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**Multiscale impacts of armoring on Salish Sea shorelines:
Evidence for cumulative and threshold effects**

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27 **Abstract**

28 Shoreline armoring is widespread in many parts of the protected inland waters of the
29 Pacific Northwest, U.S.A, but impacts on physical and biological features of local nearshore
30 ecosystems have only recently begun to be documented. Armoring marine shorelines can alter
31 natural processes at multiple spatial and temporal scales; some, such as starving the beach of
32 sediments by blocking input from upland bluffs may take decades to become visible, while
33 others such as placement loss of armoring construction are immediate. We quantified a range of
34 geomorphic and biological parameters at paired, nearby armored and unarmored beaches
35 throughout the inland waters of Washington State to test what conditions and parameters are
36 associated with armoring. We gathered identical datasets at a total of 65 pairs of beaches: 6 in
37 South Puget Sound, 23 in Central Puget Sound, and 36 pairs North of Puget Sound proper. At
38 this broad scale, demonstrating differences attributable to armoring is challenging given the high
39 natural variability in measured parameters among beaches and regions. However, we found that
40 armoring was consistently associated with reductions in beach width, riparian vegetation,
41 numbers of accumulated logs, and amounts and types of beach wrack and associated
42 invertebrates. Armoring-related patterns at lower beach elevations (further vertically from
43 armoring) were progressively harder to detect. For some parameters, such as accumulated logs,
44 there was a distinct threshold in armoring elevation that was associated with increased impacts.
45 This large dataset for the first time allowed us to identify cumulative impacts that appear when
46 increasing proportions of shorelines are armored. At large spatial and temporal scales, armoring
47 much of a sediment drift cell may result in reduction of the finer grain-size fractions on beaches,
48 including those used by spawning forage fish. Overall we have shown that local impacts of
49 shoreline armoring can scale-up to have cumulative and threshold effects -- these should be
50 considered when managing impacts to public resources along the coast.

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52

53 **1. Introduction**

54 Anthropogenic alteration of shorelines is a worldwide phenomenon as a significant
55 proportion of population growth is in coastal communities. Types of shoreline development are
56 diverse, ranging from simply building houses overlooking the water to completely altering the
57 shore by covering it with fill or structures. The Salish Sea, which includes all the inland marine
58 waters of British Columbia (Canada) and of Washington State (USA), has shorelines that range
59 from virtually pristine beaches to concrete-covered commercial ports. In the face of increasing
60 coastal urban growth and sea level rise, effective management of our shorelines requires
61 understanding both functions of natural beaches and the scales at which we are impacting them
62 (Arkema et al., 2013; Harris et al., 2015).

63 One of the most prevalent forms of coastal development in the Salish Sea and worldwide
64 is shoreline armoring, comprising various artificial means of stabilizing banks and bluffs that
65 might otherwise erode and endanger infrastructure. A recent conservative estimate of armored
66 shoreline in the continental US is 14% (Gittman et al., 2015). Local, mostly biological, effects of
67 shoreline armoring are well known for some types of embayments and marshes (e.g., Bozek and
68 Burdick, 2005; Chapman and Underwood, 2011) and open-coast sandy beaches (e.g., Dugan et
69 al., 2008; review by Nordstrom, 2014), and recently for the gravel-sand beaches of Puget Sound
70 (Sobocinski et al., 2010; Heerhartz et al., 2014). Armoring locally reduces retention of logs and
71 wrack (algae, seagrass, leaf litter, and other organic and inorganic debris left by ebbing tides) and
72 the invertebrate communities that inhabit this detritus. It can also have indirect effects on seabird
73 and shorebird use (Dugan et al., 2008) as well as abundance and diversity of large mobile
74 invertebrates (Chapman, 2003). Potential spawning locations for beach-spawning forage fish,
75 such as surf smelt (*Hypomesus pretiosus*), are reduced when armoring covers the high shore, and
76 egg mortality increases when beach temperatures are raised by shoreline modifications (Rice,
77 2006). These trophically important fish may also be negatively impacted in cases where
78 armoring coarsens the sediment due to local winnowing of finer grain sizes (Penttila, 2007;
79 Quinn et al., 2012; Fox et al., 2015; Greene et al., 2015). By changing the nearshore habitats
80 encountered by juvenile migrating salmon, armoring affects their diets (Munsch et al., 2015) and
81 possibly residence time (Heerhartz and Toft, 2015).

82 Considerable study of physical impacts of armoring on beaches has been conducted,
83 although the results are contradictory. In some circumstances, interactions of sediment

84 impoundment, wave reflection, and alterations to nearshore water currents may alter beach scour,
85 mobilization of sediment, and recovery from storms. In theory, these processes may result in
86 narrower, steeper, and coarser-grained beaches (Pilkey and Wright, 1988; Bozek and Burdick,
87 2005; Nordstrom, 2014). One clear effect is that passive erosion (e.g., caused by relative sea
88 level rise) causes narrowing of armored shorelines because the upper beach is prevented from
89 migrating inland. In contrast, whether active erosion is induced by seawalls is still argued
90 (reviews by Kraus and McDougal, 1996; Ruggiero, 2010); few long-term studies have been
91 attempted but generally do not show a definitive armoring effect (e.g., Griggs et al., 1994;
92 Griggs, 2010). Modeling work (e.g., Ruggiero, 2010) suggests that contradictions seen in the
93 literature may stem from variation among study systems in key physical parameters, in particular
94 the relative elevation of the seawall and the morphology of the beach and nearshore, including
95 their slopes.

96 Even for the more consistent biological impacts of armoring, translating local effects to a
97 landscape scale is challenging because of the myriad other natural and anthropogenic factors that
98 affect shoreline processes. The signal to noise problem is particularly large in inland waters such
99 as the Salish Sea because of the complexities of underlying geology, shoreline shape, freshwater
100 input, wave fetch, orientation to prevailing winds, nearshore bathymetry, and sources of
101 sediments, vegetation, and organisms. In most of the world, beach sediments derive
102 predominantly from rivers. On sandy shorelines, these sediments are jealously retained with
103 groins, and millions of dollars are spent annually to replenish beaches where natural sources
104 have been locked up by dams (Berry et al., 2013). Although numerous rivers empty into the
105 Salish Sea and a few of them create large deltas, much of the riverine sediment is deposited in
106 deep fjord-like basins rather than building beaches. Instead, most beach-building sediment comes
107 from erosion of bluffs (Keuler, 1988). It follows that “locking up” these sediments by armoring
108 shorelines should have large-scale and long-term impacts, including cumulative effects if few
109 sediment sources are left unaltered (reviewed by Berry et al., 2013; Nordstrom, 2014). However,
110 demonstrating cumulative effects, e.g. changes that continue to worsen with additional armoring,
111 is notoriously difficult -- especially if changes appear gradually, as is likely with many
112 geomorphic processes. In Europe, extensive coastal armoring is thought to have contributed to
113 broad-scale steepening of the shoreline (Taylor et al., 2004), but many other processes could be
114 important.

115 In the southern part of the Salish Sea (in Washington State), which includes Puget Sound,
116 extensive shoreline armoring has accompanied the last 100 years of development along the
117 greater Everett-Seattle-Tacoma urban corridor, and is thought to significantly impair nearshore
118 ecosystem processes (Simenstad et al., 2011). While local effects have recently been documented
119 (e.g., Sobocinski et al., 2010; Heerhartz et al., 2014), broader or cumulative impacts have not.
120 This uncertainty stymies managers and regulators who lack compelling data that would provide
121 the “best available science” to inform guidelines. Pressures to relax armoring regulations stem
122 from the need to protect valuable infrastructure from erosion, especially with risk exacerbated by
123 sea level rise. Sociological studies show that decisions by a few homeowners to armor their
124 shoreline often triggers neighbors to do the same, leading to cascading local impacts (Scyphers et
125 al., 2015). In addition to such possible cumulative effects, regulators are particularly interested in
126 which types or locations of armoring have greater impacts than others, and whether there are
127 thresholds that trigger these impacts. Samhoury et al. (2010) define an ecological threshold as a
128 point at which small changes in environmental conditions produce large (non-linear) responses in
129 ecosystem state. For example, ecological thresholds have been associated with habitat
130 fragmentation (e.g., Andr n, 1994) and edge effects (Toms and Lesperance, 2003). One possible
131 threshold that may apply to shoreline armoring is the extent that structures encroach on the
132 beach. In addition, slow and delayed “latent impacts” (Coverdale et al., 2013) may exist but are
133 very difficult to detect, especially given signal-to-noise problems.

134 Previous studies by our research team have focused on local impacts of shoreline
135 armoring in central and southern Puget Sound (Heerhartz et al., 2014 and 2015). We dealt with
136 among-site ‘noise’ by use of a paired sampling design, focusing our surveys on nearby,
137 physically-paired, armored and unarmored beaches. Here we broaden our geographic scale to test
138 whether the documented biological effects of armoring exist on beaches in the Salish Sea north
139 of Puget Sound. We also test whether any physical impacts are detectible, because our previous
140 work in central and southern Puget Sound found few differences in quantified physical
141 parameters that were correlated with armoring. The northern region has more bedrock shorelines
142 and different oceanographic characteristics, so we anticipated that there would be some regional
143 differences in beach parameters. Based on our own localized studies and on literature from other
144 systems (e.g., open-coast beaches), we hypothesized that: 1) Armoring-associated reduction of
145 logs, wrack, and invertebrates would be consistent across regions in paired-beach analyses; 2)

146 These associations would be increasingly clear when armoring is lower on the beach face; 3) By
147 examining a large range of sites, the predicted pattern of armoring altering beach slope and
148 sediment coarseness might be detectible; and 4) Such geomorphic signals would be most distinct
149 where extensive stretches of armoring have “locked up” more sediment sources in an area. To
150 address these questions, we discuss regional patterns but ignore the huge beach-to-beach
151 variation in geomorphic conditions, to be discussed elsewhere (A.N. McBride, pers. comm.).

152

153 **2. Methods**

154 *2.1 Sites*

155 Our analyses include data from 65 pairs of armored and unarmored beaches in the inside
156 marine waters of Washington State, from the southern extent of Puget Sound to the Canadian
157 border (Figure 1). The data thus encompass three oceanographic regions: South (6 site pairs),
158 landward of a sill at the Tacoma Narrows; Central (23 site pairs), inside Puget Sound proper,
159 south of a sill at Admiralty Inlet; and North (36 site pairs), outside of the Sound but within the
160 Salish Sea. The south sites and to a lesser extent the central ones are influenced by constrained
161 water exchange caused by the sills, and by freshwater input from several rivers. The north sites
162 have greater oceanic flushing but have substantial seasonal freshwater input from several large
163 rivers, especially the Fraser in Canada and the Skagit in Washington. The primary sediment
164 composition on our study beaches was a mix of sand and gravel predominantly derived from
165 glacial and interglacial deposits, delivered to beaches via episodic bluff erosion, and distributed
166 by longshore transport (Shipman, 2010). Wave energy regime and local geology are then the
167 primary drivers of beach sediment character and gradient in the Salish Sea. Pairs of beaches were
168 within the same drift cell (independent zone of littoral sediment transport from source to
169 deposition area) and same component of that drift cell (erosional or depositional). The 65 pairs
170 were within 49 different drift cells (out of over 600 in the Washington state portion of the Salish
171 Sea). These 49 cells ranged from 1.8 to 60.4 km long, and varied from 0 to 99% armored.

172 Sites had armoring at different elevations and of different types (e.g., concrete seawalls,
173 stone riprap, retaining walls of wood pilings). Paired beaches were matched as closely as
174 possible in terms of geomorphic setting and geology of the bluff, aspect to prevailing winds and
175 sun, wave exposure, and nearshore bathymetry. Beaches in a pair were always nearby; mean
176 distance between members of a pair was 383 m, maximum distance was 1 km. All field data

177 reported here were collected in summer (June to Aug.); central and south sites were surveyed in
178 2010-2012, north sites in 2012-2013.

179

180 *2.2 Biological Surveys*

181 Data collection followed procedures described in Heerhartz et al. (2014). Briefly, at all
182 sites we placed a 50 meter shore-parallel transect high on the shore near the wrack zone; this line
183 was used for both biological and sediment sampling. We define beach wrack as organic matter
184 consisting of detached and stranded algae, seagrass, and terrestrial debris. We surveyed the most
185 recent line of beach wrack and avoided older and usually more desiccated wrack. Armored
186 beaches lacking wrack and logs were surveyed at the highest elevation where natural beach
187 sediments were present (i.e., at the toe of armoring). At 10 randomly selected points we
188 estimated the percent cover of each type of wrack (i.e., seagrass, algae, or terrestrial-source), and
189 noted the most abundant types of algae. At 5 of these points we collected samples of wrack and
190 the top 2.5 cm of sediment using a 15-cm diameter benthic corer, and quantified the number of
191 logs (less or greater than 2 m length). We also measured the width of the log line perpendicular
192 to shore. In the lab, wrack samples were sorted into types, dried, and weighed. All invertebrates
193 were extracted (using 106 micron sieves) from the wrack, and identified and counted using a
194 dissecting microscope; talitrid “beach-hopper” amphipods and other crustaceans were identified
195 to genus, and other invertebrates to family (except oligochaetes, which were not identified
196 beyond class). Invertebrate-dense samples were split with a Folsom Plankton Splitter and
197 abundances were back-calculated. For analyses, all parameters were averaged (percent covers) or
198 summed (biomasses, invertebrate counts) across the transect (n = 5 for wrack core and log
199 samples, n = 10 for wrack percent covers).

200

201 *2.3 Geomorphic Survey Methods*

202 We characterized sediment grain sizes from the wrackline from three to five of the core
203 samples by sieving dried sediments smaller than 16 mm through progressively finer sieves (1/2
204 phi intervals) using a RoTap shaker, and weighing the amount retained in each sieve. Coarser
205 sediments (cobbles) were individually measured. Elevations of wracklines were measured;
206 because these differed within and among pairs, sediments were not all collected from the same
207 elevation on the beach. In addition, we assessed grain sizes, with lower precision, along a

208 transect at Mean Low Water (MLW: ca. +1 m above MLLW). At three randomly selected points
209 we used a 50×50 cm quadrat to estimate percent cover of cobbles (> 6 cm), pebbles (4 mm - 6
210 cm), granules (2-4 mm), sand (< 2 mm), and mud (smooth) at the surface and at 5 cm subsurface.
211 The two sets of estimates were averaged for per-quadrat proportions.

212 Beach profiles were obtained on low tides using a laser level and stadia rod or RTK-GPS,
213 measuring from the top of the berm or toe of the eroding bluff (on unarmored beaches) to
214 elevations approaching mean lower low water (MLLW), depending on the tide. On armored
215 beaches the profiles were measured from the lowest elevation on the armoring structure to
216 MLLW. Beach slope was calculated for the upper portion of each beach from the wrack line to
217 ~0.6 vertical meters above local MLW. This section was consistently in the active sediment
218 transport zone of the foreshore (an area of similar energy) of our beach transects. See
219 Supplementary Material for additional methods and data sources.

220 Due to the fjord-like shape and complex bathymetry of the Salish Sea, the magnitude of
221 the vertical tidal range varied greatly from our northern to southern sites. Mean tidal range varied
222 from 1.39 to 3.19 m, and the elevation of the mean higher high water (MHHW) datum varied
223 from 2.39 (in the north) to 4.32 m (far inside Puget Sound) above MLLW. To standardize our
224 elevation measurements in relation to tidal range and enable us to meaningfully assess impacts of
225 armoring emplaced at various elevations, we calculated a “relative encroachment” (RE) metric
226 by subtracting the elevation of armoring or toe of bluff from the MHHW datum for each beach.
227 Datum information for nearby sites was obtained from: <http://tidesandcurrents.noaa.gov/>; in
228 some cases it was necessary to interpolate between distant stations. Positive RE values indicated
229 that the toe (of armoring or bluff) was lower than MHHW, and negative values were higher. RE
230 at our study sites are reported in vertical feet, and ranged from -5.1 ft (= -1.55 m) to +7.0 ft (=
231 2.14 m), with a mean of -0.33 ± 0.16 ft SEM (standard error of the mean) (= - 0.10 m \pm 0.05
232 SEM).

233 We tested whether the proportion of the drift cell that was armored (hereafter referred to
234 as DCA: data from various sources) would generate cumulative armoring impacts, for example
235 by blocking increasing proportions of sediment sources. Variables that could be affected by
236 large-scale and long-term impacts of armoring might show these effects, including some
237 parameters where local and short-term impacts were not seen. Of particular interest was testing
238 our hypotheses of a correlation between sediment grain size or beach slope and DCA.

239

240 2.4 Statistical Analyses

241 We assessed local impacts of armoring using paired t-tests, taking advantage of our
242 sampling design to compare the differences between mean values of each measured response
243 parameter at each pair of beaches. Parameters tested are listed in Table 1.

244 We tested larger-scale effects of RE and DCA on response variables of interest using a
245 mixed effects model. For all analyses “Site” was defined as a random effect and RE or DCA as a
246 fixed effect. Each “Site” had two sampled beaches, the armored beach and its unarmored pair. In
247 this setting the model is allowed to vary the intercept for each “Site,” therefore accounting for
248 both within site and among site variation, i.e. acknowledging that sites are representative of
249 Salish Sea beaches and were randomly selected. For models testing counts of wrack invertebrates
250 (either summed, or separately for particular taxa) or components of wrack mass we used a
251 generalized mixed effects model with a quasi-Poisson distribution using the glmmPQL function
252 in the MASS package in R (Venables and Ripley, 2002; R Development Core Team, 2014). A
253 quasi-Poisson distribution was chosen over a Poisson distribution to account for overdispersion
254 and to adequately fit biological count data. For all model fits, residual plots and fitted values
255 were examined, and all appeared reasonable considering the inherent variability of the dataset.
256 For models testing the effect of RE or DCA on percent cover or proportion data (including wrack
257 cover and sediment grain sizes) we used a normal linear mixed effects model with “Site” as the
258 random effect on arcsine-square root transformation of the response variable, as is common with
259 such data to improve normality. The mixed effects models testing the effect of DCA all showed
260 high correlation between the fixed effect and the random effect of Site (Supplemental Table 1).
261 This was expected since each member of a Site existed in the same drift cell by design. Because
262 of the difficulty of deciding what constitutes an independent test, and lack of agreement in the
263 literature on adjusting alpha levels for multiple testing (e.g., Hurlbert and Lombardi, 2003,
264 2012), we present p values as reported by individual tests, and interpret our results
265 conservatively.

266 Some regression analyses showed non-linear changes in the response variable, suggesting
267 a threshold or breakpoint. For these we applied segmented (piecewise) regression to search for
268 statistically significant two-segment relationships; these can be common in ecological systems
269 and are characterized by an abrupt change in a response variable at some point (“threshold”) in

270 an independent variable (Toms and Lesperance, 2003; Samhouri et al., 2010). Our analyses used
271 an approach based on Crawley (2007) (see Supplementary Material). All univariate analyses
272 were run in R (R Development Core Team, 2014).

273 We used permutational multivariate analysis of variance (PRIMER v6 with
274 PERMANOVA+; Clarke and Gorley, 2006; Anderson et al., 2008) to test for differences in
275 sediment grain sizes between armored and unarmored beaches (type as fixed factor) with sites as
276 replicates (pair as random factor). Multivariate relationships between environmental predictor
277 variables and wrack sample invertebrate assemblages were investigated using distance-based
278 linear modeling (DISTLM) conducted using the step-wise selection procedure to minimize the
279 Akaike information criterion (AIC). These analyses partition the multivariate variability of the
280 invertebrate assemblages along best-fit axes and then test the environmental variables that are
281 most closely related to these axes.

282

283 **3. Results**

284 *3.1 Regional Differences*

285 Although we were interested in testing for armoring effects on beach parameters that
286 might exist despite regional variation, the physical backdrop for testing such local impacts
287 includes regional differences in bluff geology and shoreline geomorphology. There are
288 fundamental geologic differences among regions that result in variation in bluff material (Fig. 2).
289 The north region experienced advance and retreat of glaciers so that surface morphology reflects
290 the zone of ice grinding on bedrock; the exposed sediment in the central region transitions to a
291 glacial outwash zone; and the sediment deposits in the south are dominated by outwash that was
292 at the front edge of the ice. These influences are also seen in sediment grain sizes at the
293 wrackline of the study beaches (Fig. 2). Grain size distributions were quite consistent between
294 armored and unarmored beaches within a region, but some differed among regions; in particular,
295 very coarse gravel and very coarse sand were more abundant at the central sites (in the outwash
296 zone), medium sand was particularly abundant at the north sites, and coarse gravel in the south.
297 Sediments at MLW also showed no obvious armoring effect in the paired analysis but had some
298 regional differences, with pebbles and granules more abundant in the south (Suppl. Fig. 1).

299 Characteristics of the chosen sites varied among regions for some physical parameters but
300 not others. The large-scale parameter of wave fetch impacting each beach did not vary with

301 treatment but was lower in the south (Fig. 3). Drift cell lengths were greatest in the north and
302 shortest in the south, but with substantial variance within all regions (Suppl. Fig. 2). The DCA
303 (proportion of the drift cell armored: Suppl. Fig. 2) of the drift cells containing our study sites
304 were very different among regions, highest in the urbanized and heavily-armored central region
305 (mean $69\% \pm 5\%$ SE armored) and much lower in the north ($24 \pm 3\%$) and south ($25 \pm 15\%$).
306 Beach width was reduced consistently by armoring but showed no regional pattern (Fig. 3).
307 Elevation of the toe of the bluff/armoring showed both a treatment effect, with armoring moving
308 the toe to a lower elevation, but also a regional effect because of the much greater tidal range in
309 the south region. The toes of unarmored bluffs are much higher above MLLW in the south than
310 the north (Fig. 3) because of this factor. Our calculated relative encroachment metric (RE)
311 accounted for this background difference (see Methods). In all regions armored beaches
312 'encroached' upon Mean Higher High Water relative to unarmored beaches (Fig 3); this
313 difference was least in the north, showing that armoring is generally not emplaced as low on the
314 shore in that region, and greatest in the south. RE values for unarmored beaches were similar
315 among regions.

316 Some of the abiotic and biotic parameters that respond locally to armoring (see below)
317 also varied among regions (Fig. 3, 4). Numbers of logs stranded on the beach were much higher
318 at unarmored beaches but were most abundant in the central region and least in the south; the
319 same pattern was seen in width of the log line (data not shown). Shade from overhanging
320 vegetation likewise was always higher at unarmored beaches but was most abundant (and most
321 different with armoring) at the south sites (Fig. 3). Algal and seagrass wrack biomasses were
322 much greater at the north sites, and there was some variation in the types of wrack found there;
323 seagrass was much more common (Fig. 4), reflecting the local abundance of large seagrass
324 meadows in the region. Algal types were not weighed separately in the wrack samples, but we
325 did record the most common component in each; in all regions, ulvoid algae were the most
326 common (in 80- 85% of samples in all regions), but *Fucus* spp. was more common in the north
327 samples (most common alga in 13% of the samples, versus only 1% in the central and south).
328 The north part of the Salish Sea contains a high proportion of bedrock, the preferred substrate of
329 *Fucus*, whereas there is little such habitat in the central and south regions (Fig. 2).

330 Invertebrates in the wrackline samples showed surprising and largely unexplained
331 differences among regions. This was seen especially in the abundances of talitrid amphipods,

332 oligochaetes, and nematodes, all of which were very patchy at the north sites but often 2-10
333 times more abundant than at the central or south sites (Fig. 4). For the amphipods, these
334 differences stemmed largely from very abundant juveniles (unidentified talitrids) and adults in
335 the genus *Traskorchestia*, with lower numbers of adults in the genus *Megalorchestia* (Fig. 4).
336 Factors affecting amphipod assemblages (all three groups) were examined with multivariate
337 analyses, testing how well a wide range of ‘independent’ variables (grain sizes, amounts and
338 types of wrack, RE, shade, etc.) can predict the types and abundances of amphipods. The best
339 DISTLM analysis produced an r^2 of only 0.32, with 11 predictor variables included. More
340 amphipods of all types were found with more wrack mass and fewer with high RE and high
341 DCA, but all correlations between individual amphipod taxa and individual factors were very
342 weak (r^2 values <0.05). Total wrack mass was correlated with total amphipods over all beaches,
343 but not strongly ($r^2 = 0.19$). When the wrack was mostly terrestrial there were almost no
344 amphipods, but when the wrack was mostly marine the numbers ranged from zero to over 10,000
345 among five core samples.

346 Of the other wrackline arthropods, Collembola varied highly within and among regions;
347 in particular some of the south sites had very large numbers, but these showed no correlations
348 with amounts of any type of wrack. Insects (primarily Diptera larvae and adults) tended to be
349 more abundant in the north (Fig. 4). Insect numbers showed no correlations with algal mass but a
350 positive correlation with terrestrial wrack mass ($r^2 = 0.16$ for all beaches), especially for armored
351 beaches ($r^2 = 0.25$).

352

353 3.2 Sound-Wide Patterns in Paired Analyses

354 As was found in the central-south regions (Heerhartz et al., 2014, 2015), when data from
355 the north sites were included in a 65-pair sound-wide analysis, armoring had clear Sound-wide
356 impacts on a number of parameters on the upper shore (Figs. 3 and 4, Table 1). Unarmored
357 beaches within a pair were wider (overall means and SE Armored 27.3 ± 1.8 m, Unarmored 33.7
358 ± 1.9 m), and extended higher up the shore (Armored 3.03 ± 0.07 m above MLLW, Unarmored
359 3.77 ± 0.08 m), but we found no paired differences in slope of the upper shore (Armored $0.115 \pm$
360 0.0002 , Unarmored 0.110 ± 0.0001). This slope metric is not sensitive to small-scale armor-
361 induced scour. Unarmored beaches had far more shade from overhanging vegetation (Armored
362 $12.5 \pm 3.2\%$, Unarmored $41.7 \pm 4.8\%$), more stranded drift logs (Armored 0.7 ± 0.2 , Unarmored
363 6.7 ± 0.5), and a wider log line (Armored 0.6 ± 0.1 m, Unarmored 5.2 ± 0.4 m). More wrack also

364 accumulated on unarmored beaches, with this pattern holding true for all the measured
365 components of algae, seagrass, and terrestrial plant material (visible by region in Fig. 4). All
366 these differences except slope were significant in paired t-tests (p values < 0.01 , Table 1).

367 However, for the invertebrates found in the wrack, some of these patterns were not
368 consistent with the more localized study of Heerhartz et al. (2015). Armored beaches had
369 reduced numbers of amphipods and insects only in the central and south regions (Heerhartz et
370 al., 2015); when the north beaches were included, neither of these paired t -tests was significant
371 (Fig. 4, Table 1). The exceptions were numbers of Collembola, which varied highly among
372 regions but overall were more common at unarmored beaches, and the relatively uncommon
373 talitrid amphipod genus *Megalorchestia*, also more abundant at unarmored beaches (Fig. 4).
374 Worms involved in decomposition of the wrack (oligochaetes and nematodes) showed no overall
375 armoring effect (Fig. 4, Table 1).

376 Grain size distributions of sediment at the wrack line were generally consistent between
377 pairs of sites (visible by region in Fig. 2), even though the “wrackline” sediments were usually
378 sampled from lower elevations at armored beaches (with wrack stranded at the toe of the
379 armoring). We found no differences in any sediment grain sizes in paired t-tests (p values > 0.15 ,
380 Table 1). In the mid shore (MLW), where sediments were collected at the identical elevation on
381 armored and unarmored beaches, there was again no effect of armoring on grain sizes (Table 1,
382 Suppl. Figure 1).

383

384 *3.3 Thresholds and Cumulative Impacts*

385 We tested for the relative roles of armoring emplaced lower on the shore and of
386 increasing amounts of armoring within drift cells by regressing RE and DCA against the suite of
387 dependent variables (amounts of wrack of different kinds, counts of invertebrates in the
388 wrackline, numbers of logs, etc.). These regressions generally had the form expected from the
389 pairwise analyses, for example declines in logs, wrack, and invertebrates occurred with larger
390 encroachment of armoring on the beach, but few had r^2 values > 0.10 (data not shown). Often the
391 scatterplots were ‘wedge-shaped’ (e.g., Fig. 5, Suppl. Fig. 3). For example, Fig. 5 shows that
392 low-shore-armored beaches always had few logs or little wrack, whereas unarmored or high-
393 shore-armored beaches had highly variable amounts. These plots thus were indicative of the
394 large number of interdependent parameters causing variation in the measured shoreline variables,

395 e.g., wrack abundance at the time of measurement was affected by many factors other than
396 encroachment.

397 Based on the appearance of some scatterplots, we used segmented regression to test for
398 thresholds in the number of logs on a beach in relation to relative encroachment (Fig. 5). Our
399 analysis found that there was a breakpoint in the relationship at a relative encroachment of 1.44
400 feet (SE +/- 1.37 ft), where the regression changed from a non-significant slope of -0.31 (+/-
401 0.70) to a significant slope of -1.34 (+/- 0.27) ($p < 0.0001$) (Fig. 5). In other words, beaches with
402 armoring low on the shore had far fewer logs than expected based on the relationship between
403 number of logs and RE for beaches where the armoring was farther up the shore. Thus RE = 1.44
404 ft constitutes a threshold of relative encroachment below which logs are virtually excluded from
405 a beach. This model was compared to a simple linear regression of total logs against RE using
406 AIC and r-squared values; these were almost identical, with segmented AIC at 723.6 ($r^2 = 0.26$)
407 and simple regression at 722.2 ($r^2 = 0.25$). These comparisons suggest that both models are
408 similar in their ability to describe the data, but in terms of data useful to managers, it is helpful to
409 present the segmented model and threshold. Other scatterplots and segmented regressions
410 suggested similar relationships for the amount of wrack (Suppl. Fig. 3) but were not significant.

411 Our 65 pairs of sites varied greatly in the degree of armoring present (DCA) in the drift
412 cells where they were located (Suppl. Fig. 2). Of particular interest at this larger spatial scale was
413 testing our prediction that there would be an effect of alongshore extent of armoring on sediment
414 grain sizes or beach slope, which showed no armoring signal at the local, paired t-test scale. Our
415 mixed-effects regressions showed clear effects of DCA on a number of grain sizes (Table 1).
416 Figure 6 illustrates these patterns with DCA-extent binned (4 categories) so that the whole grain
417 size spectrum can be shown at once. Regardless of their local armoring status, beaches in the
418 more-extensively armored drift cells (“High” in Fig. 6) had significantly higher proportions of
419 coarse sediments, especially very coarse gravel (32-64 mm), and significantly lower proportions
420 of fine sand (125-250 μm) and medium sand (250-500 μm) (Table 1). Multivariate analyses
421 testing the suite of all grain sizes together showed a highly significant relationship with DCA (1-
422 way PERMANOVA, $p = 0.0001$). To be certain that these grain-size differences were not biased
423 by the generally lower elevation of “wrackline” samples at armored beaches, we ran a simple
424 linear model (without the Site term and thus not mixed-effects) on DCA versus grain sizes
425 (arcsin sqrt transformed) using data from just the unarmored beaches (visible in Fig. 6). Even

426 with the smaller sample sizes (half the N beaches), there was still a significant association of
427 higher DCA with an increased fraction of very coarse gravel ($p = 0.0002$), and decreased
428 medium sand ($p = 0.006$). Other grain sizes were not statistically related to armoring.

429 We also analyzed sediments from the mid-shore (MLW) at all beaches, although for this
430 elevation we had less precise data on grain sizes (from estimates using quadrats in the field) (Fig.
431 6). Mixed-effects regressions showed no significant effects of DCA on any grain sizes at this
432 lower elevation.

433 Because DCA varied with region (Suppl. Fig. 2), as did the proportions of different
434 sediment sizes on the beaches (Fig. 2), we were concerned that the relationship between DCA
435 and grain size might be biased, i.e. driven by some other independent variable that differed
436 among regions. To address this, we examined the underlying geological material (categories in
437 Fig. 2) of the bank or bluff in each drift cell and found that not surprisingly, DCA varied with
438 bluff material -- the most armoring occurred in drift cells dominated by loose sediment (that
439 would presumably require more stabilizing), and the least armoring in bedrock areas. A 2-factor
440 ANOVA on the proportion of very coarse gravel (the fraction with the strongest relationship to
441 degree of armoring) with factors of underlying material (6 types) and DCA (4 levels, binned as
442 in Fig. 6) showed that gravel was significantly associated with DCA ($p < 0.0001$), but not with
443 underlying bluff material ($p = 0.21$), with no significant interaction ($p = 0.08$). Thus one
444 interpretation of this analysis is that although underlying geological material in the bluff must
445 ultimately affect the amount of gravel on the beach, the regional pattern is more closely related
446 to the degree of armoring in the drift cell.

447 The slope of the upper beach also varied with DCA. This test was run with only 54 pairs
448 of beaches (see Suppl. Methods). Beaches on more-armored drift cells (regardless of local
449 armoring) had slightly but significantly steeper slopes than those in less-armored drift cells
450 (Table 1, $p = 0.0028$, data not illustrated); mean slope was 10% at low-DCA beaches and 15% at
451 high-DCA areas. Unexpectedly, relative encroachment had a small but significant ($p = 0.045$)
452 effect in the opposite direction; beaches with greater encroachment of armoring were slightly
453 flatter than those with less encroachment. This may relate to heavily-encroached beaches having
454 scour in front of armoring, which would lead to a reduced near-armor slope (A.N. McBride pers.
455 comm.).

456 The proportion of the drift cell armored was also directly or indirectly associated with
457 several biological parameters. Mixed-effects regressions showed that DCA had a significant
458 negative effect on some wrack mass parameters, and also on total numbers of wrack
459 invertebrates and Collembola (Table 1).

460

461 **4. Discussion**

462 *4.1 Local and Regional Effects*

463 As was found in our previous sampling over a smaller geographic region (Sobocinski et
464 al., 2010; Heerhartz et al., 2014, 2015), a variety of response variables, especially those
465 associated with the upper shore, differed between paired armored and unarmored beaches across
466 the southern part of the Salish Sea. Parameters locally reduced by armoring included width and
467 shadiness of the beach, and log and wrack accumulation on the upper shore. Many of the
468 invertebrate taxa that inhabit the wrack or live under logs were also less abundant with armoring.
469 Most of these patterns were visible throughout our large study area even though there were
470 substantial underlying regional differences. Northern beaches had more algal and seagrass
471 wrack; the abundance of bedrock in the north that supports large algal populations likely
472 contributed to the available algal wrack mass, as did seagrass from very large seagrass beds in
473 Padilla Bay and on large river deltas. The lesser encroachment of armoring in the north also
474 presumably allowed more wrack to accumulate. Northern shorelines have been settled and
475 altered more recently than the central region, and regulation of armoring elevation has become
476 stricter with time. The northern sites also had lower overall proportions of their drift cells
477 armored (DCA); this could be due to the greater awareness of shoreline impacts in this later-
478 developed region, and/or to the larger proportion of bedrock in the drift cells reducing the need
479 for shoreline stabilization.

480 For some biotic parameters there was an association with armoring either on local or
481 broader spatial scales, while for others the regional, geomorphic, or other sources of variation
482 obscured such potential patterns. The larger masses of wrack (especially algal) in the north were
483 occupied by higher densities of amphipods, nematodes, and oligochaetes, while more insects
484 were associated with larger amounts of terrestrial wrack. Somewhat surprisingly, numbers of
485 insects showed no relationship with the amount of overhanging vegetation (percent shade).

486 Collembolans showed a regional pattern driven by high densities in a few southern sites, but did
487 not correlate with any type of wrack.

488 The very large regional variation in talitrid amphipod abundances and their inconsistent
489 response to armoring likely relate to unexplored behavioral responses of these important wrack
490 inhabitants. *Megalorchestia* was the only amphipod taxon to show a consistent sensitivity to
491 armoring across regions (seen also by Dugan et al., 2003), and to respond significantly to relative
492 encroachment of armoring on the beach. This genus tends to burrow in sand high on the shore
493 (Pelletier et al., 2011; Dugan et al., 2013) and to be sensitive to sediment textures (Viola et al.,
494 2013). *Traskorchestia*, in contrast (likely including most of the “juvenile talitrids” counted)
495 burrows less and is more likely to move around the beach, shelter in wrack, and survive
496 submersion for extended periods (Koch, 1989). They may concentrate in lower wrack when the
497 tide is out but (in the absence of armoring) move to higher elevations to avoid being submerged
498 at high tide. Our wrack samples were taken at variable times relative to the tidal level and under
499 many different weather conditions, and we did not track age or field-moisture content of the
500 wrack; such static sampling may have affected our ability to accurately measure these highly
501 mobile organisms.

502 Our tests of armoring-associated effects lower on the shore were inconclusive. Sediment
503 analyses at Mean Low Water showed no differences between paired beaches. We also tested for
504 a biotic response to hypothesized changes in sediment texture in the abundance or species
505 richness of juvenile clams, but found no patterns (Dethier, unpubl.). Our mid-shore samples were
506 physically removed from armoring by an average of ~30 m across the beach face, meaning that
507 direct armor effects such as from wave reflection were unlikely. Long-term indirect effects such
508 as gradual loss of finer sediments from the beach face could impact the mid shore but were not
509 detected in our data.

510

511 *4.2 Broader-Scale Patterns*

512 For some parameters that we hypothesized would be affected by armoring, the local,
513 paired-beach scale was mismatched to the larger-scale processes that likely control these
514 parameters. This was particularly true for geomorphological parameters such as beach profiles
515 (e.g., slope). One likely explanation is that armoring impacts “smear” among members of a
516 sampled pair; for example, if an unarmored beach has sediment naturally eroding onto the shore,

517 some of the sediment is likely to get carried to the nearby armored beach even if that beach is, on
518 average, “updrift”. Conversely, changes to wave energies and sand supply caused by a large
519 stretch of armoring could impact sediment processes on a nearby unarmored beach. Important
520 contextual parameters such as age of the armoring often could not be ascertained due to poor
521 historical record keeping. Even with our large sample sizes we had insufficient replication to test
522 hypotheses related to different types of armoring, for example vertical concrete versus sloped
523 riprap.

524 Our ability to test for relationships between armoring and various response parameters
525 was also compromised by the interactions between space and time, and geomorphology and
526 biology. Testing hypotheses about armoring effects on sediment grain sizes and beach slopes, for
527 example, was not possible until we had data encompassing drift cells with a large range of
528 armoring – and for those conditions to have been present for long enough that finer sediments
529 had time to gradually winnow out of armored beaches. In addition, there are potential
530 geomorphic factors not considered in our analysis, and a definitive cause and effect relationship
531 is yet to be determined. Differences in upper-shore sediment grain sizes were not detected at the
532 paired beach scale but were significant when examined at a large geographic scale. We interpret
533 this regional-scale analysis to suggest that there is a reduction of sediment input resulting from
534 armoring large proportions of drift cells. In turn, this appears to have long-term, cumulative
535 geomorphological effects, such that proportions of fine sediments are reduced, leaving behind
536 coarser ones. Even at this scale, cumulative armoring effects were relatively subtle, and
537 statistically significant only for the grain sizes at the ends of the size spectrum. As in sandy
538 beach ecosystems (Berry et al., 2014), this cumulative effect may reduce the ecological resilience
539 of Puget Sound beaches, where sediment supply is already episodic. Grain sizes then affect
540 numerous biological parameters such as suitability for spawning surf smelt (Penttila, 2007), and
541 the numbers and types of invertebrates in the wrack zone and elsewhere on the beach (e.g., Viola
542 et al., 2013; Heerhartz et al., 2015). The predominance of wedge-shaped plots in our analyses
543 (e.g., Fig. 5, Suppl. Fig. 3) attests to the large numbers of factors affecting all of our measured
544 parameters; abundances of wrack invertebrates, for example, are likely influenced not only by
545 the amount and type of wrack itself, but the elevation of the wrack on the shore, the porosity of
546 the sediment, and the region.

547 Lower elevations of shoreline armoring, calculated as relative encroachment over the
548 beach, showed a clear negative association with most beach parameters at both local and larger
549 spatial scales. For some of these parameters, such as number of accumulated logs, segmented
550 regressions demonstrated that there was a distinct threshold elevation below which armoring
551 seemed to have dramatic impacts; a similar pattern was seen in total wrack mass. In each case,
552 the threshold was ca. 1 - 2 vertical feet below MHHW. Armoring below this elevation, which is
553 no longer permitted for new construction in Washington State, was associated with substantially
554 greater differences in measured parameters than armoring higher on the beach. This elevation
555 thus may constitute a “utility threshold” (Samhoury et al., 2010) to be targeted by management
556 actions or restoration to obtain the most significant beneficial changes in ecosystem functions.

557 Our study has documented both obvious and more subtle effects of armoring on Salish
558 Sea shorelines, including those detectable at diverse spatial and temporal scales as summarized
559 in Figure 7. Some differences, such as reduced wrack accumulation on armored beaches, could
560 be seen at local spatial scales (paired beaches) and probably would be observable within days of
561 armor installation. Wrack is delivered to beaches on almost every high tide, and stranding of this
562 material is clearly reduced in front of armoring, especially when the structure is relatively low on
563 the shore. At the other end of the spatial and temporal spectrum, the hypothesized geomorphic
564 responses such as slope and grain size distributions were not visible at the paired-beach scale,
565 where they are obscured by the numerous processes that impact local beaches on both short and
566 long-term time scales. These responses likely require both a large extent of armoring and
567 substantial time of sediment reworking to create a signal that is detectable over the natural
568 geomorphic variability.

569 Exact positions of responses related to armoring on the space and time axes in Figure 7
570 are only approximate, and some are context-dependent or only weakly supported. Forage fish
571 spawning, placed at intermediate scales of space and time (Fig. 7), could actually be affected
572 rapidly and locally if armoring covers spawning beaches, or slowly and only at very broad scales
573 in the case of gradual population decline due to large-scale loss of appropriate sand grains for
574 egg attachment. Our previous studies have suggested that shoreline armoring has some effect on
575 abundance and behavior of terrestrial birds (Heerhartz, 2013) and juvenile salmon (Toft et al.,
576 2013; Heerhartz and Toft, 2015), but since these organisms are highly mobile and use large
577 stretches of shoreline, distinguishing population responses to armoring is very difficult. Mobile

578 organisms in general present similar problems with regard to conservation (Runge et al., 2014;
579 Rolet et al., 2015). Juvenile salmon migrating alongshore on their way to the ocean encounter the
580 entire spectrum of armored and natural beaches, so attributing effects on diet, fitness, and
581 survival to one factor such as armoring requires manipulative studies such as holding fish in a
582 small area to measure local feeding rates (Toft et al., 2007; Toft et al., 2013). Armoring located
583 in juvenile fish habitats likely changes the character of the wrack and invertebrates therein, as
584 well as overhanging vegetation and insects, all of which may alter behavior or feeding of the
585 fish.

586 While our study did not directly address restoration efforts, our observations combined
587 with site-specific data from armor-removal projects within the Salish Sea (e.g., Toft et al., 2014)
588 suggest that many of the armoring impacts we observed may be reversible. In some cases, beach
589 functions may be at least partially restored by modification of shore structures and may not
590 require complete removal (Berry et al., 2013; Nordstrom, 2014). Our data suggest that moving
591 armoring higher on the shore may restore some ecological functions while still protecting
592 infrastructure. Recovery of beach characteristics and functions may follow the same temporal
593 patterns illustrated in Figure 7. Wrack can return quickly when armoring has been removed and
594 there is physical space on the upper shore for it to accumulate; colonization by arthropods and
595 other decomposers is likely to follow quickly if there are local sources of colonists. Terrestrial
596 birds will probably visit restored spaces quickly, once invertebrate food becomes available, and
597 rapid juvenile salmonid use of a restored beach has already been demonstrated at a site in Puget
598 Sound (Toft et al., 2013). If sediments are appropriate for spawning forage fish, or if armoring
599 removal is accompanied by beach nourishment with appropriate sediment, then egg-laying may
600 occur during the next spawning season; but even spawning on appropriate sediment is
601 unpredictable in space and time (e.g., surf smelt: Penttila, 2007). These biotic changes may
602 happen on relatively short temporal scales, for example seasonally, rather than taking years over
603 which some armoring impacts may develop. Recovery of geomorphic parameters such as beach
604 shape and pre-armoring sediment grain sizes will depend on sediment sources, whether from
605 updrift, upslope, or artificial delivery.

606 Multiscale spatial and temporal impacts of armoring are also likely to be seen on open-
607 coast sandy beaches or other systems such as armored estuarine marshes. On sandy beaches, the
608 effects of armoring on wrack accumulation and on other trophic levels have been well studied

609 (e.g., Dugan et al., 2008). A relatively unique feature of Pacific Northwest beaches is extensive
610 windrows of beach logs, but these may have some parallel in marsh vegetation that can only
611 develop when armoring is absent or very high on the shoreline (Bozek and Burdick, 2005). As in
612 the Salish Sea, on both sandy beaches and marshes the direction of drift (e.g., longshore currents,
613 estuarine outflow) should affect the location and spatial scale of armoring impacts because the
614 accumulation of both sediments and organic matter are important in those ecosystems.
615 Geomorphic effects of armoring on open beaches or marshes are similarly likely to be slow or
616 highly episodic, depending on types of sediment sources and their proximity, as well as
617 variations in wave energy. The degree to which sediment sources are locked up, either by
618 extensive alongshore armoring or by dams on riverine sources, may have cumulative effects;
619 investigating possible thresholds in the interactions between sediment budgets and marsh health
620 or beach geomorphology would be useful but temporally challenging.

621 In conclusion, our broad study covering a wide range of beaches and drift cells with
622 different types, elevations, and degrees of armoring has allowed us to quantify hitherto elusive
623 patterns of impacts of armoring on beach processes. Armoring alters beach conditions from the
624 local to the sound-wide scale, with its effects likely emerging on time scales that range from
625 immediate to years or decades. In the Salish Sea, there is great variation among beaches and
626 regions in upper-shore parameters such as logs, wrack, and invertebrates, but in many cases an
627 armoring signal overrides these complex processes, and broad associations are visible. The
628 changes in the geomorphic character of beaches towards steeper and coarser conditions appear to
629 be slow and subtle, but ultimately can ramify to impact beach functions, including supporting
630 forage fish use and altering the infauna. The elevation of armoring on the shore clearly does
631 make a difference to numerous functional characteristics, and at least in the case of log
632 accumulation, there is a threshold for this effect. Our data also suggest that adding more
633 armoring within drift cells may lead to cumulative impacts on several geomorphic and biological
634 parameters. The mechanisms that might cause these cumulative effects, for example starving the
635 beaches of sediment supply or altering local hydrodynamics, require further investigation.

636

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654

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Table 1. Summary of statistical tests.

<u>Description</u>	<u>Paired t-tests</u>		<u>RE</u>			<u>DCA</u>		
	<u>pvalue</u>	<u>direction</u>	<u>pvalue</u>	<u>Direction</u>	<u>Test type</u>	<u>pvalue</u>	<u>Direction</u>	<u>Test type</u>
Beach Width	0.0104	U > A						
Beach Slope	ns		0.0453	neg	C	0.0028	pos	C
Shade on upper shore	<0.0001	U > A						
Number of logs	<0.0001	U > A						
Width of log line	<0.0001	U > A						
Wrack Terrestrial Percent Cover	<0.0001	U > A	< 0.0001	neg	B	ns		B
Wrack Algae Percent Cover	0.0012	U > A	0.0002	neg	B	ns		B
Wrack Total Percent Cover	<0.0001	U > A	< 0.0001	neg	B	ns		B
Wrack Total Mass	<0.0001	U > A	< 0.0001	neg	A	0.0074	neg	A
Wrack Algae Mass	0.00773	U > A	< 0.0001	neg	A	0.0247	neg	A
Wrack Terrestrial Mass	0.00013	U > A	< 0.0001	neg	A	ns		A
Wrack Total Invertebrates	ns		0.0001	neg	A	0.0051	neg	A
Wrack Total Amphipods	ns		0.003	neg	A	ns		A
Wrack Total Insects	ns		< 0.0001	neg	A	ns		A
Wrack Total Collembola	0.00759	U > A	0.0001	neg	A	0.0293	neg	A
Wrack Oligochaeta + Nematoda	ns							
Wrack Megalorchestia	0.0002	U > A						
Very Coarse Gravel	ns		ns		B	0.0001	pos	B
Coarse Gravel	ns		ns		B	ns		B
Medium Gravel	ns		ns		B	ns		B
Fine Gravel	ns		ns		B	ns		B
Very Fine Gravel	ns		ns		B	ns		B
Very Coarse Sand	ns		ns		B	ns		B
Coarse Sand	ns		ns		B	ns		B
Medium Sand	ns		ns		B	0.0042	neg	B
Fine Sand	ns		ns		B	0.0008	neg	B
Fines	ns		ns		B	ns		B

Notes: Type of test: A = Mixed-effect ANOVA on quasi-Poisson data; B = Mixed-effects on arcsin sqrt transformed data; C = normal linear mixed effect model. 'ns' = non-significant. 'neg' and 'pos' refer to the direction of effect of the parameter on the response variable, e.g. large RE is associated with low wrack cover.

Figure Captions

Figure 1. Map of the Washington State portion of the Salish Sea, showing study site locations and major cities. Each pair of beaches (armored and unarmored) is represented by a dot. North sites are represented by letters, Central by #1-25, and South by #26-31. Basemap data courtesy of Washington Dept. of Ecology (WA State Basemap, Place Names)

<http://www.ecy.wa.gov/services/gis/data.htm> and Washington State Dept. of Transportation (Shoreline) <http://www.wsdot.wa.gov/mapsdata/geodatacatalog/>

Figure 2. Upper panel: Regional differences in geological materials comprising bluffs at the study sites. Lower panel: averaged sediment grain-size distributions (proportions) in samples from the wrack line (samples sieved in the lab). Sample sizes: North = 36 pairs of beaches, Central = 23 pairs, South = 6 pairs.

Figure 3. Physical parameters measured at all beaches. Bars are means and one SE. Sample sizes as in Fig. 2.

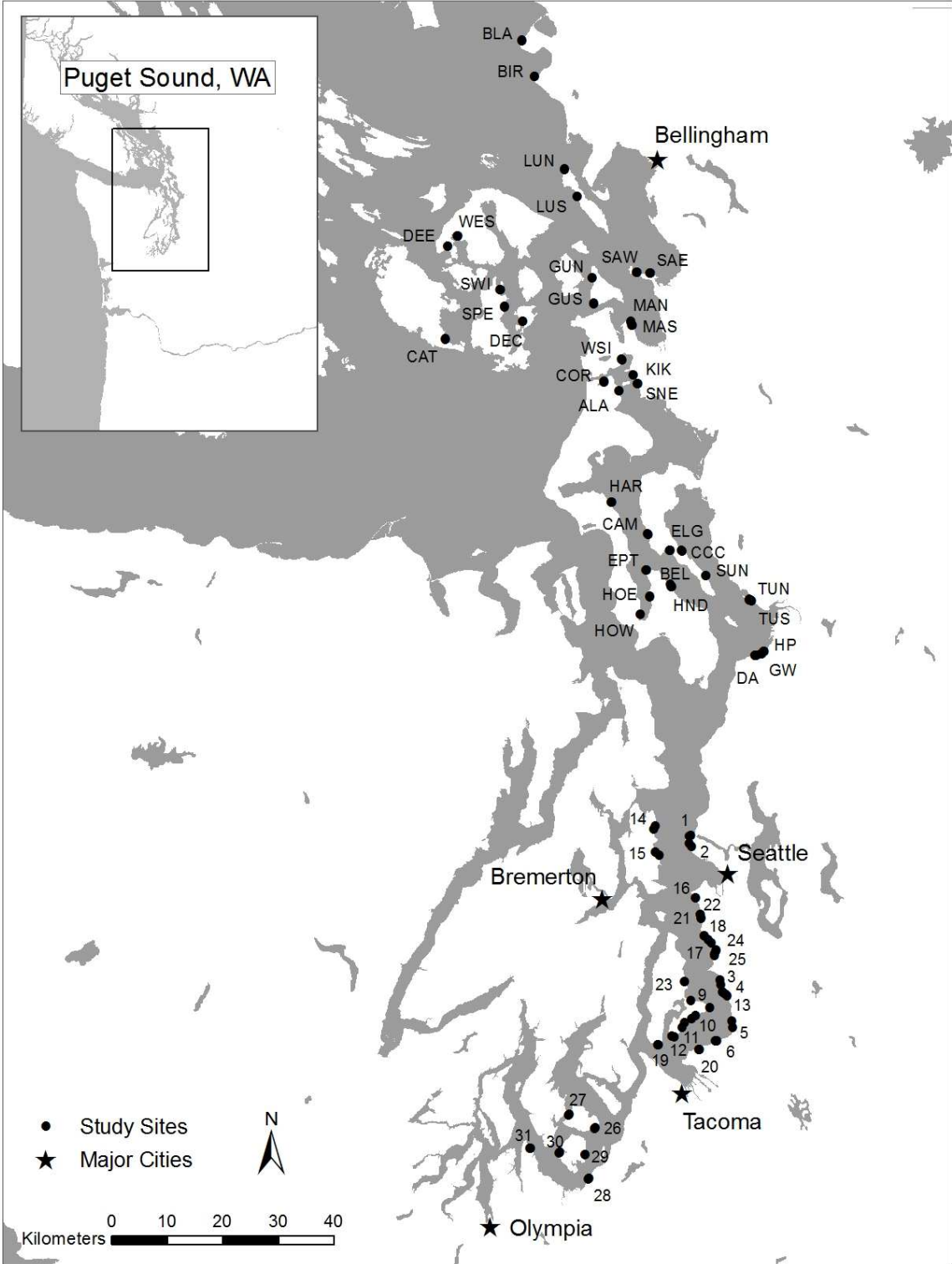
Figure 4. Abundances of types of wrack and organisms in wrack cores by region and treatment (armored vs unarmored). Bars are means and one SE of the summed elements. North = 36 pairs of beaches, Central = 23 pairs, South = 6 pairs.

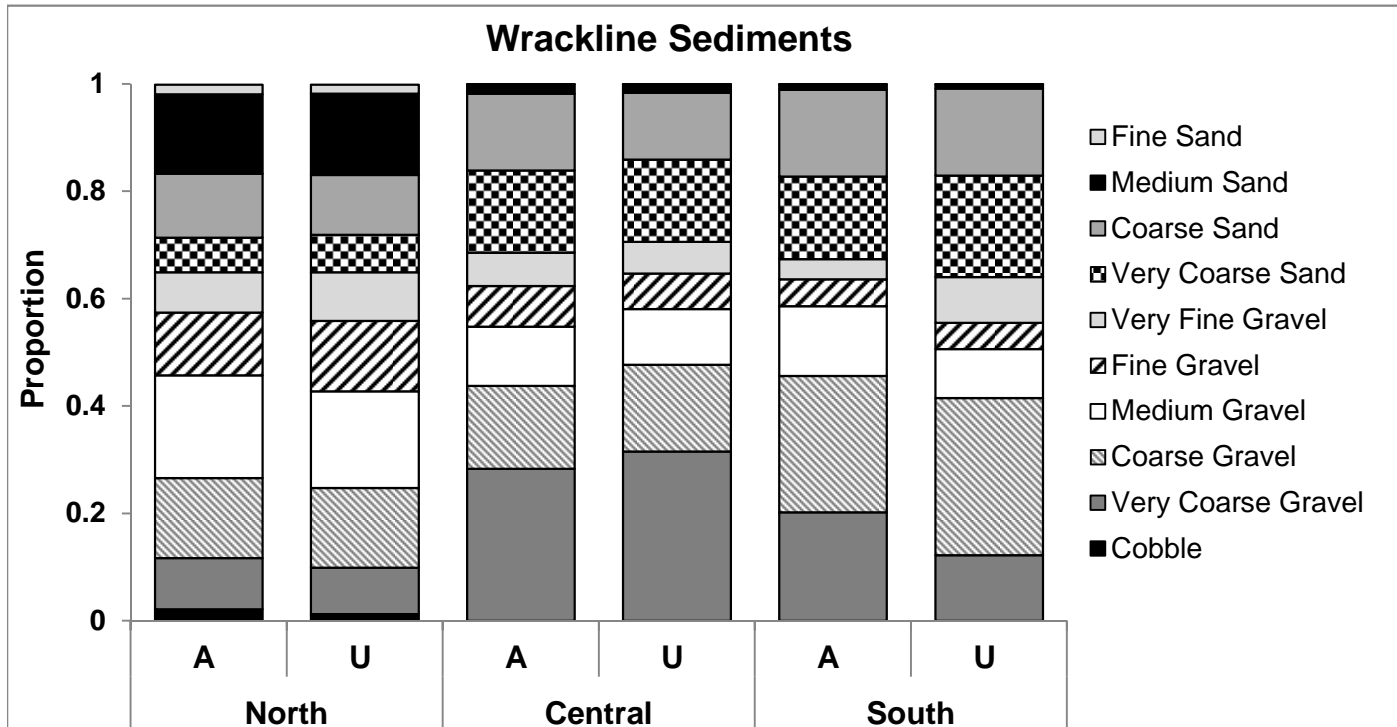
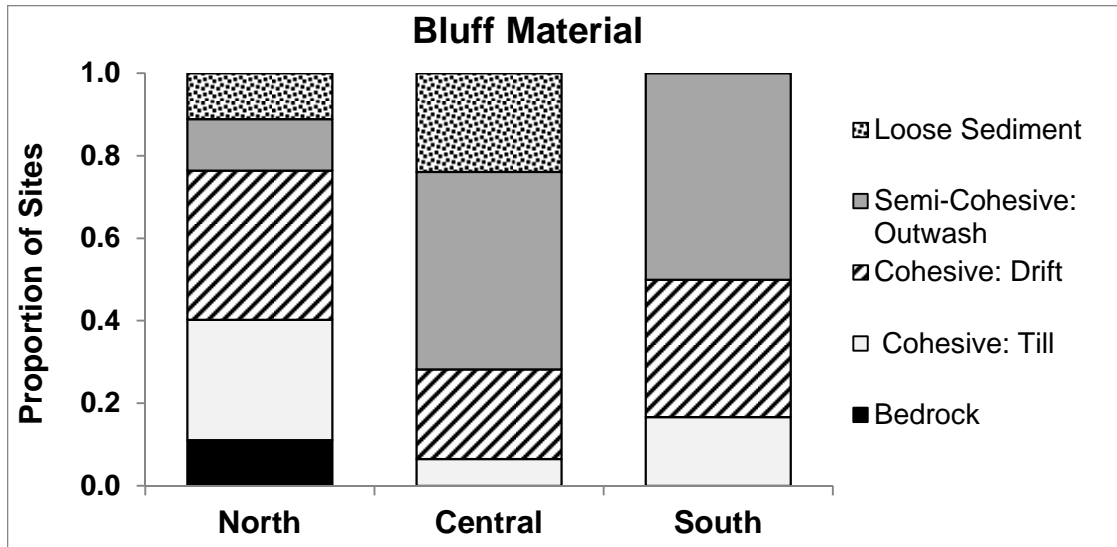
Figure 5. A segmented regression of the number of logs per transect relative to encroachment of bluff or armoring below MHHW. Regression lines incorporate both unarmored beaches (open circles), and armored beaches (filled circles).

Figure 6. Proportions of sediment grain sizes at the Wrack line (upper panel) and at Mean Low Water (lower panel) in drift cells with different degrees of armoring (Low DCA = 0-0.2 proportion armored, Medium Low = 0.2-0.5, Medium High = 0.5-0.8, and High = 0.8-1.0). We split these proportions somewhat unevenly to allow for similar site replication within each bin, and also to highlight the impacts of particularly low and particularly high amounts of armoring in the drift cell. Sediments from the Wrack line were dry-sieved in the lab; sediments at MLW were

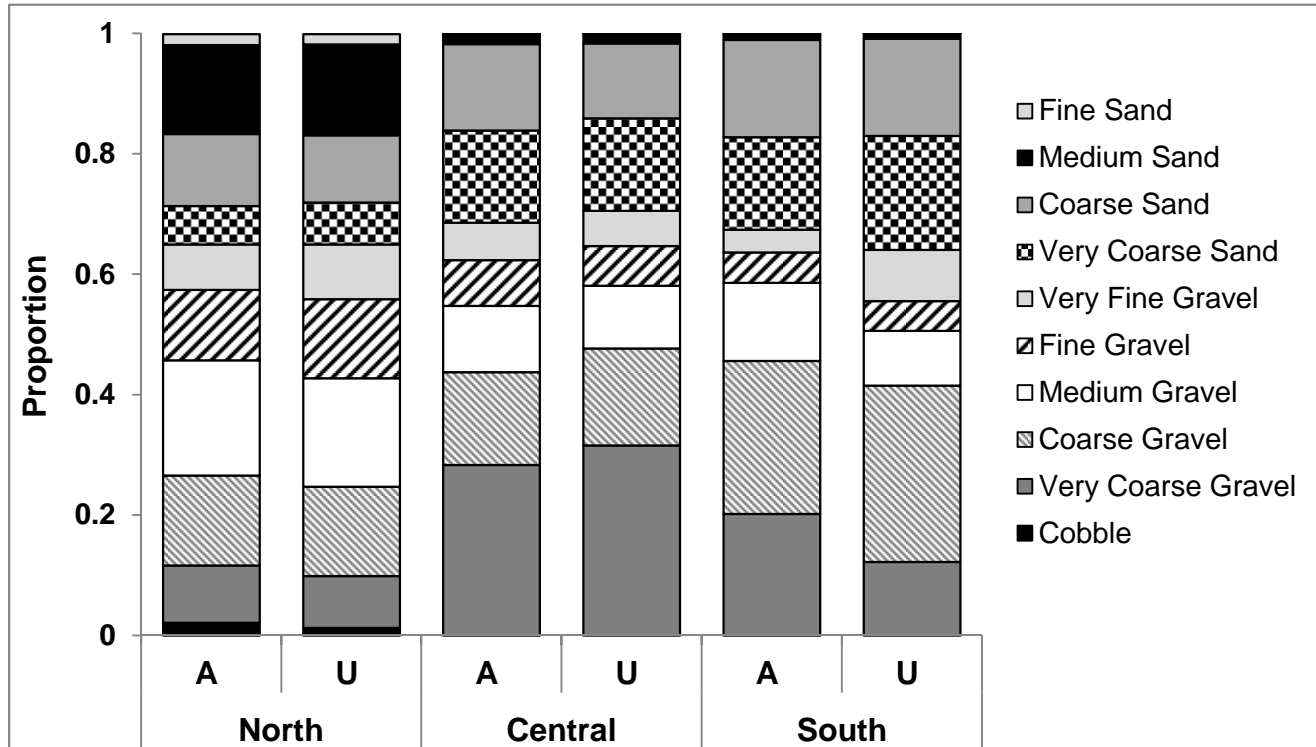
estimated in quadrats on the beach. Some of the MLW bars do not sum to 1.0 because of small amounts of hardpan or mud present. For each of the bars (A and U) in each DCA category, the N beaches = Low 20, Med Low 24, Med High 12, High 9.

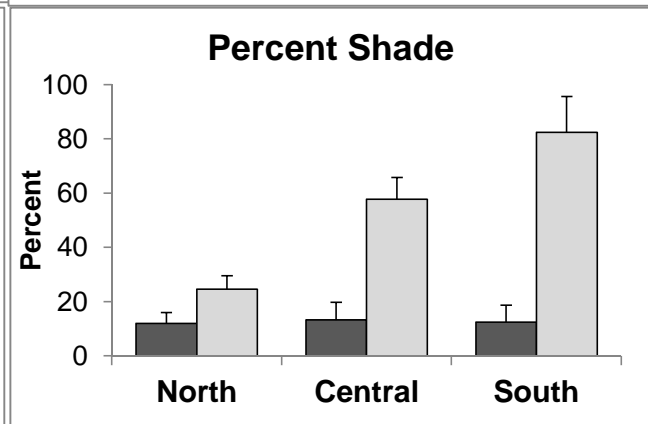
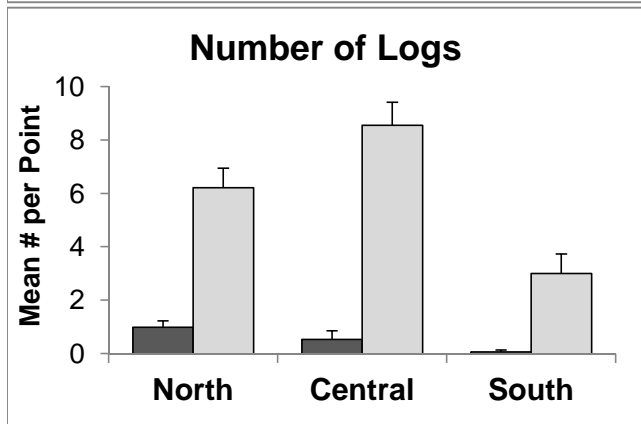
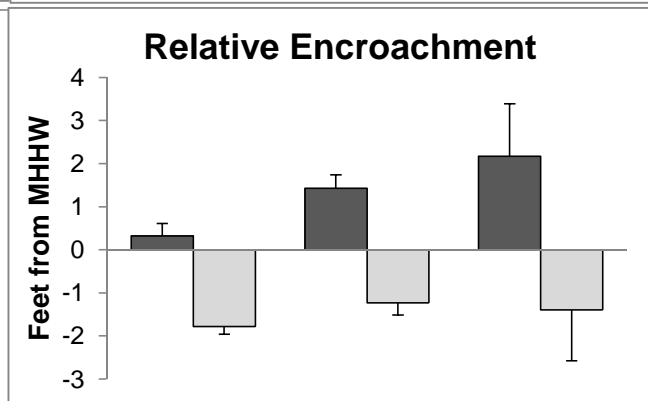
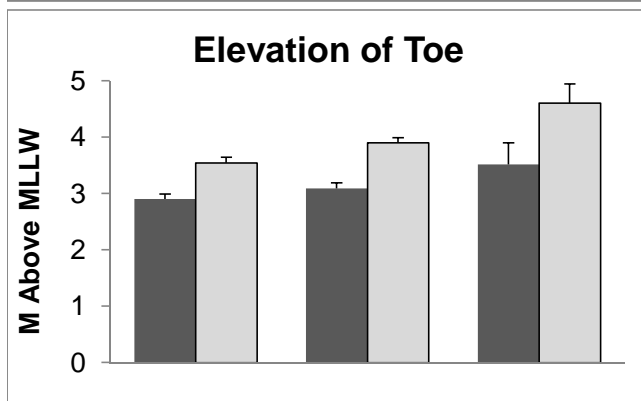
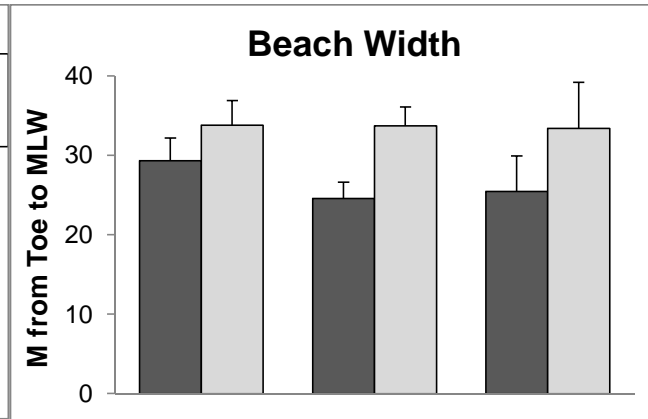
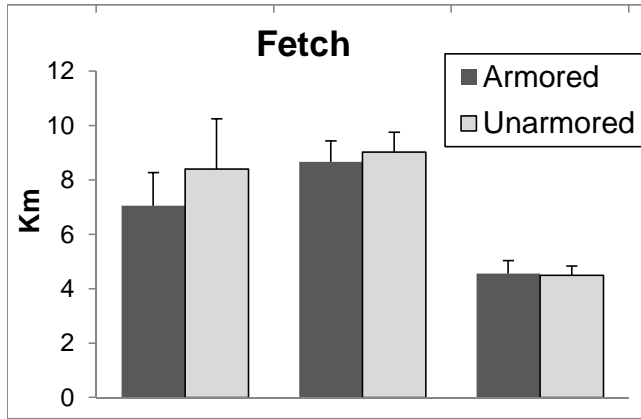
Figure 7. Temporal and spatial scales at which different types of impacts of armoring can be detected. Impacts in dashed boxes are hypothesized but not thoroughly demonstrated. Speed of responses following restoration (armor removal) may follow the same temporal and spatial patterns.

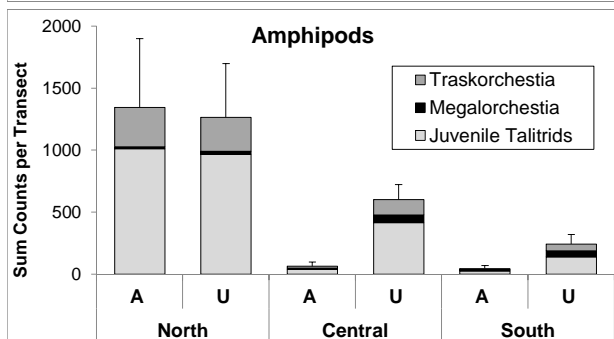
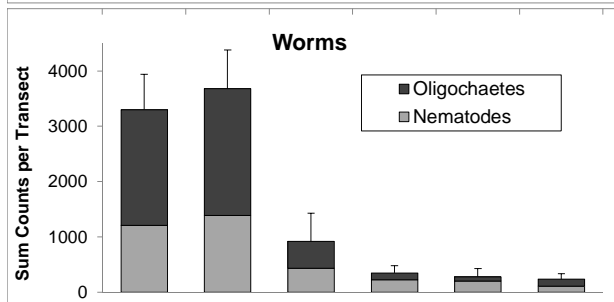
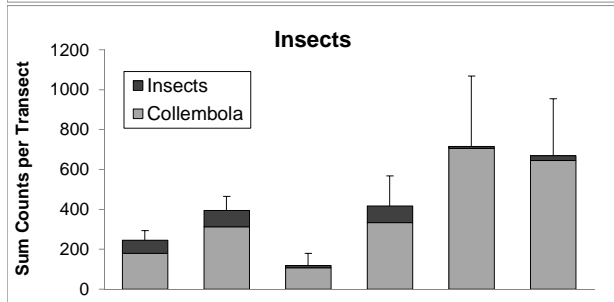
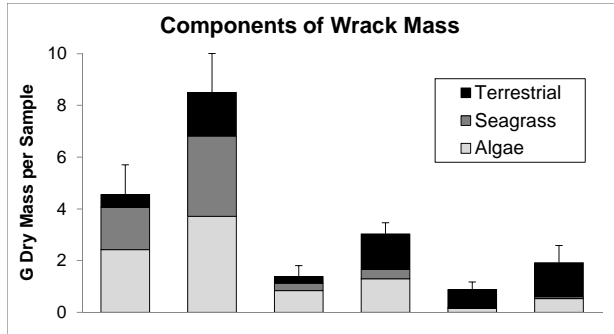




		Cobble	Very Coars	Coarse Gra	Medium G	Fine Gravel	Very Fine G	Very Coars	Coarse San	Medium Sa	Fine Sand
North	A	0.021488	0.095282	0.149155	0.191365	0.116851	0.075414	0.064057	0.119268	0.148119	0.017663
	U	0.01284	0.086154	0.148256	0.180283	0.131376	0.09069	0.069545	0.111773	0.151088	0.016959
Central	A	0	0.283364	0.154279	0.110103	0.075864	0.0623	0.152821	0.143533	0.017358	0.000318
	U	0	0.315728	0.161272	0.103893	0.066134	0.058702	0.153456	0.124483	0.015947	0.000316
South	A	0	0.202206	0.254028	0.129773	0.050142	0.037794	0.153949	0.161581	0.009792	0.000591
	U	0	0.122422	0.292684	0.091002	0.049337	0.084941	0.189321	0.161718	0.008178	0.000292

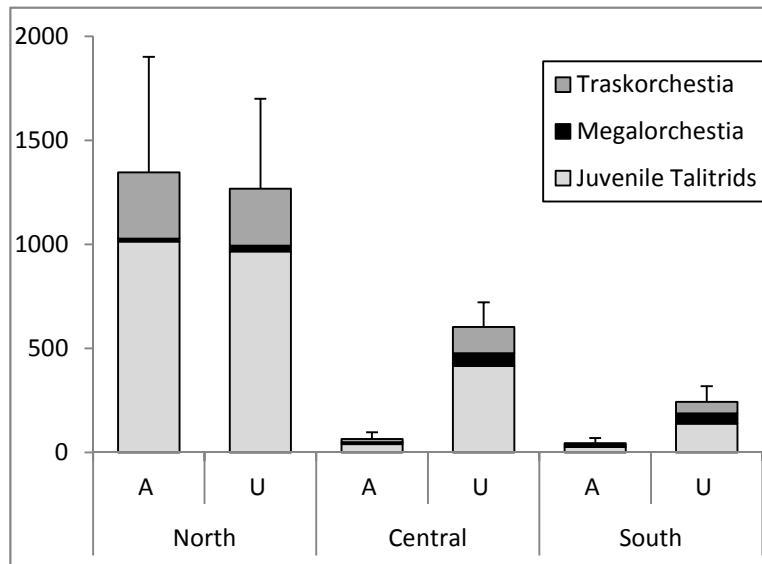


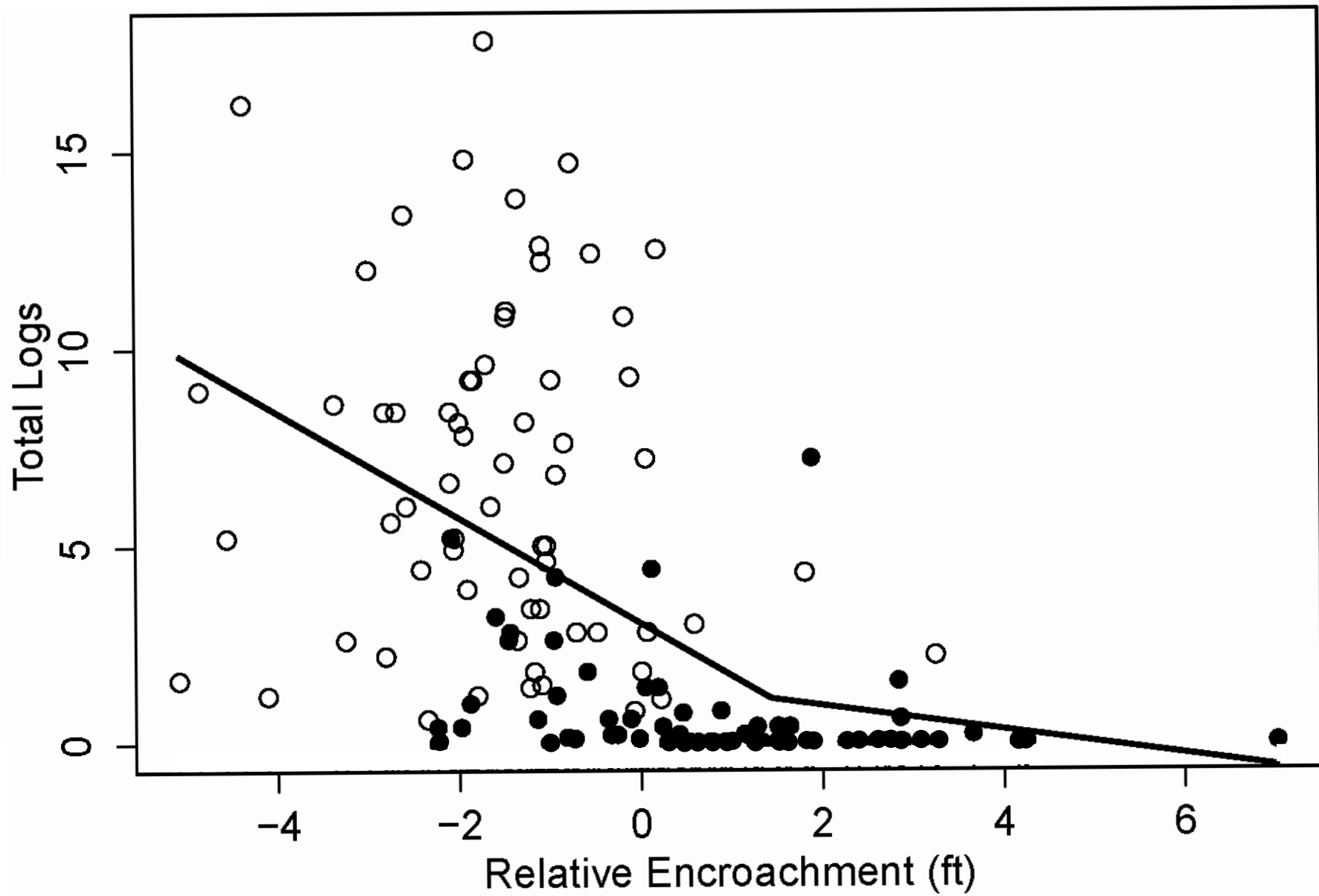


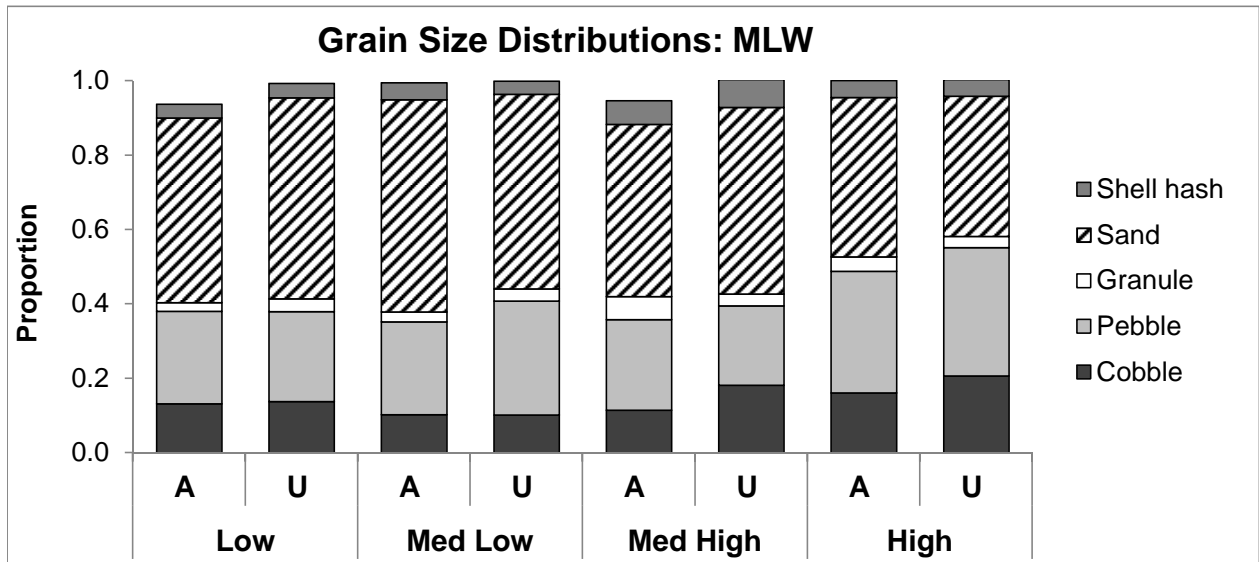
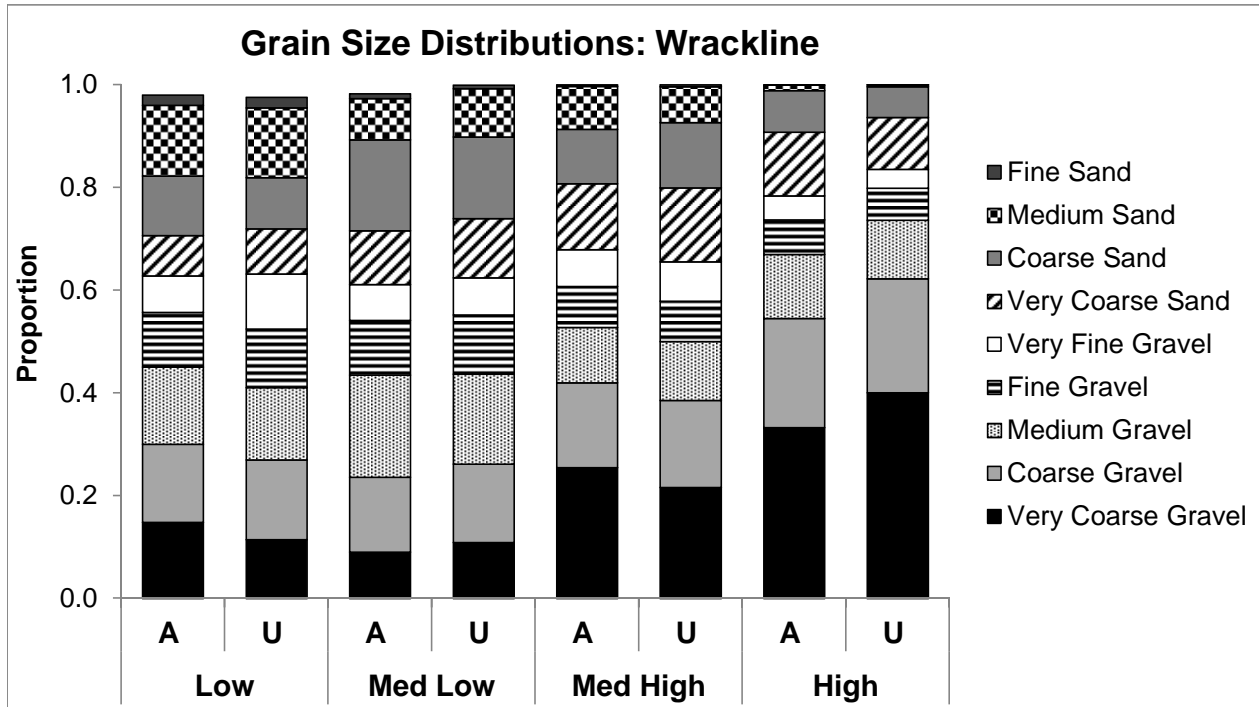


se of summed

		Juvenile Ta	Megalorch	Traskorche	amphipods
North	A	1013.556	14.55556	317.0556	555.049
	U	966.1111	27	273.9444	432.0206
Central	A	40.2971	10.01449	14.65217	32.55651
	U	416.2536	60.74457	125.538	119.734
South	A	26.08333	11.83333	7	24.45452
	U	138.7083	49.79167	55.33333	75.61202





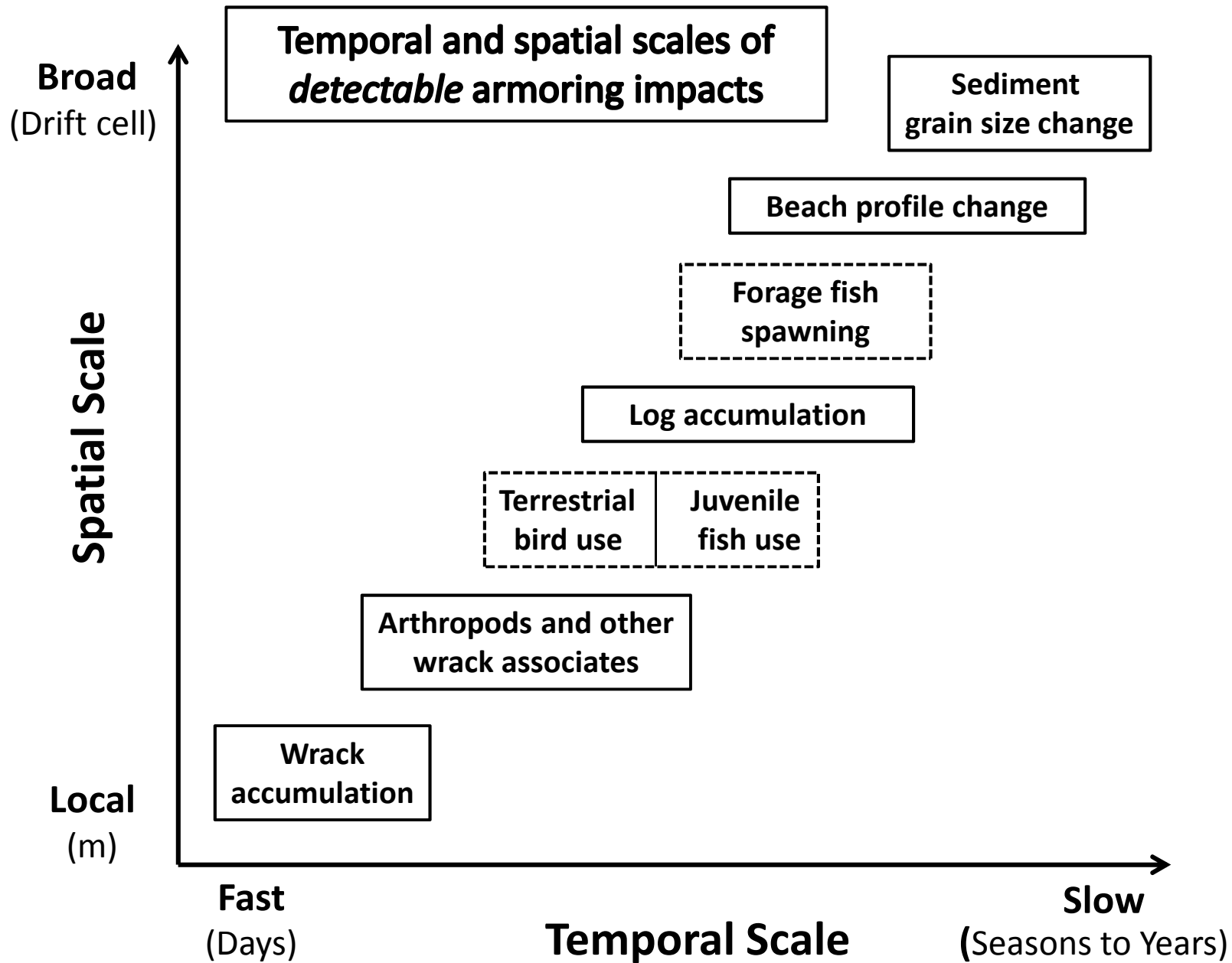


Sediments at Wrackline

		Very Coars	Coarse Gra	Medium Gr	Fine Gravel	Very Fine G	Very Coars	Coarse San	Medium Sa	Fine Sand
Low	A	0.147945	0.15229	0.150261	0.106661	0.070686	0.078953	0.115797	0.137712	0.019542
	U	0.114452	0.155417	0.140489	0.114065	0.107334	0.087705	0.09964	0.135933	0.020694
Med Low	A	0.089794	0.145919	0.199104	0.106303	0.069984	0.104732	0.176887	0.080753	0.009099
	U	0.108878	0.152796	0.175193	0.115191	0.0721	0.115617	0.158459	0.094424	0.006778
Med High	A	0.254745	0.16495	0.107665	0.0801	0.071897	0.127873	0.106228	0.083483	0.002895
	U	0.216148	0.169108	0.115094	0.0786	0.076468	0.143977	0.127238	0.069655	0.003509
High	A	0.332212	0.21277	0.124939	0.067408	0.046495	0.124167	0.080937	0.010688	0.000312
	U	0.400213	0.221983	0.114468	0.061955	0.036658	0.10138	0.059399	0.003812	9.73E-05

Sediments at MLW

		Cobble	Pebble	Granule	Sand	Shell hash
Low	A	0.13	0.25	0.02	0.50	0.04
	U	0.14	0.24	0.03	0.54	0.04
Med Low	A	0.10	0.25	0.03	0.57	0.05
	U	0.10	0.31	0.03	0.52	0.04
Med High	A	0.11	0.24	0.06	0.46	0.06
	U	0.18	0.21	0.03	0.50	0.08
High	A	0.16	0.33	0.04	0.43	0.05
	U	0.21	0.35	0.03	0.38	0.05



Highlights for Dethier et al., "Multiscale impacts..."

- Logs, wrack, and high-shore invertebrates decline with shoreline armoring.
- Armoring emplaced further down the beach face has progressively greater impacts.
- Documented threshold responses with elevation are relevant for habitat managers.
- In drift cells with extensive armoring, beaches have coarser sediment.
- Detection of geomorphic beach changes requires large spatial scales of observation.

