

The relative influence of abundance and priority effects on colonization success in a coral-reef fish

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Abstract The sequence of species colonization is increasingly recognized as an important determinant of community structure, yet the significance of sequence of arrival relative to colonizer abundance is seldom assessed. We manipulated the magnitude and timing of coral-reef fish settlement to investigate whether the competitive dominance of early-arriving Ambon damselfish (i.e., a priority effect) decreased in strength with increasing abundance of late-arriving lemon damselfish. Sequence of arrival had a stronger effect on survival than the number of competing individuals. Relative to when both species arrived simultaneously, lemon damselfish were less aggressive, avoided competitive interactions more frequently and experienced depressed survival when they arrived later than Ambon damselfish, with these effects occurring independently of lemon damselfish abundance. These results suggest priority effects are more important than colonizer abundance and should motivate the

integration of priority effects into future studies of density dependence to determine their relative importance.

Keywords Competition · Coral-reef fish · Damselfish · Density dependence · *Pomacentrus* · Settlement

Introduction

Priority effects are increasingly recognized as an important determinant of alternative stable states (Fukami et al. 2010; Fukami 2015) and species coexistence via the storage effect (Chesson 2003). Yet the importance of priority effects relative to other processes governing community structure is seldom assessed. Priority effects occur in concert with fluctuations in the number of young-of-year arriving at a site due to environmental and demographic stochasticity (Lande et al. 2003; Wilson and Lundberg 2006). These numeric fluctuations have been advocated as a major determinant of population and community structure (Gaines and Roughgarden 1985; Foster and Tilman 2003). For example, competitively inferior species can overwhelm competitively superior species simply through numeric abundance (i.e., a ‘mass effect’; Leibold et al. 2004). Ecological theory predicts that variable colonizer abundance and priority effects can have significant impacts on the transient dynamics of ecological systems (Fukami and Nakajima 2011). Such transient dynamics may operate on time scales relevant to conservation and management actions (Collie et al. 2013); however, the short-term influence of priority effects, variation in colonizer abundance and their relative contribution to population and community structure have seldom been empirically evaluated in demographically open systems.

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A coral-reef fish assemblage is a good system for examining short-term inhibitory priority effects. Recent work has shown that individual survival of reef fish settlers can depend on the timing of priority effects (Geange and Stier 2009) and that the strength of priority effects can vary dramatically within populations (Almany 2003; Munday 2004; Geange and Stier 2009; Poulos and McCormick 2014, 2015) and between habitats (Geange and Stier 2010; Adam 2011). However, these studies need to be reconciled against often large variation in settler abundance and earlier work showing that the abundance of settlers can determine their persistence by influencing density-dependent interactions (e.g., Steele 1997; Shima 2001; Shima and Osenberg 2003). Here, we factorially manipulated the abundance and sequence of settlement-stage damselfish arrival onto patch reefs to assess their relative contribution to reef fish survival.

Methods

We conducted this study in January 2015 in the lagoon of Lizard Island, Great Barrier Reef (41°41.23'S, 145°27.17'E), using a grid of 42 coral patch reefs consisting of a 50:50 mixture of live and dead *Pocillopora damicornis* with a volume of approximately 0.07 m³. Below we provide a brief overview of our experimental methodology; for a full description, see electronic supplementary material.

Due to the lack of settlement-stage Ambon damselfish (*Pomacentrus amboinensis*) captured in light traps, our experimental design focused on the effects of Ambon damselfish (11.32 ± 0.34 mm SL [mean ± SD]) on lemon damselfish (*P. moluccensis*) (11.30 ± 0.35 mm SL) rather than a fully reciprocal experimental design. We experimentally manipulated the abundance and time of arrival of settlement-stage lemon damselfish relative to settlement-stage Ambon damselfish in four treatments: (1) low settler abundance without a priority effect (two Ambon and two lemon damselfish introduced simultaneously); (2) low settler abundance with a priority effect (two Ambon damselfish introduced 3 h before two lemon damselfish individuals); (3) high settler abundance without a priority effect (two Ambon and six lemon damselfish introduced simultaneously); and (4) high settler abundance with a priority effect (two Ambon damselfish introduced 3 h before six lemon damselfish). Simulated settlement pulses of 4–8 individuals are representative of the numbers of post-settlement fishes observed on reefs of this size. A 3-h 'priority period' is ecologically relevant since reef fishes settle over several hours overnight (Dufour and Galzin 1993; Holbrook and Schmitt 1997) and competitive interactions are strongest in the hours immediately following

settlement (McCormick 2009; Poulos and McCormick 2014). The sizes of fish added to each reef did not differ by more than 0.3 mm SL. Each fish was tagged below the dorsal fin with subcutaneous fluorescent elastomer (VIE; Northwest Marine Technology, Shaw Island, WA, USA), and on each reef one individual of each species was tagged a different color and acted as the focal individual for behavioral and survival analysis. We ran the experiment in five temporal blocks between 21 and 28 January 2015. Because temporal blocks overlapped, replicates were randomized across available reefs within the grid of 42 patch reefs, yielding 13 replicates for treatments 1 and 2, 11 replicates for treatment 3 and 12 replicates for treatment 4.

To understand the mechanisms underlying the strength of variation in colonizer abundance and priority effects, we conducted 3-min behavioral observations approximately 20 min after fish were added to reefs, recording five response variables for the lemon damselfish that interacted the most frequently with other individuals: (1) the number of displays; (2) the number of chases; (3) the number of avoidances; (4) the distance ventured from the reef; and (5) relative height on the reef. Because there were strong correlations between chases and displays, we used principal components analysis (PCA) to create a composite aggression score. PCA 1 accounted for 60% of the total variation in aggressive interactions and was driven by high positive loadings of total displays and chases (71%). Thus, we interpret PCA 1 as a gradient of overall aggression.

We surveyed reefs at approximately 0800, 1200 and 1600 hrs for 3 d after we introduced fish, recording survival of all individuals and removing any new, non-tagged Ambon or lemon damselfish settlers using eugenol and hand nets. We used Cox proportional hazard regression (Cox 1972) to model survival against time of arrival, the abundance of lemon damselfish settlers and their interaction. A strata term in the model was included to account for temporal block. We right-censored data, used the Efron approximation to handle tied data, used partial likelihood to derive regression coefficients and converted these into estimated changes in mortality rate (CMR) by subtracting 1 from the hazard ratio and multiplying by 100. We used Schoenfeld residuals scaled by the Kaplan–Meier estimate to test the assumption of proportional hazards and DFBETA values to test for influential data. We selected a best-fit model based on Chi-squared tests that omitted an interaction between time of arrival and abundance ($P = 0.937$). We therefore present results for the main effects of abundance and time of arrival only. We also calculated response ratios and 95% confidence intervals for average survivorship after 68 h as per Borenstein et al. (2009).

We analyzed the behavior data to determine whether priority effects or settler abundance resulted in lemon

damsel fish displaying changes in: (1) aggressive interactions; (2) the avoidance of competitors; (3) changes in the distance ventured from the reef; or (4) relative height on the reef. For each response variable, we fit separate linear mixed-effects (LME) models that included settler abundance, time of arrival and their two-way interaction as fixed effects, and reef nested within temporal block as a random effect. We used maximum likelihood to estimate variance components (Venables and Ripley 2002) and selected the best-fit models as determined by AIC. In all cases, the best-fit model omitted an interaction between time of arrival and abundance ($P > 0.05$ in all cases) and we therefore present results for the main effects of time of arrival and abundance only. All data met assumptions of homoscedasticity (in all cases $P > 0.05$ for Levene's test of homogeneity of variance).

All statistical analyses were conducted using the statistical software R version 3.0.1 (R Development Core Team 2013). We used the survival library to fit Cox proportional hazard regressions and the library nlme to fit LMEs.

Results and discussion

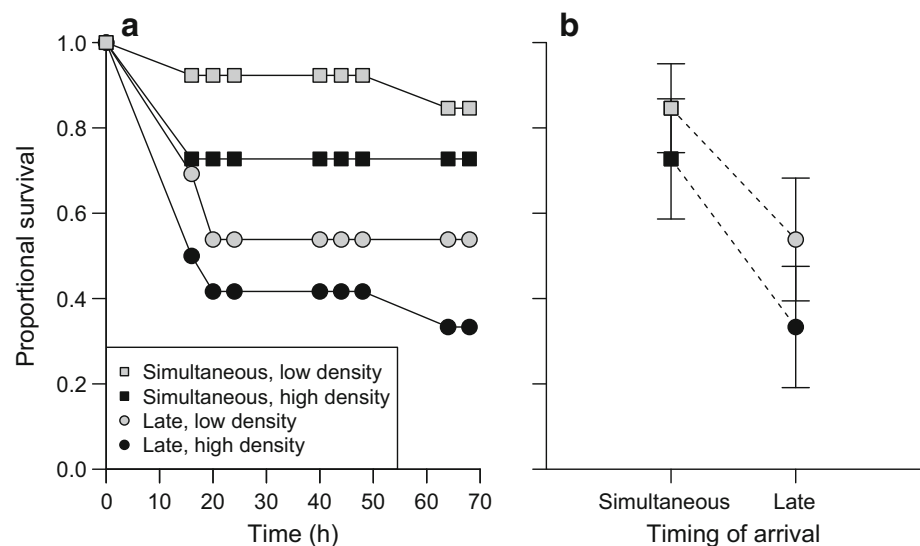
By manipulating both priority effects and settler abundance, we showed that priority effects can have a larger influence on individual survival than the number of individuals arriving at a site. We found a strong inhibitory priority effect—lemon damselfish arriving 3 h later than Ambon damselfish experienced depressed survival ($P = 0.015$, CMR = 76%; Fig. 1a, b), were less aggressive ($F_{(1,42)} = 5.12$, $P = 0.029$; Fig. 2a) and were 2.7 times more likely to avoid competitive interactions ($F_{(1,42)} = 8.011$, $P = 0.007$; Fig. 2b) relative to when they arrived at the same time as Ambon damselfish. In contrast,

increasing the abundance of lemon damselfish settlers from 2 to 6 individuals did not result in statistically significant effects on lemon damselfish mortality ($P = 0.131$, CMR = 52%; Fig. 1a, b), aggression by lemon damselfish ($F_{(1,42)} = 1.108$, $P = 0.299$; Fig. 2a) or avoidance by lemon damselfish ($F_{(1,42)} = 0.664$, $P = 0.419$; Fig. 2b); however, these results may have partly be due to insufficient statistical power. After 68 h, average survivorship of lemon damselfish at high abundance was 75% ($CI = 46$ –125%) of that at low abundance, whereas average survivorship of late-arriving lemon damselfish was 56% ($CI = 33$ –92%) of that when lemon damselfish arrived simultaneously with Ambon damselfish.

Aggression of Ambon damselfish did not differ between treatments (timing: $F_{(1,42)} = 1.923$, $P = 0.173$; density: $F_{(1,42)} = 0.713$, $P = 0.403$), suggesting that reduced aggression of late-arriving lemon damselfish was the result of submission to Ambon damselfish with concurrent decreases in survival. Previous work has found that competition increases the exposure of subordinate fish to predators due to displacement from refuge habitat (Carr et al. 2002; Holbrook and Schmitt 2002; Almany 2003). The inability to find (and defend) shelter sites may therefore explain decreased survival of late-arriving lemon damselfish in this study.

The influence of priority effects on the survival of lemon damselfish was approximately 1.5 times greater than that of variability in settler abundance, suggesting that for the species combinations and abundances used in this experiment, timing of arrival outweighs variation in settler abundance as the critical driver of early recruitment success. However, the general applicability of these results to other species and contexts requires further investigation. Although we had planned to examine for reciprocal effects between Ambon and lemon damselfish, we were limited in

Fig. 1 **a** Temporal trends in survival of focal lemon damselfish relative to timing of arrival (simultaneous or late for lemon damselfish) and the density of lemon damselfish (two individuals: gray symbols; six individuals: black symbols). Error bars are omitted for clarity. **b** Survival of lemon damselfish after 68 h. Error bars are 95% confidence intervals



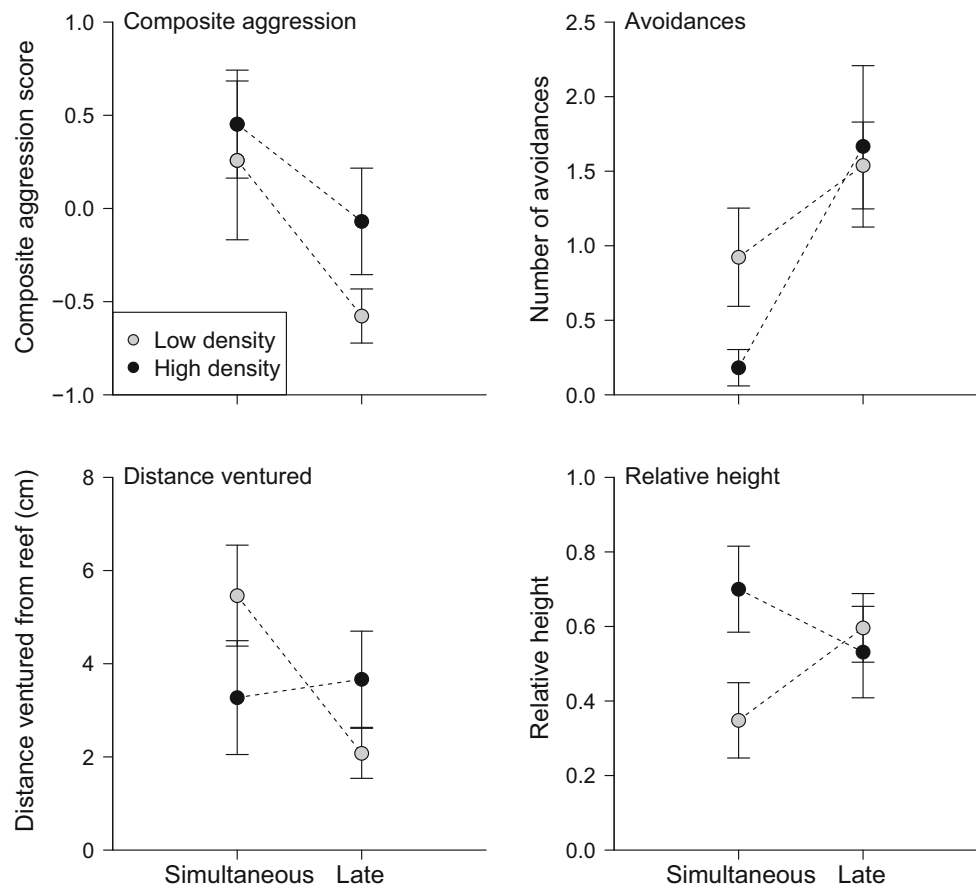


Fig. 2 Composite aggression scores (modeling chases and displays: see electronic supplementary material), number of avoidances, distance ventured from the reef and relative height on the reef (mean \pm SE) of lemon damselfish relative to timing of arrival

(simultaneous or late for lemon damselfish) and the density of lemon damselfish (two individuals: *gray symbols*; six individuals: *black symbols*)

our ability to do so due to low numbers of Ambon damselfish caught in light traps. If the results from this study are generally applicable, and priority effects have a strong influence on colonizer establishment, focusing on the number of colonizers without considering interactions between colonizers and established residents will likely reduce the accuracy of predictions of colonizer success, with potential implications for the management of fisheries yield, invasive species or restoration initiatives (e.g., Grman and Suding 2008; Dickson et al. 2012; Collie et al. 2013). Alternatively, if abundance has a stronger influence on colonizer success than priority effects, higher abundances of late-arriving individuals may either overwhelm established residents and mitigate the competitive advantage of early arrival (e.g., Gilpin and Case 1976) or enhance priority effects when predatory effort is focused on highly abundant prey (e.g., Webster and Almany 2002).

There is an emerging suggestion that the density of early-established populations is likely to have a strong impact on priority effects (e.g., Waters et al. 2013), and the results from our study highlight the requirement for

additional research into the general importance of priority effects relative to the abundance of late-arriving individuals. Our results give rise to a number of future questions that relate to determining the role of reciprocal effects (including the sequence [who arrives first]) and timing [by how much] of arrival), the relationship between the intensity of priority effects and phylogenetic relatedness, and the effect that variation in the numbers of both early and late settlers has on individual survival. Answering these questions could lead to a better mechanistic understanding of population and community dynamics than numbers of colonizers alone.

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References

- Adam TC (2011) High quality habitat and facilitation ameliorate competitive effects of prior residents on new settlers. *Oecologia* 166:121–130
- Almany GR (2003) Priority effects in coral reef fish communities. *Ecology* 84:1920–1935
- Borenstein M, Hedges LV, Higgins JPT, Rothstein HR (2009) Introduction to meta-analysis. John Wiley & Sons, United Kingdom
- Carr MH, Anderson TW, Hixon MA (2002) Biodiversity, population regulation, and the stability of coral reef fish communities. *Proc Natl Acad Sci U S A* 99:11241–11245
- Chesson P (2003) Quantifying and testing coexistence mechanisms arising from recruitment fluctuations. *Theor Popul Biol* 64:345–357
- Collie J, Rochet MJ, Bell R (2013) Rebuilding fish communities: the ghost of fisheries past and the virtue of patience. *Ecol Appl* 23:374–391
- Cox DR (1972) Regression models and life-tables. *J R Stat Soc Series B Stat Methodol* 34:187–220
- Dickson TL, Hopwood JL, Wilsey BJ (2012) Do priority effects benefit invasive plants more than native plants? An experiment with six grassland species. *Biol Invasions* 14:2617–2624
- Dufour V, Galzin R (1993) Colonisation patterns on reef fish larvae to the lagoon at Moorea Island, French Polynesia. *Mar Ecol Prog Ser* 102:143–152
- Foster BL, Tilman D (2003) Seed limitation and the regulation of community structure in oak savanna grassland. *J Ecol* 91:999–1007
- Fukami T (2015) Historical contingency in community assembly: integrating niches, species pools, and priority effects. *Annu Rev Ecol Syst* 46:1–23
- Fukami T, Nakajima M (2011) Community assembly: alternative stable states or alternative transient states? *Ecol Lett* 14:973–984
- Fukami T, Dickie IA, Wilkie JP, Paulus BC, Park D, Roberts A, Buchanan PK, Allen RB (2010) Assembly history dictates ecosystem functioning: evidence from wood decomposer communities. *Ecol Lett* 13:675–684
- Gaines S, Roughgarden J (1985) Larval settlement rate: a leading determinant of structure in an ecological community of the marine intertidal zone. *Proc Natl Acad Sci U S A* 82:3707–3711
- Geange SW, Stier AC (2009) Order of arrival affects competition in two reef fishes. *Ecology* 90:2868–2878
- Geange SW, Stier AC (2010) Priority effects and habitat complexity affect the strength of competition. *Oecologia* 163:111–118
- Gilpin ME, Case TJ (1976) Multiple domains of attraction in competition communities. *Nature* 261:40–42
- Grman E, Suding KN (2008) Within-year soil legacies contribute to strong priority effects of exotics on native California grassland communities. *Restoration Ecology* 18:664–670
- Holbrook SJ, Schmitt RJ (1997) Settlement patterns and processes in a coral reef damselfish: in situ nocturnal observations using infrared video. *Proc 8th Int Coral Reef Symp* 2:1143–1148
- Holbrook SJ, Schmitt RJ (2002) Competition for shelter space causes density-dependent predation mortality in damselfishes. *Ecology* 83:2855–2868
- Lande R, Engen S, Saether BE (2003) Stochastic population dynamics in ecology and conservation. Oxford University Press, Oxford, UK
- Leibold MA, Holyoak M, Mouquet N, Amarasekare P, Chase JM, Hoopes MF, Holt RD, Shurin JB, Law R, Tilman D, Loreau M, Gonzalez M (2004) The metacommunity concept: a framework for multi-scale community ecology. *Ecol Lett* 7:601–613
- McCormick MI (2009) Behaviourally mediated phenotypic selection in a disturbed coral reef environment. *PLoS One* 4:e7096
- Munday PL (2004) Competitive coexistence of coral-dwelling fishes: the lottery hypothesis revisited. *Ecology* 85:623–628
- Poulos DE, McCormick MI (2014) Who wins in the battle for space? The importance of priority, behavioural history and size. *Anim Behav* 90:305–314
- Poulos DE, McCormick MI (2015) Asymmetries in body condition and order of arrival influence competitive ability and survival in a coral reef fish. *Oecologia* 179:719–728
- Development Core Team R (2013) R: a language and environment for statistical computing. R Foundation for Statistical Computing, Vienna, Austria
- Shima JS (2001) Regulation of local populations of a coral reef fish via joint effects of density- and number-dependent mortality. *Oecologia* 126:58–65
- Shima JS, Osenberg CW (2003) Cryptic density dependence: effects of covariation between density and site quality in reef fish. *Ecology* 84:46–52
- Steele MA (1997) Population regulation by post-settlement mortality in two temperate reef fishes. *Oecologia* 112:64–74
- Venables WN, Ripley BD (2002) Modern applied statistics with S, 4th edn. Springer, New York
- Waters JM, Fraser CI, Hewitt GM (2013) Founder takes all: density-dependent processes structure biodiversity. *Trends Ecol Evol* 28:78–85
- Webster MS, Almany GR (2002) Positive indirect effects in a coral reef fish community. *Ecol Lett* 5:549–557
- Wilson WG, Lundberg P (2006) Non-neutral dynamics: empirical predictions for ecosystem function and diversity from linearized consumer–resource interactions. *Oikos* 114:71–83