CLONAL VARIATION FOR PHENOTYPIC PLASTICITY IN THE CORAL MADRACIS MIRABILIS

JOHN F. BRUNO1 AND PETER J. EDMUNDS

Department of Biology, California State University, 18111 Nordhoff Street, Northridge, California 91330 USA

Abstract. Morphological plasticity is common among clonal organisms, including scleractinian corals, yet the role of phenotypic plasticity in coral ecology and evolution is largely unexplored. Additionally, it is unclear how much variation in plastic responses exists among individuals, populations, and species, and thus how much potential there is for natural selection to act on coral reaction norms. In the branching coral Madracis mirabilis, corallite architecture and density, branch diameter and spacing, and overall aggregate morphology all vary among environments. To examine the role of phenotypic plasticity in generating these patterns, clonal replicates of five genotypes of M. mirabilis were transplanted from each of two source populations into four treatment environments on the north coast of Jamaica. Flow rate, sedimentation, irradiance, water temperature, and salinity all varied among these environments. DNA fingerprinting was used to ensure that the 10 transplanted genotypes were genetically distinct. Six morphological traits (intersepta area, septa length, columella area, corallite area, corallite spacing, and branch tip diameter) were measured after transplantation to determine whether they were altered in response to environmental conditions. Because these traits were correlated, principal components analysis was used to define new, uncorrelated traits for analysis. Four of the five corallite traits and branch diameter were significantly affected by the environment, demonstrating that morphological variation among environments in M. mirabilis is due in large part to phenotypic plasticity. No difference was detected between the two source populations in the magnitude or direction of their plastic responses, but there was substantial variation among genotypes (genotype × environment interaction). Many of the phenotypic changes of both populations resulted in the transplants becoming morphologically similar to resident conspecifics in each treatment environment. Genotypes from both populations were able to maintain similar growth rates under diverse environmental conditions. Such morphological convergence by phenotypic plasticity may expand the ecological range of this species by enabling genotypes to tolerate spatially and temporally variable environments.

Key words: genotype × environment interaction; Madracis mirabilis; phenotypic plasticity; reaction norm; Scleractinia; skeletal structure.

INTRODUCTION

Scleractinian corals typically display a striking degree of morphological variation in colony shape and corallite structure along environmental gradients (Foster 1980, Brown et al. 1985), and among geographic regions (Veron 1981, Veron and Wallace 1984). Such variation can be caused by genetic differences among individuals and populations (Ayre and Willis 1988), by the environment (Foster 1979), or by both. Environmental control of morphological traits is termed phenotypic plasticity and a reaction norm describes the relationship between the phenotype and the environment (Bradshaw 1965, Schlichting 1986, Stearns 1989). Phenotypic plasticity is well known among numerous other taxa, including plants (Scheiner and Goodnight 1984, Schmitt 1993), sponges (Palumbi 1984), bacteria (Forst and Inouye 1988), fish (Meyer 1987), barnacles (Lively 1986), mollusks (Martín-Mora et al. 1995, Trussell 1996), and bryozoans (Harvell 1986), where it is thought to be ecologically important because it confers broad adaptability (Bradshaw 1965). Plasticity can also have important evolutionary implications (Sultan 1987). For example, within a single environment, plasticity can maintain genetic diversity under stabilizing selection by enabling a variety of genotypes to display a similar advantageous phenotype (Bradshaw 1965, Sultan and Bazzaz 1993). In addition, there are a number of mechanisms through which phenotypic plasticity might accelerate the appearance of novel phenotypes and the rates of speciation and macroevolution (West-Eberhard 1989).

A variety of life history traits indicate that plasticity could be important in corals as well. For example, corals are sessile and many are clonal (Hughes 1983, 1989), relying heavily on asexual modes of reproduction, including fragmentation (Tunnicliffe 1981, High-
of the effects of reduced genetic diversity (Bradshaw 1985). As a branch extends, the tissue at the base of the branch recedes and the newly exposed portion of the skeleton can become colonized by boring sponges and algae. Thus, each branch is a functional colony of physiologically integrated polyps (Connell 1973), and each aggregation is likely to be a genotype made up of numerous clonal replicates (branches).

Environment

All field work took place on the north coast of Jamaica at the Discovery Bay Marine Laboratory (DBML). The reefs near DBML were originally described by Goreau (1959) and subsequently in numerous other studies (e.g., Goreau and Goreau 1973, Liddell and Ohlhorst 1987, 1992, Edmunds and Bruno 1996). Four reef sites were used as treatment environments. Two were located on the exposed forereef at Dairy Bull cove, at 10 m (DB10) and 20 m depth (DB20). The other two were <3 km west of Dairy Bull at Columbus Park (10 m depth), which is a protected lagoon environment within Discovery Bay. Columbus Park Springs (CPS, 10 m depth) was adjacent to underwater, freshwater springs, which are numerous at this site (D’Elia et al. 1981). The other lagoon environment (CP10) was 25 m from the nearest spring, but also at 10 m depth. Differences in the physical con-
October 1997  PHENOTYPIC PLASTICITY IN MADRACIS  2179

Fig. 1. (A) Madracis mirabilis aggregation from Conch Reef, Florida Keys (25 m depth); (B) branches of M. mirabilis after transplantation from Dairy Bull at 20 m to (from left): Columbus Park Springs, Columbus Park at 10 m, Dairy Bull at 10 m, Dairy Bull at 20 m (scale bar = 1 cm); (C) scanning electron micrograph of corallites of M. mirabilis from Dairy Bull at 10 m; (D) scanning electron micrograph of corallites of M. mirabilis from Columbus Park Springs at 10 m. Note the arrow in C pointing to the hexagonally shaped ridges (pseudocostae) surrounding the corallite from Dairy Bull at 10 m, which are absent in the Columbus Park Springs at 10 m corallite.

ditions among the four treatment environments have been described in previous papers and include temperature, salinity, irradiance (Foster 1979, Edmunds 1989, Bruno 1995), nutrient concentration (D’Elia et al. 1981), and flow rate (Helmuth and Sebens 1993). These physical characters are summarized in Table 1. In this paper, the corals naturally living in each of the treatment environments are referred to as residents (as opposed to those that were experimentally transplanted there) and all of the conspecific corals from each environment are loosely termed a population. However, it is likely that our populations are all part of a larger metapopulation, and that they are genetically homogeneous due to gene flow between each other and external sources.

Natural morphological variation

A number of morphological traits in Madracis mirabilis vary along environmental gradients (Schindler 1985, Fenner 1993, Bruno 1995, Sebens et al. 1997). For example, branches from forereef habitats (e.g., DB10 and DB20) are generally cylindrical, while in lagoon habitats (e.g., CP10 and CPSP) branches have a larger diameter and bulbous, flattened tips (Fig. 1B; Schindler 1985, Fenner 1993, Bruno 1995). Interconnecting, hexagonally shaped ridges (pseudocostae) often surround the corallites of forereef branches but are absent in the lagoon (Fenner 1993; Fig. 1C, D). Additionally, the spacing among corallites is larger in forereef habitats than in lagoon environments (Schindler 1985, Bruno 1995; Fig. 1C, D). Corallites on lagoon branches and on the terminal ends of forereef branches have 10 thin primary septa and the columella is formed from junction of the septa (Fig. 1D). Corallites on the sides of branches from the forereef have a prominent columella, 10 thicker primary septa, and much smaller intersepta areas than lagoon corallites. Aggregate-level traits including branch spacing and aggregate shape also vary among environments (Bruno 1995, Sebens et al. 1997).

Four corallite traits (intersepta area, ISA; columella area, CLA; corallite area, CA; and corallite spacing, CS; Fig 2), and two branch traits (branch tip diameter, BD; and branch spacing, BSP) were measured on Mad-
**Table 1.** Relative differences in environmental characteristics of the four treatment environments. Values in parentheses were measured during June 1994 (Bruno 1995). Site abbreviations: DB10, Dairy Bull at 10 m; DB20, Dairy Bull at 20 m; CP10, Columbus Park at 10 m; CPS, Columbus Park Springs at 10 m.

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Source</th>
<th>DB10</th>
<th>DB20</th>
<th>CP10</th>
<th>CPS</th>
</tr>
</thead>
<tbody>
<tr>
<td>Water velocity</td>
<td>Helmuth and Sebens 1993; J. F. Bruno, unpublished data</td>
<td>high</td>
<td>med</td>
<td>low</td>
<td>high</td>
</tr>
<tr>
<td>Wave energy</td>
<td>Brakel 1976; J. F. Bruno, unpublished data</td>
<td>high</td>
<td>med</td>
<td>low</td>
<td>low</td>
</tr>
<tr>
<td>Light intensity (%)</td>
<td>Edmunds 1989, Bruno 1995</td>
<td>high (26%)</td>
<td>med (20%)</td>
<td>med (18%)</td>
<td>med (18%)</td>
</tr>
<tr>
<td>Temperature (°C)</td>
<td>D'Elia et al. 1981</td>
<td>low (29.5)</td>
<td>high (29.5)</td>
<td>med (28.5)</td>
<td>low (27.1)</td>
</tr>
<tr>
<td>Nutrient level (nitrogen)</td>
<td>Bruno 1995</td>
<td>low (35.0)</td>
<td>high (35.0)</td>
<td>med (30.0)</td>
<td>high</td>
</tr>
<tr>
<td>Salinity (g/kg)</td>
<td>Bruno 1995</td>
<td>high (35.0)</td>
<td>high (35.0)</td>
<td>med (30.0)</td>
<td>low (26.5)</td>
</tr>
</tbody>
</table>

*Madracis mirabilis* aggregates from the four treatment environments (residents). These measurements were used to quantify field observations of morphological variation among environments and for comparison with transplanted corals (refer to Fig. 2 for further description of measured traits). Corallite area was included in the analysis so that the other corallite traits could be normalized to corallite size. To quantify the corallite traits, three branches were collected from each of five genotypes selected at random (each separated by >25 m) from each treatment environment. Three haphazardly selected corallites, 2 cm from the terminal branch end, were analyzed on each branch. Two measurements were made from each corallite for ISA (the largest and opposing area between adjacent septa), and CS (distance to the two nearest septa), and a single measurement was made on each corallite for CLA and CA.

To prepare the corals for analysis, the skeletons were bleached (with 50% bleach) to remove the tissue, rinsed in distilled water, and dried. Planar images of the skeleton surface (100×) were made of the branches with a video camera fitted to a dissecting microscope, and these were analyzed with imaging software (NIH Image 1.43). The two branch traits (BD and BSP) were measured in situ on five *M. mirabilis* aggregates (the same aggregates from which branches had been collected) from each of the four treatment environments. Branch diameter was measured with calipers 1 cm from the end of 10 branches on each aggregate. Branch spacing was measured as the distance between adjacent branches (10 pairs/aggregate).

Because the six measured skeletal traits were correlated, a one-factor MANOVA (MGHL platform, SYSTAT 3.0) was used to analyze multivariate variation among environments. After establishing the significance of the environment effect with MANOVA, univariate comparisons were performed with ANOVA. Genotype mean values for each trait were the statistical replicates, and analysis was performed on log-transformed data after all statistical assumptions had been met (Zar 1996). Environment was considered a fixed effect in all analyses because the four environments were chosen based on: (1) a priori assessment of site-specific variation in the morphology of *Madracis mirabilis*, and (2) data describing differences in environmental factors known to affect coral morphology (Table 1).

Principal components analysis (PCA) was carried out on the correlation matrix of log-transformed genotype means to investigate multivariate associations among the traits and to examine the nature of the multivariate differences among groups. Principal components (PC) that explained >10% of the total variance or that had an eigenvalue of >1.0 were interpreted (Nichols 1977, Jolliffe 1986). However, the final decision as to which components to use in subsequent analyses was based...
TABLE 2. Morphometric comparisons (means ± 1 SE) of Madracis mirabilis aggregations naturally living (resident corals) in each of the four treatment environments. F values are from univariate comparisons (ANOVA, one fixed factor = site; df = 3, 16) of each trait among the four environments. Multivariate comparison (fixed-effect MANOVA) among sites was significant (Wilks’ Λ = 0.012; F = 6.70; df = 18, 31; P < 0.001), DB10, Dairy Bull at 10 m; DB20, Dairy Bull at 20 m; CP10, Columbus Park at 10 m; CPSP, Columbus Park Springs at 10 m.

<table>
<thead>
<tr>
<th>Character</th>
<th>DB10</th>
<th>DB20</th>
<th>CP10</th>
<th>CPSP</th>
<th>F</th>
</tr>
</thead>
<tbody>
<tr>
<td>Corallite area (mm²)</td>
<td>1.75 ± 0.25</td>
<td>1.56 ± 0.17</td>
<td>1.58 ± 0.09</td>
<td>1.69 ± 0.10</td>
<td>0.29</td>
</tr>
<tr>
<td>Columella area (mm²)</td>
<td>0.35 ± 0.04</td>
<td>0.30 ± 0.05</td>
<td>0.23 ± 0.03</td>
<td>0.20 ± 0.03</td>
<td>3.05†</td>
</tr>
<tr>
<td>Interepsta area (mm²)</td>
<td>0.04 ± 0.01</td>
<td>0.04 ± 0.01</td>
<td>0.07 ± 0.01</td>
<td>0.09 ± 0.01</td>
<td>11.98***</td>
</tr>
<tr>
<td>Corallite spacing (mm)</td>
<td>0.44 ± 0.03</td>
<td>0.44 ± 0.04</td>
<td>0.38 ± 0.10</td>
<td>0.18 ± 0.03</td>
<td>6.28</td>
</tr>
<tr>
<td>Branch tip diameter (mm)</td>
<td>6.92 ± 0.37</td>
<td>5.71 ± 0.28</td>
<td>7.03 ± 0.36</td>
<td>12.68 ± 1.27</td>
<td>25.09***</td>
</tr>
<tr>
<td>Branch spacing (mm)</td>
<td>7.90 ± 0.20</td>
<td>11.30 ± 0.80</td>
<td>9.60 ± 0.40</td>
<td>6.00 ± 0.30</td>
<td>16.92***</td>
</tr>
</tbody>
</table>

* P < 0.05; *** P < 0.001; † P < 0.075.

on an examination of associated eigenvectors (component loadings) to ascertain their utility in providing biologically meaningful information (Nichols 1977, Gauch 1982).

Transplant experiment

Twelve branches (3–6 cm in length) were collected from each of five genotypes selected at random from the DB20 (DB20 population; genotypes 1–5) and the CP10 (CP10 population; genotypes 6–10) environments. These 10 genotypes were distinct from those sampled to quantify resident morphology in each environment (i.e., previous section). All genotypes sampled within a population were separated by >25 m to increase the probability that they were genetically distinct. Each branch was glued into a short piece of plastic pipe (1 cm in diameter) with marine epoxy (Z-Spar Splash Zone Compound A-788, Kop-Coat, Los Angeles, California; epoxy did not appear to adversely affect the corals, as many grew over the dried epoxy within 4 wk), and a nylon screw was set in the opposite end (Fig. 1B). The 12 branches from each genotype were randomly assigned to 12 different Plexiglas plates (15 × 25 cm) and were randomly assigned positions within each plate. Eight additional corals, which were not used in the analysis for this study, were also attached to the same plates to form small, artificial aggregations. Branches were placed as close as possible (10–20 mm between adjacent branches) to mimic natural spacing (Table 2), but were not placed in direct contact to avoid aggressive interactions (Lang and Chornesky 1990). Three randomly selected plates were transplanted to each of the four treatment environments (DB10, DB20, CP10, and CPSP) for 96 d, beginning in February 1994. Corals were kept in a flowing seawater table during the preparation of plates and were transplanted to their respective environments within 48 h of collection. The plates within each environment were separated by >10 m.

Morphological plasticity was quantified by measuring six skeletal traits at the conclusion of the transplant experiment: intersepta area, ISA; septa length, SL; columella area, CLA; corallite area, CA; corallite spacing, CS; and branch tip diameter, BD (Fig. 2). The morphological traits of the transplanted corals were quantified in the same manner as the resident corals (3 corallites/branch). Two measurements of SL were made from each corallite (the longest and opposing septa). Quantification of ISA, CLA, CA, CS, and BD is described above.

The results of the transplant experiment were analyzed in two separate sets of analyses. The purpose of the first was to examine the effects of the population (origin), genotype, and environment on coral morphology, and to compare plastic responses between populations and among genotypes. Because the six measured skeletal traits were correlated, PCA was performed on log-transformed measurements of the six traits from both populations (statistical replicates were mean branch values for each trait) and the resulting uncorrelated variables were used in ANOVA. PCA results also aided in an examination of multivariate plastic responses. Log-transformed scores from the first three PCs (see above for component selection criteria) were analyzed with three-way ANOVA (Scheffé mixed model sensu Fry 1992, fixed factor = environment, random factor = population, nested factor = genotype).

The second set of analyses was carried out to compare the morphologies of the corals transplanted into each environment with the resident corals naturally living in those environments. An additional PCA was performed on the genotype means (log transformed) of both the transplanted and the resident corals using the five traits that had been quantified on both groups: CA, CLA, ISA, CS, and BD. The log-transformed scores from the first three principal components were analyzed with two-way ANOVA (fixed-factor 1 = environment, fixed-factor 2 = population). In this analysis the population factor consisted of three levels, including the resident corals and the transplanted corals from each of the two source populations, and was considered a fixed factor because the resident level was specifically chosen to determine whether the transplants adopted the morphology of their resident conspecifics.

Reaction norms are a graphic portrayal of the mean response of individual genotypes to the environment, and allow a visual comparison of magnitudes and patterns of plasticity to be made among genotypes and
populations (Stearns 1989). The environment can consist of two or more levels of a single manipulated factor as would be controlled in a laboratory experiment (continuous environments), or two or more natural environments known to differ in a number of factors (discrete environments), as is usually the case in a field transplant experiment (Via et al. 1995). In the present study, reaction norms were plotted for both the original measured traits and for the first three principal components. For the measured traits CLA, SL, and ISA, reaction norms were plotted as proportions of corallite size, as this is likely to be more biologically important than their absolute size (Patterson 1992). In all reaction norms, the four environmental treatments are plotted in an arbitrary order and do not represent a single environmental gradient (Table 1).

**Transplant growth and survival**

The growth (linear extension and skeletal accretion) and survival of the transplanted corals were measured as estimates of fitness. In colonial corals, growth is closely linked to fecundity because the reproductive capability of modular organisms increases with living surface area (Connell 1973, Hughes 1989). Furthermore, larger colony size decreases the risk of whole-colony mortality (Hughes and Jackson 1985, Hughes and Connell 1987). Therefore, growth rate is likely to be an important component determining overall colony fitness (Jackson 1979, Hughes and Jackson 1985).

Linear extension was determined by measuring the length of each branch, before and after transplantation. Skeletal accretion was quantified as an increase in dry skeletal mass using the buoyant weighing technique (Davies 1989). At the end of the transplant period, both measures of skeletal growth were normalized to surface area of living tissue as determined with the methylene blue dye method (described in Hoegh-Guldberg 1988).

Colony survival was measured by scoring each coral as alive or dead at the completion of the transplant period.

**DNA fingerprinting**

Because *Madracis mirabilis* can reproduce asexually through fragmentation (Bak and Criens 1981, Highsmith 1982), DNA fingerprints of the genotypes used in the transplant experiment were prepared to ensure that all 10 genotypes were genetically distinct (Jeffreys et al. 1985a). Animal DNA was extracted by crushing a branch of each genotype in 2 mL of ice-cold guanidine hydrochloride buffer (8 mol/L GHCL, 0.1 mol/L sodium acetate, 5 mmol/L dithiothreitol, 0.5% N-lauryl sarcosine) and storing the resulting slurry at 4°C. The samples were cleaned with SS phenol/chloroform/isoamyl alcohol (25:24:1), followed by a chloroform/isoamyl alcohol (24:1) cleaning. DNA was then precipitated by adding 7.5 mol/L ammonium acetate and ice-cold isopropanol and chilling to −20°C overnight. DNA samples were dissolved in Tris/EDTA buffer to 1 µg/µL, treated with RNase, digested with *Hae* III, and separated on two agarose gels for 16 h at 30 V. The DNA was then transferred to nylon membranes by southern blotting.

Membranes were probed with the oligonucleotide GTG$_6$, which was 3'-end-labeled with DIG-ddUTP (Genius System, Boehringer, Mannheim, Germany). Membranes were hybridized for 6 h at 38°C and detected with chemiluminescence. The migration distance of resolvable bands between 1.6 and 10.0 kb (kilobase) was measured from autoradiograms. For quantitative analysis, bands were assigned to bins that were calculated from the error in estimating band size (after Gibbs et al. 1991) and the resulting banding patterns were used for comparisons among genotypes. Previous experiments have demonstrated that these methods can produce banding patterns from coral host DNA that are repeatable for a single colony and are not confounded by the DNA of the symbiont zooxanthellae (P. J. Edmunds, unpublished data).

**RESULTS**

**Natural morphological variation**

MANOVA revealed that the multivariate morphology of *Madracis mirabilis* resident corals varied significantly among the four treatment environments (based on Wilks’ λ, which was 0.012; $F = 6.70; df = 18, 31; P < 0.001$). Additionally, ANOVAs found that four of the six univariate traits varied significantly (ISA, CS, BD, and BSP), one varied marginally (CLA, $P < 0.075$), and one was not significant (CA, Table 2). For PCA on resident coral genotype means, the first principal component (PC1) explained 53% of the multivariate variance and was characterized by high positive loadings (eigenvectors) for CS and CLA, and high negative loadings BD and ISA (Table 3). PC2 explained 24% of the variance and was characterized mainly by positively weighted CA, while PC3 explained 24% of the variance and was defined by BSP.

**Transplant experiment**

After transplantation to CPSP the proportion of the corallite occupied by the columella decreased, while the SL and ISA (proportional to corallite size) increased (Fig. 3). This resulted in the CPSP corallites adopting a much more porous or open corallite morphology similar to that of the resident corals (Fig. 1D). CS decreased in the lagoon environments (CP10 and CPSP), while BD increased in CPSP (Fig. 3). In the PCA of transplanted corals, PC1 explained 50% of the variance and was defined by high positive loadings for ISA and SL, and negative loadings for CS and CLA (Table 3). PC2 explained 22% of the variance and was characterized mainly by positively weighted CA and CLA, and PC3 was defined mainly by BD, which was weighted negatively (refer to Table 3 for eigenvalues and eigenvectors). ANOVA of the first three principal com-
Table 3. Principal components analysis of: five skeletal traits measured on Madracis mirabilis collected from the four treatment environments (residents), six traits measured after transplantation to the treatment environments (transplants), and five traits of both transplants and the resident corals in each environment (residents and transplants).

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Residents</th>
<th>Transplants</th>
<th>Residents and transplants</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>PC1</td>
<td>PC2</td>
<td>PC3</td>
</tr>
<tr>
<td>Eigenvalue</td>
<td>3.18</td>
<td>1.46</td>
<td>0.87</td>
</tr>
<tr>
<td>Explained variance (%)</td>
<td>52.99</td>
<td>24.37</td>
<td>14.51</td>
</tr>
</tbody>
</table>

Component loadings

- **CA**
  - -0.04
  - 0.98
  - 0.04
  - 0.65
  - 0.70
  - 0.09
  - 0.49
  - 0.82
  - -0.22
- **CS**
  - 0.92
  - -0.05
  - -0.27
  - -0.73
  - 0.41
  - -0.03
  - -0.76
  - 0.20
  - 0.24
- **BD**
  - -0.85
  - 0.13
  - -0.32
  - 0.44
  - -0.21
  - -0.86
  - 0.69
  - -0.17
  - 0.70
- **CLA**
  - 0.71
  - 0.59
  - -0.37
  - -0.47
  - 0.74
  - -0.35
  - -0.55
  - 0.66
  - 0.38
- **ISA**
  - -0.84
  - 0.36
  - 0.25
  - 0.90
  - 0.16
  - 0.05
  - 0.88
  - 0.26
  - 0.02
- **SL**
  - ...
  - ...
  - ...
  - 0.91
  - 0.15
  - 0.11
  - ...
  - ...
  - ...
- **BSP**
  - 0.64
  - 0.15
  - 0.70
  - ...
  - ...
  - ...
  - ...
  - ...
  - ...

Notes: All PCA was performed on log-transformed data. Refer to Fig. 2 for trait abbreviations. Ellipses denote traits not included in the analysis.

Components found a significant environmental effect for PC1 and PC3 but not for PC2 (Table 4), which was strongly weighted by CA. No significant effect was detected for the source population or the environment \( \times \) population interaction, which is illustrated by the similarity of reaction norms of the DB10 and CP10 populations (Figs. 3 and 4). The main genotype effect, however, was highly significant for all three components, and there were significant genotype \( \times \) environment interactions (G \( \times \) E) for the first two components, and marginally significant G \( \times \) E for PC3 (Table 4). These effects are demonstrated, respectively, by differences in elevation of reaction norms and by crossing or discordant reaction norms (e.g., PC2). In the statistical design used in this analysis, comparisons are made among all 10 genotypes. However, further analysis (a separate ANOVA for each population) found the same patterns of significant genotype and G \( \times \) E effects among the five genotypes within each population (J. Bruno, unpublished data).

The PCA that included genotype trait means of both transplanted and resident corals produced three PCs that had component loadings that were very similar to those from the other two PCs (Table 3). The mean scores of transplants and residents were broadly similar for PC1 and PC2, but not for PC3 in the CPSP environment (Fig. 5). In the two-way ANOVAs used to analyze these PCs, a significant environment \( \times \) population interaction indicated that the morphology of the three populations differed significantly within environments. This interaction term was not significant for PC1 and PC2, but was highly significant for PC3, which was heavily weighted by BD (Table 5). The main environment factor had a highly significant effect on PC1 but not on PC2 (the significance of the two main effects was not tested in PC3 due to the significant interaction term; Zar 1996).

Transplant growth and survival

Most of the transplanted corals doubled their dry skeletal mass and length during the 96-d transplant period. Neither measure of growth varied significantly among environments, populations, or genotypes (Fig. 6, Table 6). However, there was a nonsignificant trend for corals from the DB10 population to extend faster than those from the CP10 population. There was no among-genotype variation for linear extension and skeletal accretion, suggesting that no one genotype grew faster in all environments than any other genotype. There was also no significant G \( \times \) E for linear extension and skeletal accretion (Table 6), indicating that rankings of genotype by growth rates did not vary across environments. Although the corals in the CP10 environment grew as fast as the corals transplanted to the other three environments, they experienced more frequent partial overgrowth by sponges, bryozoans, and bivalves. The corals transplanted to the forereef that experienced partial overgrowth were overgrown by brown algae, especially Dictyota spp. Only 4 branches (out of 120 branches) experienced whole-colony mortality during the transplant period (2 branches from each population), all of which had been transplanted to CP10. Survivorship frequencies were not independent of the environmental treatments (chi-square analysis: \( \chi^2 = 12.42; df = 3; P < 0.01 \)).

DNA fingerprinting

Scorable banding patterns were successfully produced from 8 of the 10 transplanted genotypes (four patterns from each source population). In the fingerprints from the other two genotypes, the bands were not well defined and could not be confidently scored. The eight scorable bands consisted of 4–15 bands ranging in size from 1.6 to 10.0 kb. Assigning all scorable bands to 12 bins produced 7.1 \( \pm \) 0.9 bands/colony (mean \( \pm \) 1 se, Table 7). Banding patterns of a single colony were reproducible when duplicate samples were run on separate gels (J. Bruno, unpublished data). None of the eight genotypes that were successfully fingerprinted had matching banding patterns (Table 7). This result, in addition to the fact that each of the 5 geno-
Fig. 3. Reaction norms of five traits in response to the environmental treatments. (A) Dairy Bull (DB10) genotypes; (B) Columbus Park (CP10) genotypes. Values are untransformed means. Refer to Table 1 for a summary of environmental characteristics of each treatment environment and to Fig. 2 for trait abbreviations. DB10, Dairy Bull at 10 m; DB20, Dairy Bull at 20 m; CP10, Columbus Park at 10 m; CPSP, Columbus Park Springs at 10 m.
TABLE 4. Results of three-factor nested ANOVA testing the effects of the treatment environment, origin population, and genotype on the first three principal components (PC1–3).

<table>
<thead>
<tr>
<th>Source</th>
<th>df</th>
<th>PC1</th>
<th></th>
<th>MS</th>
<th>PC1</th>
<th></th>
<th>F</th>
<th>PC2</th>
<th></th>
<th>MS</th>
<th>PC2</th>
<th></th>
<th>F</th>
<th>PC3</th>
<th></th>
<th>MS</th>
<th>PC3</th>
<th></th>
<th>F</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Environment (E)§</td>
<td>3</td>
<td>17.251</td>
<td></td>
<td>13.996*</td>
<td>3.977</td>
<td></td>
<td>2.254</td>
<td>6.488</td>
<td></td>
<td>13.491*</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Population (P)§</td>
<td>1</td>
<td>5.494</td>
<td></td>
<td>2.722</td>
<td>0.208</td>
<td></td>
<td>0.065</td>
<td>2.554</td>
<td></td>
<td>0.82</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Genotype (G)</td>
<td>8</td>
<td>2.018</td>
<td></td>
<td>6.516***</td>
<td>3.226</td>
<td></td>
<td>5.339***</td>
<td>3.115</td>
<td></td>
<td>5.235***</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>E × P§</td>
<td>3</td>
<td>1.233</td>
<td></td>
<td>1.727</td>
<td>1.764</td>
<td></td>
<td>1.544</td>
<td>0.481</td>
<td></td>
<td>0.501</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Error</td>
<td>80</td>
<td>0.31</td>
<td></td>
<td>0.31</td>
<td>2.304**</td>
<td></td>
<td>1.892*</td>
<td>0.96</td>
<td></td>
<td>1.613†</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Note: PCA was of six traits of transplanted corals. The genotype factor is nested within the population factor. Refer to Table 3 for component loadings. * P < 0.05; ** P < 0.01; *** P < 0.001; † P < 0.075.

types sampled within a population were separated by >25 m, strongly suggests that all 10 genotypes were genetically unique (Jeffreys et al. 1985b, Lewontin and Hartl 1991).

**DISCUSSION**

The goals of this study were to determine whether the polymorphism demonstrated by *Madracis mirabilis* is due to phenotypic plasticity, and whether there is variation in reaction norms among genotypes and between populations of this species. The results demonstrate numerous morphological changes in response to transplantation to a new environment as well as significant genotype × environment interactions. The phenotypic changes of both populations resulted in the transplants becoming similar to resident conspecifics in each treatment environment. Analysis of the PCA results with ANOVA found a significant environmental...
**Table 5.** Results of ANOVA testing the effects of the treatment environment and “population” (transplanted or resident corals) on the first three principal components (PC1–3) from PCA of five skeletal characters of both resident and transplanted corals.

<table>
<thead>
<tr>
<th>Source</th>
<th>d.f.</th>
<th>PC1 (MS)</th>
<th>PC2 (MS)</th>
<th>PC3 (MS)</th>
<th>F</th>
<th>F</th>
<th>F</th>
</tr>
</thead>
<tbody>
<tr>
<td>Environment (E)</td>
<td>3</td>
<td>13.836</td>
<td>0.798</td>
<td>0.798</td>
<td>0.754</td>
<td>2.740</td>
<td>...</td>
</tr>
<tr>
<td>Population (P)</td>
<td>2</td>
<td>0.825</td>
<td>0.477</td>
<td>0.486</td>
<td>0.450</td>
<td>4.139</td>
<td>...</td>
</tr>
<tr>
<td>E × P</td>
<td>6</td>
<td>0.291</td>
<td>0.496</td>
<td>0.806</td>
<td>0.761</td>
<td>2.954</td>
<td>5.722*</td>
</tr>
<tr>
<td>Error</td>
<td>48</td>
<td>0.294</td>
<td>0.990</td>
<td>1.059</td>
<td>0.516</td>
<td>0.516</td>
<td>0.516</td>
</tr>
</tbody>
</table>

*Note:* The purpose of this analysis was to compare the morphologies of the corals transplanted into each environment with the resident corals naturally living in those environments. Refer to Table 3 for component loadings. Significance tests of the two main effects were not performed for PC3 due to the significance of the interaction term.

* P < 0.05.
this study did not attempt to determine which environmental factors caused the observed plastic changes, previous studies suggest that water velocity and irradiance are two particularly important factors in determining coral morphology (Dustan 1975, Graus and Macintyre 1982, Sebens and Done 1992). Both of these factors vary among the four treatment environments used in the present study.

Coral genotype also significantly affected all six measured traits, demonstrating that both the genotype and the environment act synergistically to determine morphology in Madracis mirabilis. This finding may explain field observations of substantial morphological variation between neighboring genotypes of M. mirabilis, which appeared to have been experiencing identical environmental conditions (J. F. Bruno, personal observations). As there was no population-level genetic effect and no population × environment interaction, the origin of each population had little effect on coral morphology. This apparent lack of genetic variation between the experimental populations is further supported by the similarity of their reaction norms (Fig. 3) and is most likely due to gene flow between them.

Significant genotype × environment interactions for PC1 (combined corallite traits) and PC2 (corallite area), and a marginally significant G × E for PC3 (branch diameter), indicate among-genotype variation in the directions and/or magnitudes of plastic responses. Such variation is most likely due to genetic differences among the genotypes. However, the significance of the genotype effect and the G × E effect may also have been influenced by residual effects of the origin environment. This G × E interaction is illustrated by the reaction norm of genotype 8, which was qualitatively different than the other four CP genotypes for both interspeta area and branch diameter (Figs. 3 and 4). Intraspecific variation in plasticity among genotypes has also been reported in the corals Montastraea annularis and Siderastrea siderea (Foster 1979) and in numerous terrestrial plants and other organisms (Miller and Fowler 1993, Schmitt 1993).
Madracis mirabilis branch spacing is inversely related to branch distribution of M. mirabilis (Bruno 1995). For example, in M. mirabilis branch spacing is inversely related to branch diameter (Tables 2 and 3) and to water velocity (Table 1; Bruno 1995, Sebens et al. 1997), as it is in other coral species (e.g., Lesser et al. 1994). Previous laboratory flume studies have demonstrated that branch spacing and flow can interact to affect both the feeding efficiency (Sebens et al. 1997) and respiration rate of M. mirabilis (Bruno 1995). In low-flow conditions, increased branch spacing results in higher respiration and particle capture rates (Bruno 1995, Sebens et al. 1997), while in high-flow conditions decreased branch spacing increases feeding efficiency (Sebens et al. 1997). Therefore, the phenotype appears to “match” local environmental conditions in a way that laboratory studies and biomechanical theory (e.g., Patterson 1992) predict would be beneficial. The results of the fitness estimates of the transplanted corals are concordant with these predictions: corals from both populations were able to maintain similar growth rates and experienced a relatively high survivorship in four very different environments (Table 6, Fig. 6). In other words, they displayed flat fitness reaction norms characteristic of generalist genotypes (Schlichting and Pigliucci 1995). Thus, phenotypic plasticity could facilitate the widespread distribution of M. mirabilis (Schindler 1985) through the convergence of skeletal morphology to that of the local conspecifics. However, a more direct experimental approach would be necessary to determine unequivocally whether plasticity in M. mirabilis is beneficial.

A better understanding of phenotypic plasticity might help resolve a number of long-standing problems in coral biology. One example is coral taxonomy, which has long been plagued by confusion arising from polymorphisms of the characteristics used to distinguish species (Ayre et al. 1991). Experimental taxonomy, where transplant experiments are used to determine the effect of the environment on coral morphology, could help to unambiguously distinguish closely related species. The large amount of morphological variation seen in Madracis mirabilis exemplifies this problem (Schindler 1985, Fenner 1993). The discrete separation of each morphotype could be mistaken as an indication that M. mirabilis is composed of a complex of sibling species (sensu Knowlton 1993), without the evidence of plasticity from this study which suggests otherwise. The use of reaction norms to examine how traits change in response to environmental factors provides a valuable tool to better understand the functional morphology of polymorphic skeletal characteristics in corals and other clonal organisms. Investigations of plasticity may also aid in our understanding of how coral populations might respond to anthropogenic disturbances that are currently degrading many reefs (Grigg and Dollar 1990, Grigg 1993).

The results of this study demonstrate that a relatively long-lived, sessile organism employing both sexual and asexual reproduction can display substantial phenotypic plasticity and that reaction norms can vary significantly among clones. Such plasticity is likely to be an important component of the life history strategies of clonal populations, alleviating the negative effects of low genetic variation (Bradshaw 1965, Schlichting 1986). Morphological plasticity might also enable clonal species (and individual clones themselves) to occupy a broad range of habitats and allow those that are sessile to adapt to temporal changes in their environment. Therefore, investigations of phenotypic plasticity in predominantly asexual clonal taxa that have not previously received such attention could prove invaluable in providing insight into their evolution, speciation, population dynamics, and survival strategies in heterogeneous environments.

Acknowledgments

We especially wish to thank Mark Bertness, Robert Carpenter, Nancy Emery, Ruth Gates, Sean Grace, Mary-Bestor Grant, Rick Grosberg, Sally Hacker, Shane Heschel, Amy Melendy, Johanna Schmitt, Dione Swanson, Jon Witman, and two anonymous reviewers for their comments, which greatly improved earlier drafts of this paper. Numerous dive buddies assisted us in the field in a variety of adverse conditions, including Marco Calavetta, Amy Melendy, Matt Mills, Sean Grace, Mark Warner, and many students of the East/West Marine Biology Program. We wish to thank the staff of the Discovery Bay Marine Laboratory (DBML) for their assistance. J. F. Bruno also thanks Sandy and Frank Bruno for their support. This research represents partial fulfillment for the Master of Science degree to J. F. Bruno at California State University, Northridge (CSUN) and was financially supported by a grant to P. J. Edmunds from the Office of Naval Research (number N00014-93-1-0440) and grants to J. F. Bruno from Sigma Xi, the CSUN office of Graduate Studies and International Programs, and the CSUN student projects committee (number 323430403). This is DBML contribution number 583.

Literature Cited


Bak, R. P. M., and M. S. Engel. 1979. Distribution abundance
and survival of juvenile hermatypic corals (Scleractinia) and the importance of life history strategies in the parent coral community. Marine Biology 54:341–352.


