Two developmentally temporal quantitative trait loci underlie convergent evolution of increased branchial bone length in sticklebacks

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In convergent evolution, similar phenotypes evolve repeatedly in independent populations, often reflecting adaptation to similar environments. Understanding whether convergent evolution proceeds via similar or different genetic and developmental mechanisms offers insight towards the repeatability and predictability of evolution. Oceanic populations of three-spine stickleback fish, *Gasterosteus aculeatus*, have repeatedly colonized countless freshwater lakes and streams, where new diets lead to morphological adaptations related to feeding. Here, we show that heritable increases in branchial bone length have convergently evolved in two independently derived freshwater stickleback populations. In both populations, an increased bone growth rate in juveniles underlies the convergent adult phenotype, and one population also has a longer cartilage template. Using F2 crosses from these two freshwater populations, we show that two quantitative trait loci (QTL) control branchial bone length at distinct points in development. In both populations, a QTL on chromosome 21 controls bone length throughout juvenile development, and a QTL on chromosome 4 controls bone length only in adults. In addition to these similar developmental profiles, these QTL show similar chromosomal locations in both populations. Our results suggest that sticklebacks have convergently evolved longer branchial bones using similar genetic and developmental programmes in two independently derived populations.

1. Introduction

Independent populations that converge on similar evolved phenotypes may do so by using similar genetic and developmental mechanisms, suggesting that evolution is, at times, constrained and predictable [1,2]. When convergent phenotypic evolution is caused by parallel genetic mechanisms, the parallelism may occur on a number of different hierarchical levels, including changes in the same nucleotide, gene, genetic pathway or genomic region (reviewed in [2–4]; for examples, see [5–8]). While numerous cases of convergent evolution have been documented across natural and experimental populations of animals, plants and microbes, fewer studies have investigated whether these convergently evolved phenotypes arise in the same way during development [9]. Furthermore, most studies of convergent evolution have focused on traits with a simple genetic architecture, and less is known about whether more complex traits, which are more common in nature, convergently evolve via parallel developmental genetic features.

In vertebrates, the skeleton contributes to organismal form and function, and evolved changes in skeletal elements occur repeatedly as populations adapt to new environments. The skeleton forms largely from two types of bone: endochondral, which develops from a cartilage template, and dermal, which ossifies directly without a cartilage intermediate [10]. Atchley & Hall [11] proposed that skeletal evolution may proceed through a number of cellular
mechanisms (e.g. the size of the cartilage template or the rate of bone growth). In support of this proposal, a dramatic difference in jaw size between quails and ducks results from several differences in the specification and morphogenesis of the midbrain and midbrain neural crest cells from which the jaw is derived [12]. In Anolis lizards, however, a smaller number of cellular mechanisms appear to underlie convergent skeletal evolution. On at least four islands of the Caribbean, Anolis limb morphologies have repeatedly evolved in different ecological habitats [13]. While multiple pathways in pre- and post-embryonic development could contribute to differing limb length, increased adult limb size in four different long-limbed species arose from an increase in the size of the embryonic limb template, followed by growth rates equal to those in shorter-limbed species [14]. Whether this mechanism of evolved bone length differences is seen in other convergently evolved skeletal changes is largely unknown.

The threespine stickleback (Gasterosteus aculeatus) provides an excellent model system for studying both the developmental and genetic basis of convergent skeletal evolution. Ancestral marine sticklebacks have colonized thousands of freshwater environments throughout the Northern Hemisphere and have evolved numerous adaptations to these new freshwater environments [15]. For example, freshwater sticklebacks have repeatedly evolved changes to their head skeletons to improve feeding efficiency on new foods in freshwater environments, including convergent decreases in gill raker number [16–18], as well as increases in jaw width [17,19] and suction feeding index [20].

Here, we hypothesized that other trophic skeletal elements may also differ between marine and freshwater sticklebacks. The branchial skeleton (figure 1, adapted from [21]) is primarily made up of bilateral, segmentally reiterated bones: five ventral pairs (ceratobranchials, CB1–CB5) and four dorsal pairs (epibranchials, EB1–EB4). In fish, these bones arise from neural crest cells in the pharyngeal arches during development, and the dorsal and ventral bones are segmental homologues of the upper and lower jaw, respectively [22,23]. These long dorsal and ventral bones of the branchial skeleton are endochondral and resemble mammalian long bones (e.g. the femur) in appearance [24]. In fish, the branchial cartilages start to form late in embryogenesis, just before hatching [23,25]. As development continues, this cartilage is mostly replaced with bone deposited by osteoblasts that originate both outside and within the cartilage template [26]. The bones then elongate as the fish grows larger [24]. Thus, two key developmental processes contribute to the length of the bone: the establishment of the cartilage template early in embryonic development and the rate of subsequent bone growth.

A previous genome-wide linkage mapping study of the genetic basis of skeletal variation in sticklebacks identified 14 quantitative trait loci (QTL) with significant effects on the length of branchial bones in a marine-by-freshwater F2 cross, including two QTL on chromosomes 4 and 21 with large effects [27]. Combined, these two QTL explain approximately 27% of the variance in length of the dorsal EB1 and approximately 25% of variance in length of the ventral bones. Most (11) of these QTL, including both of the large-effect QTL, had effects in the same direction, with freshwater alleles conferring longer bones [27]. However, this study did not measure the bone length phenotypes of the parental populations. Orr [28] proposed that a concerted sign of QTL effect indicates a trait is under natural selection, as similar directions of effect would be unlikely to be observed by chance. Here, we test the hypothesis that the two previously identified large-effect QTL on chromosomes 4 and 21 are used in a second independently derived freshwater population. By studying the developmental trajectories of evolved increases in bone length, as well as the developmental timing of two bone length QTL, we also test whether similar developmental and genetic effects contribute to these evolved increases in bone length.

2. Material and methods

(a) Wild collections

Wild anadromous marine fish were collected from the Little Campbell River (LITC) in British Columbia under a fish
collection from the Washington Department of Fish and Wildlife (permit #08-284). Wild sticklebacks were collected in the summer of 2008. All wild and laboratory-reared fish were euthanized with 0.08% Tricaine and stored in 100% ethanol until staining and dissection.

(b) Fish husbandry and crosses
For the FTC × LITC cross, a wild male FTC fish was crossed to a wild female LITC. For the Paxton Benthic (PAXB, British Columbia, Canada) × LITC cross, a laboratory-reared male offspring of wild PAXB fish was crossed to two wild LITC females. Adult F1 fish were then intercrossed to their siblings to create F2 families, which were grown to ages of 20, 40 and 80 days post-fertilization (dpf), or adults (see the electronic supplementary material, table S1). All fish were raised in 3 ppt salinity (approx. 10% seawater) at 18°C in 1101 (29 gallon) tanks. Fish were fed a diet of live Artemia as young fry, live Artemia and frozen Daphnia as juveniles, and frozen bloodworms and Mysis shrimp as adults.

(c) Phenotyping, genotyping and quantitative trait loci analysis
Detailed descriptions of phenotyping, genotyping and QTL analysis can be found in the electronic supplementary material.

(d) Statistical analysis
All statistical analyses were performed using the R statistical software package (www.r-project.org). QTL analysis was performed using R/qtl (www.rqtl.org).

3. Results

(a) Population differences in bone length
To test the hypothesis that wild marine and freshwater fish differ in branchial bone length, we analysed wild-caught marine (LITC) and freshwater (FTC) sticklebacks for differences in length of the dorsal (epibranchial, EB1) and ventral (ceratobranchial, CB1–CB5) branchial long bones (figure 1). All six branchial long bones differed significantly in length (figure 2a), with freshwater fish having longer bones relative to standard length than marine fish (ventral bones were 8.8–17.1% longer; dorsal bone was 23.8% longer in 60 mm to standard length than marine fish (ventral bones were

(b) Developmental basis of bone length differences
We hypothesized that stickleback bone length differences, like evolved Anolis limb length [14], would manifest during development as transposition of the y-intercept, but not slope, of a
We found that both cartilage templates were the CB4 cartilage and stage 28 (approx. 13–14 dpf) to measure and LITC fry to stage 26 (approx. 10 dpf) [31] to measure strong genetic effects in a previous cross [27]. We raised FTC because it had a large marine–freshwater difference and had tive to marine fish. For ventral cartilages, we focused on CB4 prefigure branchial bones may be larger in freshwater fish rela-
table S5) led us to hypothesize that the cartilage templates that development time courses (electronic supplementary material, [31]). Thus, although the convergent evolution of increased bone growth rates in FTC and PAXB, we observed significant differences in the slopes (population \times standard length interaction term) of dorsal and posterior ventral bone lengths relative to standard length between marine and freshwater. ANCOVA statistics are shown in the electronic supplementary material, figure S2. Red, LITC; light blue, FTC; dark blue, PAXB. (c) EB1 and (d) CB4 cartilages are longer in FTC relative to LITC and PAXB fry (Tukey HSD \( p < 0.05 \) for FTC-PAXB comparisons of both cartilages). In (c), the FTC fish were slightly shorter in total length than the LITC and PAXB fish (Tukey HSD test \( p < 0.05 \)), which makes the cartilage size increase even greater relative to body size. Error bars = standard deviation. Asterisks indicate Tukey HSD \( p \)-values: n.s., not significant; \( * p < 0.05 \), \( ** p < 0.01 \), \( *** p < 0.001 \).

Figure 3. Developmental basis of dorsal and ventral bone length differences. (a,b) Developmental time course of (a) dorsal (EB1) and (b) ventral (CB4) bone lengths plotted against total length of laboratory-reared fish under five months of age. Both bones show statistically significant differences in slope (bone growth rate) as well as \( y \)-intercept between marine and freshwater. ANCOVA statistics are shown in the electronic supplementary material, table S5; additional bones are shown in the electronic supplementary material, figure S2. Red, LITC; light blue, FTC; dark blue, PAXB. (c) EB1 and (d) CB4 cartilages are longer in FTC relative to LITC and PAXB fry (Tukey HSD \( p < 0.05 \) for FTC-PAXB comparisons of both cartilages). In (c), the FTC fish were slightly shorter in total length than the LITC and PAXB fish (Tukey HSD test \( p < 0.05 \)), which makes the cartilage size increase even greater relative to body size. Error bars = standard deviation. Asterisks indicate Tukey HSD \( p \)-values: n.s., not significant; \( * p < 0.05 \), \( ** p < 0.01 \), \( *** p < 0.001 \).

regression of bone length against standard length. We col-
collected laboratory-reared fish from each population at regular development time points, resulting in fish varying in total length from 10 to 40 mm. We looked for differences in bone growth rate and initial bone size using an ANCOVA with standard length as the covariate and population as an interacting factor. Contrary to our prediction, we observed significant differences in the slopes (population \times standard length interaction term) of dorsal and posterior ventral bone lengths relative to standard length between marine and freshwater fish, suggesting that freshwater bones grow more rapidly relative to body size (figure 3a,b). Thus, unlike in Anolis lizards, the convergent evolution of increased bone length in two derived freshwater stickleback populations appears to use a similar faster bone growth rate in both populations.

The significant differences in \( y \)-intercepts in the bone development time courses (electronic supplementary material, table S5) led us to hypothesize that the cartilage templates that prefigure branchial bones may be larger in freshwater fish relative to marine fish. For ventral cartilages, we focused on CB4 because it had a large marine–freshwater difference and had strong genetic effects in a previous cross [27]. We raised FTC and LITC fry to stage 26 (approx. 10 dpf) [31] to measure the CB4 cartilage and stage 28 (approx. 13–14 dpf) to measure EB1 cartilage. We found that both cartilage templates were longer in FTC relative to both LITC and PAXB (figure 3c,d). Thus, despite the convergent increased bone growth rates, one unique developmental difference contributes to the convergent evolution, with one freshwater population (FTC) but not a second (PAXB) evolving a longer cartilage template early in development.

(c) Genetic basis of bone length differences

QTL mapping provides a powerful first test of possible parallel genetic mechanisms underlying convergent evolution. We hypothesized that previously identified bone length QTL might be re-used in multiple freshwater stickleback populations due to extensive sharing of the genetic basis of evolved traits in stickleback populations [32,33], and the similar increased bone growth rates in FTC and PAXB. Because there are probably multiple developmental mechanisms that can be altered to change bone length [11], we further predicted that these QTL might exert different effects at specific points in development.

We focused on the two largest-effect QTL controlling adult bone length in a previous cross (chromosomes 4 and 21 [27]) and observed strikingly similar developmental profiles of these QTL in our two crosses. We raised F2 fish to four time points (20, 40 and 80 dpf and adults, see the electronic supplementary material, table S1), and tested for the
Figure 4. Similar developmental effects of chromosome 21 QTL in two independently derived freshwater populations. F₂ fish from two marine × freshwater F₂ crosses ((a,c) FTC × LITC; (b,d) PAXB × LITC) were raised to four time points (given in days post-fertilization, dpf; or adults > 150 dpf) and tested for effects of chromosome 21 genotype on size-corrected bone length residuals. See the electronic supplementary material, table S1 for a summary of the fish included in each time point. (a,b) Chromosome 21 controlled dorsal (EB1) bones at all time points in the FTC cross and at all time points except 20 dpf in the PAXB cross, and the effect was strongest at 80 dpf in both crosses. (c,d) The effect of chromosome 21 on ventral bone length (CB4) was significant at 80 dpf in the FTC cross and nearly significant at 80 dpf in the PAXB cross. ANOVA p-values for the marker Stn489 are indicated with asterisks when significant (p < 0.05; see the electronic supplementary material, table S6 for complete listing of ANOVA results for all branchial bones). Error bars = standard deviation.

4. Discussion
(a) A heritable increase in branchial bone length in two freshwater stickleback populations is likely to be a trophic adaptation

A previous QTL mapping study found that most (11/14) freshwater alleles controlling stickleback branchial bone length produced longer bones [27], suggesting increased branchial bone length is under natural selection in freshwater environments. Supporting this prediction, we show that marine and freshwater bone lengths differ in the wild, and that two populations of freshwater stickleback show strongly heritable increases in branchial bone length. This elongation of branchial bones may facilitate the processing of larger prey items in freshwater by providing a larger buccal cavity for...
food to pass through, generating greater crushing force and/or offering increased muscle attachment area for the crushing of freshwater prey. While many studies have focused on evolutionary loss, these evolved increases in bone length demonstrate that despite the predictable loss of several skeletal elements (including gill rakers, dorsal spines and armour plates) in freshwater environments [18,32,34], other parts of the skeleton (i.e. the branchial bones) increase in size despite the much lower environmental calcium concentration in freshwater. In both freshwater populations studied here, the increased bone length differences are most pronounced in the dorsal (EB1) and posterior ventral bones (CB4 and CB5 demonstrate larger marine–freshwater differences than the more anterior three CBs). These findings suggest that the entire branchial skeleton is not uniformly enlarged relative to standard length in freshwater fish, but rather that independent genetic and developmental mechanisms have led to modular changes in the relative sizes of bones in the branchial skeleton. Heritable and similarly modulated increases in bone length in two independent freshwater populations suggest that this trait may be adaptive in these environments [35].

(b) A convergent increase in bone growth rate underlies bone elongation in freshwater sticklebacks

Here, we find two developmental mechanisms of evolved bone elongation (increased cartilage template size and bone growth rate) are at work in freshwater stickleback populations. Relative to marine fish, both freshwater populations have evolved an increased bone growth rate. All PAXB branchial bones and dorsal and posterior ventral FTC branchial bones have an increased growth rate relative to marine bones. Early cartilage template size is also increased in FTC freshwater fish. Therefore, the convergent evolution of these independent stickleback populations uses one shared developmental feature (increased bone growth rate) as well as at least one unique feature (increased cartilage template size in only one freshwater population). Differences in juvenile bone growth rates have been observed in the limbs of large and small mouse strains [36], and the elongated craniofacial bones of needlefish [37]. Multiple aspects of chondrocyte hypertrophy (the enlargement of chondrocytes that promotes bone growth) are altered to produce elongated digits in bats [38] and elongated limbs in jerboas [39]. Thus, developmental modulation of bone growth rates seems to be a shared mechanism of altering skeletal proportions in multiple taxa, including sticklebacks.

(c) Shared quantitative trait loci on chromosomes 4 and 21 suggest a parallel developmental genetic basis for freshwater bone length increase

Consistent with the convergent increased bone growth rate in two freshwater populations, these populations also share two overlapping QTL with strong effects on bone length at various stages of development. These two QTL, initially identified in the PAXB freshwater population [27], were successfully replicated here by crossing a different PAXB fish to a different marine background, and also were observed in a second freshwater population, Fishtrap Creek. The development profiles of the QTL are remarkably similar between the two crosses. The effect of chromosome 4 is only seen in adult bones in both crosses. This QTL may only act late in development, or its earlier effects may only be apparent when fish reach a larger size. By contrast, chromosome 21 seems to exert its effects earlier than chromosome 4 in both crosses. Thus, similar developmental genetic features underlie the convergent evolution of longer
branchial bones, suggesting that even complex traits can have at least partially predictable genetic bases.

Although QTL in vertebrates are typically mapped in adults, a handful of studies have linked adult phenotypes to changes in juveniles. QTL for stickleback juvenile pigmentation and standard length have been identified [40]. In a finding similar to ours, a QTL controls shank growth rate in beach chickens during a specific time period of juvenile development [41]. Additionally, a QTL controlling adult hair colour in beach mice can be traced to differential expression of the Agouti gene early in development [42]. These studies demonstrate that genetic changes that manifest at specific developmental stages can contribute to differences in final adult phenotype.

Future fine-mapping work will determine whether FTC and PAXB share the alleles on chromosomes 4 and 21 that control bone size. Since freshwater stickleback populations are derived from a large oceanic ancestral population and often share alleles controlling evolved morphological changes [32,33,43,44], we parsimoniously hypothesize that the same alleles are at work in the FTC and PAXB populations. This recycling of ancestral alleles to produce a convergent phenotype has been called ‘collateral’ evolution [3]. This hypothesis is supported by the similar developmental profiles of the QTL, and the similarities in chromosomal location and dominances of the QTL in each cross. However, sticklebacks have also been shown to independently evolve alleles in multiple populations in the case of pelvic reduction [45], so the possibility remains that unique alleles have evolved in each population.

In conclusion, we find evidence of similar genetic and developmental properties underlying evolved increases in bone length in two independently derived freshwater stickleback populations. Both derived freshwater populations share an increased rate of growth of some bones relative to the bones of their marine counterparts, and the two QTL on chromosomes 4 and 21 demonstrate strikingly similar effects throughout development in crosses of each population. Our developmental genetic evidence supports a model that the same chromosome 4 and 21 genomic regions were selected independently in two freshwater populations to produce quantitative changes in a convergently evolved trophic phenotype. Future studies of other freshwater populations and crosses will test whether this evolved gain trait and the use of bone length QTL on chromosomes 4 and 21 are predictable features of freshwater adaptation.

All animal work was approved by the Institutional Animal Care and Use Committees of the University of California-Berkeley or Stanford University (protocol numbers R350 and 13834).

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Data accessibility. All data are available as an electronic supplementary material file.

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