation of habitat variables (sediment characteristics of carbon, sand, silt, and clay, as well as elevation, eelgrass shoot count and percent cover) to the ordination was examined using the Envfit function within the package 'Vegan' (Oksanen et al. 2017). Using continuous environmental data, Envfit fits vectors to show the direction of increase for that environmental variable. The strength of the correlation to the ordination is reflected in the length of the arrow, with a stronger correlation being displayed with a longer arrow (Oksanen et al. 2017). This can help to determine what environmental factors might be driving potential differences in community composition between groups.

To investigate further the relationship between invertebrate community composition and the environmental variables, I used a gradient forest analysis (Ellis et al., 2012; Pitcher et al. 2012; Stephenson et al., 2018). Using taxa abundances and continuous environmental data, a gradient forest analysis evaluates compositional community changes along the environmental gradients of interest, in this case: elevation, eelgrass cover, eelgrass shoot count, sediment carbon content, and sediment size. I conducted this analysis using the R package "GradientForest" (Ellis et al. 2012). This package creates regression trees to group sites based upon the community response to the environmental predictors. Regression trees use the taxa dissimilarity data at each site to split different sites into two groups (or 'branches') based upon the community response to an environmental predictor variable. The diverging branches are organized relative to a split value, 's', such that one branch is composed of sites having predictor values less than s, while the other contains sites having predictor values greater than s. In many cases, a set of taxa sensitive to an environmental gradient react to a threshold that sorts taxa composition above and below that threshold level; in these cases, the first split value would be the most informative and would occur at an 's' value close to that of the threshold. The gradient forest method uses an aggregation of many individual regression trees, created using bootstrapped samples. This method evaluates which environmental variables have the strongest overall impact on the invertebrate community, as well as

where along the environmental gradient the response occurs (Ellis et al. 2012; Pitcher et al. 2012).

RESULTS

Eelgrass and Sediment Metrics

Comparison of eelgrass shoot count and percent cover revealed similar effects of oyster aquaculture on both eelgrass metrics. Two-way ANOVA comparison using the model: Percent Cover ~ Season * Region * Aquaculture (present or absent), resulted in a significant result for the aquaculture comparison only, with less eelgrass occurring when longlines were present (p< 0.001; F= 31.31; Figure 4); all other comparisons and interactions were found to be non-significant. Shoot counts, using the same model factors as above, were found to be higher in the winter season (p= 0.002; F= 10.42; Figure 4), although the difference between count averages was less than one eelgrass shoot. There was also significantly lower shoot counts (p= 0.012; F= 6.76) when oyster longlines were present. Although a significant interaction existed between season and region (p= 0.004; F= 6.21), for the purpose of comparing the impacts of oyster culture on eelgrass cover and count, it was considered to be of low importance and is not considered further. For both analyses, the assumptions of homogeneity of variance and normality were met.



Figure 4. Comparison of eelgrass percent cover (left) and shoot count (right) in habitats with and without longline oyster aquaculture. Because both seasons resulted in significantly less eelgrass when longlines were present, the seasons were pooled for simplicity of visual comparison. Error bars indicate standard error of the mean.

Due to their potential to drive community differences, the other environmental characteristics of site elevation, percent carbon, sand, silt, and clay were summarized by average per habitat type per region for each season (Table 2). Carbon content of the sediment was highest in the summer, with Mad River region having the highest percent carbon in the winter months, and East Bay having the highest carbon content during the summer. Of the three regions, Bird Island sediment had the highest sand content and the lowest silt and clay composition. Eelgrass habitats, both with and without aquaculture were located in the lowest elevations, followed by mudflat with aquaculture, with mudflat only habitats located in the highest elevations.

		Bird Is	land Reg	gion		East Ba	ay Regio	on		Mad R	iver Re	gion	
		AE	AM	NE	NM	AE	AM	NE	NM	AE	AM	NE	NM
Eelgrass Percent Cover	Summer Winter	22.04 19.15	0 0	45.1 54.3	0.08 0	22.44 29.21	2.2 0	51.42 47.55	0.04 0	24.68 14.1	0.2 0.5	45.92 34.16	0.04 0
Eelgrass Shoot Count	Summer Winter	1.94 1.89	0 0	2.52 3.02	0 0	2.32 2.09	0.1 0	2.58 2.8	0.02 0	1.68 0.95	0.04 0.02	1.55 1.71	0.02 0
Elevation (m to MLLW)	Summer Winter	-0.22 -0.16	0.13 0.15	-0.25 -0.12	0.27 0.36	-0.15 -0.14	-0.08 -0.09	-0.24 -0.27	0.53 0.09	-0.28 -0.21	0.32 0.16	-0.12 -0.21	0.36 0.26
Sediment Percent Carbon	Summer Winter	3.66 3.07	3.47 4.1	3.93 2.96	3.35 3.28	5.45 3.35	5.44 4.1	5.11 3.79	5.864 3.25	6.58 5.84	4.68 3.97	4.69 4.73	5.86 3.45
Percent Sand	Summer Winter	50.2 72.29	46.29 35.6	44.2 46.4	55.83 44	28.2 21.83	20.2 35.6	33.83 36.4	22.4 27	11.83 15.2	31 22.8	28.67 28.67	22.4 24
Percent Silt	Summer Winter	33.4 48.2	43.43 45.2	44.8 39	34.33 42.6	47 50.33	54.8 45.2	49.17 42.2	52.2 47.67	47.83 45.6	49.2 51.2	48.17 47.5	52.2 54.2
Percent Clay	Summer Winter	16.4 18.2	10.29 19.2	11 14.6	9.83 13.4	24.8 27.83	25 19.12	17 21.4	25.4 25.33	40.33 39.2	19.8 26	23.17 23.83	25.4 21.8

Table 2. Seasonal mean environmental characteristics for each habitat type within the three regions. AE= eelgrass with aquaculture, AM= mudflat with aquaculture, NE= eelgrass without aquaculture, NM= mudflat without aquaculture.

Benthic Macrofauna

Taxa accumulation curves

Taxa accumulation curves evaluate the sufficiency of the invertebrate sampling protocol. The taxa accumulation curves for both the summer and winter seasons indicate that community analysis would benefit from additional samples. For either season, although some habitats were close, none completely achieved an asymptote, indicating that a complete census of the invertebrate community did not occur (Figure 6), and additional taxa may have been identified had more samples been collected. A two-way ANOVA of the number of taxa, with main factors Season, Habitat, and an interaction term resulted in significantly different numbers of taxa between seasons (F= 57.536; P<0.001) and habitat types (F=12.017; P<0.001). For all habitat types, more taxa were encountered during the winter season (Figure 5). Comparing the abundance of invertebrates in eelgrass habitats with and without aquaculture revealed no difference in abundances, during either season (summer: p= 0.40, F= 0.708; winter: p= 0.60, F= 0.263). The top five most abundant taxa for each habitat type within each region were summarized by season (Appendix B).



Figure 6. Taxa accumulation curves for each habitat type for both the summer (left) and winter (right) seasons.



Figure 5. Comparison of the mean number of taxa encountered within each habitat type during the summer and winter seasons. All three regions are compared, from left to right: Bird Island, East Bay, Mad River. For each habitat type, there were significantly more taxa encountered during the winter season. Although there were exceptions, generally there were no strong trends or differences between habitat types. Error bars indicate standard error of the mean.

Multivariate analyses

Summer NMDS results

Using NMDS ordination and PERMANOVA analysis, invertebrate communities within the four habitats were statistically and visually compared. For all three regions, permutation tests revealed equal dispersions (Appendix C). Following Hellinger transformation, the Bray-Curtis distance was used to create regional dissimilarity matrices. For the Bird Island region, PERMANOVA and post hoc analyses resulted in differences between community composition of mudflat habitats with and without aquaculture (Appendix D and Appendix E). The ordination for the summer Bird Island sites had a stress level of 0.122 with three dimensions, and Envfit analysis resulted in no environmental variables being significantly correlated to the ordination. PERMANOVA analysis of the East Bay dissimilarity matrix resulted in differences in the community composition between mudflat without aquaculture and both eelgrass habitats (Appendix D and Appendix E). The ordination for East Bay had a stress value of 0.124 with three dimensions, and elevation relative to MLLW was found to be correlated to the ordination. For the Mad River region, PERMANOVA analysis resulted in differences in community composition between mudflat with aquaculture and both eelgrass habitats (with and without aquaculture), as well as differences between eelgrass with aquaculture and mudflat without aquaculture (Appendix D and Appendix E). The three-dimensional ordination for Mad River had a stress value of 0.144, with eelgrass percent cover, shoot count, carbon, clay, and elevation relative to MLLW significantly correlated to the ordination. Each region, regardless of significant differences between habitat types, had

significant indicator taxa (Table 3). For all regions, ordinations for axes 1 and 2 are displayed below (Figure 7), with the additional axis comparisons in Appendix F and stress plots in Appendix G.

Winter NMDS results

Community analysis of the regional winter samples revealed differences in community composition between several habitat types. Permutational tests resulted in equal dispersion for Bird Island, East Bay, and Mad River regions (Appendix C). For all regions, taxa abundances were Hellinger transformed, and the Bray-Curtis distance was used to create a dissimilarity matrix for each region. For the Bird Island region, post-hoc testing of a significant PERMANOVA result found differences in the community composition of eelgrass without aquaculture to both mudflat habitats (Appendix D and Appendix E). The three-dimensional ordination had a stress value of 0.150, with eelgrass percent cover, shoot count, and elevation relative to MLLW reported by Envfit to be significantly correlated to the ordination. East Bay winter PERMANOVA results showed differences in the community composition of mudflat without aquaculture and all other habitat types (Appendix D and Appendix E). The East Bay ordination had a stress value of 0.140 with three dimensions, and Envfit analysis found no environmental variables to be significantly correlated to the ordination. PERMANOVA and post-hoc analysis of the Mad River winter sites resulted in a significantly different result between mudflat without aquaculture and both eelgrass habitats (Appendix D and Appendix E). Three-dimensional ordination of this region had a stress value of 0.140, with Envfit finding eelgrass shoot count, percent cover, and elevation relative to MLLW to be significantly correlated to the

ordination. Ordination solutions for axes 1 and 2 are displayed below (Figure 7), with additional axes comparisons in Appendix F and stress plots in Appendix G.



Figure 7. NMDS ordinations of the invertebrate samples collected from Humboldt Bay's North Bay. Habitat codes within ellipses indicate habitat types: AE= aquaculture and eelgrass, AM= aquaculture and mudflat, NE= eelgrass no aquaculture, and NM= mudflat no aquaculture. Overlapping ellipses indicate similarity in community composition, while clearly separated ellipses represent habitat types with differing invertebrate communities. Arrows indicate the strength and direction of increase of the significantly correlated environmental variables.

Table 3. Indicator taxa analysis and the associated functional feeding group for the summer and winter sampling seasons for the three sampled regions of Humboldt Bay. Superscript indicates functional feeding group for that taxa (D= deposit, Su= suspension, Sc= scavenger, P= predator, M= mixed.

	Bird	Bird Island		Bay	Mad River		
Habitat Type	Summer	Winter	Summer	Winter	Summer	Winter	
Eelgrass with aquaculture	Caprellidae ^M Oligochaete ^D Oweniidae ^D Phyllodocidae ^P	Ampharetidae ^D	Ostracoda ^{Su} Oligochaete ^D	Isopoda ^{Sc}	Ostracoda ^{Su}	Cirratulidae ^D Bivalvia ^{Su}	
Mudflat with aquaculture	Oligochaete ^D Oweniidae ^D Phyllodocidae ^P	Ampharetidae ^D	-	Oweniidae ^D	-	Cirratulidae ^D Oweniidae ^D	
Eelgrass no aquaculture	Caprellidae ^M Oligochaete ^D Oweniidae ^D Phyllodocide ^P	Ampharetidae ^D	Ostracoda ^{Su} Oligochaete ^D	Nephytidae ^P Oweniidae ^D	Ostracoda ^{Su} Bivalvia ^{Su}	Cirratulidae ^D Bivalvia ^{Su} Oweniidae ^D	
Mudflat no aquaculture	-	Chironomidae ^D	Oligochaete ^D	Oweniidae ^D	Bivalvia ^{Su}	Bivalvia ^{Su} Oweniidae ^D	

Gradient Forest results

The gradient forest analysis was used to determine which of the seven environmental variables (elevation relative to MLLW, eelgrass percent cover, eelgrass shoot count, percent sand, silt, clay, and sediment carbon content) had the largest influence on the invertebrate community in each region, and where along the gradient of each of the variables the invertebrate community had the largest response. The overall importance of each of the environmental variables was expressed as R² weighted importance (Appendix H). To evaluate the response of the invertebrate community along those environmental gradients, the two most important variables were analyzed further. Because eelgrass shoot count and percent cover revealed similar trends in invertebrate response, in situations where these eelgrass measures were the two most important environmental parameters, the second eelgrass measure was excluded and the next most informative environmental factor was used in its place. The selected habitat factors were plotted to evaluate where along the environmental gradient the split density, and therefore community response, was greatest (Figure 8). These plots delve into the community response, displaying the cumulative importance of an environmental factor on individual taxa, where shallow slopes are indicative of a slow rate of compositional change, and a steep slope indicates a relatively high rate of change in community composition.

For most regions, I found substantial invertebrate compositional change to be related to changes in elevation, while the other factors that influenced invertebrate compositional change varied by region and season. In the Bird Island region, the gradient forest analysis indicated that the primary factor influencing the invertebrate community

was elevation, while silt was the secondary factor, for both summer and winter. During the summer in Bird Island, at an elevation of approximately -0.2 m, and at 25% silt content, there were considerable changes in the importance of Lumbrineridae, a predatory polychaete. During the winter, the composition of bivalvia varied at an elevation of approximately 0.4m and the composition of Pholoidae, another predatory polychaete, varied at approximately 30% silt content. In the East Bay region, the important factors influencing the invertebrate community differed between the summer (percent silt and carbon) and winter (eelgrass cover and elevation). During the summer, the major change was in the composition of the taxa Maldanidae, a deposit feeding polychaete, at carbon content of approximately 4.0%. In the winter, the largest observed changes were also in polychaete taxa. For eelgrass cover, it was a predatory taxa that had the largest change (Nephytidae), whereas for elevation it was an omnivore (Nereididae). Finally, elevation and eelgrass metrics were the factors that had the largest influence on invertebrate community composition in the Mad River region in summer and winter. During the summer, a predatory polychaete (Glyceridae) was the taxa whose composition varied the most for both elevation (at approximately -0.2 m) and eelgrass count (at approximately 1.5 shoots). During the winter, the composition of another predatory polychaete (Pholoidae) varied with cover (at approximately 30%) and the composition of bivalvia varied with elevation (at approximately 0.2 m).



Figure 8. Graphical outputs of GradientForest analysis for the Bird Island, East Bay, and Mad River regions of Humboldt Bay. Plots display taxa which were most impacted by changes along that gradient; top three most impacted taxa are indicated in legend.

DISCUSSION

Benthic Invertebrate Habitat Use

Based on the results of this study, benthic invertebrate assemblages are most strongly impacted by the presence of structure. Although the trend was stronger for some regions than others, throughout both seasons the communities of mudflat without aquaculture differed from the invertebrate communities of aquaculture and eelgrass habitats. The slight exception to this trend was found in the Mad River summer and Bird Island winter samples, where there was also a significant difference between mudflat with aquaculture and eelgrass communities. In general, the results from the NMDS ordinations, in which the confidence intervals for the oysters and eelgrass habitats overlap, support the findings of other studies that have also found that invertebrate communities are similar in various types of structures habitats (Dumbauld 2003; Hosack 2003; Dealteris et al. 2004; Rumrill and Poulton 2004; Ferraro and Cole 2007). This is true regardless of whether, or not, that structure was native eelgrass beds or commercial oyster aquaculture.

In the otherwise barren bottom of the intertidal mudflats in Humboldt Bay, oyster longlines contribute broken shell pieces as well as complete, living oysters to the bottom habitat. These fallen oysters add heterogeneity to the bottom environment, providing substratum for boring and attachment and a refuge from predation and tidal currents (Gutierrez et al. 2003). The production of habitat heterogeneity, complexity, and structure

appears to support a similar suite of invertebrates, whether that habitat is eelgrass or aquaculture. However, in addition to creating physical habitat, oysters can also increase deposition to the sediment. Although organic inputs are necessary for a thriving invertebrate community, an overabundance of organics can have considerable impacts to the local area. Soft sediment communities are commonly characterized by relatively large filter feeders, though as organic inputs increase, a shift towards smaller deposit-feeding organisms often occurs (Pearson and Rosenburg 1978; Forrest et al. 2009; Mckindsey et al. 2011). The indicator taxa analysis generally associated deposit feeders with aquaculture habitats (both eelgrass and mudflat), as well as eelgrass without aquaculture. Suspension feeders, however, were associated with habitats lacking aquaculture, as well as eelgrass with aquaculture (Table 3). If a shift towards deposit feeders (which are associated with organically enriched habitats) were occurring, additional field studies would be needed to investigate this further. However, as evidenced by the taxa accumulation curves, this study would have benefitted from additional samples, which might support or contradict this potential trend in functional feeding group shifts.

Seasonality

Both the summer and winter seasons were characterized by distinctions between habitat types, particularly between habitats with structure and the barren mudflats. Interestingly, there were, regardless of habitat type, more taxa encountered during the winter season than the summer season. This is in contrast to the findings of other local studies, which found either fewer or no difference in taxa in the winter months compared to the summer season (Rumrill and Poulton 2004; Osborn 2017). Differences in seasonal patterns could have been impacted by the timing of the winter sample collection. Additionally, pump collection occurred in the early morning during the summer, and in the evening in the winter. This introduced variability with the potential for diurnal movement from mobile epifauna to impact the invertebrate community collected via the pump sampler.

In contrast to the expected seasonal differences, eelgrass percent cover was not found to be significantly different between seasons, and shoot count was actually higher in the winter season. However, it has been well documented that eelgrass undergoes seasonal fluctuations, with the active growing season ranging from May to September, and eelgrass coverage often decreasing in the winter months (Rumrill and Poulton 2004; Gilkerson and Merkel 2014). If the winter of 2017-2018 had a late start, as might be indicated by the lack of eelgrass decline, then perhaps the invertebrate community was still in transition during the winter sampling season. However, despite this slight seasonal inconsistency with some published literature, the overall patterns of community composition between habitat types are consistent with similar studies.

Ecosystem Implications

This study did not find significant differences in the invertebrate community of habitats with and without longline oyster aquaculture. In the case of foraging shorebirds and wading birds in Humboldt Bay, the invertebrate community within longline beds may be preferred. A study by Connolly and Colwell (2005) found greater abundances of these birds within longline plots than in control areas. Pinnix (2004) likewise found similar numbers of fish species between habitats with and without longline culture, although this study was primarily to investigate the effectiveness of sampling equipment rather than comparison of community assemblages. Because this study was focused on the invertebrate community, the predator-prey connections can only be speculated upon, and additional studies focusing on the foodweb implications would be instrumental in drawing connections between invertebrate community and the Humboldt Bay ecosystem.

I found that longline aquaculture habitats had reduced eelgrass densities, which are an important component of the Humboldt Bay ecosystem. Comparison of eelgrass habitats with and without aquaculture resulted in significantly less eelgrass, both shoot count and cover, when aquaculture was present. This is consistent with previous research investigating the impacts of off-bottom bivalve culture on eelgrass. A recent metaanalysis by Ferriss et al. (2019) reported that longline off-bottom culture methods resulted in a 44% decrease in eelgrass density and a 61% decrease in eelgrass reproduction. Although the presence or absence of aquaculture in eelgrass habitats may not have a strong effect on the number of invertebrates present, there may still be impacts to the broader ecosystem. Semmens (2008) found smolts of the ecologically and economically important Chinook salmon to exhibit a strong preference for native eelgrass habitats in Willapa Bay. Pacific Herring, an important forage fish, preferentially spawn on the blades of eelgrass (Barnhart 1988). Most herring spawning in Humboldt Bay occurs in the North Bay (Rabin and Barnhart 1986), and although there has been a recent dearth of research into Humboldt Bay herring populations, anecdotal reports of declines through the 1990s may link Herring and eelgrass populations in Humboldt Bay (Watters et al. 2001).

Results from the gradient forest analysis suggest that the factors that influence invertebrate composition varies regionally throughout Humboldt Bay, but is generally related to elevation. In addition to elevation, the factors that had the largest impact on invertebrate concentration were related to the sediment composition or a measure of the eelgrass density (cover or shoot count). The Bird Island region had percent silt concentration as the second most important factor influencing the invertebrate community in both summer and winter. In contrast, the Mad River region in both the summer and winter seasons resulted in at least one eelgrass measure as a primary factor influencing changes in community composition. The East Bay region was a mix, with sediment composition in the summer and eelgrass cover in the winter. Based on the gradient forest analysis, the taxa which were shown to be most strongly impacted in all regions were predators in the polychaete family. Predators have been found to be an important factor in structuring soft-bottom communities (Ambrose 1984; Wilson 1990). Based on these findings, future research is warranted to examine how invertebrates in the predator functional feeding group respond to aquaculture.

Benthic invertebrates, as well as eelgrass habitats, are also vital to the survival of wintering waterbirds in Humboldt Bay. While invertebrate consumption by waterfowl can be varied, many shorebirds feed primarily upon aquatic invertebrates (Afton et al. 1991; Euliss and Grodhaus 1991; Skagen and Oman 1996) In addition, the dynamics of piscivorous waterbird populations in Tomales Bay, CA, were found to be tied to the availability of herring roe, with pulses of herring spawning leading to increases in the abundance of waterbirds for the next three winters (Kelly et al. 2018). Dabbling waterbirds, including *Branta bernicla nigricans*, the Black Brant goose, are highly dependent upon eelgrass success. Moore et al. (2004) found variability in Brant numbers across flyway sites to be strongly correlated to the presence of high eelgrass abundance. Although I measured that longline aquaculture habitats had reduced eelgrass densities, a resource important to a variety of estuarine species, the impacts appear to be localized to the area encompassed by the longline bed. This indicates that careful management of the oyster aquaculture industry could conserve eelgrass beds for use by dependent members of the broader ecosystem.

Management Implications

This study of benthic invertebrate habitat use provides insight into one component of the complex ecosystem of Humboldt Bay. The increasing importance of oyster aquaculture to feeding the earth's growing population suggests that this and similar studies should be applied to developing options for management. As the need for global protein production grows, efforts should be focused on methods which can achieve a balance between production and environmental sustainability.

With the balance between providing resources for sustaining human populations while also conserving the natural environment, the available research should be used to inform best management practices for oyster aquaculture in Humboldt Bay. Rumrill and Poulton (2004) found that small changes to longline culture methods can have large impacts on eelgrass success. For example, they found that increasing line spacings from 2.5 feet (which is standard in Humboldt Bay) to five or ten feet resulted in significantly increased eelgrass cover and shoot count. With the potential for positive impacts to other eelgrass-associated species, and negligible impacts to the benthic invertebrate community (Dumbauld 2003), increased line spacing of longline oyster aquaculture beds within Humboldt Bay could be considered.

When viewing the benthic invertebrate community in isolation, the results of this study are encouraging for the lack of impacts of longline oyster culture. However, I did find that the eelgrass cover was lower in longline oyster culture habitats relative to habitats without oyster culture. These effects can have implications throughout the ecosystem, and these should be considered when planning placement of oyster aquaculture in Humboldt Bay.

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APPENDIX A

Appendix A. Summer and winter sampling sites for the Bird Island, East Bay, and Mad River regions of Humboldt Bay's North Bay.







Appendix A, continued. Summer and winter sampling sites for the Bird Island, East Bay, and Mad River regions of Humboldt Bay's North Bay.

APPENDIX B

	Habitat	Bird Island		East Bay		Mad River	
	Туре	Taxa	Count	Таха	Count	Taxa	Count
		Capetellidae	67	Tanaid	87	Tanaid	418
	ith ure	Cirratulidae	57	Capetellidae	78	Gammarid	127
	iss w cultu	Tanaid	55	Syllidae	66	Syllidae	124
	elgra vqua	Bivalvia	55	Bivalvia	53	Capetellidae	88
	Ε	Cumacea	45	Cirratulidae	40	Spionidae	32
		Gammarid	45				
	4 0	Cumacea	114	Tanaid	70	Tanaid	56
	: witl	Capetellidae	92	Syllidae	64	Capetellidae	45
	dflat uacu	Tanaid	88	Capetellidae	63	Cirratulidae	30
r	Mu Aq	Cirratulidae	79	Bivalvia	45	Syllidae	11
nme		Syllidae	50	Cirratulidae	30	Gammarid	8
Sur							
	out	Tanaid	120	Gammarid	120	Tanaid	59
	Vith	Syllidae	108	Bivalvia	90	Capetellidae	53
	ass V uacu	Cumacea	74	Capetellidae	83	Gammarid	33
	elgra Aq	Bivalvia	70	Tanaid	71	Syllidae	21
	Ц	Gammarid	51	Ostracod	66	Bivalvia	21
	out	Gammarid	209	Gammarid	230	Tanaid	220
	Vitho Iture	Tanaid	99	Syllidae	102	Gammarid	100
	lat V uacu	Capetellidae	86	Bivalvia	62	Syllidae	70
	1udf Aqı	Cirratulidae	75	Tanaid	54	Bivalvia	50
	2	Bivalvia	46	Capetellidae	27	Capetellidae	26

Appendix B. Top five most abundant taxa per habitat type per region, divided into the summer and winter sampling seasons.

	Habitat	Bird Islan	d	East Bay		Mad River	
	Туре	Taxa	Count	Taxa	Count	Taxa	Count
	e ih	Syllidae	113	Syllidae	183	Syllidae	293
	wit	Capetellidae	111	Tanaid	162	Tanaid	154
	rass acu	Cirratulidae	108	Capetellidae	154	Capetellidae	105
	lelg Aqu	Spionidae	98	Cirratulidae	124	Gammarid	70
	_ ш `	Tanaid	84	Spionidae	65	Ostracod	58
	ë	Tanaid	186	Tanaid	555	Tanaid	273
	wit ltur	Cirratulidae	148	Syllidae	520	Gammarid	174
	lf1at Iacu	Syllidae	130	Gammarid	171	Syllidae	88
	Muđ Aqu	Spionidae	125	Capetellidae	152	Spionidae	56
er		Capetellidae	124	Bivalvia	129	Capetellidae	47
Vint							
>	grass Without Aquaculture	Capetellidae	142	Capetellidae	200	Capetellidae	155
		Syllidae	126	Tanaid	185	Tanaid	140
		Cumacea	83	Syllidae	170	Syllidae	136
		Cirratulidae	78	Cirratulidae	145	Spionidae	56
	Eel	Spionidae	64	Spionidae	119	Cumacea	36
	iout e	Tanaid	386	Bivalvia	183	Bivalvia	220
	Vith	Capetellidae	145	Tanaid	165	Tanaid	219
	at V acu	Bivalvia	141	Gammarid	96	Gammarid	117
	adfl Aqu	Spionidae	113	Syllidae	81	Capetellidae	76
	/ Mı	Syllidae	86	Capetellidae	52	Syllidae	37

Appendix B, continued. Top five most abundant taxa per habitat type per region, divided into the summer and winter sampling seasons.

APPENDIX C

Appendix C. Results of permutational analysis tests to evaluate equality of dispersion prior to PERMANOVA analysis of regional community composition. Significance values greater than 0.05 indicate failure to reject the null hypothesis of equal dispersion, indicating fulfillment of this assumption. Rejection of the null hypothesis indicates a need to further evaluate equality of dispersion using visual plotting of habitat dispersions.

Season	Region	Degrees of Freedom	F-value	Significance
	Bird Island	3	0.4039	0.7552
Summer	East Bay	3	0.4441	0.7333
	Mad river	3	0.6084	0.6374
r	Bird Island	3	1.0827	0.3736
Winte	East Bay	3	1.6335	0.2008
	Mad River	3	0.1101	0.958

APPENDIX D

Appendix D. Results of permutational analysis of variance (PERMANOVA) for each region for each sampling season. Significant results were evaluated using a post-hoc test, found in Appendix E.

Season	Region	Degrees of Freedom	F-value	Significance
	Bird Island	3	1.5889	0.0170*
mmer	East Bay	3	1.2001	0.1389
Su	Mad River	3	2.1444	0.0021*
s	Bird Island	3	1.7859	0.0051*
Winte	East Bay	3	2.4454	<0.001*
	Mad River	3	2.1552	< 0.001*

APPENDIX E

Appendix E. Results of post-hoc testing of significant PERMANOVA results for the summer (top) and winter (bottom) seasons. Dashes indicate those habitats were not significantly different for any region. Regional entries indicate those habitats were found to be significantly different within that region.

Habitat Type				
Summer season	Eelgrass with	Mudflat with	Eelgrass no	
	aquaculture	aquaculture	aquaculture	
Mudflat with aquaculture	Mad River	-	-	
Eelgrass no aquaculture	-	Mad River	-	
Mudflat no aquaculture	Mad River	Bird Island	-	

Habitat Type				
Winter season	Eelgrass with	Mudflat with	Eelgrass no	
	aquaculture	aquaculture	aquaculture	
Mudflat with	-	-	-	
aquaculture				
Eelgrass no aquaculture	-	Bird Island	-	
Mudflat no aquaculture	East Bay Mad River	East Bay	Bird Island East Bay Mad River	

APPENDIX F

Appendix F. Additional axis comparisons of NMDS ordinations for the invertebrate samples collected from Humboldt Bay's North Bay. Habitat codes within ellipses indicate habitat types: AE= eelgrass with aquaculture, AM= mudflat with aquaculture, NE= eelgrass no aquaculture, and NM= mudflat no aquaculture. Arrows indicate the strength and direction of increase of the significantly correlated environmental variables



Appendix G. Stress plotted against number of dimensions to determine the correct number of dimensions for ordination plotting



APPENDIX H

Appendix H. R^2 weighted importance of environmental variables based on GradientForest analysis. The top two variables for each season and region combination were used for further analysis of the relationship between individual variables and invertebrate taxa, except in the case of the top two variables being eelgrass shoot cover and count, in which case the next best variable was used, due to the similarity in invertebrate response to the two eelgrass metrics.



APPENDIX I

Appendix I. Plots of trends in compositional change of invertebrate communities within the Bird Island, East Bay, and Mad River regions of the North Bay. The x-axis indicates the environmental parameters most influential to driving changes in the invertebrate community. The black line indicates the density of regression tree splits at that level of the environmental gradient (how much community sorting occurred at that point in the gradient) and the red line indicates the density of samples taken at various points along the gradient. The blue line displays the ratio of the black line to the red; peaks in the blue line indicate gradient locations where the compositional change of the invertebrate community occurred.



Appendix I, continued. Plots of trends in compositional change of invertebrate communities within the Bird Island, East Bay, and Mad River regions of the North Bay. The x-axis indicates the environmental parameters most influential to driving changes in the invertebrate community. The black line indicates the density of regression tree splits at that level of the environmental gradient (how much community sorting occurred at that point in the gradient) and the red line indicates the density of samples taken at various points along the gradient. The blue line displays the ratio of the black line to the red; peaks in the blue line indicate gradient locations where the compositional change of the invertebrate community occurred.

