

Figure S1| Schematic of the experimental design. Each treatment, which included a control and the effect of reduced pH and/or elevated temperature, was performed relative to the background variance of each habitat (Low variance (LV) outer-reef and high variance (HV) seagrass). *Acropora palmata* sampled from the LV habitat ($n= 40$), and *Porites astreoides* from both the HV ($n= 40$) and LV ($n= 40$) habitats) were simultaneously subjected to each of the eight treatments. Each vessel held one fragment of *A. palmata* (LV), *P. astreoides* (LV) and *P. astreoides* (HV) as shown in the close-up image of the vessel.

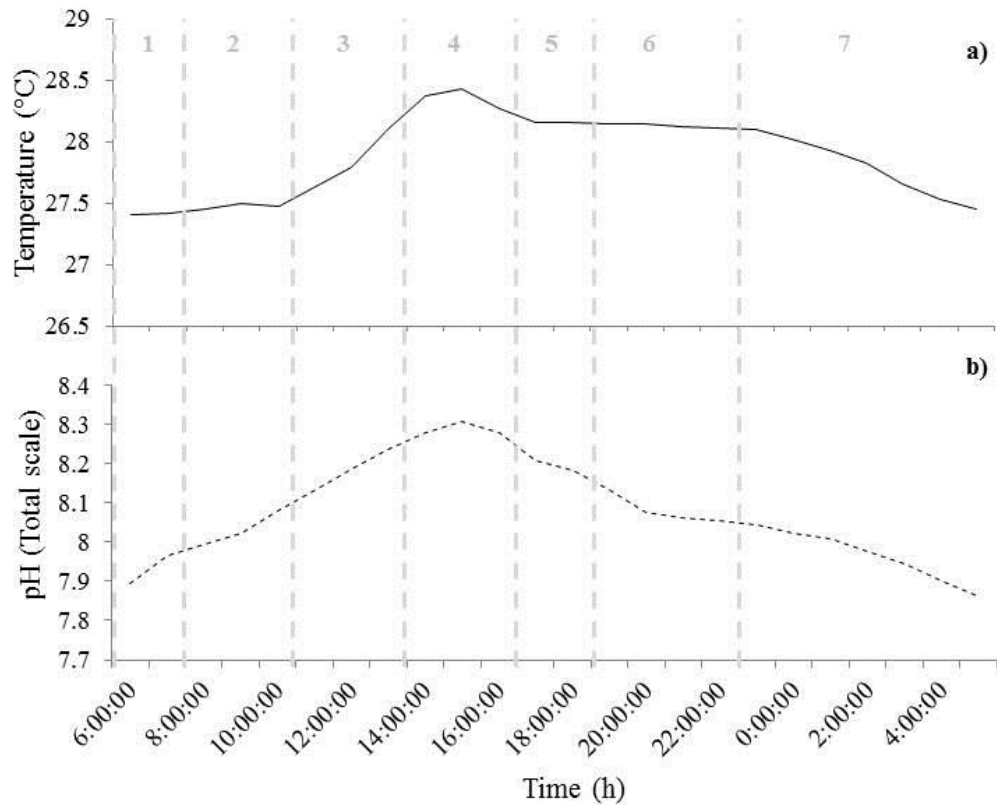


Figure S2| The natural diurnal oscillations in: a) temperature and b) pH for the high-variance seagrass habitat on Little Cayman, Cayman Islands, BWI. From the natural diurnal cycles, seven time periods were selected to manipulate the temperature and pH to conditions predicted in 2100 under the IPCC A1B scenario. The seven time periods were selected to try and best-represent the natural diurnal conditions.

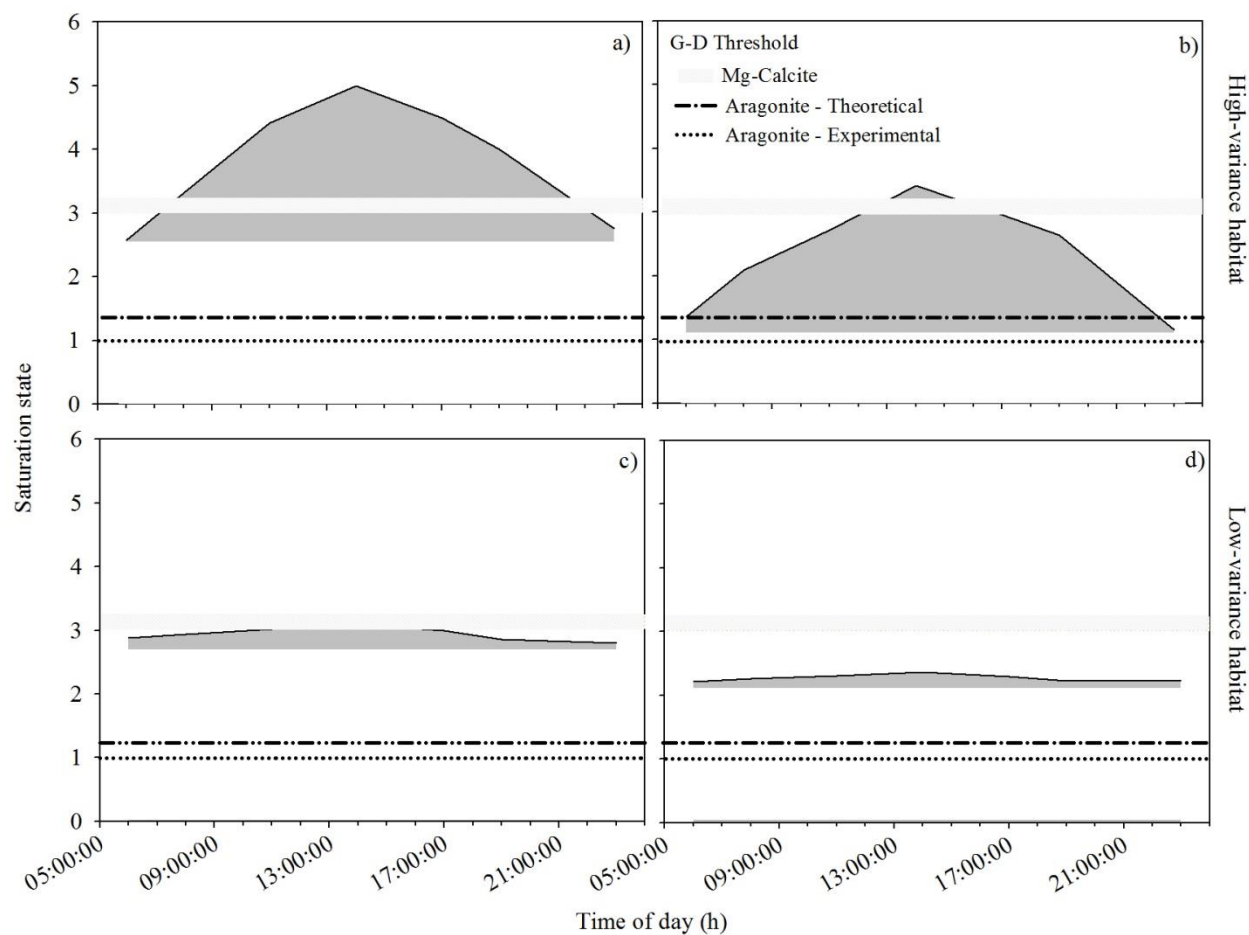


Figure S3| The time each habitat (outer-reef, low-variance and seagrass, high-variance) was exposed to a given saturation state. Data is averaged for each level of variability (high or low). a & c show current ambient seawater conditions, b & d show the effect a change of -0.3 pH units and a 2.0 °C temperature increase has on the saturation states of each habitat as predicted under IPCC scenario A1B. Calcification-to-dissolution thresholds (G-D) are shown for Mg-calcite, aragonite theoretical (e.g. the calculated level of one) and experimental aragonite (levels experiments have measured the aragonite threshold to occur at).

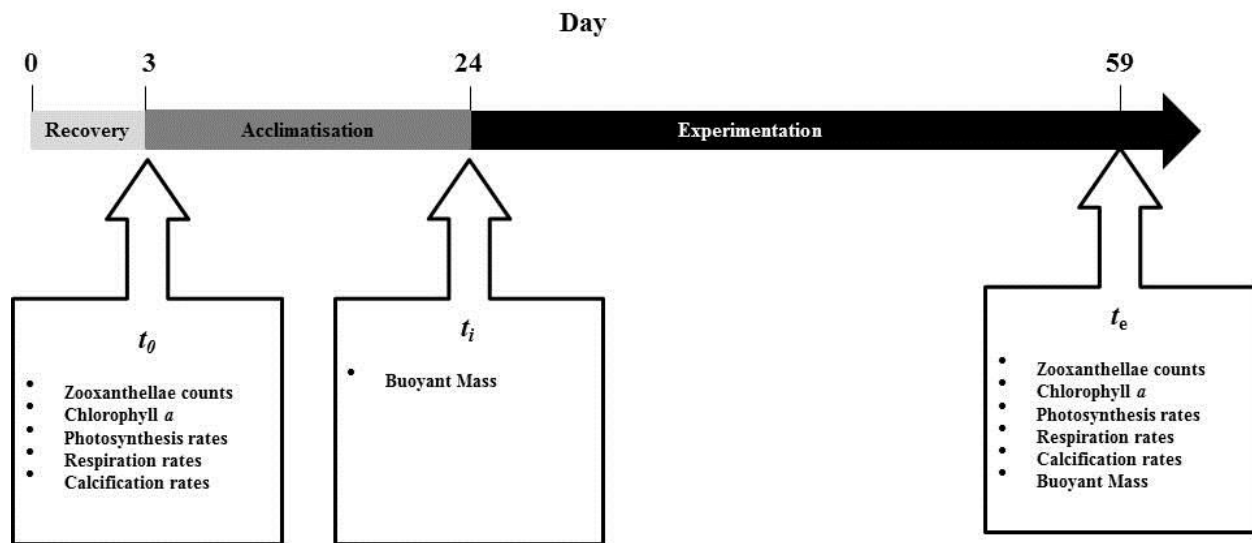


Figure S4| Sampling timeline demonstrating the experimental design and times of physiological measurements. Corals were removed from their native environment at day 0 and allowed to recover in the laboratory within tanks set to their ambient conditions for three days (Recovery period). At the end of the recovery period (t_0) a series of metabolic measurements were taken. Corals were then given a 21-day acclimatisation period to the *ex situ* treatment conditions. At the end of the acclimatisation period (t_i) buoyant mass was measured for each colony. The experimentation conditions were then run for 35-days. At the end of the experimentation period (t_e) the metabolic parameters measured at t_0 along with buoyant mass were re-measured.

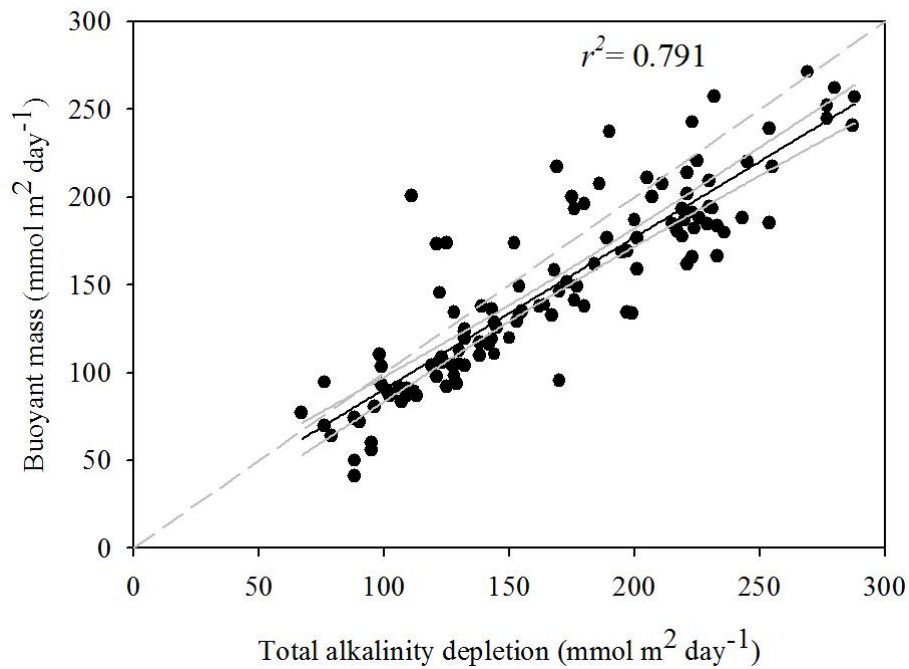


Figure S5| The relationship between growth rates based on the total alkalinity depletion and buoyant mass methods. Each point represents measurements from both methods on an individual coral fragment. Data is shown for the high-variance and low-variance treatments. The grey-dotted line illustrates a 1:1 ratio, the solid black line represents the best fit line (Buoyant Mass ($\text{mmol m}^{-2} \text{ day}^{-1}$) = $4.32 + 0.864 * \text{Alkalinity depletion (mmol m}^{-2} \text{ day}^{-1}$), $r^2 = 0.791$, $n = 120$, $p < 0.001$) and the grey-solid lines represent the 95% confidence intervals. Rates were obtained from 8 x 3h incubations conducted over a 24h period at t_0 and t_e .

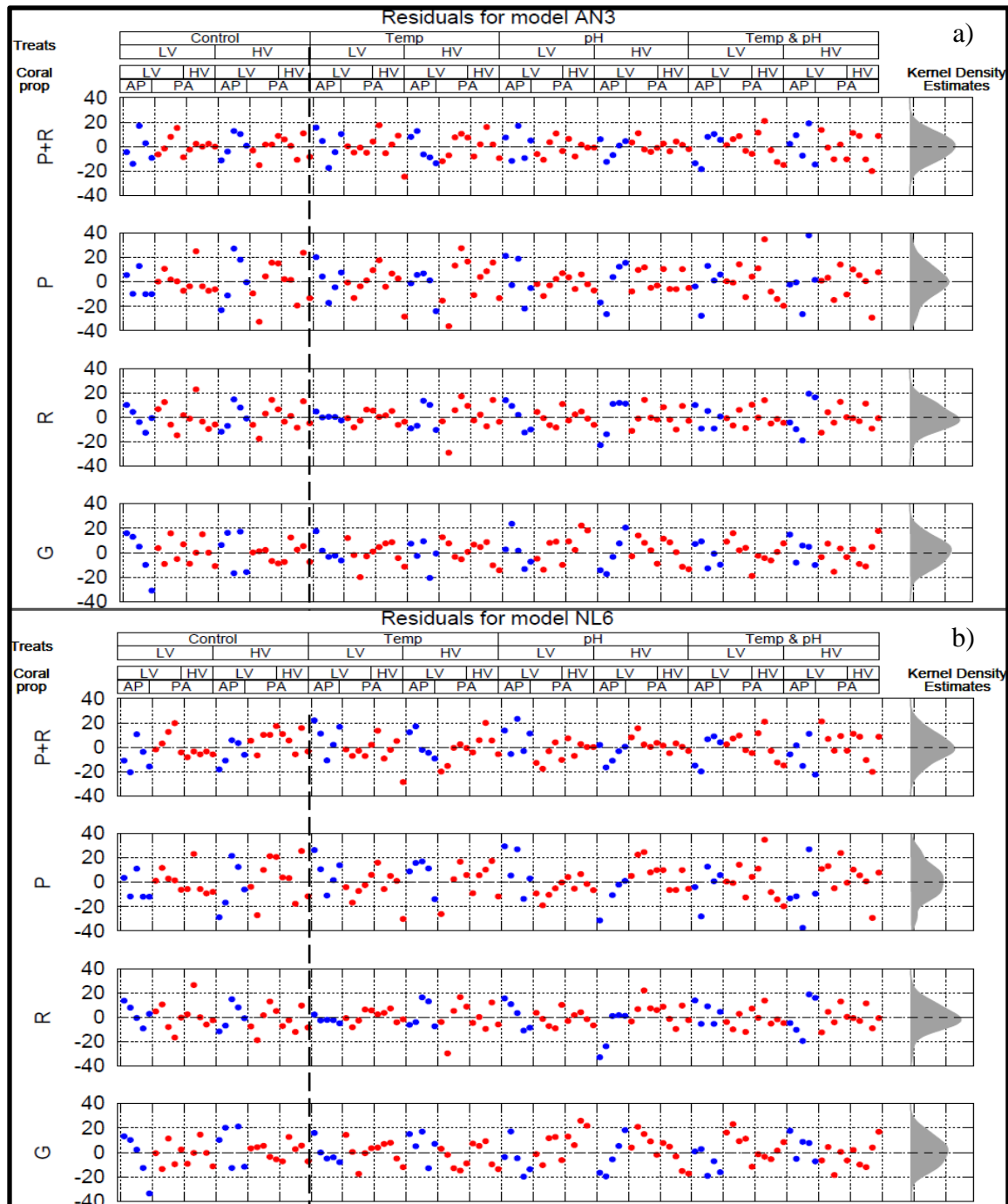


Figure S6| Residuals of models: a) AN3 and b) NL6, showing those for controls (left of the dashed line) and treatments (right of the dashed line.) The issue is whether residuals from the controls (left of the dashed line) are differently distributed from the treatments (right of dashed line). This was quantified by the Kolmogorov-Smirnoff test (table S6).

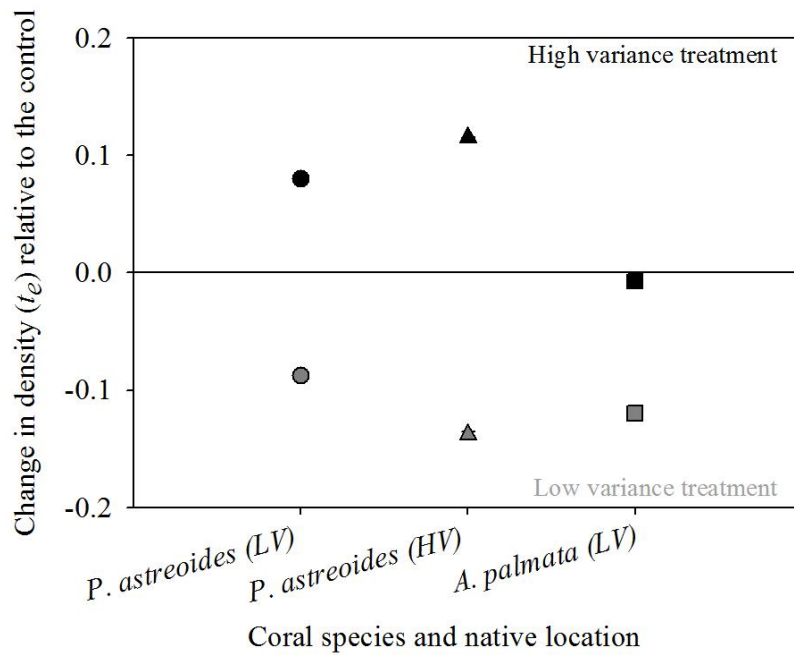


Figure S7| The changes in coral density over the experimental period. The average (\pm standard error) changes in density for each coral species, from each habitat (outer-reef, low-variance (LV), seagrass, high-variance (HV)) for both the HV and LV treatments, relative to the controls at the end of the experiment (t_e). Density was determined on a sub-sample of corals from each experimental treatment, ($n= 3$ per treatment, total $n= 24$) using a 3D scanning process following the protocol of [27].

Electronic Supplementary Material- Statistical Analysis

Summary of models:

Saturating model: SAT

This is a somewhat crude model that gives each experiment treatment a mean and assumes a common error. It corresponds to maximum number of variables that can be used to describe the results. It serves as an upper bound on model complexity.

Linear models / ANOVA: AN1-3

These models were validations against the R package. As the experiment was not fully factorial it was necessary to eliminate species-habitat interaction terms from the ANOVA. In R these were written as:

$$\text{AN1: } v \sim 1 + s + h + v + T + pH$$

$$\text{AN2: } v \sim 1 + (s + h + v + T + pH)^2 - s : h$$

$$\text{AN3: } v \sim 1 + (s + h + v + T + pH)^3 - s : h : (1 + v + T + pH)$$

Non-linear models : NL1-6

A set of nested non-linear models were fitted to the data (table S2). The form for each model was chosen from initial exploratory analyses that suggested the major effects were pH, temperature and species. The general form these non-linear models were:

$$V = \underbrace{V_0}_{\text{Baseline}} \times \underbrace{\left[1 + \begin{matrix} T & (b_T + b_{T,v} v + b_{T,h} h) \\ + pH & (c_{pH} + c_{pH,v} v + c_{T,h} h) \\ + T \cdot pH & (d_{T,pH} + d_{T,pH,v} v + d_{T,pH,h} h) \end{matrix} \right]}_{\text{Treatment effects}} .$$

The general form of the model (NL6) was then simplified and alternative non-linear models derived by setting various coefficients to zero. The nested models NL1-6 are listed in table S2. The values of the coefficients were found by likelihood maximisation, which the case of normally distributed errors is equivalent to least squares fitting. 95% confidence intervals were calculated by the log-profile method. The linear models (ANOVA) and non-linear models were compared using their corrected Akaike Information Criteria (AICc).

Motivation for model choice:

There were concerns with interpretation of ANOVA since:

- The data structure was non-factorial, which might make results of the ANOVA unreliable.
- It was difficult to rationalise the behaviour of the terms in the linear model / ANOVA. For example, take the ANOVA series of models for photosynthesis in table S3.

In table S3 the following can be seen:

- In the 1st order ANOVA the habitat variable is significant and has a positive effect.
- At 2nd order the habitat is not significant and has changed sign. However, there is a significant habitat:temperature term and an almost significant habitat:pH term.
- The significant 2nd-order terms containing habitat become less significant at 3rd order.
- Some terms, especially the variability:pH term, change dramatically between 2nd-order and 3rd-order, suggesting that they may not have converged.

This makes it difficult to assess:

- Whether there is confounding or aliasing of the main effects with higher interactions;
- If there are artefacts arising from not having fully factorial data.

For these reasons a set of non-linear models were constructed. Usually non-linear models are used to build more complicated functional dependence in main effect. In this case we have built non-linearity into the interactions by assuming a multiplicative structure in the terms of the model. This allows sparseness to be enforced through model specification. The nonlinear models have significantly fewer terms (23 for 3rd order ANOVA vs 13 for NL6), yet comparable performance in terms of AICc scores – NL6 has a lower relative AIC for 4 out of the 5 observations (table S4). NL6 was used as it allows the role of habitat to be decomposed into multiple contributions; the differences in the ‘baseline’ condition, and whether habitat affects the response to temperature stress, pH stress and the combined stress of temperature and pH.

The nested models NL1-6 also allow the effect of each additional degree of flexibility to be introduced incrementally. It is reassuring to see the stability of parameters and that, for example, the baseline habitat parameter changes only when addition flexibility related to habitat is added. While perhaps not important in this case, in general there are multiple solutions for nonlinear models so it is prudent to use the optimised parameters from a simpler models as initial parameters for more complex models (table S5).

Comparison of AN3 and NL6 models

The parameters in the ANOVA/linear model can be compared to a non-linear model by performing a power series expansion (ie. expanding out the non-linear model and collecting the terms order-by-order). This is shown for AN3 and NL6 in table S6. The arrows highlight how the second order habitat terms come from multiple sources in NL6.

From table S6 it can be seen that:

- The models are in general agreement, of particular interest are the habitat:temperature and habitat:pH terms.
- In the expansion of the NL6 model the habitat:temperature and habitat:pH terms primarily come from differences between the stress response due to habitat. This suggests that confounding/aliasing of main effects of habitat and pH/temp are unlikely to be a problem in ANOVA / linear models.

In conclusion:

The linear and non-linear models are broadly in agreement, with the non-linear models allowing some of the subtler points to be tested. Consistently across all the observations (P,R,G) habitat gave a small positive effect (<+4%) which in only a single case (photosynthesis) rose above the threshold for statistical significance. The small benefit of exposure to higher variability conditions is unable to offset the larger negative effects of temperature, pH and temperature:pH (ranging from -20 to -40%). Overall the influence of native habitat was therefore negligible.

Testing for heteroskedasticity

Concerns were raised about the heteroskedasticity of errors in the dependent variables as the variance in the independent variables was slightly higher in the future temperature and pH manipulations. We tested the analysis of residuals for model NL6 and AN3. Residuals were plotted and grouped into control and treatment. Kolmogorov–Smirnov tests were performed to verify that residuals for the control and treatment appear to come from the same distribution (table S7). The Bruesch-Pagan test was used on the ANOVA/linear models with no significant effects (except for respiration whose *p*-value 0.047). The non-linear model NL6 was extended to have two error terms (one error term for control, one temperature, and pH manipulations). Based on AIC difference, the increased complexity of an additional error term was not justified by better fitting the data (table S8). We therefore concluded there was no evidence of heteroskedasticity in this data set.

Supplementary Tables

Table S1| Physio-chemical tank and *in situ* conditions.

Physio-chemical variable	<i>In situ</i>		High-variance (HV)				Low-variance (LV)			
	HV	LV	Control	Temp.	pH	Temp. & pH	Control	Temp.	pH	Temp. & pH
pH (total scale)	8.154 ± 0.1	8.121 ± 0.1	8.148 ± 0.2	8.145 ± 0.2	7.809 ± 0.2	7.817 ± 0.3	8.115 ± 0.1	8.119 ± 0.1	7.827 ± 0.2	7.841 ± 0.3
Temperature (°C)	28.0 ± 0.3	27.4 ± 0.1	27.7 ± 0.2	30.1 ± 0.2	27.9 ± 0.3	30.3 ± 0.3	27.6 ± 0.3	29.6 ± 0.2	27.8 ± 0.4	29.5 ± 0.3
Aragonite saturation state	4.0 ± 0.1	3.0 ± 0.1	3.8 ± 0.1	3.9 ± 0.1	2.2 ± 0.1	2.4 ± 0.2	2.9 ± 0.1	2.8 ± 0.2	2.2 ± 0.2	2.1 ± 0.2
<i>p</i> CO ₂ (µatm)	280 ± 8.9	310 ± 12.1	290 ± 9.6	297 ± 13.2	700 ± 18.9	719 ± 14.1	311 ± 6.5	303 ± 8.0	740 ± 23.1	722 ± 16.6
Total Alkalinity (µmol Kg/SW)	2280.7 ± 0.2	2250.9 ± 0.1	2255.3 ± 0.4	2295.3 ± 0.2	2265.6 ± 0.3	2245.4 ± 0.3	2228.9 ± 0.2	2226.0 ± 0.1	2243.7 ± 0.3	2209.0 ± 0.4
Salinity (ppm)	35.5 ± 0.2	35.1 ± 0.1	35.2 ± 0.3	35.3 ± 0.2	35.2 ± 0.1	35.1 ± 0.1	35.0 ± 0.2	35.1 ± 0.2	35.2 ± 0.1	35.0 ± 0.1
Light (µmol photons m ⁻² s ⁻¹)	406.5 ± 46.9	458.8 ± 28.7	435.2 ± 30.1	440.7 ± 25.9	450.3 ± 15.0	463.2 ± 35.0	445.1 ± 18.1	463.9 ± 15.2	470.1 ± 30.8	450.8 ± 26.1
Nitrates (µM)	0.85 ± 0.1	1.10 ± 0.1	0.95 ± 0.1	0.90 ± 0.1	0.92 ± 0.1	0.90 ± 0.1	1.04 ± 0.1	0.99 ± 0.1	1.05 ± 0.1	1.01 ± 0.1

In situ discrete water samples were taken weekly over the study duration ($n= 8$) to obtain all of the physio-chemical conditions. Temperature, salinity, pH, and total alkalinity experimental conditions were determined from discrete water samples collected at the start of every water exchange ($n= 531$). Total alkalinity and pH were used with temperature and salinity to calculate *p*CO₂ and aragonite saturation state ($n= 531$). Light and nitrates were measured daily within each experimental treatment ($n= 59$).

Table S2| The parameters included in the non-linear models (NL1-NL6)

	NL1	NL2	NL3	NL4	NL5	NL6
V_0	•	•	•	•	•	•
a_s	•	•	•	•	•	•
a_h	•	•	•	•	•	•
b_T	•	•	•	•	•	•
$b_{T,v}$			•	•	•	•
$b_{T,h}$					•	•
c_{pH}	•	•	•	•	•	•
$c_{pH,v}$			•	•	•	•
$c_{pH,h}$					•	•
$d_{pH,T}$		•	•	•	•	•
$d_{pH,T,v}$				•	•	•
$d_{pH,T,h}$						•

Model parameters for the first six non-linear models. The parameters included were: species (s), habitat (h), experimental treatment variability (v), temperature (t) and pH (pH).

Table S3| Point estimates for the parameters in the ANOVA, linear models.

(Intercept)	AN1		AN2		AN3	
	322.6	***	298.3	***	297.1	***
Species	17.9	***	21.0	***	23.1	***
Habitat	6.7	*	-1.7		-2.8	
Variance	-1.5		-4.1		-0.1	
Temperature	-16.3	**	28.9	***	31.0	***
pH	-97.4	***	-52.8	***	-50.7	***
species:variance			3.7	*	2.1	
species:temp			-5.7		-9.8	*
species:pH			-0.6		-4.7	
habitat:varince			-1.3		-0.2	
habitat:temp			6.6	.	8.9	.
habitat:pH			10.2	**	12.5	*
variance:temp			15.9	***	12.0	**
variance:pH			-14.2	***	-24.6	***
temp:pH			-82.1	***	-86.4	***
species:variance:temp					-4.2	
sppecies:variance:pH					7.4	*
sppecies:temp:pH					8.3	
habitat:variance:temp					2.9	
habitat:variance:pH					-5.0	
habitat:temp:pH					-4.2	
variance:temp:pH					12.6	*
Studentized Bruesch-Pagan <i>p</i> -value	0.3		0.3		0.2	

Terms relating to the habitat variable are highlighted in bold. (*: $p < 0.05$, **: $p < 0.01$, ***: $p < 0.001$)

Table S4| Comparison of the corrected Akaike Information Criteria (AICc) for all models.

Model	K	Photosynthesis (G)		Respiration		Photosynthesis (N)		Calcification	
		AICc	AICc rel	AICc	AICc rel	AICc	AICc rel	AICc	AICc rel
NL1	6	1144.6	209.9	915.5	0.7	1160.0	136.1	994.6	29.1
NL2	7	1026.0	91.3	916.5	1.7	1079.0	55.1	994.8	29.3
AN1	7	1141.1	206.4	914.8	0.0	1156.6	132.7	1009.4	43.9
NL3	9	987.1	52.5	918.1	3.4	1046.4	22.5	986.7	21.2
NL4	10	988.4	53.7	915.9	1.2	1044.5	20.6	984.4	18.9
NL5	12	939.8	5.1	917.6	2.9	1023.9	0.0	965.5	0.0
NL6	13	939.0	4.3	916.4	1.6	1026.4	2.5	968.0	2.5
AN2	16	946.3	11.6	924.1	9.3	1023.9	0.0	973.0	7.5
AN3	23	934.7	0.0	930.2	15.5	1029.3	5.4	979.6	14.1
SAT	25	938.9	4.2	936.3	21.5	1034.0	10.1	983.1	17.6

Non-linear (NL) models 1-6 and Analysis of Variance (ANOVA) 2-3 were compared using their AICc with decreasing number of interaction terms (k) for the parameters: Respiration, Photosynthesis (Net and Gross) and Calcification. AICc rel. is the difference in AICc relative to best performing model (lowest AICc score). NL6 was used as it allows the role of habitat to be decomposed into multiple contributions; the differences in the ‘baseline’ condition, and whether habitat affects the response to temperature stress, pH stress and the combined stress of temperature and pH.

Table S5| Point estimates for the variables in the non-linear models.

		NL1	NL2	NL3	NL4	NL5	NL6
Baseline	Intercept	321.1	301.2	301.4	301.4	299	299.1
	Species	0.066	0.065	0.064	0.064	0.066	0.065
	Habitat	0.021	0.021	0.021	0.021	-0.005	-0.004
Temp	Temperature	-0.050	0.080	0.080	0.080	0.084	0.083
	Temperature:Variance			0.047	0.034	0.034	0.034
	Temperature:Habitat					0.014	0.012
pH	pH	-0.296	-0.183	-0.183	-0.183	-0.172	-0.173
	pH: Variance			-0.050	-0.063	-0.063	-0.063
	pH:Habitat					0.040	0.038
Temp & pH	Temperature& pH		-0.266	-0.266	-0.266	-0.266	-0.265
	Temperature& pH:Variance				0.037	0.037	0.037
	Temperature& pH:Habitat						0.005

Table S6| Comparison between the parameters of the AN3 model and NL6 model by power expansion of NL6.

		AN3			NL6 expanded
		Linear Variable	Est	sig	Est
1st order	(Intercept)		297.2	***	299.1
	Species (spe)		23.1	***	19.4
	Habitat (hab)		-2.9		-1.2
	Variance (var)		-0.1		NA
	Temp		31.0	***	24.8
	pH		-50.7	***	-51.7
2nd order	spe:var		2.1		
	spe:temp		-9.8	*	1.6
	spe:pH		-4.7		-3.4
	hab:var		-0.2		NA
	hab:temp		8.9	.	10.1
	Hab:pH		12.5	*	11.6
	var:temp		12.0	**	10.2
	var:pH		-24.6	***	-18.8
	temp:pH		-86.4	***	-83.6
	3rd order	spe:var:temp		-4.2	
spe:var:pH			7.4	*	-1.2
spe:temp:pH			8.3		-5.4
hab:var:temp			2.9		-0.1
hab:var:pH			-5.0		0.1
hab:temp:pH			-4.2		1.8
var:temp:pH			12.6	*	11.1

		NL6 Non-linear model			
		sig	Variable		
299.1	***		V0	baseline	
6.5	***	Spe	a		
-0.4		Hab			
8.3	***	Temp	b	Temp effect	
3.4	***	Temp:var			
1.2		Temp:hab			
-17.3	***	pH	c	pH effect	
-6.3	***	pH:var			
3.8	**	pH:hab			
-26.5	***	Temp:pH	d	Temp & pH	
3.7	*	T:pH:var			
0.5		Temp:pH:hab			

The parameters in the ANOVA/linear model can be compared to a non-linear model by performing a power series expansion (ie. expanding out the non-linear model and collecting the terms order-by-order). This is shown for AN3 and NL6 in table S5. The arrows highlight how the second order habitat terms come from multiple sources in NL6

Table S7| Kolmogorov-Smirnov tests of whether the residuals in the control and treatments appear to come from different distributions.

	Kolmogorov-Smirnov		Bruesh-Pagan	
	<i>p</i> -values (two sided)		<i>p</i> -value	
	NL6	AN3	AN3	
Photosynthesis (P)	0.74	0.74	0.22	
Respiration (R)	0.65	0.82	0.05	
P&R	0.27	0.82	0.06	
Calcification	0.94	0.89	0.14	

Table S8| Comparison between the NL6 model with a single error term and NL6 with two error terms (one for control, one for treatment).

		AICc	sigma	
			control	treatment
Photosynthesis	one error	299.09		15.41
	two error	298.96	14.26	15.78
Respiration	one error	150.97		9.74
	two error	150.96	9.99	9.66
PR	one error	148.79		10.71
	two error	148.64	10.24	10.86
Calcification	one error	248.58		12.08
	two error	248.59	11.70	12.21

Table S9| Physio-chemical tank and *in situ* conditions.

Physio-chemical variable	<i>In situ</i>		High-variance (HV)				Low-variance (LV)			
	HV	LV	Control	Temp.	pH	Temp. & pH	Control	Temp.	pH	Temp. & pH
pH (total scale)	8.154 ± 0.1	8.121 ± 0.1	8.148 ± 0.2	8.145 ± 0.2	7.809 ± 0.2	7.817 ± 0.3	8.115 ± 0.1	8.119 ± 0.1	7.827 ± 0.2	7.841 ± 0.3
Temperature (°C)	28.0 ± 0.3	27.4 ± 0.1	27.7 ± 0.2	30.1 ± 0.2	27.9 ± 0.3	30.3 ± 0.3	27.6 ± 0.3	29.6 ± 0.2	27.8 ± 0.4	29.5 ± 0.3
Aragonite saturation state	4.0 ± 0.1	3.0 ± 0.1	3.8 ± 0.1	3.9 ± 0.1	2.2 ± 0.1	2.4 ± 0.2	2.9 ± 0.1	2.8 ± 0.2	2.2 ± 0.2	2.1 ± 0.2
<i>p</i> CO ₂ (µatm)	280 ± 8.9	310 ± 12.1	290 ± 9.6	297 ± 13.2	700 ± 18.9	719 ± 14.1	311 ± 6.5	303 ± 8.0	740 ± 23.1	722 ± 16.6
Total Alkalinity (µmol Kg/SW)	2280.7 ± 0.2	2250.9 ± 0.1	2255.3 ± 0.4	2295.3 ± 0.2	2265.6 ± 0.3	2245.4 ± 0.3	2228.9 ± 0.2	2226.0 ± 0.1	2243.7 ± 0.3	2209.0 ± 0.4
Salinity (ppm)	35.5 ± 0.2	35.1 ± 0.1	35.2 ± 0.3	35.3 ± 0.2	35.2 ± 0.1	35.1 ± 0.1	35.0 ± 0.2	35.1 ± 0.2	35.2 ± 0.1	35.0 ± 0.1
Light (µmol photons m ⁻² s ⁻¹)	406.5 ± 46.9	458.8 ± 28.7	435.2 ± 30.1	440.7 ± 25.9	450.3 ± 15.0	463.2 ± 35.0	445.1 ± 18.1	463.9 ± 15.2	470.1 ± 30.8	450.8 ± 26.1
Nitrates (µM)	0.85 ± 0.1	1.10 ± 0.1	0.95 ± 0.1	0.90 ± 0.1	0.92 ± 0.1	0.90 ± 0.1	1.04 ± 0.1	0.99 ± 0.1	1.05 ± 0.1	1.01 ± 0.1

In situ discrete water samples were taken weekly over the study duration ($n= 8$) to obtain all of the physio-chemical conditions. Temperature, salinity, pH, and total alkalinity experimental conditions were determined from discrete water samples collected at the start of every water exchange ($n= 531$). Total alkalinity and pH were used with temperature and salinity to calculate *p*CO₂ and aragonite saturation state ($n= 531$). Light and nitrates were measured daily within each experimental treatment ($n= 59$).

Table S9| Raw data measurements before experimental manipulation (t_0) and at the end (t_e) of the experimental period for the corals *Acropora palmata* and *Porites astreoides* from a high-variance (HV) seagrass habitat and a low-variance (LV) outer-reef location on Little Cayman, Cayman Islands, BWI. Measurements include: daily net photosynthesis (P_N), daily respiration (R), daily calcification (G), and growth rates. Note: buoyant mass rates were calculated from the end of the acclimatization period (t_i) to t_e .

Variability	Treatment	Species	P_N (mmol m ⁻² d ⁻¹)		R (mmol m ⁻² d ⁻¹)		G (mmol m ⁻² d ⁻¹)		Zooxanthellae density (cells/cm ⁻²)		Chlorophyll a (µg cm ⁻²)		Buoyant Mass (mg d ⁻¹ g ⁻¹ coral)
			t_0	t_e	t_0	t_e	t_0	t_e	t_0	t_e	t_0	t_e	Rate (t_i to t_e)
High	Control	<i>A.palmata LV</i>	132.0 ± 1.7	138.3 ± 0.3	142.1 ± 2.1	139.5 ± 3.3	274.1 ± 1.3	272.2 ± 2.1	1.1 × 10 ⁶ ± 3.1 × 10 ⁴	1.2 × 10 ⁶ ± 3.4 × 10 ⁴	14.2 ± 2.1	15.0 ± 3.2	6.08 ± 1.2
		<i>P.astreoides HV</i>	157.2 ± 0.6	161.2 ± 1.7	165.2 ± 1.9	159.1 ± 1.7	226.1 ± 1.6	231.6 ± 2.6	2.0 × 10 ⁶ ± 7.5 × 10 ⁴	2.2 × 10 ⁶ ± 6.3 × 10 ⁴	12.4 ± 1.8	12.0 ± 1.5	3.93 ± 2.1
		<i>P.astreoides LV</i>	162.1 ± 2.2	157.8 ± 2.8	169.9 ± 3.1	163.8 ± 2.5	230.1 ± 1.0	221.4 ± 7.7	1.3 × 10 ⁶ ± 3.8 × 10 ⁴	1.2 × 10 ⁶ ± 1.9 × 10 ⁴	13.1 ± 2.0	13.6 ± 4.0	3.60 ± 2.1
	Temperature	<i>A.palmata LV</i>	129.9 ± 3.2	173.7 ± 1.2	140.7 ± 3.2	138.1 ± 5.1	280.1 ± 0.7	227.3 ± 0.9	1.9 × 10 ⁶ ± 7.5 × 10 ⁵	5.1 × 10 ⁵ ± 5.0 × 10 ⁴	13.9 ± 2.2	7.5 ± 0.9	4.71 ± 0.8
		<i>P.astreoides HV</i>	142.1 ± 2.2	193.8 ± 2.8	150.1 ± 0.8	159.1 ± 3.7	231.9 ± 0.6	189.3 ± 1.1	1.9 × 10 ⁶ ± 2.9 × 10 ⁴	2.0 × 10 ⁶ ± 3.2 × 10 ⁴	12.8 ± 1.4	11.4 ± 2.0	2.71 ± 2.3
		<i>P.astreoides LV</i>	160.4 ± 2.1	179.3 ± 4.9	152.1 ± 5.9	160.9 ± 7.9	223.1 ± 0.2	169.5 ± 1.3	1.1 × 10 ⁶ ± 3.8 × 10 ⁴	9.9 × 10 ⁵ ± 2.1 × 10 ⁴	12.9 ± 2.7	13.0 ± 1.6	2.22 ± 3.2
	pH	<i>A.palmata LV</i>	127.8 ± 1.9	72.1 ± 2.2	137.2 ± 2.6	130.9 ± 3.6	277.3 ± 1.7	185.4 ± 2.7	1.9 × 10 ⁶ ± 9.8 × 10 ⁴	7.7 × 10 ⁵ ± 4.9 × 10 ⁴	15.2 ± 3.1	12.8 ± 1.9	3.58 ± 1.1
		<i>P.astreoides HV</i>	155.1 ± 3.8	100.1 ± 1.7	155.6 ± 0.7	160.6 ± 3.7	233.1 ± 0.2	140.6 ± 2.7	2.1 × 10 ⁶ ± 3.5 × 10 ⁴	2.2 × 10 ⁶ ± 6.0 × 10 ⁴	12.2 ± 2.9	11.0 1.6	2.38 1.7
		<i>P.astreoides LV</i>	162.9 ± 0.5	188.5 ± 2.3	151.1 ± 0.3	162.7 ± 4.1	228.1 ± 4.3	138.2 ± 3.1	1.1 × 10 ⁶ ± 5.6 × 10 ⁵	9.9 × 10 ⁵ ± 2.3 × 10 ⁴	13.1 ± 1.4	12.7 ± 1.1	1.85 ± 0.9
	Temperature & pH	<i>A.palmata LV</i>	139.2 ± 2.4	28.1 ± 2.8	139.1 ± 0.9	138.5 ± 4.4	265.9 ± 6.4	118.6 ± 2.0	2.0 × 10 ⁶ ± 1.0 × 10 ⁵	3.7 × 10 ⁵ ± 4.8 × 10 ⁴	14.0 ± 2.3	5.3 ± 1.9	2.56 ± 2.0
		<i>P.astreoides HV</i>	141.2 ± 6.4	68.8 ± 4.1	151.6 ± 1.1	158.6 ± 3.3	225.5 ± 0.2	122.2 ± 0.2	1.0 × 10 ⁶ ± 4.6 × 10 ⁵	1.1 × 10 ⁶ ± 5.3 × 10 ⁴	13.8 ± 2.8	11.3 ± 2.6	1.78 ± 1.1
		<i>P.astreoides LV</i>	146.2 ± 1.9	42.1 ± 3.1	158.3 ± 5.5	164.9 ± 4.2	225.1 ± 2.4	115.3 ± 6.9	1.1 × 10 ⁶ ± 6.8 × 10 ⁴	9.9 × 10 ⁵ ± 3.6 × 10 ⁴	12.7 ± 0.9	12.1 ± 3.1	1.76 ± 0.6

Variability	Treatment	Species	P _N (mmol m ⁻² d ⁻¹)		R (mmol m ⁻² d ⁻¹)		G (mmol m ⁻² d ⁻¹)		Zooxanthellae density (cells/cm ²)		Chlorophyll a (µg cm ⁻²)		Buoyant Mass (mg d ⁻¹ g ⁻¹ coral)
			t ₀	t _e	t ₀	t _e	t ₀	t _e	t ₀	t _e	t ₀	t _e	Rate (t _i to t _e)
Low	Control	<i>A.palmata LV</i>	129.2 ± 1.1	134.8 ± 1.9	135.1 ± 0.6	141.7 ± 1.9	272.0 ± 0.9	269.6 ± 1.7	2.0 × 10 ⁶ ± 7.7 × 10 ⁵	1.9 × 10 ⁶ ± 7.3 × 10 ⁴	14.6 ± 2.7	12.2 ± 1.2	6.21 ± 1.1
		<i>P.astreoides HV</i>	143.9 ± 2.1	149.3 ± 2.7	170.3 ± 1.3	167.3 ± 4.7	231.0 ± 3.6	229.0 ± 5.9	1.1 × 10 ⁶ ± 5.6 × 10 ⁵	1.1 × 10 ⁶ ± 6.2 × 10 ⁴	12.1 ± 2.1	10.2 ± 1.2	3.74 ± 2.1
		<i>P.astreoides LV</i>	160.1 ± 2.4	158.8 ± 4.9	160.1 ± 0.3	163.2 ± 1.8	221.1 ± 6.1	218.6 ± 0.9	1.2 × 10 ⁶ ± 3.5 × 10 ⁵	1.2 × 10 ⁶ ± 2.6 × 10 ⁴	11.8 ± 2.0	14.5 ± 1.8	3.46 ± 2.1
	Temperature	<i>A.palmata LV</i>	133.8 ± 2.1	164.5 ± 4.5	134.2 ± 0.9	135.0 ± 1.7	225.1 ± 0.4	203.8 ± 1.3	1.8 × 10 ⁶ ± 9.0 × 10 ⁴	9.1 × 10 ⁵ ± 4.9 × 10 ⁴	14.4 ± 1.6	7.9 ± 1.1	4.88 ± 1.1
		<i>P.astreoides HV</i>	149.3 ± 2.4	170.6 ± 6.2	167.7 ± 7.8	163.5 ± 2.0	272.0 ± 0.9	183.0 ± 1.4	1.1 × 10 ⁶ ± 1.9 × 10 ⁴	1.0 × 10 ⁶ ± 3.3 × 10 ⁴	11.3 ± 2.1	9.8 ± 2.6	2.64 ± 1.2
		<i>P.astreoides LV</i>	140.1 ± 0.1	163.9 ± 2.2	171.9 ± 1.6	163 ± 2.7	231.0 ± 0.6	168.8 ± 1.1	1.3 × 10 ⁶ ± 8.7 × 10 ⁴	1.0 × 10 ⁶ ± 3.0 × 10 ⁵	11.9 ± 2.9	13.0 ± 2.1	2.07 2.2
	pH	<i>A.palmata LV</i>	163.9 ± 1.5	111.8 ± 3.4	144.3 ± 2.9	137.8 ± 5.2	221.0 ± 1.6	152.9 ± 8.7	1.5 × 10 ⁶ ± 1.0 × 10 ⁵	1.3 × 10 ⁶ ± 3.7 × 10 ⁴	15.0 ± 2.1	9.4 ± 3.4	3.53 ± 1.3
		<i>P.astreoides HV</i>	147.3 ± 3.1	125.3 ± 1.1	168.1 ± 4.1	168.8 ± 3.9	270.0 ± 1.2	159.0 ± 1.7	1.0 × 10 ⁶ ± 2.1 × 10 ⁵	1.2 × 10 ⁶ ± 3.4 × 10 ⁴	11.9 ± 2.8	10.1 ± 1.6	2.5 ± 1.2
		<i>P.astreoides LV</i>	139.9 ± 3.4	103.2 ± 5.5	152.3 ± 7.5	139.5 ± 3.5	235.0 ± 6.1	133.1 ± 3.9	1.5 × 10 ⁶ ± 6.7 × 10 ⁵	1.3 × 10 ⁶ ± 1.6 × 10 ⁵	13.2 ± 1.2	14.4 ± 3.2	2.06 ± 1.1
	Temperature & pH	<i>A.palmata LV</i>	137.9 ± 3.4	21.6 ± 2.2	144.7 ± 2.6	139.5 ± 3.8	229.6 ± 6.3	87.4 ± 7.9	2.3 × 10 ⁶ ± 9.0 × 10 ⁴	6.8 × 10 ⁵ ± 4.6 × 10 ⁴	14.2 ± 1.4	6.0 ± 1.5	2.4 ± 0.9
		<i>P.astreoides HV</i>	153.1 ± 2.8	63.4 ± 2.1	162.1 ± 6.7	164.8 ± 3.5	275.0 ± 3.1	76.4 ± 2.9	1.4 × 10 ⁶ ± 9.9 × 10 ⁴	1.5 × 10 ⁶ ± 2.6 × 10 ⁵	10.1 ± 1.0	9.7 ± 2.3	2.09 ± 1.3
		<i>P.astreoides LV</i>	146.2 ± 2.1	28.8 ± 2.9	150.3 ± 1.9	158.9 ± 3.7	230.0 ± 2.1	87.9 ± 1.6	1.1 × 10 ⁶ ± 6.0 × 10 ⁵	1.0 × 10 ⁶ ± 1.2 × 10 ⁴	13.3 ± 1.6	12.7 ± 0.7	1.61 ± 1.6