



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

2

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

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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) 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 ≥ 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 (± 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

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729

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	https://myfwc.com/research/gis/regional-projects/unified-reef-map/
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	https://oceanwealth.org
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	https://geodata.dep.state.fl.us/
Septic/Sewer Areas: proxy for potential sewage pollution	Septic area (area in km ² 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 ² 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	https://www.floridahealth.gov/environmental-health/onsite-sewage/research/fiwm/index https://www10.doh.state.fl.us/pub/bsis/inventory/FloridaWaterManagementInventory
Land Use: degree of urbanization (affects coastal runoff & pollution)	High intensity land use area (in km ²) = Constructed materials account for 80 – 100% of the total cover. Vegetation, if present, occupies less than 20 % of the landscape.	0	289	https://coast.noaa.gov/digitalcoast/data/geobase.html
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	http://secr.flu.edu/wqmnetwork/
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	https://www.census.gov/geographies/mapping-files/2010/geo/tiger-data.html

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 ²) 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 ²) within 21 km	1.5
total organic carbon (um/L) in bottom waters within 21 km	1.3
degree of urbanization (m ²) 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.

759

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).

769

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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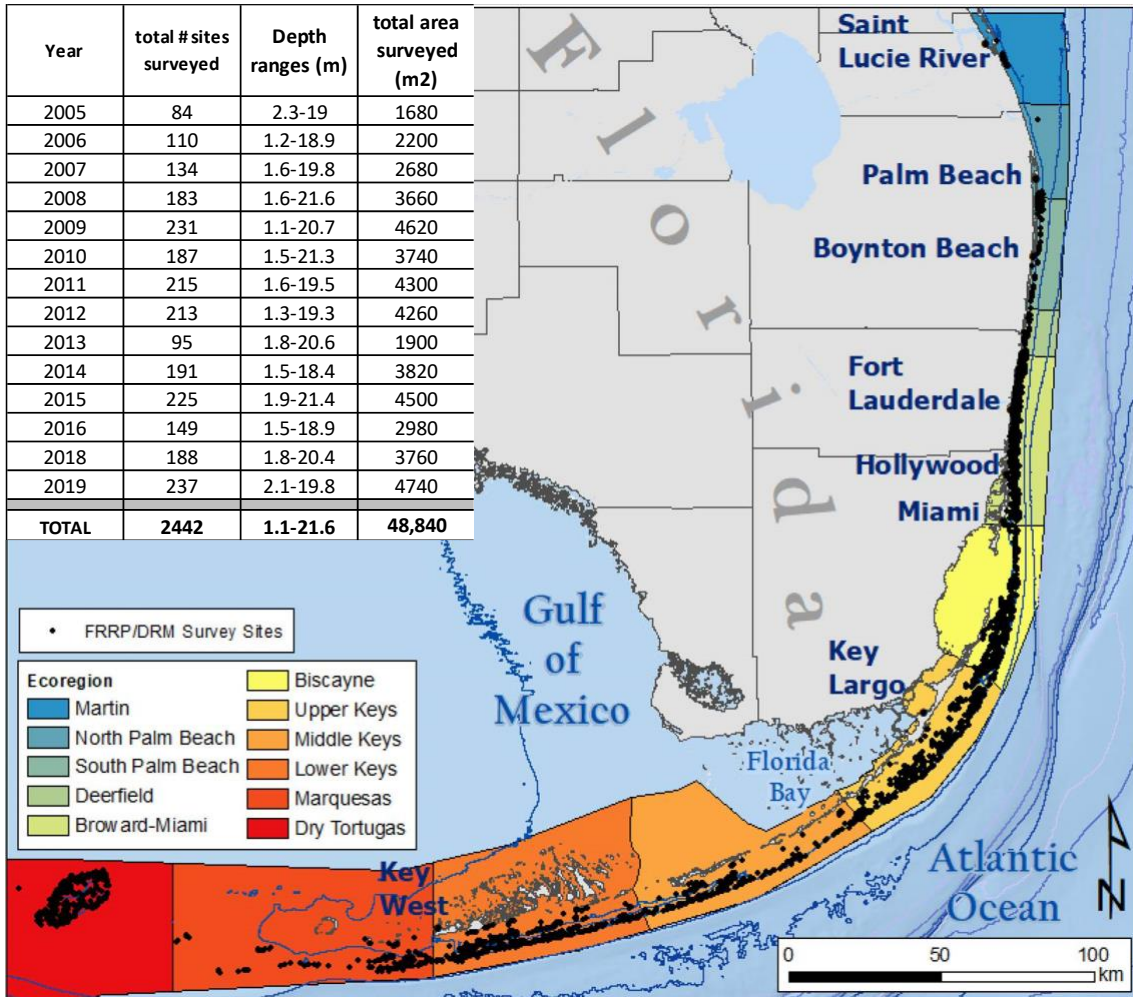
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802 Figure 2

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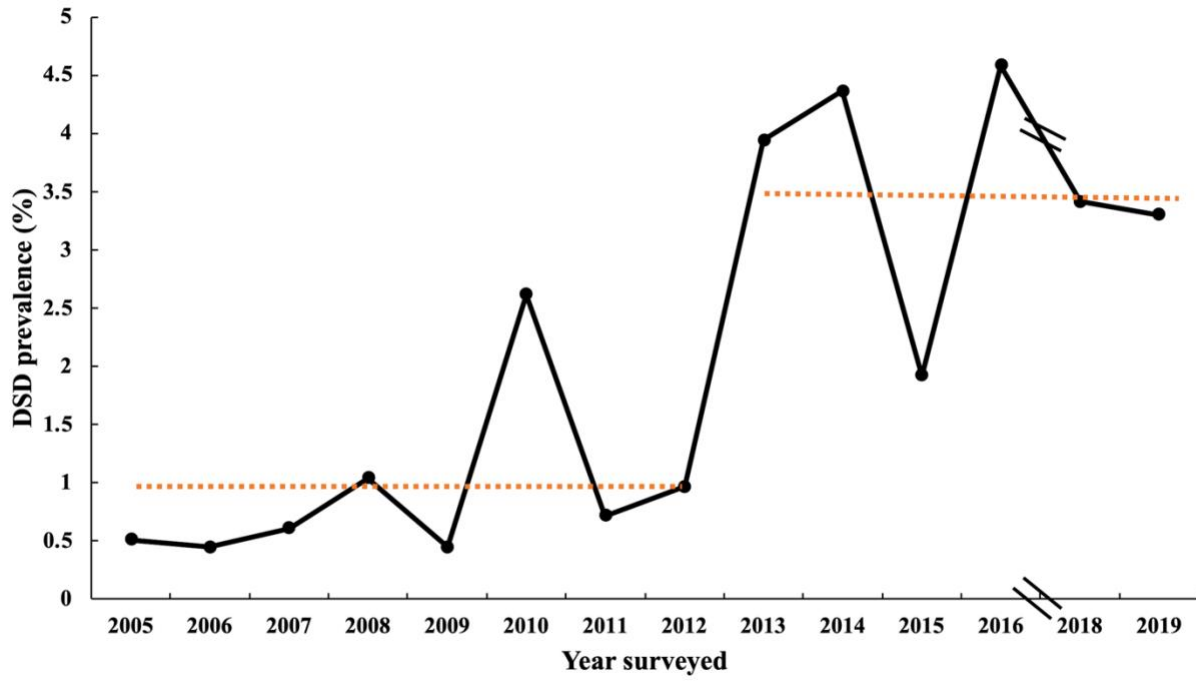
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Figure 3

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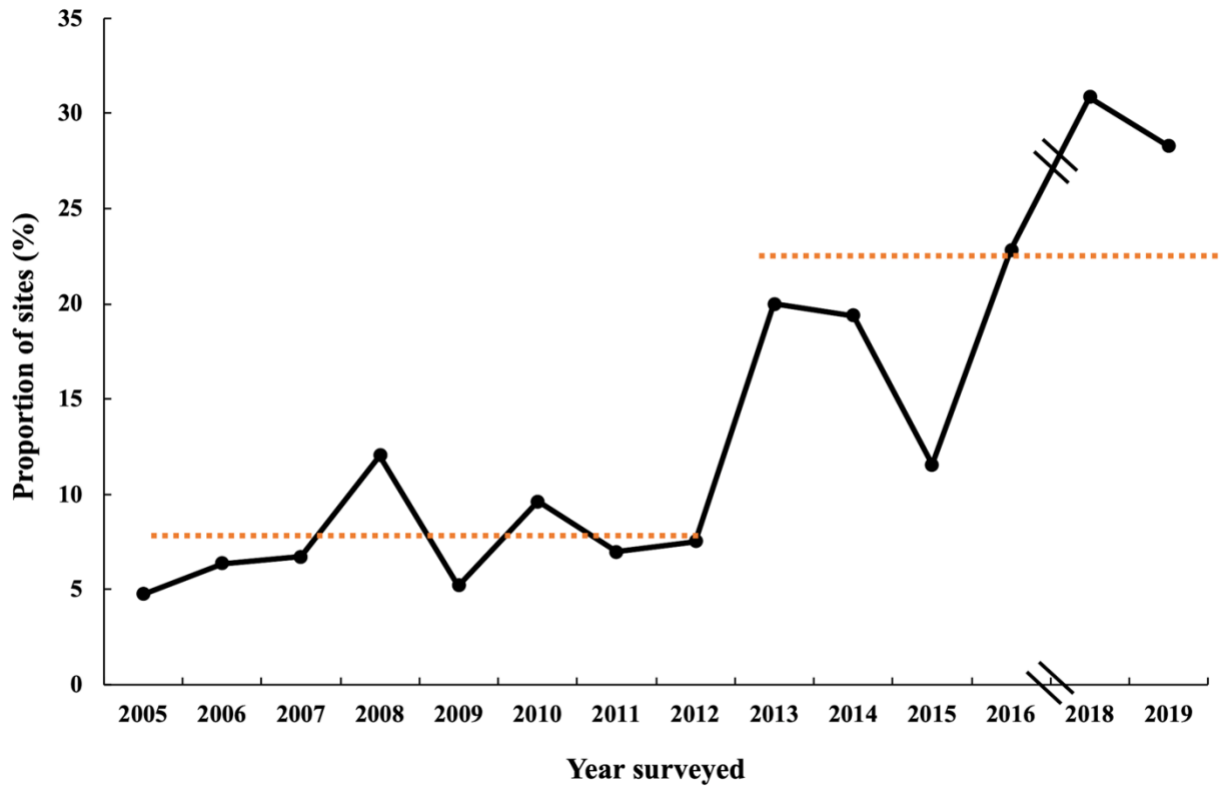
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825 Figure 4

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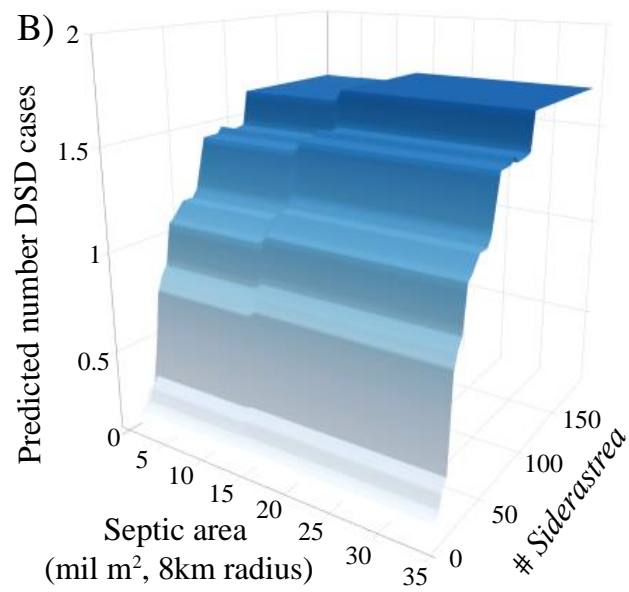
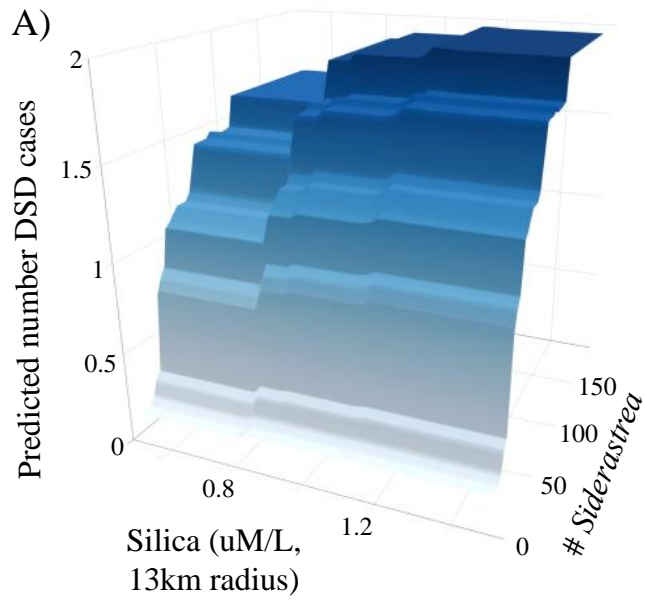
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840 Figure 5. A & B.

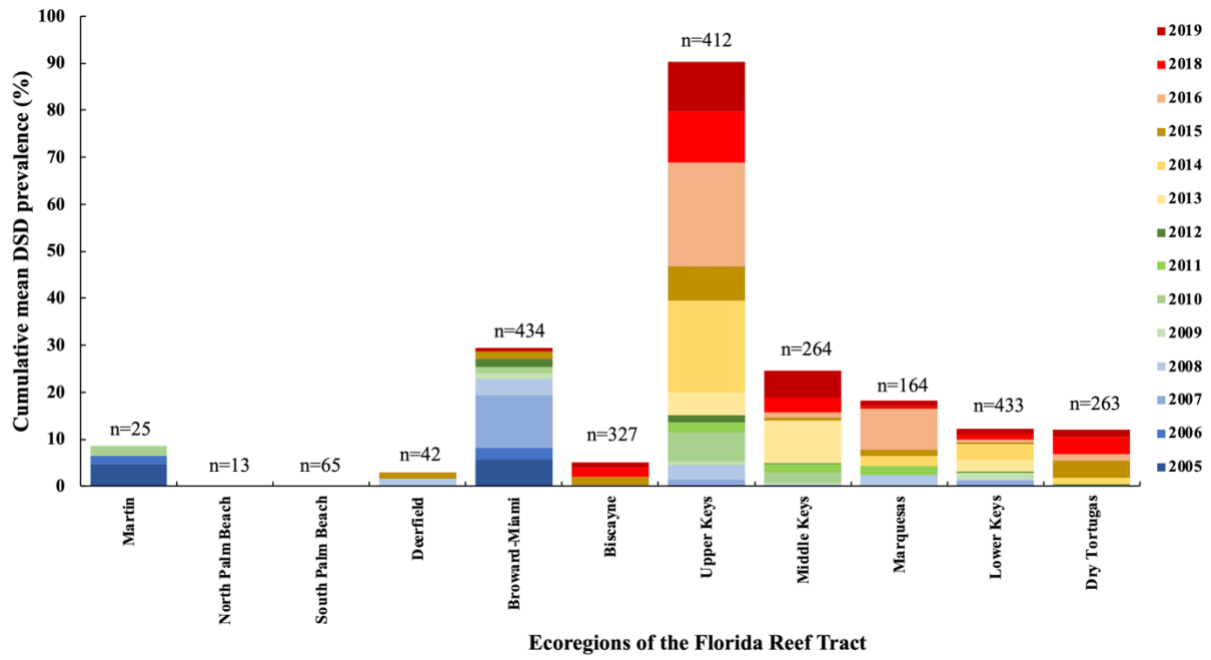
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847 Figure 6

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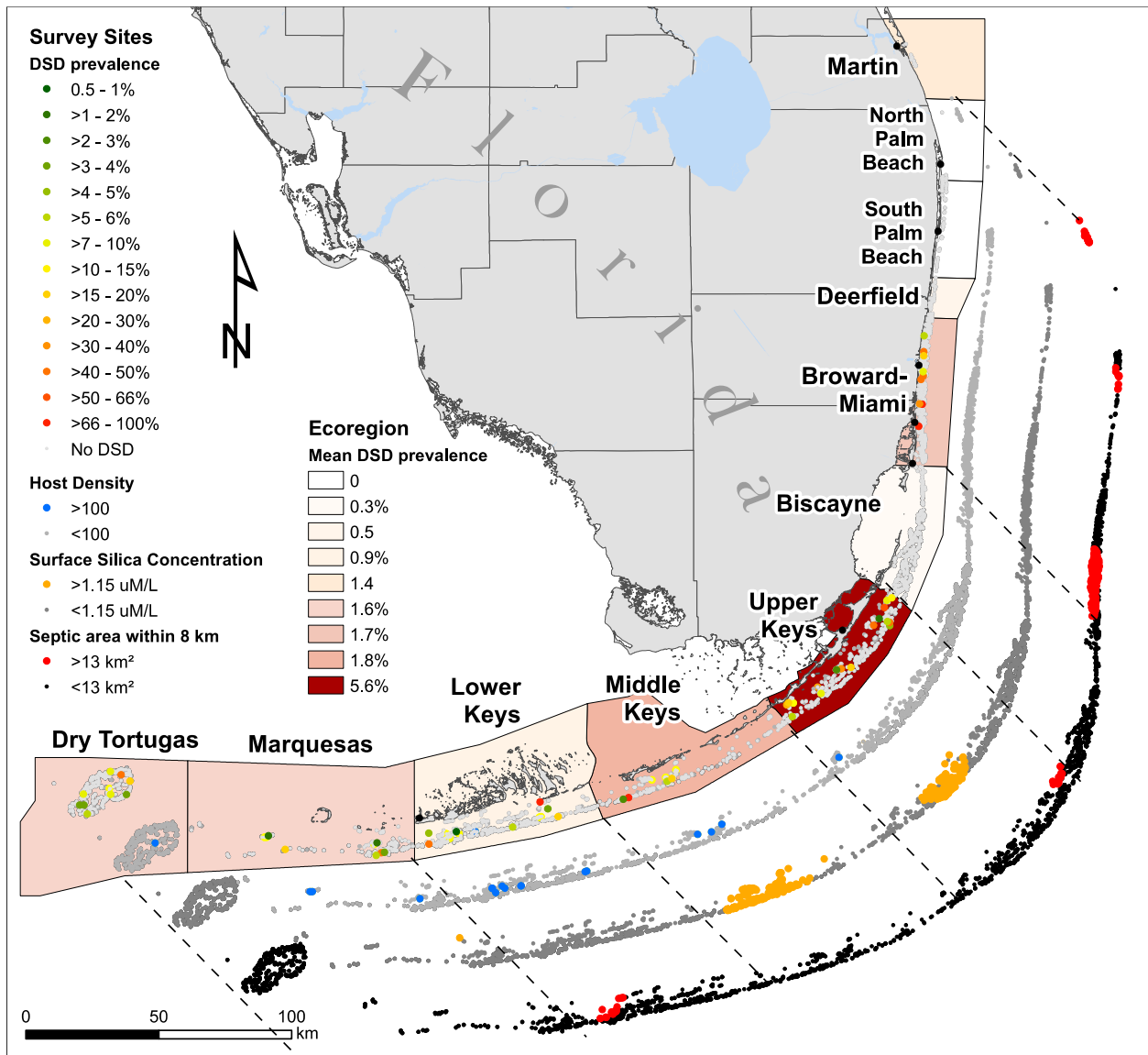
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862 Figure 7

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