



## Host density and anthropogenic stress are drivers of variability in dark spot disease in *Siderastrea siderea* across the Florida Reef Tract

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**Bulletin of Marine Science**

E-pub ahead of print: 27/06/2023

Peer reviewed version

[Cyswllt i'r cyhoeddiad / Link to publication](#)

*Dyfyniad o'r fersiwn a gyhoeddwyd / Citation for published version (APA):*

Aeby, G. S., Williams, G. J., Whittall, D., Davies, A., Fromuth, E., & Walker, B. K. (2023). Host density and anthropogenic stress are drivers of variability in dark spot disease in *Siderastrea siderea* across the Florida Reef Tract. *Bulletin of Marine Science*. Advance online publication.

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1 **Running title:** Host density and anthropogenic stress drive patterns of dark spot disease

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3 **Title:** Host density and anthropogenic stress are drivers of variability in dark spot disease in

4 *Siderastrea siderea* across the Florida Reef Tract

5

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## ABSTRACT

Dark spot disease (DSD) was first reported within Florida’s coral reefs in the 1990s but factors affecting its spatial distribution have not been well studied. We used a 14-year (2005-2019) coral monitoring data set, utilizing 2,242 surveys collected along Florida’s coral reefs (~530 linear km) to explore the spatial and temporal patterns of DSD occurrence. We built predictive statistical models to test for correlations between a suite of environmental and human impact factors and the occurrence of DSD in the reef coral, *Siderastrea siderea*. DSD in *S. siderea* is a chronic disease which occurred in all 14 years of the study. Annual DSD prevalence ranged from 0.45% to 4.4% and the proportion of survey sites that had DSD ranged from 4.8% to 30.9%. During the study period, DSD became more widespread across Florida’s coral reefs and affected a higher proportion of *S. siderea* populations. Spatial variations in DSD correlated with environmental and human factors which together explained 64.4% of the underlying variability. The most influential factors were concentration of silica in the surface waters (a proxy for freshwater input), the total number of coral hosts, and distance to septic areas. DSD occurred in all regions, but the highest cumulative prevalence occurred in the upper Keys on reefs around major urban centers with links to coastal water discharges. Our results support the hypothesis that coastal water quality is a key component of DSD disease dynamics in Florida and provides motivation for addressing land-sea connections to ameliorate disease occurrence in the region.

## INTRODUCTION

47           Disease is a normal component of host populations, but increased levels of disease can  
48 indicate changes in host-pathogen ecology often due to shifting environmental conditions  
49 (Schrag and Wiener 1995, Daszak et al. 2000). Climate change and environmental degradation  
50 from local human impacts are cited as major influences contributing to disease outbreaks in  
51 wildlife populations (Daszak et al. 2000, Dobson and Foufopoulos 2001, Lafferty and Holt 2003)  
52 including reef corals (Harvell et al. 1999, 2007, Carpenter et al. 2008, Maynard et al. 2015).  
53 Diseases in corals have become a problem worldwide with outbreaks reported from the Indo-  
54 Pacific (Willis et al. 2004, Myers and Raymundo 2009, Aeby et al. 2011a, 2015, 2016, Haapkylä  
55 et al. 2011), Persian Gulf (Howells et al. 2020, Aeby et al. 2020), Indian Ocean (Thinesh et al.  
56 2009, Raj et al. 2016) and western Atlantic (Patterson et al. 2002, Miller et al. 2009, Voss and  
57 Richardson 2006, Croquer and Weil 2009, Brandt et al. 2012, Croquer et al. 2021). The most  
58 devastating coral disease in recent history, stony coral tissue loss disease, emerged in Florida in  
59 2014 (Precht et al. 2016) and continues to spread across the Caribbean, causing significant  
60 mortality on affected coral reefs (Walton et al. 2018, Alvarez-Filip et al. 2019, Sharp et al. 2020,  
61 Heres et al. 2021, Estrada-Saldivar et al. 2021). Coral disease is a major threat to the survival of  
62 coral reefs and identifying the primary factors affecting coral disease dynamics is key to  
63 designing effective local mitigation strategies, prioritizing disease intervention resources, and  
64 identifying areas suitable for reef restoration.

65           Environmental factors can affect host-pathogen dynamics by impairing host immune  
66 responses and enhancing pathogen abundance, transmission, or virulence (Daszak et al. 2001,  
67 Dobson and Foufopoulos 2001, Harvell et al. 2007) and several environmental conditions have  
68 emerged as important in driving increased disease levels in corals. Thermal stress has been  
69 linked to numerous disease outbreaks (Bruno et al. 2007, Miller et al. 2009, Bruno 2015,

70 Brodnicke et al. 2019, Howells et al. 2020) with higher ocean temperatures thought to impair  
71 coral immune responses as well as enhance pathogen abundance and virulence (Mydlarz et al.  
72 2009, Maynard et al. 2015, Ushijima et al. 2016, Vega Thurber et al. 2020). Lamb et al. (2018)  
73 found that disease establishment was facilitated via abrasion of corals by plastics, allowing  
74 pathogen invasion, reducing coral resources for immune function due to wound healing and by  
75 the colonization of plastics by pathogens. Poor water quality due to dredging (Pollock et al.  
76 2014), land-based runoff (Haapkylä et al. 2011, Sheridan et al. 2014, Aeby et al. 2016) and  
77 eutrophication (Bruno et al. 2003, Vega Thurber et al. 2014) have also been implicated in  
78 increasing disease prevalence. A common component of all these studies is the link between  
79 increases in coral disease and changes in environmental conditions due to anthropogenic actions.

80         Dark spot disease (DSD), also called dark spot syndrome (DSS), is a disease reported on  
81 coral reefs across the wider Caribbean (23 countries) affecting 16 coral species (Work and Weil  
82 2016). DSD manifests as multi-focal or coalescing spots of dark discolored tissue (Fig. 1) that  
83 can spread across the coral surface causing slow, progressive tissue loss (Work and Weil 2016).  
84 DSD lesions can also resolve and appear elsewhere on the colony (Borger et al. 2005, Gochfeld  
85 et al. 2006, Porter et al. 2011) and it is unclear what determines the trajectory of these lesions.  
86 Corals with similar disease signs from the Indo-Pacific (Work et al. 2008, 2014), and Red Sea  
87 (Aeby et al. 2021) are termed ‘endolithic hypermycosis’ as histopathology showed the lesions to  
88 be associated with endolithic fungal infections (Work et al. 2008, 2014, Aeby et al. 2021). In the  
89 Caribbean, the etiology of DSD is less certain, although Renegar et al. (2008) also found  
90 endolithic fungal infections in DSD-affected *Siderastrea siderea*. Cervino et al. (2001) showed  
91 that zooxanthellae from dark spot lesions are swollen and darker in pigment and suggested DSD  
92 was primarily a disease of the symbiotic zooxanthellae found within the coral host. The lesions



116 We used data collected as part of the Florida Reef Resilience Program's (FRRP)  
117 Disturbance Response Monitoring program (DRM)(<https://frrp.org/coral-monitoring/>). The  
118 DRM uses a stratified random sample design across 28 discrete reef zones in 10 sub-regions. At  
119 each site, two independent 1x10 m belt transects were surveyed with all coral colonies with  
120 maximum diameter >4 cm identified to species, enumerated and their health condition noted.  
121 The data are available from Florida Fish and Wildlife Conservation Commission  
122 (<https://ocean.floridamarine.org/FRRP/>). For our analysis, we used a subset of the data, limiting  
123 it to the coral species, *Siderastrea siderea*, and the health condition noted as dark spot disease  
124 (DSD). *Siderastrea siderea* is a widespread species occurring across the Caribbean and is the  
125 most common host of DSD (Gil-Agudelo et al. 2001, Weil 2004, Borger 2005, Gochfeld et al.  
126 2006, Voss and Richardson 2006). The dataset used in our analyses included 2,442 individual  
127 surveys from 2005-2019, excluding 2017 due to low and inconsistent sampling effort that year  
128 after hurricane Irma (Fig. 2). From these data, we calculated DSD prevalence, defined as the  
129 percent of colonies surveyed at each site that had a lesion consistent with DSD. Overall  
130 prevalence was the proportion of all colonies that had a DSD lesion (all surveys combined).  
131 Frequency of disease occurrence (FOC) is a measure of spatial distribution of disease and was  
132 calculated as the percentage of survey sites having one or more coral colonies manifesting DSD.  
133 There was a large increase in DSD prevalence and FOC between the time periods 2005-2012 and  
134 2013-2019. Thus, differences in DSD prevalence and FOC between these two time periods were  
135 analyzed using a non-parametric Wilcoxon 2-group test (JMP vers. 16.1). Regional patterns of  
136 DSD prevalence were explored with the data split by ecoregions (Walker 2012). Differences in  
137 DSD prevalence among ecoregions was examined by calculating the cumulative average DSD  
138 prevalence for surveys between 2005 and 2019 (excluding 2017).

139

140 ***Modeling the association between DSD and multiple human and environmental factors***

141         Several predictor variables were hypothesized to be linked to spatial variations in coral  
142 disease dynamics, including 1) The Nature Conservancy’s (TNC) Ocean Wealth Index (as a  
143 proxy of reef “use” by people), 2) impacts from wastewater, 3) septic tanks (potential sewage  
144 pollution), 4) local land use (as indicators of local coastal development), 5) water quality, and 6)  
145 local human populations. These were quantified using multiple data sources and across a range  
146 of scales and combined with measurements of survey depth, susceptible coral host abundance  
147 (i.e., the abundance of *Siderastrea siderea* at each site), survey year (to account for the  
148 temporally variable nature of the disease data), and a range of metrics that captured variations in  
149 reef habitat (Table 1). For our predictor groups 1– 6 above, we quantified each of them within a  
150 radial distance of each survey location over a range of spatial scales using a Fibonacci sequence  
151 (1, 2, 3, 5, 8, 13 and 21 km). We examined different spatial scales for predictor variables as the  
152 influence each predictor might have on our disease response variable could vary depending on  
153 their distance from the survey locations. In some cases, there was a high proportion of missing  
154 data values over the smaller scales for some predictor variables (e.g., 1 – 8km for water quality  
155 estimates and wastewater treatment facilities) so these scales were excluded prior to model  
156 fitting (see Table 1 for the final scales included for each predictor). Despite our knowledge that  
157 some of the predictor variables were likely collinear, given the large replication in the response  
158 variable, and the fact that our chosen modeling framework (see below) is robust to the inclusion  
159 of spurious predictors, we included all predictors in the model fitting process.

160         Statistical models were built using a boosted regression tree (BRT) framework to test the  
161 ability of the predictor variables to explain variation in the number of diseased coral hosts across



162 the Florida Reef Tract. Unlike many modeling techniques that aim to fit a single parsimonious  
163 model, BRT incorporates machine learning decision tree methods (Breiman et al. 1984) and  
164 boosting, a method to reduce predictive error (Elith et al. 2008), to build an additive regression  
165 model in which individual terms are regression trees, fitted in a forward stage-wise manner (i.e.,  
166 sequentially fitting each new tree to the residuals from the previous ones). In summary, BRT  
167 gives two crucial pieces of information, namely the underlying relationship between the response  
168 and each predictor, and the strongest statistical predictor (among the simultaneously tested  
169 predictors) of the response variable in question. Due to their flexible use and improved  
170 predictive power, the use of BRTs has increased over recent years to model non-linear ecological  
171 relationships at a range of spatial scales including coral reef-environment associations (e.g.,  
172 Williams et al. 2010, Gove et al. 2015, Aston et al. 2019).

173 BRTs were constructed using the *gbm.step* routine (Elith et al. 2008) in the *dismo* package  
174 (Hijmans et al. 2017) for R ([www.r-project.org](http://www.r-project.org)) and all model outputs were visualized in *ggplot2*  
175 using *ggBRT* (Jouffray et al. 2019). The data were modeled using a Poisson distribution, as our  
176 response variable was the number of corals in each survey location showing signs of DSD (i.e.,  
177 number of disease cases). Number of disease cases was used rather than disease prevalence (#  
178 cases/total # colonies surveyed) as host abundance was already included in the model. We used  
179 a 10-fold cross-validation approach to test the model against withheld portions of the data  
180 (iterated thousands of times) and the cross-validated percentage deviance explained, calculated  
181 as  $(1 - (\text{cross-validated deviance}/\text{mean total deviance}))$ , as our measure of model performance  
182 (Jouffray et al. 2019). To optimize model predictive performance, we varied three core  
183 parameters of the BRT algorithm: the *bag-fraction* (bf, proportion of data to be selected at each  
184 step), the *learning rate* (lr, used to shrink the contribution of each tree as it is added to the

185 model), and the *tree complexity* (tc the number of terminal nodes in a tree). Using a customized  
186 loop routine (Richards et al. 2012), we identified the combination of these three parameters that  
187 resulted in the lowest cross-validation deviance (CVD) over bf-values 0.5, 0.7, and 0.8, lr-values  
188 0.001, 0.0001, and 0.00001, and tc-values 1–5, while maintaining a minimum of  $\geq 1000$  fitted  
189 trees and a maximum of 50,000 trees. This identified the following optimal model parameter  
190 settings: bf = 0.8, lr = 0.001, tc = 2. We calculated the relative importance of each predictor  
191 based on the number of times a variable was selected for splitting, weighted by the squared  
192 improvement to the model as a result of each split, and averaged over all trees (Friedman and  
193 Meulman 2003, Elith et al. 2008). To assess the relative contribution of each predictor, we only  
194 considered predictors with a relative influence above that expected by chance (100/number of  
195 variables) (Muller et al. 2013) and then rescaled their influence to 100%. The interactive effect  
196 of the most influential predictors was then visualized using the *ggInteract\_3D* function within  
197 *ggBRT*. All input data (DSD.csv) and R code (DSD\_Cases\_BRT\_Florida.R) used to execute our  
198 BRT models are provided as Supplementary Materials.

199

200

## RESULTS

201

### *DSD prevalence through time within S. siderea populations across the Florida Reef Tract*

202

Between 2005 and 2019, mean ( $\pm 1$ SE) DSD prevalence among all sites and all years was

203

2% ( $\pm 0.4\%$ ). DSD was observed in every year of the survey period (14 years) and mean DSD

204

prevalence ranged from a low of 0.45% in 2006 to a high of 4.4% in 2016. Overall, DSD

205

prevalence (all sites combined) increased through time, with mean DSD prevalence before 2013

206

(2005-2012) equaling 1.1% ( $\pm 0.17\%$ ) compared to 2.9% ( $\pm 0.27\%$ ) from 2013-2019 (Fig. 3)

207

(Wilcoxon 2-group test,  $Z=2.9$ ,  $p=0.0037$ ).

208

209 ***Distribution of DSD through time across the Florida Reef Tract***

210           Region-wide distribution of DSD within our study period was 14.6% with 357 of the  
211 2,442 survey sites having colonies with DSD signs (all years combined). Frequency of  
212 occurrence ranged from a low of 4.8% (n=84 sites) in 2005 to a high of 30.9% (n=188 sites) in  
213 2018 (Fig. 4). Through time, DSD occurrence increased from 7.4% ( $\pm 0.8\%$ ) of survey sites  
214 between 2005 and 2012 to an average of 22.1% ( $\pm 2.8\%$ ) of survey sites between 2013-2019  
215 (Wilcoxon 2-group test,  $Z=2.9$ ,  $p=0.0037$ ).

216

217 ***Association between DSD and suspected human and environmental drivers***

218           Our predictive model explained 64.4% of the underlying spatial deviance in the number of  
219 DSD cases across the entire dataset (Table 2). The top three predictors contributed 45.5% of the  
220 relative influence within the model and included the concentration of silica in the surface waters  
221 within a 13 km radius (proxy for freshwater input), the total number of susceptible coral hosts (*S.*  
222 *siderea*), and the number of septic areas within an 8 km radius. When predicting to new data from  
223 the training data, 47.1% of the cross-validated percentage deviance was explained, suggesting a  
224 reasonable ability to predict to new data. The probability of DSD occurrence increased as both  
225 surface water silica concentration and coral host abundances increased and became maximized  
226 where silica exceeded 1.15  $\mu\text{M/L}$  and where there were  $>100$  coral hosts within the survey area  
227 (Fig. 5A). DSD occurrence was also higher in areas with  $>13 \text{ km}^2$  septic areas within 8 km  
228 regardless of coral host abundance but was highest where there were also  $>100$  coral hosts (Fig.  
229 5B).

230 Other significant variables in the model were year of the study (consistent with our  
231 findings of increased DSD levels through time), survey depth, habitat characterization (URM  
232 ClassLv4, Reef Zone)(<https://myfwc.com/research/gis/regional-projects/unified-reef-map/>),  
233 measures of water quality, total carbon in surface waters within 8 km and in bottom waters  
234 within 21 km, and total phosphorous in bottom waters within 21 km, and surface waters within 8  
235 km, distance to pollution sources (number of sewer systems within 21 km and number of septic  
236 systems within 8 km) and variables associated with human impacts (amount of spending within 1  
237 and 5 km which indicates degree of human presence and degree of urbanization within 8 km  
238 which could influence coastal runoff and pollution).

239

240 ***Regional distribution of DSD and the association with predictor variables across the Florida***  
241 ***Reef Tract***

242 *S. siderea* colonies with DSD were found throughout the Florida Reef Tract but there was  
243 a distinct spatial clustering of DSD colonies on reefs around the upper Keys (Fig. 6) where there  
244 was a threshold concentration of silica in the water and a threshold area of septic systems within  
245 8 km of the survey sites (Fig. 7).

246

247 **DISCUSSION**

248 In the first long-term study examining dark spot disease (DSD) occurrence across the  
249 Florida Reef Tract, DSD in *Siderastrea siderea* was found to be a chronic disease occurring in  
250 the annual monitoring surveys in all 14 years of the study. Between 2005 and 2019, *S. siderea*  
251 DSD prevalence ranged from less than 0.5% to over 4% which is less than from prior studies in  
252 Florida and regions in the Caribbean. In Florida, DSD prevalence in *S. siderea* was found to be

253 between 4.3% to 13.3% (Porter et al. 2011), and in the Caribbean reported studies found 25% in  
254 Columbia (Gil-Agudelo & Garzon-Ferreira 2001), 53% in Bonaire, 58% in Turks and Caicos,  
255 and 42% in Grenada (Cervino et al. 2001). We only found an average DSD prevalence of 2%  
256 (all years combined) but the spatial and temporal coverage from the number of surveys we  
257 examined (2508 surveys across 15 years) far exceeded the spatial and temporal scale of other  
258 studies which might explain these differences. Indeed, we did find higher DSD prevalence at  
259 smaller spatial and temporal scales. As example, in the upper Florida Keys, DSD prevalence  
260 was 19.4% in 2014 and 22% in 2016. In addition, human and environmental variables are also  
261 important factors underlying disease occurrence which also vary in time and space.

262 DSD became more widespread across Florida's coral reefs and affected a higher  
263 proportion of *S. siderea* populations through time. DSD was found in 7% of the total surveys  
264 between 2005 and 2012 but occurred in 22% of the surveys between 2013 and 2019. Similarly,  
265 average DSD prevalence was approximately 1% of the surveyed corals from 2005 to 2012 but  
266 affected nearly 3% of *S. siderea* colonies thereafter. As with most coral diseases, pinpointing a  
267 cause to this increase in DSD occurrence through time is challenging. We ruled out changes in  
268 methods or field personnel and there were no obvious acute environmental events (bleaching,  
269 hurricanes, annual rainfall) that could explain changing disease levels. Previous studies have  
270 shown that DSD is dynamic, with colonies recovering and becoming reinfected over time  
271 (Gochfeld et al. 2006, Porter et al. 2011). However, our data indicate a persistent and increasing  
272 annual DSD prevalence from 2012 – 2019, indicative of a chronic source or stressor. Florida's  
273 coral reefs have been increasingly impacted by humans with densely populated coastlines, higher  
274 visitor numbers, and chronic polluted terrestrial run-off for many decades (Jackson et al. 2014).  
275 Florida has also suffered an acceleration of both acute and chronic thermal stress over the past 30

276 years with reefs experiencing six mass bleaching events since the late 1980s (Manzello 2015)  
277 with the most recent event occurring in 2014 (Eakin et al. 2019, van Woesik and McCaffrey  
278 2017). It is plausible that Florida's coral reefs have reached the limit of their ability to tolerate  
279 multiple, possibly additive or synergistic stressors which may be affecting the coral's ability to  
280 resist disease.

281         Statistical modeling found host density, indicators of freshwater input, and proximity to  
282 septic systems were the strongest predictors of DSD, suggesting that environmental and  
283 anthropogenic stress are important drivers of this disease across the Florida Reef Tract. It must  
284 be noted that other factors may be important in determining DSD levels in regions where  
285 pollution is less problematic. Identifying the cause of DSD will help in teasing out all relevant  
286 environmental co-factors. However, for Florida's reefs polluted waters have been identified as a  
287 major problem for decades (Zhao et al. 2013). Florida's long history of coastal pollution has  
288 resulted in harmful algal blooms, seagrass die-offs, and declining coral reefs, with wastewater  
289 and on-site sewage treatment and disposal systems (OSTDS) representing major nitrogen sources  
290 contributing to ecosystem impacts (Lapointe et al. 2004, 2015). In addition to excess nitrogen,  
291 sewage pollution also results in coastal influxes of freshwater, phosphorous, pathogens,  
292 endocrine disrupters, suspended solids, sediments, heavy metals, and other toxicants (Wear and  
293 Vega Thurber 2015). Evidence of sewage pollution is common in Florida's nearshore  
294 environments with human fecal contamination of nearshore or offshore coral reefs found along  
295 the Florida Reef Tract (Lipp et al. 2002, Futch et al. 2010, 2011, Staley et al. 2017). Sewage can  
296 also act as a reservoir for potential pathogens as evidenced by Sutherland et al. (2011) who found  
297 human wastewater to be a source of the pathogen causing white pox disease in the reef coral,  
298 *Acropora palmata*. Additionally, freshwater stress alone can increase a coral's susceptibility to

299 bacterial infections (Shore-Maggio et al. 2018) and numerous field studies show a link between  
300 rainfall, terrestrial run-off, and disease outbreaks (Haapkylä et al. 2011, Sheridan et al. 2014,  
301 Aeby et al. 2016).

302         It is not usually a single stressor that results in the emergence of disease but multiple co-  
303 factors. Coastal runoff exposes nearshore coral reefs to sediments, freshwater, excess nutrients,  
304 and other pollutants. Other studies support the importance of multiple stressors impacting coral  
305 reefs. Oberle et al. (2019) found a combination of nutrient loading and low salinities from  
306 groundwater discharge explained an outbreak of black band disease on coral reefs in Hawaii.  
307 Considering the long-term problems Florida has had with polluted waters, it is not surprising that  
308 we found a link between DSD levels, proximity to sewage sources and indicators of land-based  
309 freshwater influx. Similarly, Porter et al. (2001) showed a five-fold increase in number of survey  
310 stations with disease over time in the Florida Keys and this trend was especially evident on reefs  
311 close to the population centers of Key West and Key Largo. In additional support of our  
312 findings, Vega Thurber et al. (2014) showed a direct link between excess nitrogen and DSD  
313 development in *S. siderea*. In an *in situ* nutrient enrichment experiment, they found *S. siderea* in  
314 treatment plots, had a 100% increase in DSD compared to corals in control plots.

315         Spatial analyses among regions showed that the progressively higher DSD prevalence  
316 through time, was especially prominent in the upper Florida Keys. The upper Florida Keys were  
317 also associated with higher levels of silica indicating an influx of freshwater, and larger septic  
318 areas, two of the top predictor variables explaining DSD occurrence along the Florida Reef  
319 Tract. Several changes in local water management have occurred around the upper Keys since  
320 2012 that might have influenced the amount of freshwater, and thus levels of DSD, occurring in  
321 that region, namely the development of wastewater management systems, and changes in the

322 management of the Everglades canal system. Wastewater is recognized as a historic source of  
323 excess nutrients into local Florida waters (Barreras et al. 2019) and billions of dollars have been  
324 spent developing wastewater management systems to convert septic systems to sewer. In the  
325 upper Keys, there are two water management districts, Key Largo Wastewater Treatment District  
326 (KLWTD) and the North Key Largo Utility Corporation (Barreras et al 2019). The KLWTD  
327 advanced wastewater treatment system was completed in 2010 and effluent from this system is  
328 deep-well injected to reduce pollution in nearshore marine waters (Reich et al. 2002, Chanton et  
329 al. 2003). Implementation of these systems has shown improvements in water local quality  
330 (Barreras et al. 2019, Kelly et al. 2021) so wastewater seems a less likely cause of increased  
331 disease within this region. However, deep-well injected wastewater on the island of Maui in  
332 Hawaii was found to be a significant source of eutrophication on adjacent coral reefs,  
333 contributing to their decline (Dailer et al. 2010, Murray et al. 2019).

334         Conversely, freshwater from the heavily managed and controlled Everglades canal  
335 system has been shown to affect much of the Florida Keys coral reef system (Lapointe et al  
336 2019). This water contains runoff from both urban and agriculture areas that can result in ‘black  
337 water’ events where patches of dark colored water develop due to high concentrations of  
338 phytoplankton and colored dissolved organic matter (Hu 2004, Zhao et al. 2013). Lapointe et al.  
339 (2019) implicate this water as the main source of enriched nutrients to the reef system and a  
340 primary driver of coral reef decline in the lower Florida Keys over the past few decades. Given  
341 that increasing the freshwater flows from the Everglades has been a priority for managers for  
342 decades (NOAA 1996), it is also possible that these flows are increasing DSD. Clearly, more  
343 research is needed to understand the source and dynamics of DSD in the Florida Keys.



344 Host density was also a significant factor in explaining DSD occurrence across the  
345 Florida Reef Tract. A positive relationship between host abundance and disease has been  
346 demonstrated for numerous host-pathogen systems (Altizer and Augustine 1997, McCallum et al.  
347 2004, Lafferty 2004) including DSD (Borger and Steiner 2005) and other coral diseases (Bruno  
348 et al. 2007, Haapkylä et al. 2009, Aeby et al. 2010). High host density can result in increased  
349 transmission of disease throughout a population as distance between individuals decreases.  
350 However, this would only apply toward infectious diseases, and it is not known whether DSD is  
351 infectious. DSD is not well studied, but to date, there is no evidence that DSD is infectious  
352 (Randall et al. 2016). However, if DSD were an environmentally induced disease, then host  
353 density could still play a role in predicting disease occurrence (e.g., more corals exposed to the  
354 adverse environmental conditions would lead to higher local disease levels).

355 On the Florida Reef Tract, *S. siderea* densities have been increasing through time (Hayes  
356 et al. 2022), and it is thought that weedy species, such as *S. siderea*, are becoming more common  
357 due to the declines of historically dominant species (Burman et al. 2012, Jones et al. 2020). The  
358 species that have declined the most were also the most stenotopic species, i.e., those able to  
359 tolerate a narrower range of environmental conditions (Burman et al. 2012). Increasing thermal  
360 stress events combined with multiple disease outbreaks have led to this shift in coral community  
361 structure (Burman et al. 2012, Jones et al. 2020). Understanding disease in *S. siderea* is  
362 becoming even more important as its numerical dominance increases throughout the Florida Reef  
363 Tract.

364 Several studies have suggested DSD is caused by the overgrowth of endolithic fungi  
365 (Work et al. 2008, 2014, Renegar et al. 2008) and it is plausible that DSD is an environmentally  
366 induced disease whereby environmental stressors are disrupting the coral holobiont facilitating

367 fungal overgrowth. If so, the environmental stressors and thus underlying drivers of DSD may  
368 also differ between regions explaining why pollution may be important in driving DSD on  
369 Florida's reefs but perhaps not in other regions. Corals live in a dynamic partnership between  
370 the cnidarian host, its endosymbiotic dinoflagellates, and a suite of microbiota that includes  
371 archaea, bacteria, viruses, fungi, and endolithic algae (Rosenberg et al. 2007, Bourne et al. 2009,  
372 Boilard et al. 2020). Environmental conditions that disrupt the delicate balance among partners  
373 create imbalances or dysbiosis which can result in disease (Lesser et al. 2007, Bourne et al. 2009,  
374 Vega Thurber et al. 2020). MacKnight et al. (2021) examined the link between microbial  
375 dysbiosis and disease susceptibility of seven Caribbean coral species exposed to white plague  
376 disease. They found that coral species that showed comparatively higher microbial dysbiosis had  
377 the greatest susceptibility to infection upon exposure to white plague. Diseases in marine  
378 organisms are more often due to dysbiosis and the emergence of opportunistic pathogens rather  
379 than infection by a single pathogen (Lesser et al. 2007, Burge 2014, Egan et al. 2014. Egan and  
380 Gardner 2016). In other cases, pathogens can live as a component of a host's normal  
381 microbiome but under certain conditions switch from a commensal to a pathogen. For example,  
382 the marine bacterium *Phaeobacter gallaeciensis* BS107 is a commensal on the microalgae,  
383 *Emiliania huxleyi*, and produces compounds that are beneficial to its algal host (Seyedsayamdost  
384 et al. 2011). However, in response to p-coumaric acid, a breakdown product symptomatic of  
385 aging algae, *Phaeobacter gallaeciensis* BS107 initiates production of potent algaecides  
386 becoming an opportunistic pathogen of its algal host. Whether or not DSD is caused by a  
387 disruption of the symbiosis between the coral host and its fungal endoliths will require further  
388 research on the ecology and etiology of this disease.

389           Interestingly, human population densities were not significant in explaining DSD  
390 occurrence unlike studies of other coral diseases (e.g., growth anomalies: Aeby et al. 2011b)  
391 where human populations were associated with disease. Instead, the amount of spending was an  
392 important factor reflecting Florida’s resident population as well as its extensive visitor numbers.  
393 Florida’s resident population is around 22 million ([https://www.populationu.com/us/florida-](https://www.populationu.com/us/florida-population)  
394 [population](https://www.populationu.com/us/florida-population)) whereas it hosted 122 million visitors in 2021  
395 (<https://www.visitflorida.org/resources/research/>). For Florida, this metric (spending) appears to  
396 be a better indicator of potential degree of human impacts rather than simply resident human  
397 population. Other important variables identified in the model (measures of water quality,  
398 distance to pollution sources, degree of urbanization that influence coastal runoff and pollution)  
399 confirm anthropogenic impacts as a major influence on coral disease occurrence.

400           Florida’s coral reefs have declined over the past several decades with increased coastal  
401 populations, high visitor numbers, overfishing, and impaired water quality contributing to their  
402 decline. Florida is also considered a “coral disease hotspot” with disease outbreaks occurring  
403 repeatedly since the 1970s (Dustan 1977, Richardson 1998, Green and Bruckner 2000, Precht et  
404 al. 2016, van Woesik and McCaffrey 2017). The downward trend of Florida’s coral reefs  
405 continues with a recent study on southeast Florida coral populations by Walton et al. (2018) who  
406 found region-wide declines in coral diversity, density and live tissue area which was attributed to  
407 disease. Our study documents the long-term persistence of disease in *S. siderea*, a common reef  
408 coral across Florida, and our modeling results suggest that coastal water quality is a key  
409 component to coral reef disease management, including maximizing resiliency in the face of  
410 global climate change. Disease not only impact host populations but can also result in  
411 ecosystem-wide impacts due to mortality of keystone species (Burge et al. 2014). Diseases are

412 predicted to increase with global climate change and anthropogenic pressures (Gattuso et al.  
413 2015, Maynard et al. 2015), placing more emphasis on the need to further understand drivers of  
414 marine diseases.

415

416

## ACKNOWLEDGEMENTS

417 We thank the Florida Department of Environmental Protection’s Office of Resilience and  
418 Coastal Protection and to NOAA's National Centers for Coastal Ocean Science for supporting  
419 these efforts. Thank you to J Stein and the Florida Fish and Wildlife Conservation Commission  
420 Fish and Wildlife Research Institute’s Florida Reef Resilience Program Disturbance Response  
421 Monitoring for facilitating data requests and advice. Thanks to J Maynard and D Tracey at  
422 SymbioSeas for help with data analysis and GIS methods development. Funding was provided by  
423 the Florida Department of Environmental Protection Award No. B7B6F3. The manuscript was  
424 greatly improved by comments from C. Woodley, E. Davenport and J. Guyon. The views,  
425 statements, findings, conclusions, and recommendations expressed herein are those of the authors  
426 and do not necessarily reflect the views of the State of Florida or any of its sub-agencies.

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## LITERATURE CITED

429

430 Aeby GS, Ross M, Williams GJ, Lewis TD, Work TM. 2010. Disease dynamics of *Montipora*  
431 white syndrome within Kaneohe Bay, Oahu, Hawaii: distribution, seasonality, virulence,  
432 and transmissibility. *Dis Aquat Org* 91:1–8

433 Aeby GS, Bourne DG, Wilson B, Work TM. 2011a. Coral diversity and the severity of disease

434 outbreaks: A cross-regional comparison of *Acropora* white syndrome in a species-rich

435 region (American Samoa) with a species-poor region (Northwestern Hawaiian Islands). J  
436 Mar Biol 2011:1–8

437 Aeby GS, Williams GJ, Franklin EC, Haapkyla J, Harvell CD, Neale S, Page CA, Raymundo L,  
438 Vargas-Ángel B, Willis BL, et al. 2011b. Growth anomalies on the coral genera  
439 *Acropora* and *Porites* are strongly associated with host density and human population  
440 size across the Indo-Pacific. PLoS ONE 6:e16887

441 Aeby GS, Work TM, Runyon CM, Shore-Maggio A, Ushijima B, Videau P, Beurmann S,  
442 Callahan SM. 2015. First record of black band disease in the Hawaiian archipelago:  
443 response, outbreak status, virulence, and a method of treatment. PLoS ONE 10:e0120853

444 Aeby GS, Callahan S, Cox E, Runyon C, Smith A, Stanton F, Ushijima B, Work T. 2016.  
445 Emerging coral diseases in Kāneʻohe Bay, Oʻahu, Hawaiʻi (USA): two major disease  
446 outbreaks of acute *Montipora* white syndrome. Dis Aquat Org 119:189–198

447 Aeby GS, Howells E, Work T, Abrego D, Williams GJ, Wedding LM, Caldwell JM, Moritsch  
448 M, Burt JA. 2020. Localized outbreaks of coral disease on Arabian reefs are linked to  
449 extreme temperatures and environmental stressors. Coral Reefs 39:829–846

450 Aeby GS, Shore A, Jensen T, Ziegler M, Work T, Voolstra CR. 2021. A comparative baseline of  
451 coral disease in three regions along the Saudi Arabian coast of the central Red Sea. PLoS  
452 ONE 16:e0246854

453 Altizer SM, Augustine DJ. 1997. Interactions between frequency–dependent and vertical  
454 transmission in host–parasite systems. Proc R Soc Lond B 264:807–814

455 Alvarez-Filip L, Estrada-Saldívar N, Pérez-Cervantes E, Molina-Hernández A, González-Barrios  
456 FJ. 2019. A rapid spread of the stony coral tissue loss disease outbreak in the Mexican  
457 Caribbean. *PeerJ* 7:e8069

458 Aston E, Williams GJ, Green J, Davies A, Wedding L, Gove J, Jouffray J, Jones T, Clark J.  
459 2019. Scale-dependent spatial patterns in benthic communities around a tropical island  
460 seascape. *Ecography* 42:578–590

461 Barreras H, Kelly EA, Kumar N, Solo-Gabriele HM. 2019. Assessment of local and regional  
462 strategies to control bacteria levels at beaches with consideration of impacts from climate  
463 change. *Mar Pollut Bull* 138:249–259

464 Boilard A, Dubé CE, Gruet C, Mercière A, Hernandez-Agreda A, Derome N. 2020. Defining  
465 coral bleaching as a microbial dysbiosis within the coral holobiont. *Microorg* 8:1682

466 Borger JL. 2005. Dark spot syndrome: a scleractinian coral disease or a general stress response?  
467 *Coral Reefs* 24:139–144

468 Bourne DG, Ainsworth TD, Pollock FJ, Willis BL. 2014. Towards a better understanding of  
469 white syndromes and their causes on Indo-Pacific coral reefs. *Coral Reefs* 34:233–242

470 Bourne DG, Garren M, Work TM, Rosenberg E, Smith GW, Harvell CD. 2009. Microbial  
471 disease and the coral holobiont. *Trends Microbiol* 17:554–562

472 Brandt ME, Ruttenger BI, Waara R, Miller J, Witcher B, Estep AJ, Patterson M. 2012.  
473 Dynamics of an acute coral disease outbreak associated with the macroalgae *Dictyota*  
474 spp. in Dry Tortugas National Park, Florida, USA. *Bull Mar Sci* 88:1035–1050

475 Breiman L, Friedman, J, Olshen A, Stone C. 1984. Classification and Regression Trees.  
476 Belmont, CA: Wadsworth. International Group 432:151-166

477 Brodnicke OB, Bourne DG, Heron SF, Pears RJ, Stella JS, Smith HA, Willis BL. 2019.  
478 Unravelling the links between heat stress, bleaching and disease: fate of tabular corals  
479 following a combined disease and bleaching event. *Coral Reefs* 38:591–603

480 Bruno JF. 2015. The coral disease triangle. *Nature Clim Change* 5:302–303

481 Bruno JF, Petes LE, Drew Harvell C, Hettinger A. 2003. Nutrient enrichment can increase the  
482 severity of coral diseases. *Ecol Letters* 6:1056–1061

483 Bruno JF, Selig ER, Casey KS, Page CA, Willis BL, Harvell CD, Sweatman H, Melendy AM.  
484 2007. Thermal stress and coral cover as drivers of coral disease outbreaks. *PLoS Biol*  
485 5:e124

486 Burge CA, Eakin M, Friedman CS, Froelich B, Hershberger PK, Hofmann EE, Petes LE, Prager  
487 KC, Weil E, Willis BL, et al. 2014. Climate change influences on marine infectious  
488 diseases: implications for management and society. *Annu Rev Mar Sci* 6:249–277

489 Burman S, Aronson R, van Woesik, R. 2012. Biotic homogenization of coral  
490 assemblages along the Florida reef tract. *Mar Ecol Prog Ser*, 467, 89-96.  
491 doi:10.3354/meps09950

492 Carpenter KE, Abrar M, Aeby G, Aronson RB, Banks S, Bruckner A, Chiriboga A, Cortés J,  
493 Delbeek JC, DeVantier L, et al. 2008. One-third of reef-building corals face elevated  
494 extinction risk from climate change and local impacts. *Science* 321:560–563

495 Cervino J, Goreau TJ, Nagelkerken I, Smith GW, Hayes R. 2001. Yellow band and dark spot  
496 syndromes in Caribbean corals: distribution, rate of spread, cytology, and effects on  
497 abundance and division rate of zooxanthellae. *In*: Porter JW, editor. *The Ecology and*  
498 *Etiology of Newly Emerging Marine Diseases*. Springer Netherlands, Dordrecht, p. 53–  
499 63

500 Chanton J P, Burnett, WC, Dulaiova H, Corbett, DR, Taniguchi M. 2003. Seepage Rate  
501 Variability in Florida Bay Driven by Atlantic Tidal Height. *Biogeochem*, 66:87-202

502 Cróquer A, Weil E. 2009. Changes in Caribbean coral disease prevalence after the 2005  
503 bleaching event. *Dis Aquat Org* 87:33–43

504 Cróquer A, Weil E, Rogers CS. 2021. Similarities and differences between two deadly Caribbean  
505 coral diseases: white plague and stony coral tissue loss disease. *Front Mar Sci* 8:709544

506 Dahlgren C, Pizarro V, Sherman K, Greene W, Oliver J. 2021. Spatial and temporal patterns of  
507 stony coral tissue loss disease outbreaks in the Bahamas. *Front Mar Sci* 8:682114

508 Dailer ML, Knox RS, Smith JE, Napier M, Smith CM. 2010. Using  $\delta^{15}\text{N}$  values in algal tissue  
509 to map locations and potential sources of anthropogenic nutrient inputs on the island of  
510 Maui, Hawai‘i, USA. *Mar Pollut Bull* 60:655–671

511 Daszak P. 2000. Emerging infectious diseases of wildlife-- threats to biodiversity and human  
512 health. *Science* 287:443–449

513 Daszak P, Cunningham AA, Hyatt AD. 2001. Anthropogenic environmental change and the  
514 emergence of infectious diseases in wildlife. *Acta Tropica* 78:103–116



515 Dobson A, Foufopoulos J. 2001. Emerging infectious pathogens of wildlife. *Phil Trans R Soc*  
516 *Lond B* 356:1001–1012

517 Dustan P. 1977. Vitality of reef coral populations off Key Largo, Florida: Recruitment and  
518 mortality. *Environ Geol* 2:51–58

519 Eakin CM, Sweatman HPA, Brainard RE. 2019. The 2014–2017 global-scale coral bleaching  
520 event: insights and impacts. *Coral Reefs* 38:539–545

521 Egan S, Fernandes ND, Kumar V, Gardiner M, Thomas T. 2014. Bacterial pathogens, virulence  
522 mechanism and host defense in marine macroalgae: bacterial pathogens of macroalgae.  
523 *Environ Microbiol* 16:925–938

524 Egan S, Gardiner M. 2016. Microbial dysbiosis: rethinking disease in marine ecosystems. *Front*  
525 *Microbiol* 7:991

526 Elith J, Leathwick JR, Hastie T. 2008. A working guide to boosted regression trees. *J Anim Ecol*  
527 77:802–813

528 Estrada-Saldívar N, Molina-Hernández A, Pérez-Cervantes E, Medellín-Maldonado F, González-  
529 Barrios FJ, Alvarez-Filip L. 2021. Reef-scale impacts of the stony coral tissue loss  
530 disease outbreak. *Coral Reefs* 39:861–866

531 Friedman J, Meulman J. 2003. Multiple additive regression trees with application in  
532 epidemiology. *Statistics in medicine* 22:1365–1381

533 Futch JC, Griffin DW, Lipp EK. 2010. Human enteric viruses in groundwater indicate offshore  
534 transport of human sewage to coral reefs of the upper Florida Keys. *Environ Microbiol*  
535 12:964–974

536 Futch JC, Griffin DW, Banks K, Lipp EK. 2011. Evaluation of sewage source and fate on  
537 southeast Florida coastal reefs. *Mar Pollut Bull* 62:2308–2316

538 Gattuso J, Magnan A, Bille R, Cheung W, Howes E, Joos F, Allemand D, Bopp L, Cooley S,  
539 Eakin M, et al. 2015. Contrasting futures for ocean and society from different  
540 anthropogenic CO<sub>2</sub> emissions scenarios. *Science* 349

541 Gil-Agudelo DL, Garzón-Ferreira J. 2001. Spatial and seasonal variation of dark spots disease in  
542 coral communities of the Santa Marta area (Colombian Caribbean). *Bull Mar Sci* 69:619-  
543 629

544 Gochfeld D, Olson J, Slattery M. 2006. Colony versus population variation in susceptibility and  
545 resistance to dark spot syndrome in the Caribbean coral *Siderastrea siderea*. *Dis Aquat*  
546 *Org* 69:53–65

547 Gove J, Williams G, McManus M, Clark S, Ehses J, Wedding L. 2015. Coral reef benthic  
548 regimes exhibit non-linear threshold responses to natural physical drivers. *Mar Ecol Prog*  
549 *Ser* 522:33–48

550 Green EP, Bruckner AW. 2000. The significance of coral disease epizootiology for coral reef  
551 conservation. *Biol Cons* 96:347–61

552 Haapkylä J, Unsworth RKF, Flavell M, Bourne DG, Schaffelke B, Willis BL. 2011. Seasonal  
553 rainfall and runoff promote coral disease on an inshore reef. *PLoS ONE* 6:e16893

554 Harvell CD, Jordan-Dahlgren E, Merkel S, Rosenberg E, Raymundo L, Smith G, Weil E, Willis  
555 B. 2007. Coral disease, environmental drivers, and the balance between coral and  
556 microbial associates. *Oceanography* 20:172–195

557 Harvell CD, Kim K, Burkholder JM, Colwell RR, Epstein PR, Grimes DJ, Hoffmann EE, Lipp  
558 EK. 1999. Emerging marine diseases: climate links and anthropogenic factors. *Science*  
559 285:1505–1510

560 Hayes NK, Walton CJ, Gilliam DS. 2022. Tissue loss disease outbreak significantly alters the  
561 Southeast Florida stony coral assemblage. *Front Mar Sci* 9:975894

562 Heres MM, Farmer BH, Elmer F, Hertler H. 2021. Ecological consequences of stony coral tissue  
563 loss disease in the Turks and Caicos Islands. *Coral Reefs* 40:609–624

564 Hijmans RJ, Phillips S, Leathwick J, Elith J. 2017. Dismo: species distribution modeling. R  
565 package version 1.1–4. See <https://CRAN.R-project.org/package=dismo>

566 Howells EJ, Vaughan GO, Work TM, Burt JA, Abrego D. 2020. Annual outbreaks of coral  
567 disease coincide with extreme seasonal warming. *Coral Reefs* 39:771–781

568 Hu C. 2004. Linkages between coastal runoff and the Florida Keys ecosystem: A study of a dark  
569 plume event. *Geophys Res Lett* 31:L15307

570 Jackson EJ, Donovan M, Cramer K, Lam V. 2014. Status and Trends of Caribbean Coral Reefs:  
571 1970-2012. Global Coral Reef Monitoring Network, IUCN 306

572 Jones NP, Figueiredo J, Gilliam DS. 2020. Thermal stress-related spatiotemporal variations in  
573 high-latitude coral reef benthic communities. *Coral Reefs* 39:1661–1673

574 Jouffray JB, Wedding LM, Norström AV, Donovan MK, Williams GJ, Crowder LB, Erickson  
575 AL, Friedlander AM, Graham NAJ, Gove JM, et al. 2019. Parsing human and  
576 biophysical drivers of coral reef regimes. *Proc R Soc B* 286:20182544

577 Kelly E, Gidley M, Sinigalliano C, Kumar N, Solo-Gabriele HM. 2021. Impact of wastewater  
578 infrastructure improvements on beach water fecal indicator bacteria levels in Monroe  
579 County, Florida. *Science of The Total Environment* 763:143024

580 Lafferty KD. 2004. Fishing for lobsters indirectly increases epidemics in sea urchins. *Ecol App*  
581 14:1566–1573

582 Lafferty KD, Holt RD. 2003. How should environmental stress affect the population dynamics of  
583 disease? *Ecol Lett* 6:654–664

584 Lamb JB, Willis BL, Fiorenza EA, Couch CS, Howard R, Rader DN, True JD, Kelly LA, Ahmad  
585 A, Jompa J, Harvell CD. 2018. Plastic waste associated with disease on coral reefs.  
586 *Science* 359:460–462

587 Lapointe BE, Barile PJ, Matzie WR. 2004. Anthropogenic nutrient enrichment of seagrass and  
588 coral reef communities in the Lower Florida Keys: discrimination of local versus regional  
589 nitrogen sources. *J Exp Mar Biol Ecol* 308:23–58

590 Lapointe BE, Herren LW, Debortoli DD, Vogel MA. 2015. Evidence of sewage-driven  
591 eutrophication and harmful algal blooms in Florida’s Indian River Lagoon. *Harmful*  
592 *Algae* 43:82–102

593 Lapointe BE, Brewton RA, Herren LW, Porter JW, Hu C. 2019. Nitrogen enrichment, altered  
594 stoichiometry, and coral reef decline at Looe Key, Florida Keys, USA: a 3-decade study.  
595 *Mar Biol* 166:108

596 Lesser MP, Bythell JC, Gates RD, Johnstone RW, Hoegh-Guldberg O. 2007. Are infectious  
597 diseases really killing corals? Alternative interpretations of the experimental and  
598 ecological data. *J Exp Mar Biol Ecol* 346:36–44

599 Lipp EK, Jarrell JL, Griffin DW, Lukasik J, Jacukiewicz J, Rose JB. 2002. Preliminary evidence  
600 for human fecal contamination in corals of the Florida Keys, USA. *Mar Pollut Bull*  
601 44:666–670

602 MacKnight NJ, Cobleigh K, Lasseigne D, Chaves-Fonnegra A, Gutting A, Dimos B, Antoine J,  
603 Fuess L, Ricci C, Butler C, et al. 2021. Microbial dysbiosis reflects disease resistance in  
604 diverse coral species. *Commun Biol* 4:679

605 Manzello DP. 2015. Rapid recent warming of coral reefs in the Florida Keys. *Sci Reports* 5:1–10

606 Maynard J, van Hooidonk R, Eakin CM, Puotinen M, Garren M, Williams G, Heron SF, Lamb J,  
607 Weil E, Willis B, Harvell CD. 2015. Projections of climate conditions that increase coral  
608 disease susceptibility and pathogen abundance and virulence. *Nature Clim Change*  
609 5:688–694

610 McCallum H, Kuris A, Harvell C, Lafferty K, Smith G, Porter J. 2004. Does terrestrial  
611 epidemiology apply to marine systems? *Trends Ecol Evol* 19:585–591

612 Miller J, Muller E, Rogers C, Waara R, Atkinson A, Whelan KRT, Patterson M, Witcher B.  
613 2009. Coral disease following massive bleaching in 2005 causes 60% decline in coral  
614 cover on reefs in the US Virgin Islands. *Coral Reefs* 28:925–937

615 Muller D, Leitao P, Sikor T. 2013. Comparing the determinants of cropland abandonment in  
616 Albania and Romania using boosted regression trees. *Agricultural systems* 117:66–77

617 Murray J, Prouty NG, Peek S, Paytan A. 2019. Coral skeleton  $\delta^{15}\text{N}$  as a tracer of historic  
618 nutrient loading to a coral reef in Maui, Hawaii. *Sci Rep* 9:5579

619 Mydlarz L, Couch C, Weil E, Smith G, Harvell C. 2009. Immune defenses of healthy, bleached,  
620 and diseased *Montastraea faveolata* during a natural bleaching event. *Dis Aquat Org*  
621 87:67–78

622 Myers R, Raymundo L. 2009. Coral disease in Micronesian reefs: a link between disease  
623 prevalence and host abundance. *Dis Aquat Org* 87:97–104

624 Neely KL, Lewis CL, Lunz KS, Kabay L. 2021. Rapid population decline of the pillar coral  
625 *Dendrogyra cylindrus* along the Florida Reef Tract. *Front Mar Sci* 8:656515

626 NOAA Florida Keys National Marine Sanctuary Final Management Plan/Environmental Impact  
627 Statement. 1996. <https://www.govinfo.gov/content/pkg/CZIC-qh91-75-f56-f583-1996-v-1/html/CZIC-qh91-75-f56-f583-1996-v-1.htm>  
628

629 Oberle FKJ, Storlazzi CD, Cheriton OM, Takesue RK, Hoover DJ, Logan JB, Runyon C,  
630 Kellogg CA, Johnson CD, Swarzenski PW. 2019. Physicochemical controls on zones of  
631 higher coral stress where black band disease occurs at Mākua Reef, Kaua‘i, Hawai‘i.  
632 *Front Mar Sci* 6:552

633 Patterson KL, Porter JW, Ritchie KB, Polson SW, Mueller E, Peters EC, Santavy DL, Smith  
634 GW. 2002. The etiology of white pox, a lethal disease of the Caribbean elkhorn coral,  
635 *Acropora palmata*. *Proc Nat Acad Sci* 99:8725–8730

636 Pollock FJ, Lamb JB, Field SN, Heron SF, Schaffelke B, Shedrawi G, Bourne DG, Willis BL.  
637 2014. Sediment and turbidity associated with offshore dredging increase coral disease  
638 prevalence on nearby reefs. *PLoS ONE* 9:e102498

639 Porter JW, Dustan P, Jaap WC, Patterson KL, Kosmynin V, Meier OW, Patterson ME, Parsons  
640 M. 2001. Patterns of spread of coral disease in the Florida Keys. *In*: Porter JW. Editor.  
641 *The Ecology and Etiology of Newly Emerging Marine Diseases*. Springer Netherlands,  
642 Dordrecht, p. 1–24

643 Porter JW, Torres C, Sutherland KP, Meyers MK, Callahan MK, Ruzicka R, Colella M. 2011.  
644 Prevalence, severity, lethality, and recovery of dark spots syndrome among three  
645 Floridian reef-building corals. *J Exp Mar Biol Ecol* 408:79–87

646 Precht WF, Gintert BE, Robbart ML, Fura R, van Woesik R. 2016. Unprecedented disease-  
647 related coral mortality in southeastern Florida. *Sci Rep* 6:31374

648 Raj KD, Aeby GS, Mathews G, Bharath MS, Rajesh S, Laju RL, Arasamuthu A, Kumar D,  
649 Edward JKP. 2016. Patterns in the abundance of fish and snail corallivores associated

650 with an outbreak of acute tissue loss disease on the reefs of Vaan Island in the Gulf of  
651 Mannar, India. Proc 13th Int Coral Reef Symp, Hono., HI. 85–89

652 Randall CJ, Jordán-Garza AG, Muller EM, van Woesik R. 2016. Does dark spot syndrome  
653 experimentally transmit among Caribbean corals? PLoS ONE 11:e0147493

654 Reich CD, Shinn EA, Hickey TD, Tihansky AB. 2002. Tidal and meteorological influences on  
655 shallow marine groundwater flow in the upper Florida Keys. The Everglades, Florida  
656 Bay, and coral reefs of the Florida Keys: an ecosystem sourcebook. pp 659–676

657 Renegar D-EA, Blackwelder P, Miller JD, Gochfeld DJ. 2008. Ultrastructural and histological  
658 analysis of dark spot syndrome in *Siderastrea siderea* and *Agaricia agaricites*. Proc 11th  
659 Int Coral Reef Symp, Ft. Lauderdale, FL. Session 7.

660 Richards BL, Williams ID, Vetter OJ, Williams GJ. 2012. Environmental factors affecting large-  
661 bodied coral reef fish assemblages in the Mariana Archipelago. PLoS ONE 7:e31374

662 Richardson L, Goldberg W, Carlton R, Halas J. 1998. Coral disease outbreak in the Florida  
663 Keys: Plague Type II. Rev Biol Trop 46:187–198

664 Rosenberg E, Koren O, Reshef L, Efrony R, Zilber-Rosenberg I. 2007. The role of  
665 microorganisms in coral health, disease, and evolution. Nat Rev Microbiol 5:355–362

666 Santavy DL, Mueller E, Peters EC, MacLaughlin L, Porter JW, Patterson KL, Campbell J. 2001.  
667 Quantitative assessment of coral diseases in the Florida Keys: strategy and methodology.  
668 *In:* Porter JW, editor. The Ecology and Etiology of Newly Emerging Marine Diseases.  
669 Springer Netherlands, Dordrecht, pp 39–52



670 Schrag SJ, Wiener P. 1995. Emerging infectious disease: what are the relative roles of ecology  
671 and evolution? Trends Ecol Evol 10:319–324

672 Seyedsayamdost MR, Case RJ, Kolter R, Clardy J. 2011. The Jekyll-and-Hyde chemistry of  
673 *Phaeobacter gallaeciensis*. Nature Chem 3:331–335

674 Sharp WC, Shea CP, Maxwell KE, Muller EM, Hunt JH. 2020. Evaluating the small-scale  
675 epidemiology of the stony coral tissue loss disease in the middle Florida Keys. PLoS  
676 ONE 15:e0241871

677 Sheridan C, Baele JM, Kushmaro A, Fréjaville Y, Eeckhaut I. 2014. Terrestrial runoff influences  
678 white syndrome prevalence in SW Madagascar. Mar Environ Res 101:44–51

679 Shore-Maggio A, Aeby GS, Callahan SM. 2018. Influence of salinity and sedimentation on  
680 *Vibrio* infection of the Hawaiian coral *Montipora capitata*. Dis Aquat Org 128:63–71

681 Staley C, Kaiser T, Gidley ML, Enochs IC, Jones PR, Goodwin KD, Sinigalliano CD, Sadowsky  
682 MJ, Chun CL. 2017. Differential impacts of land-based sources of pollution on the  
683 microbiota of southeast Florida coral reefs. Appl Environ Microbiol 83:e03378-16

684 Sutherland KP, Shaban S, Joyner JL, Porter JW, Lipp EK. 2011. Human pathogen shown to  
685 cause disease in the threatened elkhorn coral *Acropora palmata*. PLoS ONE 6:e23468

686 Thinesh T, Mathews G, Edward JKP. 2009. Coral disease prevalence in Mandapam group of  
687 islands, Gulf of Mannar, Southeastern India. Ind J Mar Sci 38:444-450

688 Ushijima B, Videau P, Poscablo D, Stengel JW, Beurmann S, Burger AH, Aeby GS, Callahan  
689 SM. 2016. Mutation of the *toxR* or *mshA* genes from *Vibrio coralliilyticus* strain OCN014

690 reduces infection of the coral *Acropora cytherea*: virulence genes in *Vibrio*  
691 *coralliilyticus*. Environ Microbiol 18:4055–4067

692 van Woesik R, McCaffrey KR. 2017. Repeated thermal stress, shading, and directional selection  
693 in the Florida Reef Tract. Front Mar Sci 4:182

694 Vega Thurber RL, Burkepile DE, Fuchs C, Shantz AA, McMinds R, Zaneveld JR. 2014. Chronic  
695 nutrient enrichment increases prevalence and severity of coral disease and bleaching.  
696 Glob Change Biol 20:544–554

697 Vega Thurber R, Mydlarz LD, Brandt M, Harvell D, Weil E, Raymundo L, Willis BL, Langevin  
698 S, Tracy AM, Littman R, et al. 2020. Deciphering coral disease dynamics: integrating  
699 host, microbiome, and the changing environment. Front Ecol Evol 8:575927

700 Voss J, Richardson L. 2006. Coral diseases near Lee Stocking Island, Bahamas: patterns and  
701 potential drivers. Dis Aquat Org 69:33–40

702 Walker BK. 2012. Spatial analyses of benthic habitats to define coral reef ecosystem regions and  
703 potential biogeographic boundaries along a latitudinal gradient. PLoS ONE 7:e30466

704 Walker BK, Gilliam DS. 2013. Determining the extent and characterizing coral reef habitats of  
705 the northern latitudes of the Florida Reef Tract (Martin County). PLoS ONE 8:e80439

706 Walton CJ, Hayes NK, Gilliam DS. 2018. Impacts of a regional, multi-year, multi-species coral  
707 disease outbreak in southeast Florida. Front Mar Sci 5:323

708 Wear SL, Thurber RV. 2015. Sewage pollution: mitigation is key for coral reef stewardship. Ann  
709 NY Acad Sci 1355:15–30

710 Weil E. 2004. Coral reef diseases in the wider Caribbean. *In*: Rosenberg E., Loya Y. (eds) Coral  
711 Health and Disease. Springer Berlin Heidelberg, Berlin, Heidelberg, pp 35–68

712 Williams GJ, Aeby GS, Cowie ROM, Davy SK. 2010. Predictive modeling of coral disease  
713 distribution within a reef system. PLoS ONE 5:e9264

714 Willis BL, Page CA, Dinsdale EA. 2004. Coral disease on the Great Barrier Reef. *In*: Rosenberg  
715 E, Loya Y, editors. Coral Health and Disease. Springer Berlin Heidelberg, Berlin,  
716 Heidelberg, p. 69–104

717 Work TM, Aeby GS, Stanton FG, Fenner D. 2008. Overgrowth of fungi (endolithic  
718 hypermycosis) associated with multifocal to diffuse distinct amorphous dark  
719 discoloration of corals in the Indo-Pacific. Coral Reefs 27:663–663

720 Work TM, Aeby GS, Lasne G, Tribollet A. 2014. Gross and microscopic pathology of hard and  
721 soft corals in New Caledonia. J Invert Path 120:50–58

722 Work TM, Weil E. 2015. Dark-Spots Disease. *In*: Woodley CM, Downs CA, Bruckner AW,  
723 Porter JW, Galloway SB, editors. Diseases of Coral. John Wiley and Sons, Inc, Hoboken,  
724 NJ, p. 354–360

725 Zhao J, Hu C, Lapointe B, Melo N, Johns E, Smith R. 2013. Satellite-observed black water  
726 events off southwest Florida: implications for coral reef health in the Florida Keys  
727 National Marine Sanctuary. Remote Sensing 5:415–431

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730 Table 1. Predictor variables with their description and units used to model potential human and  
 731 environmental drivers of dark spot disease (DSD) across the Florida Reef Tract.

Predictor variable	Description and units	Min	Max	Data source
Depth	Depth (m) of coral at time of survey	1.1	21.6	Recorded in situ
Host abundance	Total number of host colonies along transect	1	193	Recorded in situ
Year	Survey year	2005	2019	
Habitat characterization	Reef Zone - Thirteen mutually exclusive zones identified from shore to shelf edge corresponding to typical insular shelf and coral reef geomorphology. Examples include Reef Flat, Lagoon, Back Reef, Reef Crest, Fore Reef, Bank/Shelf, Bank/Shelf Escarpment. (Zitello et al 2009)  URM GeoForm - Distinct and non-overlapping broad geomorphological structure types that can be mapped by visual interpretation of remotely sensed imagery. Examples include Coral Reef and Hardbottom, Unconsolidated Sediment, Other Delineations, and Unknown. (Zitello et al 2009)  URM GeoFormDet - Distinct and non-overlapping more descriptive geomorphological structure types that can be mapped by visual interpretation of remotely sensed imagery. Examples include Rock Outcrop, Boulder, Spur and Groove, Individual Patch Reef, Aggregated Patch Reefs, Aggregate Reef, Reef Rubble, Pavement, Pavement with Sand Channels, Rhodoliths, Sand, Mud, Sand with Scattered Coral and Rock, Artificial, Land, and Unknown. (Zitello et al 2009)  URM Class Lv - (Lv 0-4) The most detailed classification that incorporates GeoForm and dominant live biotic cover estimates. Examples include Aggregate Reef-Algae (Continuous), Aggregate Reef-Algae (Discontinuous), Aggregate Reef-Algae (Patchy), Aggregate Reef, Live Coral (Discontinuous), Aggregate Reef-Live Coral (Patchy), Aggregate Reef- Live Coral (Sparse), Aggregate Reef-Uncolonized	NA	NA	<a href="https://myfwc.com/research/gis/regional-projects/unified-reef-map/">https://myfwc.com/research/gis/regional-projects/unified-reef-map/</a>
The Nature Conservancy Ocean Wealth Data	Mean number of coastal tourist visits that year (within 1, 2, 3, 5, 8, 13, 21 km radius) Mean spend (\$) by coastal tourists that year (within 1, 2, 3, 5, 8, 13, 21 km radius)	0	6,305	<a href="https://oceanwealth.org">https://oceanwealth.org</a>
Wastewater: proxy of human presence/influence (impacts to coastal areas)	Number of wastewater treatment facilities (within 13 and 21 km radius) Mean design capacity (millions gallons/day) of wastewater treatment facilities (within 13 and 21 km radius) Amount of wastewater (millions gallons/day) permitted to be processed (within 13 and 21 km radius)	1	57	<a href="https://geodata.dep.state.fl.us/">https://geodata.dep.state.fl.us/</a>
Septic/Sewer Areas: proxy for potential sewage pollution	Septic area (area in km <sup>2</sup> known/likely to drain to septic within 1, 2, 3, 5, 8, 13, 21 km radius) Septic count (total number of known/likely septic 'areas' that intersect 1, 2, 3, 5, 8, 13, 21 km radius) Sewer area (area in km <sup>2</sup> known/likely to drain to sewer within 1, 2, 3, 5, 8, 13, 21 km radius) Sewer count (total number of known/likely sewer 'areas' that intersect 1, 2, 3, 5, 8, 13, 21 km radius)	0	230	<a href="https://www.floridahealth.gov/environmental-health/onsite-sewage/research/fiwm/index">https://www.floridahealth.gov/environmental-health/onsite-sewage/research/fiwm/index</a> <a href="https://www10.doh.state.fl.us/pub/bsis/inventory/FloridaWaterManagementInventory">https://www10.doh.state.fl.us/pub/bsis/inventory/FloridaWaterManagementInventory</a>
Land Use: degree of urbanization (affects coastal runoff & pollution)	High intensity land use area (in km <sup>2</sup> ) = Constructed materials account for 80 – 100% of the total cover. Vegetation, if present, occupies less than 20 % of the landscape.	0	289	<a href="https://coast.noaa.gov/digitalcoast/data/geobase.html">https://coast.noaa.gov/digitalcoast/data/geobase.html</a>
Water Quality: direct measurements at water surface and reef floor	Mean total phosphorus (um/L) (within 8, 13, 21 km radius) Mean silica (um/L) (within 8, 13, 21 km radius) Total organic carbon (um/L) TN-ANTEX 9000=tot nitrogen (um/L) Chlorophyll-a (ug/L)	0.033	0.531	<a href="http://secr.flu.edu/wqmnetwork/">http://secr.flu.edu/wqmnetwork/</a>
Human population (impacts to coastal areas)	Human population count (total number within 1, 2, 3, 5, 8, 13, 21 km radius) Housing units (total number within 1, 2, 3, 5, 8, 13, 21 km radius)	0	1,564,696 709,263	<a href="https://www.census.gov/geographies/mapping-files/2010/geo/tiger-data.html">https://www.census.gov/geographies/mapping-files/2010/geo/tiger-data.html</a>

732

733 Table 2. Relative influence (%) of the 18 significant variables that together explained 64.4% of the  
 734 variation in the number of DSD cases (cross-validated percentage deviance explained = 47.1%).  
 735 See Table 1 for a more detailed description of the predictor variables.

736

Predictor	Influence (%)
mean silica (um/L) in surface waters within 13 km	18.4
total number of <i>S. siderea</i>	16.9
septic area (km <sup>2</sup> ) within 8 km	10.2
habitat characterization (URM class Lv4)	9.4
depth	8.6
year	7.1
reef zone	5.2
total organic carbon (um/L) in surface waters within 13 km	4
Chlorophyll-a (um/L) in surface waters within 21 km	3.9
total organic carbon (um/L) in surface waters within 8 km	3.8
mean total phosphorous (um/L) in bottom waters within 21 km	2.7
mean spending within 1 km	2.4
septic areas (km <sup>2</sup> ) within 21 km	1.5
total organic carbon (um/L) in bottom waters within 21 km	1.3
degree of urbanization (m <sup>2</sup> ) within 8 km	1.3
mean spending within 5 km	1.2
number of septic systems within 8 km	1.1
mean total phosphorous (um/L) in surface waters within 8 km	1.1

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745 **Figure legends**

746

747 Figure 1. Example of dark spot disease (DSD) in *Siderastrea siderea*. Dark reddish-brown areas  
748 indicate disease.

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750 Figure 2. Benthic survey locations (black dots) from 2005 to 2019 within ecoregions across  
751 Florida's Reef Tract. The table summarizes the number of sites surveyed per year.

752

753 Figure 3. Overall DSD prevalence (all sites combined for each year) through time on reefs  
754 across the Florida Reef Tract. Dotted lines indicate mean prevalence for the two time periods  
755 2005-2012 and 2013-2019.

756

757 Figure 4. Overall frequency of occurrence of DSD (all sites combined for each year) through  
758 time. Dotted lines indicate the mean FOC for the two time periods 2005-2012 and 2013-2019.

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760 Figure 5. Predicted relationship from the BRT model between *Siderastrea siderea* abundance and  
761 (A) surface silica concentration within a 13 km radius and (B) septic area within 8 km radius, on  
762 the number of DSD cases (fitted values). See Table 1 for a more detailed description of the  
763 predictor variables.

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766 Figure 6. Cumulative DSD prevalence within different ecoregions of the Florida Reef Tract.  
767 Number above each bar indicates total number of surveys conducted in each region between  
768 2005 and 2019 (excluding 2017).

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770 Figure 7. Map of the Florida Reef Tract showing the association between DSD prevalence in *S.*  
771 *siderea* and the top three predictor variables among different ecoregions along the Florida Reef  
772 Tract. Colored dots indicate sites exceeding the threshold levels for the model's three top  
773 predictors of DSD occurrence. Blue dots indicate sites where the threshold levels of *S. siderea*  
774 densities are found, yellow dots indicate the threshold for silica concentrations and red dots  
775 indicate the threshold for septic areas within 8 km.

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792 Figure 1

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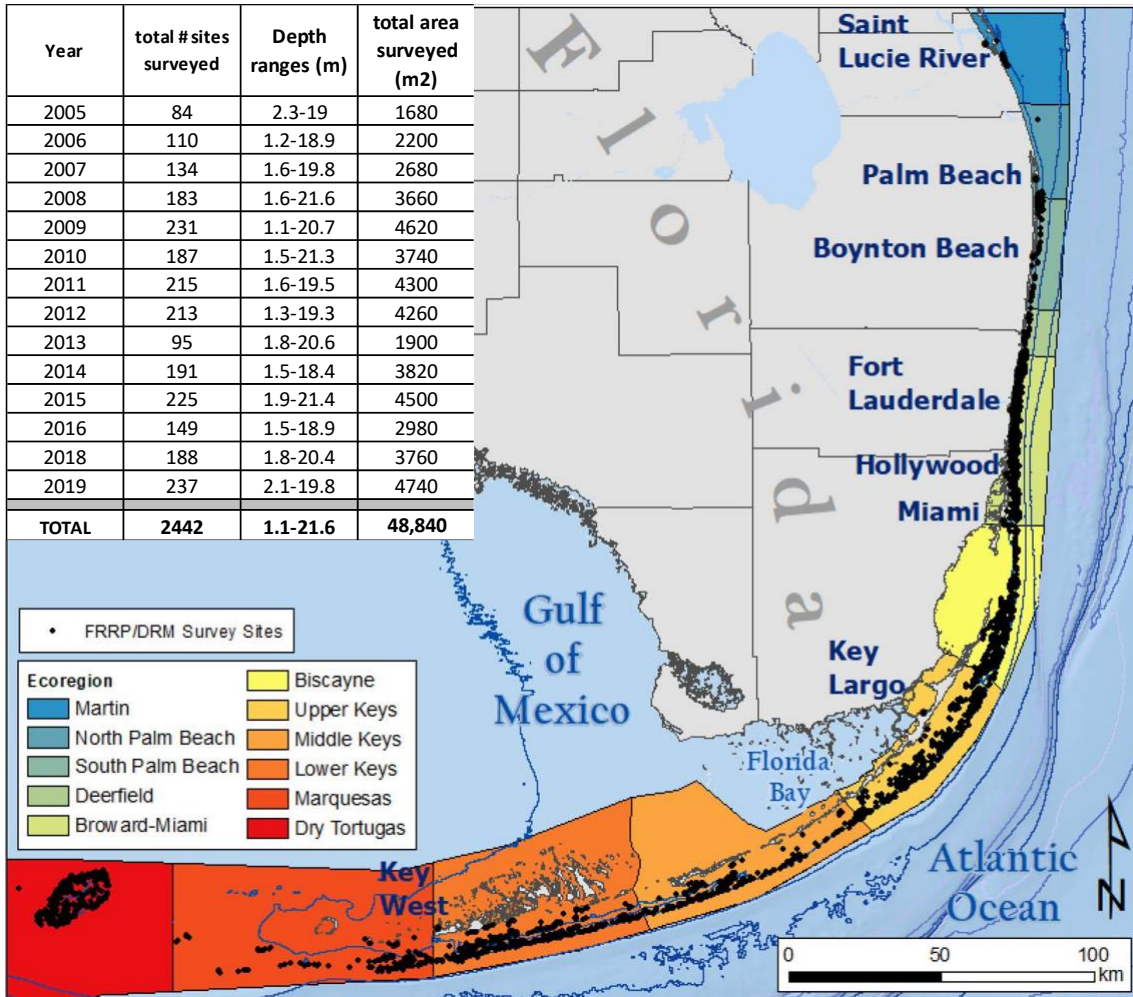
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802 Figure 2

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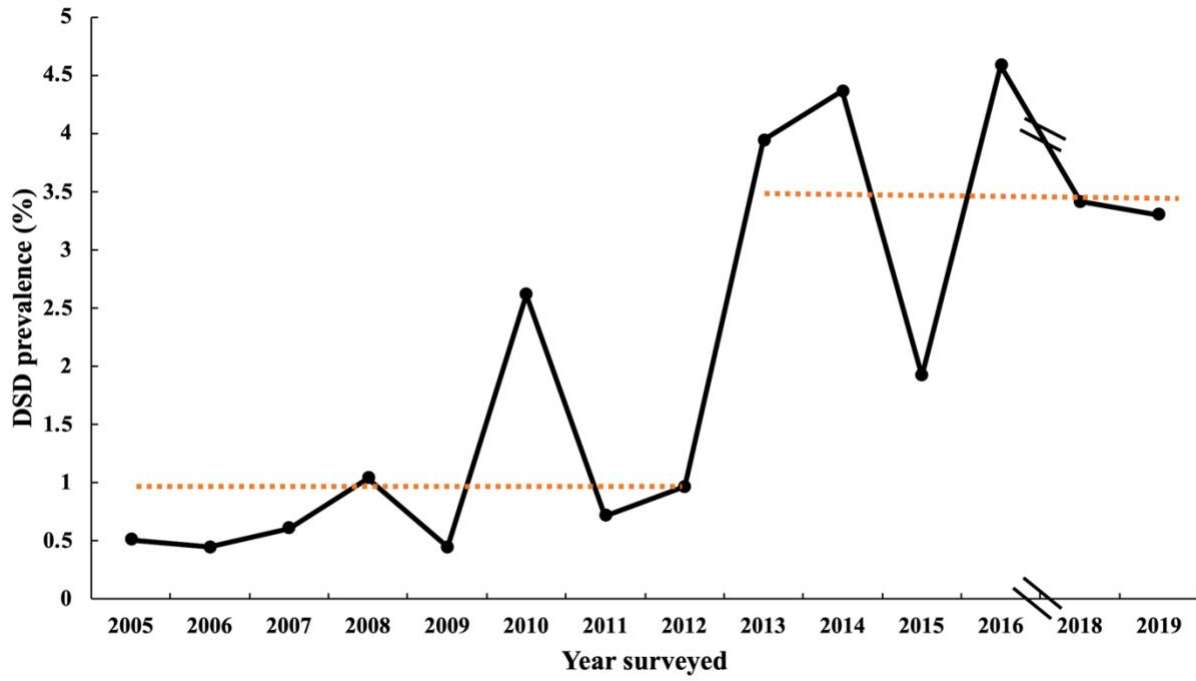
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Figure 3

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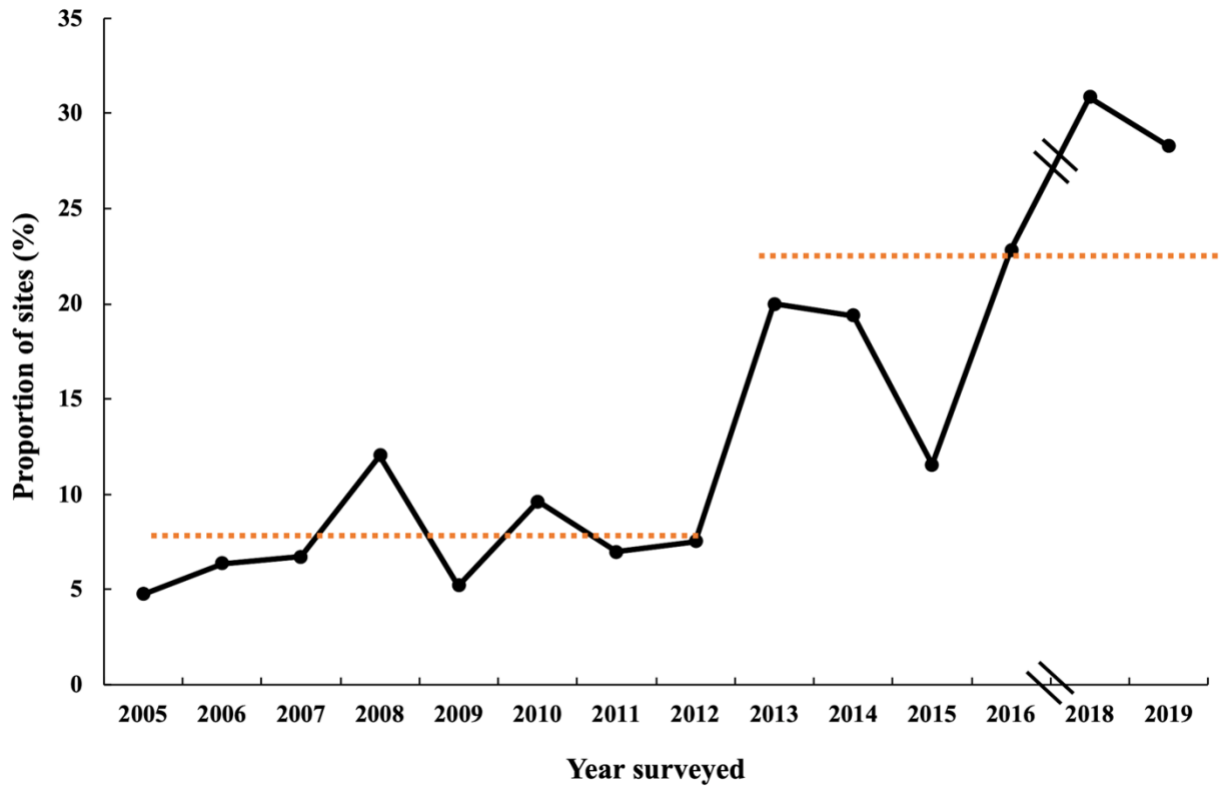
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825 Figure 4

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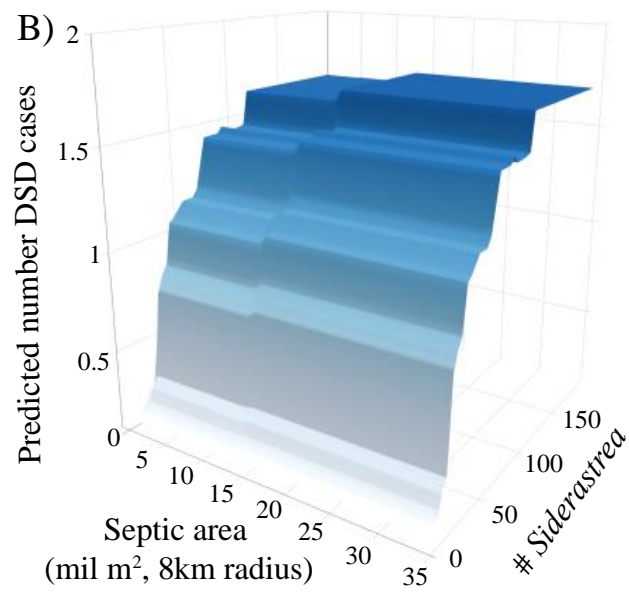
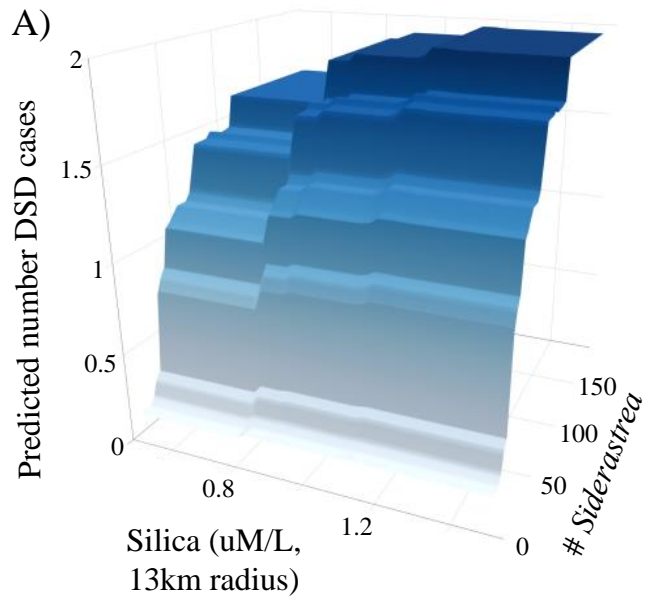
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840 Figure 5. A & B.

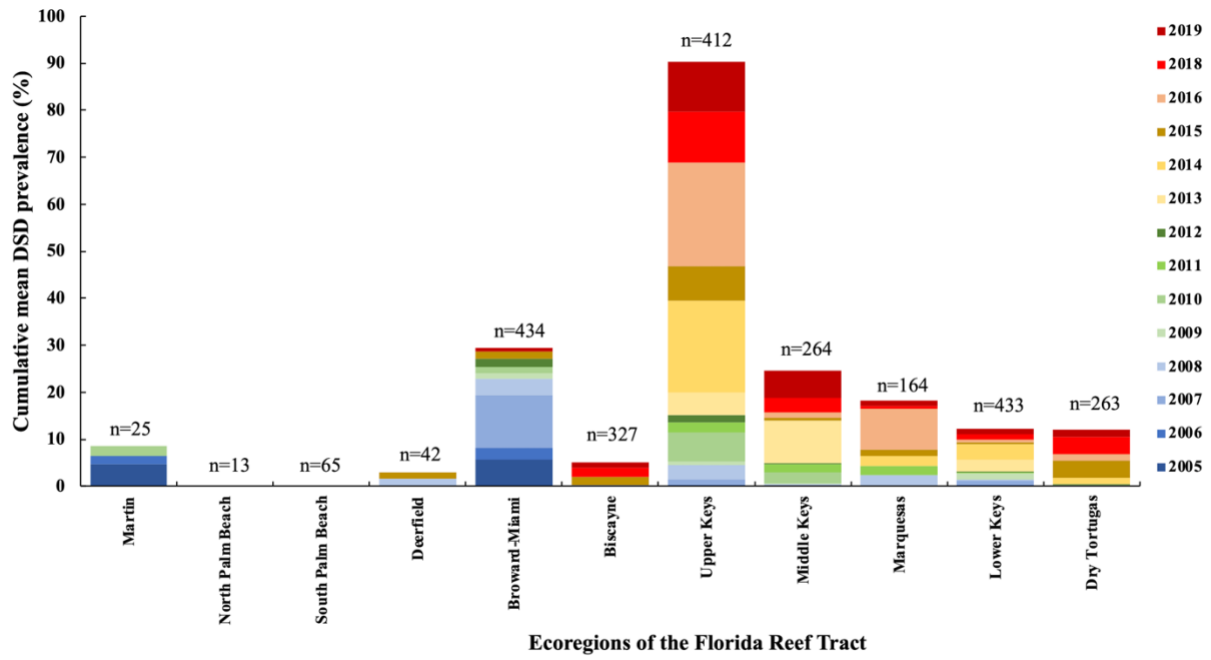
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847 Figure 6

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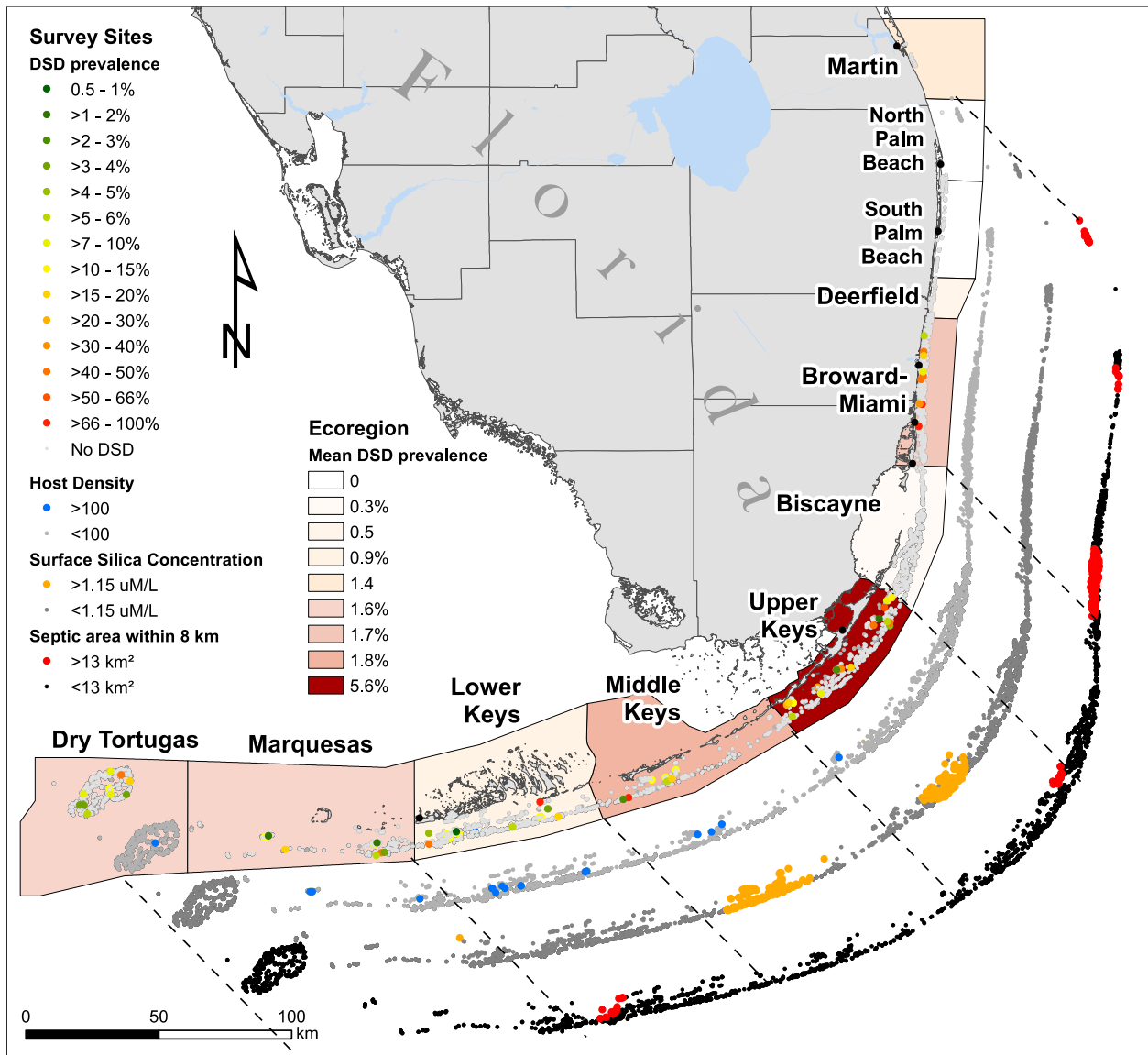
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862 Figure 7

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