**From the predictable to the unexpected: kelp forest and benthic invertebrate community dynamics following decades of sea otter expansion**

Andrew O. Shelton1, Chris J. Harvey1, Jameal F. Samhouri1, Kelly S. Andrews1, Blake E. Feist1, Kinsey E. Frick2, Nick Tolimieri1, Gregory D. Williams3, and Liam D. Antrim4

1Conservation Biology Division, Northwest Fisheries Science Center, National Marine Fisheries Service, National Oceanic and Atmospheric Administration, 2725 Montlake Blvd E, Seattle, WA 98112 USA

2Fisheries Ecology Division, Northwest Fisheries Science Center, National Marine Fisheries Service, National Oceanic and Atmospheric Administration, 2725 Montlake Blvd E, Seattle, WA 98112 USA

3Pacific States Marine Fisheries Commission, under contract to Northwest Fisheries Science Center, National Marine Fisheries Service, National Oceanic and Atmospheric Administration, 2725 Montlake Blvd E, Seattle, WA 98112 USA

4Olympic Coast National Marine Sanctuary, National Ocean Service, National Oceanic and Atmospheric Administration, 115 E. Railroad Ave. Suite #301, Port Angeles, WA 98362 USA

**Abstract**

The recovery of predators has the potential to restore ecosystems and fundamentally alter the services they provide. One of the most iconic examples of this potential comes from keystone predation by sea otters in nearshore habitats of the Northeast Pacific. Here we combine spatial time-series of sea otter abundance, canopy kelp area, and benthic invertebrate abundance from Washington State, USA, to examine the shifting consequences of sea otter introduction for kelp and kelp forest communities. We leverage the spatial variation in timing and speed of sea otter recovery to understand connections between sea otters and the kelp forest community. A small population of sea otters was re-introduced to the Washington coast in the late 1960s, which has since grown to more than 1,400 individuals. This population increase created a pronounced decline in sea otter prey - particularly kelp-grazing sea urchins – and led to an expansion of canopy kelps until roughly 2000. However, while sea otter and kelp population growth rates were positively correlated prior to 2000, this association disappeared over the last two decades. This occurred despite the fact that surveys of benthic invertebrates show that sea otter prey have continued to decline. Kelp area trends appear to be largely decoupled from both sea otter and benthic invertebrate abundance at current densities. However, variability in kelp abundance has declined in the most recent 15 years of the time series. Altogether these results suggest that initial nearshore community responses to sea otter population expansion follow predictably from trophic cascade theory, but now other factors may be as or more important in influencing community dynamics. Thus, the utility of sea otter predation in ecosystem restoration must be considered within the context of complex and shifting environmental conditions.

**Keywords**

Sea otters, sea urchins, kelp forests, top-down control, trophic cascades, predator-prey interactions, keystone predator hypothesis, nearshore ecology, community ecology, spatial ecology, marine ecosystems, *Enhydra lutris,* *Nereocystis luetkeana,* *Macrocystis pyrifera*

**Introduction**

Sustainable management and conservation of marine ecosystems hinges on understanding natural and anthropogenic pressures and structural forces that act on system stability (Knowlton 2004). While marine resources and ecosystem services in coastal zones are affected by climate and environmental variability as well as human activities like fishing, nutrient loading and habitat alteration (e.g., Sherman and Duda 1999, Möllmann et al. 2009), species interactions also play an important and more nuanced role in marine ecosystem dynamics. For example, “keystone” species affect marine community structure and function to an extent that is disproportionate to their biomass (Paine 1969, Power et al. 1996). A classic example is the sea otter, *Enhydra lutris,* in coastal waters of the North Pacific Ocean from Alaska to California. Sea otter predation can severely reduce local densities of benthic grazing invertebrates, such as sea urchins, thereby allowing kelp canopies to develop and expand (Estes and Palmisano 1974, Breen et al. 1982, Estes and Duggins 1995, Steneck et al. 2002).(Watson & Estes 2011). The indirect effect of sea otters on kelp is important, given kelp forests are among the most productive ecosystems on Earth (Mann 1973), support distinct fish, invertebrate, and understory algal communities (Duggins 1988, Ebeling and Laur 1988, Reisewitz et al. 2006, Markel and Shurin 2015) (Watson & Estes 2011) and perform ecosystem roles such as wave energy attenuation (Pinsky et al. 2013) and carbon storage (Wilmers et al. 2012). Similar community- and ecosystem-level consequences of sea otters have been noted in other coastal habitats as well (e.g. seagrass communities; Hughes et al. 2013).

Sea otters are native to the coast of the Olympic Peninsula of Washington state, USA (Fig. 1), but were hunted to extirpation by the early 1900s (Lance et al. 2004). Reestablishment efforts began in 1969-1970, when 59 sea otters were translocated to Washington from Amchitka Island, Alaska (Jameson et al. 1982). Despite high mortality in the early 1970s, the population eventually expanded (Fig. 1), surpassing 200 individuals by 1989 (Jameson 1993) and 600 by the late 1990s (Jameson and Jeffries 1999); Fig. 1). SCUBA surveys at multiple sites (Fig. 1) in 1987 indicated that otter densities were correlated with increased coverage of foliose and canopy-forming kelps, and reduced abundance and size of benthic invertebrates, including kelp-grazing sea urchins in the genera *Mesocentrotus* and *Strongylocentrotus* (Kvitek et al. 1989). Subsequent surveys in 1995 and 1999 indicated that the expanding sea otter population had produced these keystone predator effects on invertebrates and kelp at additional areas off the coast (Kvitek et al. 1989, 1998, 2000) (Kvitek et al. 2000) (Kvitek et al. 1989). Around this time, the kelp canopy reached peak surface coverage at the scale of the Olympic Coast (Fig. 1; Washington Department of Natural Resources kelp monitoring program; https://www.dnr.wa.gov/programs-and-services/aquatics/aquatic-science/kelp-monitoring; (Pfister et al. 2017))

Since the subtidal community surveys in 1999, the Olympic Coast sea otter population has more than doubled (Fig. 1; Jeffries and Jameson 2014). We expect that increased top down predation by sea otters might have further suppressed benthic macroinvertebrates and increased kelp canopy cover; however, the total kelp canopy area has declined from a peak abundance in roughly 2005 (Fig. 1). The apparent decoupling of sea otter and kelp changes warrants renewed research to understand patterns of nearshore community change at the regional and landscape scales. An intriguing possibility is that the decoupling of sea otters and kelp indicates a recovery of benthic grazers on the Olympic Coast.

While sea otters are generally assumed to play a strong top-down role in shifting North Pacific coastal ecosystems from herbivore-dominated to algae-dominated (Soulé et al. 2003), this generality has been both affirmed and challenged over the past 40 years (Watson & Estes 2011). There are numerous examples of Pacific coastal systems that are not herbivore-dominated in the absence of sea otters (Foster 1990, Lafferty 2004, Carter et al. 2007, Reed et al. 2011). For example, Reed et al. (2011) found that wave disturbance overwhelmed the effect of herbivory and nutrient availability in determining kelp forest dynamics in central and southern California. This example highlights the importance of other physical and biological interactions for structuring coastal habitats, and encourages an explicit consideration of the spatiotemporal heterogeneity of coastal kelp systems. Such a landscape perspective on the drivers of heterogeneity and complexity has been used to improve understanding of kelp forest dynamics in California (Bell et al. 2015).

Here we combine a 30-year time series of sea otter, kelp and benthic invertebrate data off the Olympic Coast in order to better understand nearshore marine community dynamics at regional and landscape scales. We conduct spatio-temporal analyses on sea otter and kelp data from publicly available surveys and conduct new field sampling to extend previous kelp forest invertebrate surveys conducted at focal sites by Kvitek et al. (1989, 1998, 2000). Our ability to understand the interactions between habitat, predators, and prey is essential for proper management and conservation of coastal species, habitats, ecosystems and services, particularly in this area where maintaining high native biodiversity and keystone species populations are explicit management objectives (Office of National Marine Sanctuaries 2008).

**Materials and Methods**

*Study locations*

Our study sites fall within the Olympic Coast National Marine Sanctuary (OCNMS, designated in 1994). We conduct analyses at landscape- (sea otter abundance and kelp coverage for the entire OCNMS-wide region) and local-scales (sea otter, kelp and invertebrate abundance by region and by site). For the landscape-scale analysis, data for sea otters and total kelp canopy surface coverage were derived from long-term monitoring surveys conducted in most years. Below we describe these data sources and detail how we connect these OCNMS-wide surveys to provide sea otter and kelp abundance at our focal sites.

For our local-scale analyses, we focus on ten kelp forest sites located within the range of OCNMS surveys for sea otters and kelp canopies (Fig. 1). Most of the sites are on the outer coast, while two sites, Chibadehl Rocks and Neah Bay, are inside the Strait of Juan de Fuca (Fig 1). All sites feature subtidal rocky reef habitat with dense stands of canopy kelp (*Nereocystis luetkeana* and/or *Macrocystis pyrifera)*, along with diverse communities of understory red, brown, and green algae. Canopy forming kelp forests generally occupy depths shallower than 10 m in OCNMS. SCUBA divers surveyed each site for benthic invertebrates in 2015 (see methods below) and in at least two of the three years (1987, 1995, and 1999) by Kvitek and colleagues (Kvitek et al. 1989, Kvitek et al. 2000) (add 1998). Six sites (Teahwhit Head, Rock 305, Cape Johnson, Cape Alava, Anderson Point, and Neah Bay; Fig. 1) were surveyed in all four years monitoring was conducted.

For regional comparisons, we divided the ten sites into three geographic groupings: Northern (Neah Bay, Chibadehl Rocks, and Tatoosh Island), Central (Andersen Point, Point of the Arches, and Cape Alava) and Southern (Cape Johnson, Rock 305, Teahwhit Head, and Destruction Island). These groupings are similar to the areas used to describe sea otter trends within OCNMS historically (Lance et al. 2004) and reflect distinct geographic patterns in kelp and sea otter trends (see Results). We also use these groupings to account for pseudo-replication in statistical analyses and to allow for regional differences in biological relationships.

Finally, we calculated an exposure metric for each site, which was a composite of potential wind driven wave energy, to use as an explanatory variable in statistical models (see wave supplement for specific details). We used the waves tool (Rohweder et al. 2012) in ArcGIS to create a gridded surface of potential wave exposure for the length of coast represented across the study sites. We ran the model on a 30-m grid extending from the coastline out to 1,000 km. We used the NOAA Continually Updated Shoreline Product (CUSP) shoreline for the United States as the boundary for the land-sea interface (<https://shoreline.noaa.gov/data/datasheets/cusp.html>; accessed 19 September 2017). We ran the model using the SPM-Restricted method, which calculates wind fetch using the average of five radials spread three degrees apart (Smith 1991). We used the weighted fetch option in the waves tool, with probabilistic wind direction inputs derived from 1981-2010 normal hourly wind direction data for the Quillayute State Airport (Arguez et al. 2010). Finally, we overlaid study site points with the resulting weighted wave exposure grid and extracted the corresponding grid cell values.

*Sea otter abundance and distribution*

We extracted sea otter location and abundance information from research reports (see e.g. Lance et al. 2004, Jeffries and Jameson 2014) to examine shifts in otter abundance and distribution over the past several decades. Sea otter surveys along the Olympic Coast have been conducted by a mix of aerial surveys and land-based observations since 1977. Surveys were approximately biennial through the 1980s (data in 1977-9, 1981, 1983, 1984, and 1987), and annual from 1989-2015 (but no surveys in 2009 or 2014). Sea otter surveys were conducted in summer and thus reflect summer distribution and abundance (Laidre et al. 2009). Available evidence does not suggest that summer and winter distributions of sea otters are substantially different in this region (Laidre et al. 2009). However, sea otters are highly mobile predators with substantial home ranges and information on seasonal patterns is notably uncertain.

To estimate trends in sea otter abundance at each focal site, we developed a kernel-smoothed distribution of otters along the coast to incorporate uncertainty about how snapshot surveys translate to effective numbers of otters present at a given location. We first developed a one-dimensional coastline for the Olympic Peninsula and identified the position of each WDNR survey location along this coastline. We generated a smooth density of otters along the coastline using kernel density estimates, which approximate the observed otter data using a mixture of Normal (Gaussian) distributions. Specifically, we placed a Gaussian distribution centered at each survey location with a standard deviation *h* (the bandwidth) that corresponds to the estimated sea otter home-range size of 40 km for the Washington coast (*h* = 10.2; Laidre et al. 2009, their Fig. 3). After calculating the smoothed kernel probability density, we calculated the proportion of the total sea otter population that was present within a radius of 10 km of each focal site by integrating the probability density and multiplying by the total sea otter population size to arrive at an estimate of sea otters at each site. Due to uncertainty in the effective home range size of sea otters, we performed sensitivity analyses using a range of bandwidths (*h* = 5 and 15). The qualitative pattern of results did not change with alternate bandwidths.

We estimated the temporal trend in sea otter abundance at each site and OCNMS-wide by regressing the natural logarithm of sea otter abundance against time. We performed this analysis on the time-series beginning with the first year kelp canopy survey (1989-2015, see below), and separately for the two halves of the time-series (1989-2001 and 2002-2015) to assess if trends shifted over time. As estimates of trends become progressively less precise with less data, we elected not to further subdivide the time series. To facilitate comparison among sites that vary substantially in the sea otter abundance, we constructed a log-index of sea otter abundance. We standardized the number of sea otters by dividing by the average number of sea otters estimated at each site during the first three years of the kelp surveys (1989-91;) and taking a natural logarithm of this ratio:. Using such an index allows for sites across a large range of abundances to be visualized on the same axes and provides a graphical interpretation of sea otter trend (linear trends are exponential changes in abundance). We explored alternate years for dividing the time-series as well, setting the break variously at 1999 to 2003; these breaks made little qualitative change to the results.

*Kelp canopy area*

To describe kelp abundance at each site, we used publicly available data from aerial overflight surveys of algae from the Washington Department of Natural Resources (WADNR 2014; WADNR 2017, survey methods described in Van Wagenen 2015 (Pfister et al. 2017)). Surveys were conducted annually between 1989 and 2015 (no data available for 1993) during peak kelp abundance for the region (late July or early August). Kelp canopies in this region consist of a mix of *Macrocystis* and *Nereocystis*. While overflight surveys differentiate between the two species, we are primarily interested in the total canopy habitat provided, and thus we focus on the total surface coverage provided by the two species; additionally, the two species’ abundances are strongly positively correlated in this region (Pearson’s *r = 0.689;* (Pfister et al. 2017)). We examined kelp abundance at two scales. First, we used kelp area within discrete area strata along the coast to provide estimates of local kelp surface coverage, , for the strata containing each of our ten sites, *j*, in each year, *t* (see Fig. 1). The strata (kelp index map regions) used by WADNR are substantially larger than the area surveyed during invertebrate surveys (see below). Unfortunately, these strata are the smallest spatial unit for which it is appropriate to generate kelp area estimates (WADNR 2017). Second, we summed kelp surface coverage in all strata between Neah Bay and Destruction Island to provide an OCNMS-wide estimate of kelp area (Fig. 1).

We estimated the temporal trend in kelp canopy coverage at each site and OCNMS-wide by regressing the natural logarithm of kelp area against time. We also calculated the standard deviation (SD) and coefficient of variation (CV = SD / mean) of observations around each trend. Thus, our measures of CV represent variability after accounting for the overall trend in area. We performed this analysis on the entire time series (1989-2015), and separately for the two halves of the time-series (1989-2001 and 2002-2015) in order to determine if trends shifted over time. As with sea otter data, to facilitate comparison among sites that vary substantially in kelp area, we constructed a log-index of kelp area; we standardized the area of kelp by dividing the average kelp area observed during the first three years of the survey (1989-91)*,* , for site *j* and taking the natural logarithm of the ratio:.

*Invertebrate SCUBA surveys*

We conducted SCUBA surveys between 3-7 August 2015, and gathered historical survey information collected by Kvitek and colleagues in 1987, 1995, and 1999 (Kvitek et al. 1989, 1998, 2000). During 2015, SCUBA divers surveyed benthic communities in kelp beds at each site (Fig. 1) at depths between 5-10 m, along visual transects (30 m x 2 m, *n* = 4 transects per site). On each transect, divers counted large, non-cryptic invertebrates >5 cm diameter (sea urchins, sea stars, sea cucumbers, crabs, bivalves, gastropods, anemones, chitons, tunicates, etc.).

For the 1987, 1995, and 1999 subtidal surveys, we extracted summary statistics on benthic invertebrate densities from Kvitek and colleagues (Kvitek et al. 1989, Kvitek et al. 2000). (Kvitek et al. 1989). We include surveys that occurred at the same sites and comparable depths (5-10m). All surveys use standard quadrat and transect sampling methods, though the sample sizes vary among years (Table 1, Table S1). For the sake of comparison, we converted data from all subtidal surveys into units of countsm-2. Not all sites were sampled in each year, and some taxonomic groups of interest were not identified in available reports (e.g., sea stars were not listed in the results for 1995, gastropod densities were only available for 1987 and 2015). We used all available data for each site and year. When necessary, we combined quadrat and transect data using a weighted average with weights corresponding to the area surveyed by each type. We include only species that are large and readily identifiable to avoid concerns about among-diver variation in detection of cryptic species (e.g., chitons; class *Polyplacophora*). We focus on the time-series of abundance for six species groups that are common, important members of the Olympic coast nearshore invertebrate community: sea urchins (genera *Mesocentrotus* and *Strongylocentrotus*), sea cucumbers (genera *Cucumaria* and *Parastichopus)*, crab (primarily genera *Pugettia* and *Cancer)*, bivalves (primarily rock scallops, *Crassadoma gigantea*), and sea stars (including genera *Pisaster, Orthasterias, Dermasterias, Henricia, Pycnopodia)*. Consistent with previous research, we identified sea urchins as the dominant invertebrate grazer in this system and contrast the trends in sea urchin abundance with the other invertebrate groups. Based on sea otter diet information provided by Jesse, we classified these groups based on their frequency of occurrence in otter diets (Walker et al. 2008). We identified sea urchins and scallops (bivalves)as common prey items (>5% of observed successful sea otter foraging dives); crabs as occasional prey items (1%-5% of foraging dives); chitons, sea stars, and sea cucumbers as rare prey items (<1%); and, anemones, tunicates, and nudibranchs as non-prey items (not observed).

*Statistical Analyses*

To ask if local changes in sea otter abundance resulted in subsequent changes in kelp area among the 10 focal sites, we regressed exponential trends in sea otter abundance against kelp area. We performed this analysis for the entire time-series (1989-2015) and separately for each half of the study period (1989-2001 and 2002-2015), using region and otter growth rate as fixed effects. In the model with two time periods we allowed for a period otter growth rate interaction to ask if the relationship between sea otters and kelp shifted between periods. We also included our measure of wave exposure as a potential covariate in all models.

To assess the effect of otter abundance on the variability of kelp cover, we used the difference in CV of kelp area at each site between periods as the response variable with region, and wave exposure, difference in otter abundance between periods at each site, and CV of kelp area in the 1989-2001 period as predictors. We explored only additive main effects due to a sample size of 10 and selected among models using AIC corrected for small sample sizes (AICc).

To examine changes in the abundance of sea urchins, bivalves, crabs, sea stars, and sea cucumbers over time, we used permutation-based multivariate analysis of variance (PERMANOVA) to compare community structure across three time periods (1987, 1999, 2015) or three regions (northern, central, and southern) using the adonis function in R. We exclude data from 1995 because sea star data were absent. The taxa-specific average densities (individuals m-2) for each site-year-region were used as the dependent variables, and converted to dissimilarity matrices using Manhattan log(*x* + 1) distances. We used Manhattan dissimilarity measures because we viewed joint absences of species as informative and wanted to include that information. The more commonly used Bray-Curtis dissimilarity excluded information about joint absences (Legendre et al. 2005)(REF… Andersen et al… JAMEAL A LITTLE HELP?). We performed randomizations within strata based on regions or time periods. We also tested whether community composition was more variable in some regions than others or in some time periods than others by examining multivariate dispersion in community composition using the betadisper function in R. To visualize differences among time periods or regions in invertebrate community structure, we used non-metric multidimensional scaling (nMDS) based on the nmds function and plotted vectors explaining how variation in the densities of individual taxa related to community dissimilarity using the envfit function. Because information about gastropod densities was not collected at some sites in 1999, we repeated all of the above analyses for 1987 and 2015 data only to determine if doing so modified our inferences about changes in the mean or variability in community composition. All multivariate analyses and visualizations were conducted in the R package vegan. We also calculated proportional declines in mean abundance and used paired *t*-tests to evaluate their significance.

**Results**

*Spatiotemporal trends of sea otters and kelp*

Sea otter density trends have followed three spatially distinct patterns along the Olympic Coast since the 1970s (Fig. 2a - c). In general, local trends in sea otters differ substantially from the OCNMS-wide trend. Near the most northerly study sites, sea otter densities increased sharply from the mid-1980s until the early 1990s before declining slightly and then remaining stable from the mid-1990s to present (Fig. 2a). Sea otter densities in the Central region of the study area including Anderson Point, Point of the Arches and Cape Alava experienced exponential growth from the late 1970s until the mid-1990s, but have remained largely stable at densities just above those observed in 1990 (Fig. 2b). This represents a longer period of increasing otter densities than the northern region. The increase in sea otter density has been strongest and most consistent in the southern region of the study area (Fig. 2c). Sea otter densities near the southern sites have increased exponentially since the late 1970s; since roughly 2000, the rate of increase in the Destruction Island area has outpaced rates of increase near Teahwhit Head and Cape Johnson / Rock 305. At present, the absolute abundance of sea otters is also greatest in the southern region; sea otter abundances in the northern region are the lowest by at least an order of magnitude (estimated 2015 sea otter abundance of 18, 207, and 439 for Tatoosh Island (Northern region), Cape Alava (Central), and Destruction Island (Southern) respectively). Cape Johnson and Rock 305 have essentially the same trend in Fig. 2c due to their proximity relative to the kernel bandwidth used for home range estimation (Fig. 1; see Methods).

Further analysis of sea otter data shows that the distribution of the Olympic Coast population has shifted over time (see also Jefferies and Jameson 2014). The population has had a bimodal or multimodal distribution for much of the study period, with the most significant modes in the area between Cape Alava and Cape Johnson, and another further south near Destruction Island (Fig. 3). The center of gravity of the population was in the vicinity of Teahwhit Head in the late 1970s, but then shifted north to the area around Cape Alava for much of the 1980s and 1990s. Starting in the late 1990s, the center of gravity rapidly shifted south to near Destruction Island, where it has remained. In recent years sea otter observations are rare inside the Strait of Juan de Fuca (Fig. 3, above dashed line) but common at most points to Point Grenville in the south (Fig. 3).

Canopy kelp area exhibited spatiotemporally distinct patterns in the three regions of the study area from 1989-2015 (Fig. 2d-f). Kelp area showed substantial interannual variation both at the individual sites and the OCNMS-wide scale (Fig. 1; see also (Pfister et al. 2017)). While the area of kelp in absolute terms varied substantially among sites within a region (Table 3; Pfister et al. 2017), kelp trends varied predominantly by region along the Olympic Coast. At the furthest north sites, kelp area indices showed no clear long-term trends but displayed notably higher interannual variability at Tatoosh Island than Neah Bay and Chibadehl Rocks inside the Strait of Juan de Fuca (Fig. 2d; note that Neah Bay and Chibadehl Rocks are in the same kelp monitoring stratum (Fig. 1), and thus share a single kelp time series). The central region also showed within-region differences among sites (Fig. 2e). Canopy area at Cape Alava increased from 1989 to 2000 before stabilizing and possibly declining in recent years, while Point of the Arches and Anderson Point decreased in the early 1990s before following a qualitative pattern similar to Cape Alava. The index of canopy area at Cape Alava was far less variable than the other two central sites. At the southern sites, canopy area generally increased until the early 2000s before stabilizing or declining slightly (Fig. 2f). As with the central region (Fig. 2e) there were some differences in the signs of short-term trends across the four southern sites early in the time series, although the degree of interannual variability was fairly consistent across the sites.

*Connections between sea otters, invertebrates, and kelp*

We detected differences in the kelp population growth rate between periods with a OCNMS-wide average growth rate of 0.056 for 2002-2015 and of -0.037 for the 1989-2001 period (difference between periods *p* =0.012; Fig. 4). While the temporal effect is intriguing, more interesting is the interaction between sea otter population growth rate and time period. Sea otter growth had a positive relationship with kelp during the earlier period (point estimate of slope = 0.285) but a negative effect of sea otters during the later period (point estimate of slope = -0.507; interaction term *p* = 0.045). There was no support for regional variation in kelp population growth after accounting for other factors (*p* = 0.128). The model considering only a single time period found no effect of sea otters on kelp (*p =* 0.40) but differences in kelp population growth rate among regions (*p =* 0.024*).* This result shows how the temporal context can substantively alter the interpretation of mechanisms driving kelp population growth. Importantly, our analyses should not be taken to suggest a discrete break in the population time-series in 2001 – using alternate breakpoints in the time series between 1999 and 2003 yield qualitatively similar results – rather, dividing the time series is a convenient way of summarizing changes in a continuous time series (see Fig. 2).

After accounting for kelp population growth rates, the variability in kelp area declined at most sites between the two time periods (Fig. 5). Specifically, bootstrapped estimates of CV showed that variability at all sites but one (Tatoosh Island) declined, though the magnitude of decline varied substantially by region. The three northern sites had virtually no change in CV (changes of less than 0.05 for all sites), the central region showed declines in CV but with variability among sites (declines of 0.033, 0.343, and 0.351, for Cape Alava, Anderson Point, and Point of the Arches, respectively), while the southern sites showed substantial declines in CV (declines of 0.175 to 0.694). For most sites, these are large and biologically significant changes in kelp variability. Linear models showed that including kelp CV in 1989-2001 alone best predicted the change in CV between periods ( *adj. R2* = 0.54). Sites with high CV in the first period showed reduced CV in the second. The only other model with a small amount of statistical support included both kelp CV in 1989-2001 and the change in the number of otters ( *adj. R2* = 0.64). In this model, the coefficient for the change in otters was negative, indicating increased sea otter abundance was associated with reduced kelp CV (point estimates correspond to an increase of approximately 13 otters leading to a decrease of 0.01 in CV). As we only have 10 sites for comparison, our statistical power and precision of these estimates is low. Estimates of wave exposure were not a significant explanatory variable for any aspect of kelp CV.

As both primary sea otter prey and the major grazers of kelp, benthic invertebrates are the mechanistic link between sea otters and kelp. While we lack continuous time series for invertebrates at OCNMS, available information shows significant variation in the benthic invertebrate community over years but not across regions (Fig. 6; Table 2). Not only was there a shift in mean community composition between 1987 and the two later survey years, but community composition among sites became less variable after 1987 (Table 3). The differences in community composition among years reflected substantial declines in all 5 major taxonomic groups from 1987-2015 (Fig. 7). Sea urchins declined precipitously with the across-site mean density falling by more than 99% between 1987 to 2015 (from 3.7 m-2 to 0.01 m-2). While the other five species groups did not decline as dramatically as urchins, they all showed substantial declines from 1987 to 2015: bivalves (decline of 90%), sea cucumbers (86%), crabs (84%), and sea stars (70%). All of these declines were significant (paired t-tests, *p*<0.01 for all species groups). Only sea urchins showed a pattern in which the highest density occurred in the three sites that Kvitek et al. (1989) defined as outside of the range of sea otters (Neah Bay, Anderson Point, Point of the Arches; Fig. 7a). For the four other species groups, densities were not notably different between sites inside and outside of the otter range in 1987. This suggests that the dramatic and immediate effect of otters on invertebrates is limited to a few species or species groups, even if over time there are substantial but gradual changes in invertebrate communities. Beyond declines in mean densities, all five species also show notable declines in the among-site variation in density; the among-site standard deviation fell by 75 to 99% for our six species groups. By all measures, the spatial variability in invertebrate densities has declined over the past 30 years.

**Discussion**

Sea otters are iconic keystone predators in coastal ecosystems of the northeastern Pacific, and their presence radically affects invertebrate and algal communities (Watson & Estes 2011) Estes and Duggins 1995, Steneck et al. 2002) (Breen et al. 1982). Here we revisit a series of historical invertebrate surveys and complement these surveys with independent spatiotemporal data on kelp and sea otters in Washington’s OCNMS since 1987 along with new *in situ* invertebrate surveys. Our analyses reveal a strong correlation between sea otter and kelp population growth at the local scale during the rapid expansion of sea otter populations. However, there is a temporal component to these associations: the relationship between kelp and sea otter growth rates shifted from positive during the 1990s to neutral or possibly slightly negative post-2000 (Fig. 4). Together our analyses demonstrate that OCNMS-wide trends in sea otter, kelp, and benthic invertebrate abundance are not necessarily evident at a finer spatial grain. They suggest that while a sea otter derived trophic cascade initially drove changes in the nearshore community along the Washington coast, additional factors in more recent years may account for fundamental shifts in invertebrate community dynamics. In addition, contrary to predictions from trophic cascade theory (Estes et al. 2011), kelp and sea otter abundance are statistically decoupled when viewed at an OCNMS wide scale and over the entirety of the 30-year period. The decoupling of otter and kelp abundance in more recent years, and the disconnect between OCNMS-wide and local scale spatial patterns, may provide insight into appropriate theoretical expectations for short- vs long-term dynamics following the reintroduction of predators more generally (Sergio et al. 2014, Stier et al. 2016), and sparks intriguing hypotheses about the relative influence of top-down and bottom-up forcing factors in temperate coastal habitats.

Classic trophic cascade theory predicts that increased sea otter abundance should reduce the abundance of their prey, including functionally important invertebrate grazers such as sea urchins. In turn, reduced grazer abundance should release kelp and other algae from top-down control and lead to increases in their abundance. Our results echo those of Kvitek et al. (1989, 1998, 2000) and demonstrate large, immediate, and persistent effects of sea otter expansion on the main invertebrate kelp-grazer and a preferred prey, sea urchins. We also show that increases in otter abundance were correlated with declines for a broad suite of invertebrate species and multi-year increases in kelp area. These reductions in invertebrate abundances suggest that the consequences of sea otter populations for kelp forest community composition are not exclusively an immediate shift in state, but can manifest gradually over the span of decades (Watson & Estes 2011, Kenner et al. 2013). Furthermore, because invertebrate densities remain far below historical levels and those associated with shifts to non-kelp dominated states (Filbee-Dexter & Scheibling 2014), the observed declines in kelp population growth rates (Fig. 4) and total area (Fig.1) in OCNMS since 2000 are unlikely to be caused by increased invertebrate grazing pressure. Rather, these results suggest that forces unrelated to otter abundance drive kelp abundance and kelp forest community structure once kelp have equilibrated to the presence of sea otters.

One hypothesis for the decoupling of kelp and otter growth rates after 2001 is an increase in the influence of abiotic factors. Oceanographic dynamics in the late 1990s and early 2000s in the Northeast Pacific have been the subject of intense study because they were characterized by major El Niño and La Niña events and shifts in productivity regimes (PDO; (Mantua et al. 1997, Mantua & Hare 2002)). It is possible that at that time the vast majority of changes in invertebrate grazer abundance due to otter predation had already occurred, and kelp dynamics became more strongly influenced by sea surface temperature, upwelling, nutrient availability and other bottom-up forces (Pfister et al. 2017). For kelp in particular, we expected that the variability in kelp area would be strongly related to wave exposure at a given site (Reed et al. 2011, Bell et al. 2015). However, while kelp CV varied substantially among sites, it was unrelated to calculated wave exposure values (Fig. S1). Surprisingly, Neah Bay, a site largely protected from wave exposure (Fig. S1), has had nearly equivalent kelp CVs since 2002 as five sites on the outer coast, including Cape Alava and Destruction Island (Fig. 5; detrended CV of approximately 0.2). We speculate that this may be driven predominantly by the fact that we only have information about kelp area during the summer, when seasonal wave energy on the coast is relatively low. An alternative prediction is that kelp CV in Neah Bay would decline if sea otters invaded Neah Bay, as kelp CV appears to be related to the number of otters present at each site (Fig. 5). While we cannot definitively identify what caused the decoupling of otter and kelp dynamics in OCNMS, shifts in factors controlling alternate states within ecological communities are not without precedent in other systems (Bellwood et al. 2006, Petraitis et al.). Our study adds richness and complexity to the classic trophic cascade explanation for the dynamics of kelp forest communities in the presence of sea otters.

Beyond the mean effects of sea otters on kelp and invertebrates, both kelp area and the benthic invertebrate community showed reduced variability as sea otter abundance increased. We know of no other study that has examined the interaction between otter abundance and kelp variability, but suggest this is a particularly interesting avenue of future research due to the connection between canopy kelps, benthic community structure (Arkema et al. 2009), and various ecosystem services (Wilmers et al. 2012, Pinsky & Fogarty 2012). Indeed, incorporating the effects of keystone species and ecosystem engineers in marine planning seems especially wise when ecological endpoints include conservation and restoration of biogenic habitats such as kelp forests. For benthic invertebrates, both multivariate and univariate analyses show that invertebrate communities cluster by year rather than spatial region, suggesting the primary driver of communities is a temporal rather than a spatial process (Fig. 6, 7). Thus both kelp and invertebrates show evidence of homogenization in concert with the expansion of sea otters, which aligns with previous suggestions that sea urchin-dominated habitats may show more variability than sea otter controlled habitats (Watson & Estes 2011). We acknowledge that there are potentially many consequences for other groups including understory algae (Watson & Estes 2011) and fish (Markel & Shurin 2015) that we cannot explore here.

Our data does not allow us to examine year to year changes in the linkages between sea otters, benthic invertebrates, and kelp, opening the possibility that invertebrate communities substantially shifted during the years between surveys in a way that can explain kelp variation. While there is ample evidence that other factors have affected the abundance of some invertebrate groups (e.g. sea star wasting disease outbreak in 2013-4; Eisenlord et al. 2016), personal observations of one of the authors (AOS) between 2004 and 2008 at two sites - Tatoosh Island and Point of the Arches - do not support radical changes in invertebrate abundances during the 1999 to 2015 gap in our invertebrate time-series. We cannot exclude the possibility of strong variability in invertebrate communities driving these patterns but we suggest that it is an unlikely driver of observed kelp patterns.

Looking to the future, sea otter numbers appear to have stabilized in much of the northern and central regions of OCNMS (Fig. 2) and may be at or near carrying capacity in this region. Invertebrate densities in these areas are very low, which begs the question of how sea otter populations are maintained with very low prey abundances. Rocky subtidal and kelp forest habitats support higher densities of sea otters in Washington than either sandy bottom or estuarine habitats (Laidre et al. 2001, 2002), suggesting that sea otters are not likely to simply be foraging in nearby sandy habitats, but they may be capturing invertebrates in deeper rocky habitats that were not included in our surveys or historical surveys. Furthermore, we note that the abundance of sea otters in 2015 (*n* > 1400) is substantially above the published estimates of carrying capacity for the Washington outer coast (922 to 1189; Laidre et al. 2002), suggesting that either the carrying capacity of otters needs to be revisited or, if carrying capacity estimates are correct, the population is predicted to decline in the coming years.

In conclusion, place-based management of resources and ecosystem services is of great importance in coastal regions, and informed, holistic management requires accounting for the dynamics of keystone species and major biogenic habitats. Along the Olympic Coast, place-based management is a priority at both the state level—for example, the recently drafted Washington state marine spatial plan (http://www.msp.wa.gov/wp-content/uploads/2017/draft\_MSP\_and\_appendices.pdf)—and the federal level, as practiced by OCNMS (Office of National Marine Sanctuaries 2008) and the adjacent Olympic National Park (National Park Service 2008). The reestablishment of a healthy sea otter population in this region has already yielded considerable ecosystem change, through trophic cascade dynamics that have enabled kelp canopy habitats to expand. However, our research shows that kelp canopy dynamics are now being influenced predominantly by drivers other than otter abundance. This apparent decoupling poses a challenge to spatial management of marine resources in the area because the new prevalent mechanisms must be identified in order to anticipate further change and understand how management actions interact with natural variation. A second challenge may be in revising management objectives for sea otters, which have been prioritized as keystone species that have major impacts on ecosystem structure and functioning, biodiversity, and other attributes (Watson & Estes 2011, Wilmers et al. 2012). Our work suggests that their keystone effect on kelp forests on the Olympic Coast has diminished over time, and also that they must be relying on prey in habitats other than kelp forests in order to maintain their population at its present level. The current ecological dynamics of sea otters at different sites along the Olympic Coast must be better understood in order to determine if their tremendous predatory demands are impacting other habitats and potentially introducing new management tradeoffs in habitats beyond the shallow kelp forest studied here.

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