

# Current Biology

## A Single Interacting Species Leads to Widespread Parallel Evolution of the Stickleback Genome

### Highlights

- Biotic selection correlated with rapid, parallel genetic differentiation
- Stickleback from the two biotic environments differed in about 600 genes
- There are about 140 discrete genomic regions potentially under biotic selection
- Genetic differentiation correlated with variation in morphology

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### In Brief

Miller et al. analyze genetic differentiation in wild populations of threespine stickleback fish in response to a single agent of biotic selection, intraguild predation by prickly sculpin. Sculpin presence was correlated with parallel, widespread genetic divergence. Ecological interactions between these species had large evolutionary consequences.



# A Single Interacting Species Leads to Widespread Parallel Evolution of the Stickleback Genome

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## SUMMARY

Biotic interactions are potent, widespread causes of natural selection and divergent phenotypic evolution and can lead to genetic differentiation with gene flow among wild populations (“isolation by ecology”) [1–4]. Biotic selection has been predicted to act on more genes than abiotic selection thereby driving greater adaptation [5]. However, difficulties in isolating the genome-wide effect of single biotic agents of selection have limited our ability to identify and quantify the number and type of genetic regions responding to biotic selection [6–9]. We identified geographically interspersed lakes in which threespine stickleback fish (*Gasterosteus aculeatus*) have repeatedly adapted to the presence or absence of a single member of the ecological community, prickly sculpin (*Cottus asper*), a fish that is both a competitor and a predator of the stickleback [10]. Whole-genome sequencing revealed that sculpin presence or absence accounted for the majority of genetic divergence among stickleback populations, more so than geography. The major axis of genomic variation within and between the two lake types was correlated with multiple traits, indicating parallel natural selection across a gradient of biotic environments. A large proportion of the genome—about 1.8%, encompassing more than 600 genes—differentiated stickleback from the two biotic environments. Divergence occurred in 141 discrete genomic clumps located mainly in regions of low recombination, suggesting that genes brought to lakes by the colonizing ancestral population often evolved together in linked blocks. Strong selection and a wealth of standing genetic variation explain how a single member of the biotic community can have such a rapid and profound evolutionary impact.

## RESULTS AND DISCUSSION

Threespine stickleback (*Gasterosteus aculeatus*) were sampled in British Columbia (Canada) from 17 geographically interspersed lakes of two types: “lakes with sculpin” (n = 9) and

“lakes without sculpin” (n = 8) (Figure 1A). Prickly sculpin fish (*Cottus asper*) are a piscivorous intraguild predator—preying upon stickleback and competing with it for food [10]. Sculpin presence or absence is associated with heritable phenotypic differences in stickleback defensive armor, body shape, and behavior [10, 11]. Lakes formed about 10,000 years ago and were colonized from the sea by marine or anadromous stickleback (hereafter marine) [12]. Study lakes are similar in size and were chosen to minimize regional differences in lake characteristics (Figure 1B; Figure S1; Table S1). The result is an unusual natural system of populations having access to shared ancestral standing genetic variation that allowed for independent and repeated adaptation to environments differing primarily in the presence or absence of sculpin.

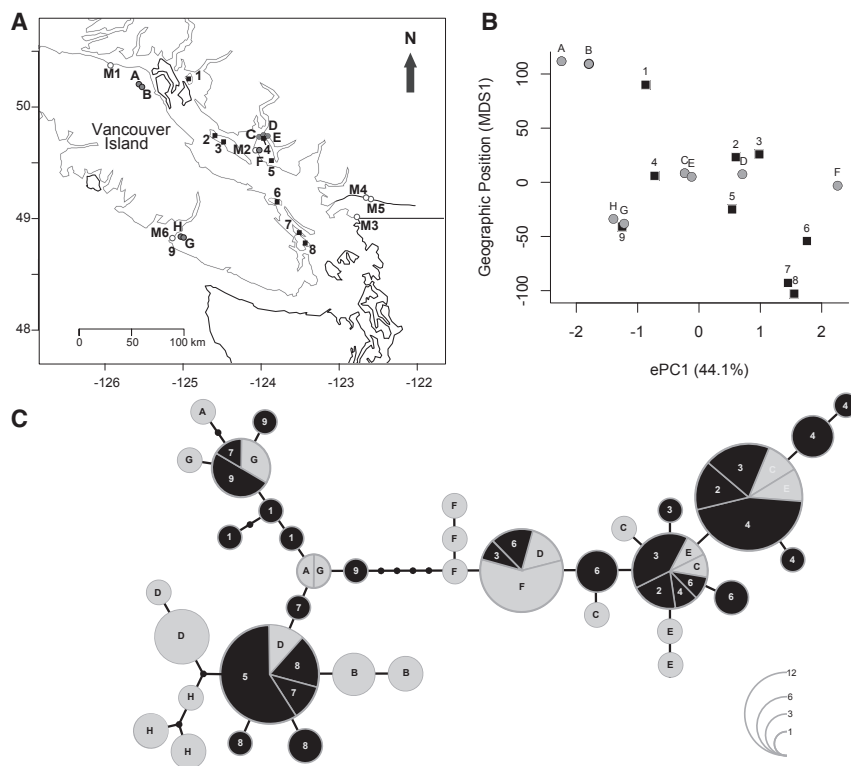
We tested the phylogenetic independence of stickleback from study lakes using a haplotype network based on putatively neutral sequences from the mitochondrial control region. Systematic structuring between population types would suggest that distinct monophyletic clades colonized lakes with and without sculpin, whereas shared haplotypes between lake types would be consistent with colonization of lakes by a common ancestral population. We found that most populations contained multiple haplotypes, many haplotypes were shared among several populations, and stickleback from lakes with and without sculpin did not cluster in the haplotype network (Figure 1C). This pattern agrees with the geological history of the region [12] and phylogenetic studies indicating that freshwater stickleback populations formed at the end of the last ice age by multiple colonization events from the sea and subsequently evolved independently [13].

### Most Genomic Variation Is Associated with Sculpin Presence or Absence

To detect parallel adaptive evolution in response to the presence or absence of sculpin, we sequenced a single representative stickleback from each freshwater population to an average coverage of 9x using Illumina 100 bp paired-end whole-genome re-sequencing. Sequences were aligned to the stickleback reference genome [14, 15]. After filtering, we generated a dataset of 6.3 million SNPs from the 17 freshwater genomes, corresponding to approximately one SNP every 73 bp.

A principal-component analysis (PCA) of all nuclear SNPs, excluding those on the sex chromosome, revealed sculpin presence or absence to be the predominant factor organizing stickleback genome-wide variation. The first genomic principal component (gPC1) explained 11.6% of the genome-wide SNP





**Figure 1. Sculpin Presence or Absence Is the Primary Difference among Lakes**

(A) Location of study populations. Lakes in which stickleback co-occur with sculpin are indicated with gray circles (A–H), and lakes without sculpin are given in black squares (1–9). Sampling locations of marine stickleback are shown with open circles (M1–M6). Lake names are provided in Table S1.

(B) The association between the first principal component of physical and chemical properties of the study lakes (ePC1) and the relative geographical position of lakes (MDS1), calculated as the principal axis from a multidimensional scaling of pairwise geographical distances among lakes. Lakes located in closer proximity to each other, as indicated by similarity in MDS1, have ePC1 values that are more similar than ePC1 values between more distant lakes. Environmental trait values are in Table S1.

(C) Genealogical haplotype network based on a putatively neutral DNA sequence from the mitochondrial control region. Circle size indicates the total number of individuals with that haplotype (see lower right corner). The length of the line segments indicates the number of mutations separating haplotypes. Haplotypes are often shared between stickleback from lakes with sculpin (gray, A–H) and stickleback from lakes without sculpin (black, 1–9). The haplotype network shows no evidence for monophyly of stickleback from lakes with sculpin or of stickleback from lakes without sculpin. See Figure S1 for more information on lake characteristics.

variation and separated stickleback from the two lake types with only slight overlap (Figure 2A;  $F_{1,15} = 22.5$ ,  $p < 2.6e-4$ ; lakes with sculpin =  $412.1 \pm 120.6$ ; lakes without sculpin =  $-366.3 \pm 108.7$ ). The second genomic principal component (gPC2) accounted for 7.8% of SNP variation and was associated with geographical position of lakes (Figure 1B). The two lake types overlapped broadly in gPC2, and their means were not significantly different ( $F_{1,15} = 0.15$ ,  $p = 0.71$ ; lakes with sculpin =  $42.8 \pm 191.5$ ; lakes without sculpin =  $-38.1 \pm 105$ ). We found a weak spatial autocorrelation in gPC1 (Moran's I:  $I = 0.08$ ,  $p = 0.08$ ) but a significant spatial autocorrelation in gPC2 (Moran's I:  $I = -0.141$ ,  $p = 0.02$ ) (Figure S2). We retested the difference between the lake types in gPC1 after controlling for spatial effects using a distance-based Moran's eigenvector map analysis (dbMEM) and again found a significant association ( $F_{4,12} = 9.2$ ,  $p = 0.001$ ). Overall, we conclude that sculpin presence or absence had a large overall effect on genomic variation in stickleback, more so than geography in this sample of populations.

We repeated these analyses including stickleback from six marine locations. The first genomic principal component explained 13.8% of the genomic variation and again differentiated stickleback from lakes with and without sculpin (Figure 2B;  $F_{6,16} = 18.9$ ,  $p < 2e-6$ ). Stickleback sympatric with sculpin were more similar to the marine populations along the gPC1 than stickleback from sculpin-absent lakes (Figure 2B), indicating that populations from lakes with sculpin have retained mainly marine alleles at the SNPs having high loadings. Since the marine population represents the putative ancestral form, the alleles at SNPs evolving in parallel are most often in the

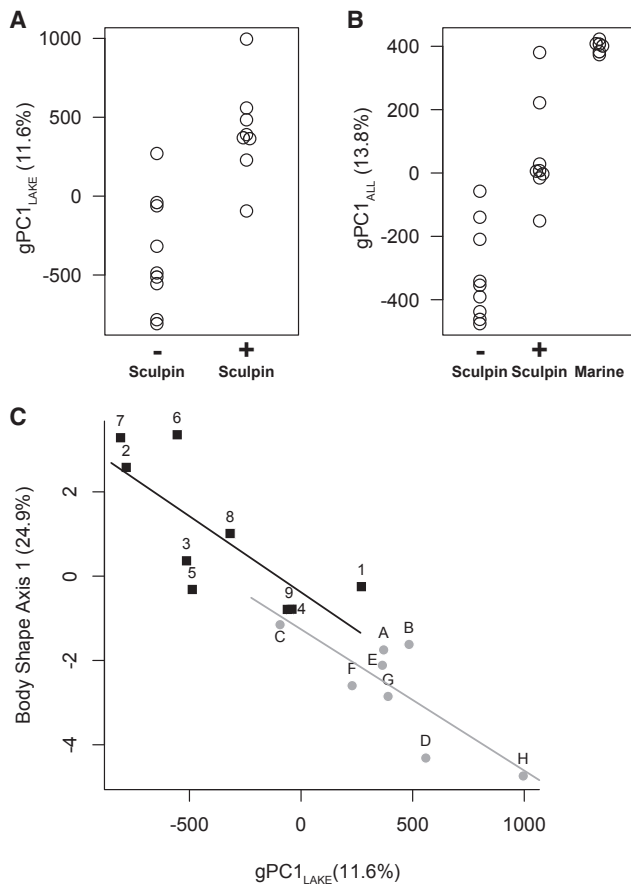
derived state in sculpin-absent lakes compared with alleles in sculpin-present lakes. This agrees with previous phenotypic trait comparisons of the three groups [16].

### Genomic and Phenotypic Differentiation Are Associated

Position along the major axis of genomic variation among lakes (gPC1) was correlated with phenotypic differentiation, both within and between lake types in body shape (Figure 2C; lakes with sculpin  $r = 0.80$ ,  $p = 0.02$ ; lakes without sculpin  $r = 0.77$ ,  $p = 0.02$ ; overall  $r = 0.91$ ,  $p = 4e-7$ ), and bony armor (Figure S2). This pattern might reflect widespread pleiotropy, with suites of phenotypic traits mapping to the same parallel diverging genetic variants, but this seems unlikely given the diversity of traits and large number of genomic regions involved (see below). A more likely cause is correlated selection on multiple traits and underlying genes along a gradient of environmental differences, chiefly sculpin presence or absence.

### Genomic Differentiation Is Extensive

Widespread genomic regions differed between stickleback from lakes with and without sculpin (Figure 3A). To quantify absolute divergence between the two groups, we calculated the raw variance between the two groups at each nucleotide ( $F_{ST\_NUM}$ ) and then took the average in 10,000 bp sliding windows across the genome. This divergence metric is the numerator of the  $F_{ST}$  statistic and has a maximum of 1 at bi-allelic SNPs and a value of 0 at invariant sites. Although lake types are not strictly geographically structured subpopulations, we also calculated  $F_{ST}$  to quantify relative differentiation between stickleback



**Figure 2. Genomic Variation Is Associated with Sculpin Presence or Absence**

Separation of stickleback from lakes with and without sculpin along the first principal component of whole-genome SNP variation.

(A) First principal-component values based on all 17 lake populations ( $gPC1_{LAKE}$ ). (B) First principal-component values based on an analysis that also includes marine (ancestral) stickleback sampled at six localities ( $gPC1_{ALL}$ ). Each point represents a single individual from a population. Stickleback from lakes with sculpin resemble marine stickleback more closely than do stickleback from sculpin-absent lakes. The percentage of SNP variation explained by each first principal component is shown in parentheses.

(C) Comparison of the first principal component of all SNPs in all lake populations ( $gPC1_{LAKE}$ ) with the first linear discriminant axis of body shape (body shape axis 1). Greater phenotypic differentiation both between lake types and within lake types is associated with more extreme genotypic differentiation along  $gPC1_{LAKE}$ . See also Figures S2 and S3.

from the two lake types.  $F_{ST}$ , allele frequency difference (AFD), and a modified version of the cluster separation score of [14] produced similar findings (Figure S3; Table S2).

Using permutation tests, and controlling false discovery rate, we identified significant differences in 1,395 of 76,109 informative genomic windows (hereafter outliers), representing  $\sim 1.8\%$  of the genome. Outlier windows frequently contained alleles not yet fixed between populations from the different lake types as only 689 SNPs (out of 6.3 million) had  $F_{ST_{NUM}} > 0.8$  between stickleback from the two lake types. Outlier windows overlapped introns or exons of 609 (3.3%) of the 18,254 annotated genes

that fell within sampled windows. Each outlier window contained an average of 0.88 genes (SD = 0.79), with individual genes often spanning multiple windows. Genes in outlier windows were enriched for Gene Ontology (GO) terms related to neuron development, synaptic transmission, heart morphogenesis, and many other diverse biological functions (Table S3), suggesting that interactions with sculpin had a comprehensive effect on stickleback evolution.

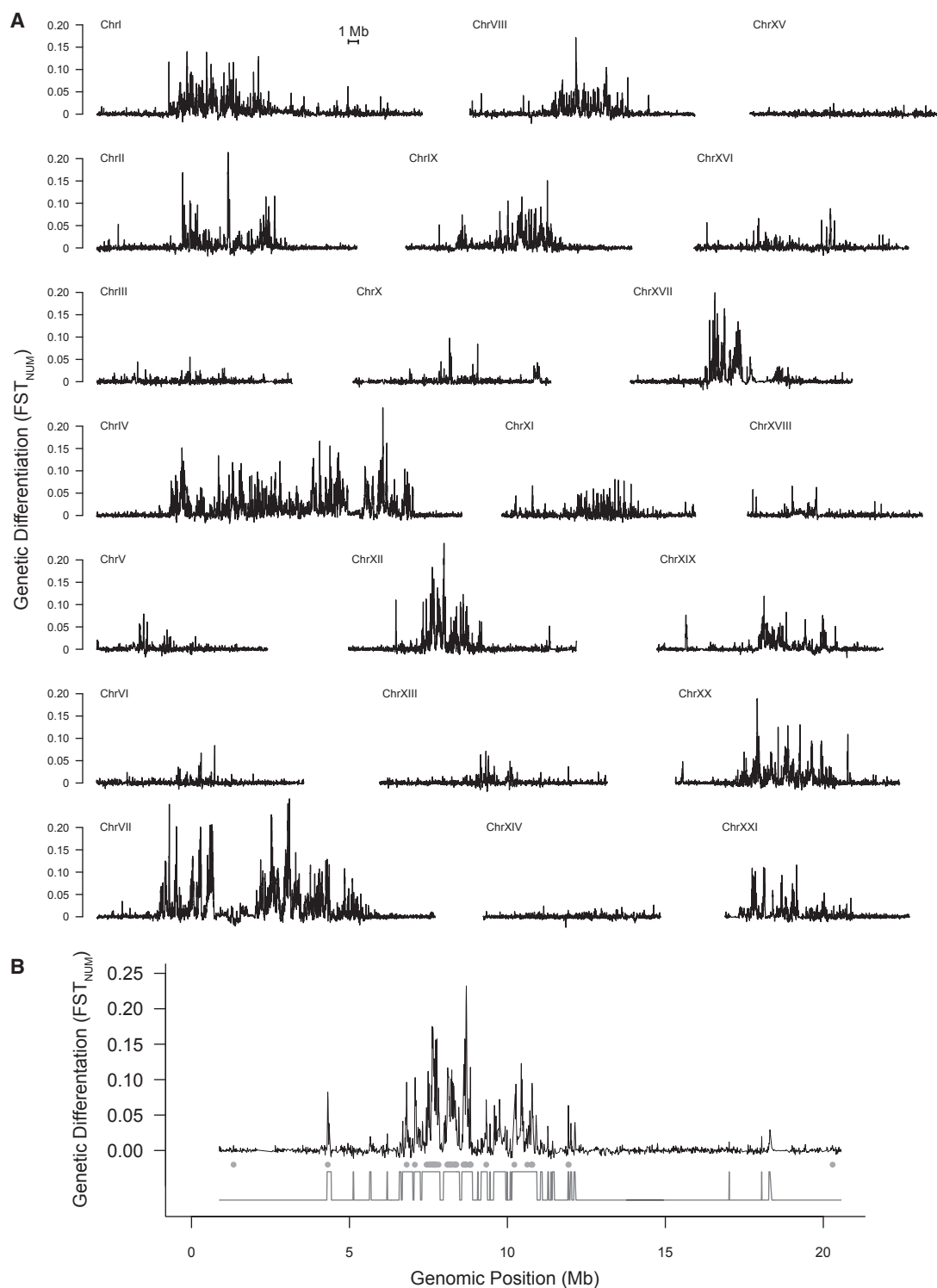
Outlier windows occurred on many chromosomes, with chromosomes IV, VII, XII, and XX displaying especially elevated levels of divergence (Figure 3A). This spatial distribution of outlier windows across chromosomes was highly non-random ( $\chi^2 = 1753$ , df = 20,  $p < 2.2e-16$ ). Outlier windows occurred most frequently in chromosomal centers, which have relatively low recombination rates [17]. Across all windows,  $F_{ST_{NUM}}$  and local recombination rate were negatively correlated (Figure 4;  $r = -0.31$ ,  $p = 1.5e-15$ ).

### Influence of Abiotic Lake Environment

Lakes with and without sculpin overlapped along axes  $ePC1$  and  $ePC2$  describing abiotic environmental differences among lakes (Figure S1;  $ePC1$ :  $U = 51$ ,  $p = 0.17$ ;  $ePC2$ :  $U = 40$ ,  $p = 0.74$ ). However, lakes without sculpin had higher average pH ( $U = 63$ ,  $p = 0.01$ ) and greater average calcium concentrations ( $U = 59$ ,  $p = 0.03$ ) than lakes with sculpin (Table S1; Figure S1). Calcium concentration and pH were strongly correlated with each other ( $r = 0.83$ ,  $p = 4e-5$ ). Nevertheless, pH and calcium are unlikely to determine the presence or absence of sculpin. Sculpin occur in lakes with higher pH and calcium concentrations than are represented in our sample of non-sculpin lakes [18, 19]. However, sculpin-absent study lakes were more variable in these parameters, in association with local differences in geological substrates: none of the study lakes with sculpin occurred on the high limestone substrates on which several of the sculpin-absent lakes occurred. We reassessed the relationship between  $gPC1_{LAKE}$  and lake type while including pH and calcium concentration as covariates and accounting for spatial effects. Stickleback from different lake types remained divergent along  $gPC1$  (type:  $t = 2.9$ ,  $p = 0.016$ ). No relationship was detected between  $gPC1_{LAKE}$  and pH or calcium concentration (pH:  $t = 0.26$ ,  $p = 0.80$ ; calcium concentration:  $t = -0.91$ ,  $p = 0.39$ ).

We investigated the influence of variation in pH and calcium separately on the stickleback genome, focusing only on lakes without sculpin, whose range of values in pH and calcium encompassed the narrower range of values in sampled lakes with sculpin (Figure S1). We calculated  $F_{ST_{NUM}}$  between stickleback sequences from sculpin-absent lakes grouped into low or high calcium concentration and low or high pH (Figure S4). The comparison of  $F_{ST_{NUM}}$  between stickleback in lakes with and without sculpin shared 23 outlier windows (12 genes) with the analysis of lakes with low and high calcium, and 12 outlier windows (10 genes) with the analysis of lakes with low and high pH.

We conclude that abiotic differences between study lakes in pH or calcium do not drive the majority of the genomic changes differentiating stickleback between lakes with and without sculpin. To minimize the possibility of influence, we excluded the 35 shared outlier windows from subsequent analyses, leaving 1,360 outlier windows and 587 genes associated with the presence or absence of sculpin.

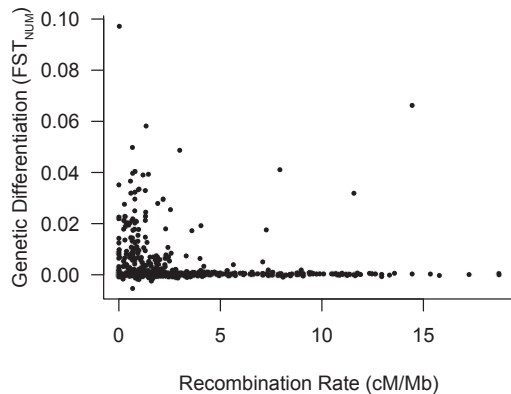


### Figure 3. Extensive Genomic Differentiation between Lake Types

(A) Genomic differentiation between stickleback from lakes with and without sculpin.  $F_{ST\_NUM}$  was measured in 10,000 bp sliding windows (step size 5,000 bp). All chromosomes are plotted on the same scale and ordered by size.

(B)  $F_{ST\_NUM}$  between stickleback from lakes with and without sculpin for an exemplary chromosome showing marked divergence (ChrXII). Estimated state changes (high compared to low differentiation) identified with the HMM are shown below. Outlier windows are indicated with gray circles.

See also [Figures S3](#) and [S4](#) and [Tables S2](#) and [S3](#).



**Figure 4. Genomic Differentiation Correlated with Recombination Rate**

The average value of  $F_{ST\_NUM}$  calculated between lakes with and without sculpin in consecutive 500 kb windows are negatively correlated with average recombination rate ( $R = 0.19$ ,  $p < 1.7e-5$ ). Recombination rate data are from [17].

### Estimating Boundaries of Divergent Regions

Outlier windows typically occurred consecutively in the genome, suggesting that genomic blocks of differentiated sequence might often have evolved together during adaptation to contrasting lake types [20]. To estimate the extent of these blocks, we used a two-state hidden Markov model (HMM) to define boundaries between genomic regions of high and low divergence. This model collapsed the 1,298 contiguous outlier windows into 141 discrete outlier blocks across the genome (Figure 3B; Table S2), with a median width of 220 kb (range: 10–1,470 kb). This number might still overestimate the actual number of independent genetic loci under selection during lake adaptation, since it is possible that linked blocks of standing genetic variation in the ancestral population included a mixture of windows with low and high divergence.

### Broader Implications

Extensive genomic divergence of stickleback was associated with the presence or absence of a single biotic agent of selection, the prickly sculpin. Parallel differentiation of genomes between stickleback from the different lake types involved ~1.8% of the genome, overlapping 587 genes with a wide diversity of biological functions. Widespread adaptation is implicated because genetic drift is unlikely to cause repeated, parallel evolution in multiple evolving populations in association with a specific environmental feature [21]. These extensive changes underscore the rapid and profound effects of a seemingly simple biotic interaction on stickleback evolution.

Study lakes formed recently (<10,000 generations), meaning that differentiation between lake types has repeatedly occurred in a remarkably short time period. On the other hand, genes in outlier windows are not evolutionarily independent, because most occur in contiguous blocks of differentiated genes whose alleles might have increased in frequency in unison during adaptation. We found that stickleback populations sympatric with sculpin have retained more marine genetic variants than populations in lakes without sculpin. Because marine stickleback co-

occur with numerous other fish species, including several other sculpin species, recent release from selection by sculpin produced the direction of most genetic divergence between stickleback populations in the two lake types.

How can a single species from the ecological community rapidly produce differentiation across so many genomic regions? One possible explanation is that traits under selection might be highly polygenic. Stickleback from the two lake types differ in body shape, defensive armor, diet, and behavior [10, 11], and variation in such traits is commonly controlled by many genetic loci [16, 22, 23]. Second, stickleback traits in lakes with sculpin might represent the cumulative outcome of a long series of reciprocal coevolutionary changes in the two species that unraveled in lakes where sculpin are absent. This hypothesis could be addressed in future studies investigating the evolutionary dynamics of sculpin.

A third potential explanation is that sculpin and stickleback are members of a larger network of interacting species. Stickleback may indirectly experience multifarious selection if the presence or absence of sculpin leads to changes in the strength of stickleback interactions with other species. The accumulated effects of “diffuse” or “indirect” selection might be as great or greater than direct selection by sculpin. For example, the lack of sculpin may allow stickleback to colonize the shallow benthic environment, changing stickleback diet and thereby leading to selection to alter behaviors and morphology to improve catching and handling performance on new food items and avoid predators.

Lastly, differentiation of a large portion of the genome might be a correlated response to selection on a smaller set of genes. Most identified differentiated genome regions are clustered in blocks of relatively old standing genetic variation brought to the lakes by colonists [20], with genes in the same block remaining in relatively high linkage disequilibrium during adaptation of a more limited set of loci [24]. In this way, neutral and even deleterious alleles could hitchhike along with linked loci under selection, overestimating the number of genes under selection [25]. It is likely that freshwater allele copies present as standing genetic variation in the colonizing marine population originated from gene flow with other nearby freshwater populations not long before lake colonization [26] and so were initially present as co-segregating blocks. By estimating where blocks of differentiated windows begin and end, we calculate that at most 141 genomic regions were involved in the process of adaptation by stickleback to the presence or absence of sculpin. This still represents a large number of genomic regions responding to biotic selection over a relatively short time span.

Although recombination would eventually break down linkage between genes in the marine population, regions of low recombination would aid the persistence of large blocks of freshwater alleles in the marine population [27]. Consistent with this, we found a negative correlation between local recombination rate and genetic divergence between stickleback from lakes with and without sculpin. We note that by using whole genome sequence data, our study is less likely to be biased toward detecting selection in regions of low recombination than reduced representation methods (e.g., restriction-site-associated DNA sequencing).

How did selection bring so many genes or blocks of genes from low to high frequency in so few generations? Selection on



many genomic regions would presumably generate a high substitution load requiring many selective deaths [28]. One possibility is that biotic interactions may lead to “soft” selection, in which the fitness of an individual is a function of the difference between its phenotype and the most fit phenotype in the population, rather than its difference from an optimum phenotype, reducing substitution load [29]. A change to a new optimum can occur quickly when initial allele frequency in a population is high [30]; therefore, a second possible answer is that adaptive genetic variants did not start at a low frequency. Colonizing populations were most likely large, since colonization occurred during post-glacial rebound of coastal lands. At this time, increased gene flow may have maintained much higher levels of standing variation in the ancestral marine population compared to the present-day marine population. As a result, differentiated genes may have begun at relatively high frequency in the colonizing population, reducing substitution load.

This study provided evidence that ecological interactions between species—even non-intimate and symbiotic interactions—can have large evolutionary consequences. Other studies have found biotic selection to be associated with substantial genomic divergence [14, 31–33], although examples remain few. It is tempting to suggest, based on our study and others, that biotic selection is likely to have more substantial effects on the genome and the phenotype than abiotic agents of selection. Testing this hypothesis will require future controlled studies on the genomic impact of different agents of selection.

## STAR★METHODS

Detailed methods are provided in the online version of this paper and include the following:

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## SUPPLEMENTAL INFORMATION

Supplemental Information includes four figures and three tables and can be found with this article online at <https://doi.org/10.1016/j.cub.2018.12.044>.

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## AUTHOR CONTRIBUTIONS

This study was initially conceived and designed by S.M. and D.S. with later contributions from M.R. Sample collection, morphometrics, DNA extraction, and library preparation were done by S.M. M.R. and S.M. collected water samples and stained fish. M.R. sequenced and constructed the mtDNA haplotype network, analyzed water chemistry, and provided recombination rate data. Genomic data analysis was done by S.M. and D.S. The manuscript was written by S.M. with contributions from D.S. and M.R.

## DECLARATION OF INTERESTS

The authors declare no competing interests.

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## STAR★METHODS

## KEY RESOURCES TABLE

REAGENT or RESOURCE	SOURCE	IDENTIFIER
Biological Samples		
23 Threespine stickleback ( <i>Gasterosteus aculeatus</i> ) tissue samples	This paper	Table S1
Chemicals, Peptides, and Recombinant Proteins		
MS-222	Argent Chemical	Cat#ARF5G
Alizarin Red	Sigma-Aldrich	Cat#A5533
Critical Commercial Assays		
TruSeq DNA PCR-Free Sample kit	Illumina	Cat#20015962
High-Sensitivity DNA Bioanalyzer kit	Agilent Technologies	Cat#5067-4626
Deposited Data		
Threespine stickleback whole genome resequencing data	This paper	NCBI SRA project PRJNA387728
Lake_morphometric_data.csv	This paper	<a href="https://doi.org/10.5061/dryad.nb37pn3">https://doi.org/10.5061/dryad.nb37pn3</a>
FSTnum_outlier_genes.csv	This paper	<a href="https://doi.org/10.5061/dryad.nb37pn3">https://doi.org/10.5061/dryad.nb37pn3</a>
FST_outlier_genes.csv	This paper	<a href="https://doi.org/10.5061/dryad.nb37pn3">https://doi.org/10.5061/dryad.nb37pn3</a>
CSprime_outlier_genes.csv	This paper	<a href="https://doi.org/10.5061/dryad.nb37pn3">https://doi.org/10.5061/dryad.nb37pn3</a>
Oligonucleotides		
L-Pro-F	[34]	N/A
TDK-D	[35]	N/A
Software and Algorithms		
CodonCode Aligner v6.0.2		<a href="https://www.codoncode.com/aligner/">https://www.codoncode.com/aligner/</a>
RAxML v8.0.0	[36]	<a href="https://cme.h-its.org/exelixis/software.html">https://cme.h-its.org/exelixis/software.html</a>
FITCHI	[37]	<a href="http://www.evoinformatics.eu/fitchi.htm">http://www.evoinformatics.eu/fitchi.htm</a>
tpsDig v2.3	[38]	<a href="http://life.bio.sunysb.edu/ee/rohlf/software.html">http://life.bio.sunysb.edu/ee/rohlf/software.html</a>
R package 'shapes'	[39]	<a href="https://www.maths.nottingham.ac.uk/personal/ild/shapes/">https://www.maths.nottingham.ac.uk/personal/ild/shapes/</a>
BWA v0.7.13	[40]	<a href="http://bio-bwa.sourceforge.net/">http://bio-bwa.sourceforge.net/</a>
GATK v3.6	[41]	<a href="https://software.broadinstitute.org/gatk">https://software.broadinstitute.org/gatk</a>
Picard v2.8.2	N/A	<a href="https://broadinstitute.github.io/picard">https://broadinstitute.github.io/picard</a>
VCFTools v0.1.14	[42]	<a href="https://vcftools.github.io/index.html">https://vcftools.github.io/index.html</a>
R script 'convertCoordinate.R': R script for converting to improved stickleback genome assembly coordinates	[15]	<a href="https://datadryad.org/resource/doi:10.5061/dryad.q018v">https://datadryad.org/resource/doi:10.5061/dryad.q018v</a>
R package 'VariantAnnotation'	[43]	<a href="http://www.bioconductor.org/packages/release/bioc/html/VariantAnnotation.html">http://www.bioconductor.org/packages/release/bioc/html/VariantAnnotation.html</a>
R package 'pcaMethods'	[44]	<a href="https://bioconductor.org/packages/release/bioc/html/pcaMethods.html">https://bioconductor.org/packages/release/bioc/html/pcaMethods.html</a>
R package 'Ape'	[45]	<a href="https://CRAN.R-project.org/package=ape">https://CRAN.R-project.org/package=ape</a>
R package 'quanteco'	[46]	<a href="https://github.com/quanteco/quanteco-tools/">https://github.com/quanteco/quanteco-tools/</a>
vcftools_custom: A modified version of vcftools that prints the numerator and denominator of FST		<a href="https://github.com/dalloliogm/vcftools_custom/">https://github.com/dalloliogm/vcftools_custom/</a>
R script 'FSTnum': custom R script for calculating FST <sub>NUM</sub> in windows	This paper	<a href="https://datadryad.org/resource/doi:10.5061/dryad.q018v">https://datadryad.org/resource/doi:10.5061/dryad.q018v</a>
R script 'CSprime': custom R script for calculating CS' in windows	This paper	<a href="https://datadryad.org/resource/doi:10.5061/dryad.q018v">https://datadryad.org/resource/doi:10.5061/dryad.q018v</a>
R package 'qvalue'	[47]	<a href="https://github.com/jdstorey/qvalue">https://github.com/jdstorey/qvalue</a>
R package: 'biomaRt'	[48]	<a href="https://www.bioconductor.org/packages/release/bioc/html/biomaRt.html">https://www.bioconductor.org/packages/release/bioc/html/biomaRt.html</a>

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**Continued**

REAGENT or RESOURCE	SOURCE	IDENTIFIER
R package: 'depmixS4'	[49]	<a href="https://CRAN.R-project.org/package=depmixS4">https://CRAN.R-project.org/package=depmixS4</a>
R package: 'topGO'	[50]	<a href="https://bioconductor.org/packages/release/bioc/html/topGO.html">https://bioconductor.org/packages/release/bioc/html/topGO.html</a>
R core team	[51]	<a href="https://www.r-project.org/foundation/">https://www.r-project.org/foundation/</a>

**CONTACT FOR REAGENT AND RESOURCE SHARING**

Further information and requests for resources and reagents should be directed to and will be fulfilled by the Lead Contact, Sara Miller ([sem332@cornell.edu](mailto:sem332@cornell.edu)).

**EXPERIMENTAL MODEL AND SUBJECT DETAILS**

Adult threespine stickleback fish (*Gasterosteus aculeatus*) specimens were collected using minnow traps from lake and marine regions in Southwestern British Columbia (Table S1). Collection protocols were approved by the British Columbia Ministry of Forests, Lands, and Natural Resource Operations (permits NA-SU12-76311, NA-SU13-85151, NA-SU14-93473). Upon sampling, stickleback were euthanized with an overdose of buffered MS-222 anesthetic (Argent Chemical Laboratories, Redmond, WA) and stored in 95% ethanol.

**METHOD DETAILS****Sampling Strategy**

Our study design, based on [14], applied whole genome re-sequencing of a single representative (diploid) stickleback individual from multiple independent lakes, maximizing the number of populations sampled rather than the number of individuals within populations. The design minimizes false positives signatures of selection produced by the effects of demographic history. The approach is mainly adept at identifying genomic regions that evolved repeatedly from shared standing genetic variation present in the common ancestral population [13, 52]. Standing genetic variation has been previously shown to be a source of adaptive alleles in stickleback [13]. The method is unlikely to detect genomic regions evolving by new mutations unless they produce a parallel signature at the same loci in different populations. Hence our approach provides a conservative estimate of genes responding to biotic environments.

Study lakes were initially identified either based on previous lake observations [11] or were found using Habitat Wizard (<http://www.env.gov.bc.ca/habwiz>), an online database maintained by the Provincial government of British Columbia. We searched for lakes with and without prickly sculpin and whose fish community otherwise contained only threespine stickleback and coastal cutthroat trout (*Oncorhynchus clarkii clarkii*). Cutthroat trout are piscivorous predators of stickleback and sculpin [53] and their presence was unavoidable as they are found in virtually every lake in this region (as are diving piscivorous birds during summer, especially loons (*Gavia immer*)). Therefore, we emphasize that our comparisons are between lakes with and without sculpin, and not between lakes with and without predation. The presence or absence of sculpin was verified in all lakes using minnow traps.

Study lakes selected were geographically interspersed to minimize the contribution of regional differences in lake characteristics. All study lakes, except North Lake (a lake with sculpin) and Klein Lake (a lake without sculpin), were in separate watersheds, and all were inaccessible from the sea, ensuring no contemporary gene flow between populations. We therefore treat each lake as an independent replicate.

We sampled up to 25 adult threespine stickleback during the breeding season (May–June 2012–2014) from nine lakes with sculpin and eight lakes without sculpin (Figure 1A). In addition, we sampled marine stickleback at six localities (Figure 1A). Present day marine stickleback are thought to be phenotypically representative of the ancestral marines that colonized freshwater habitats after the end of the last ice age [12, 54]. Modern Pacific Ocean marine populations are considered nearly panmictic, with high gene flow among nearby populations [13, 14, 52, 55].

**Mitochondrial Haplotype Network**

We used mitochondrial DNA (mtDNA) to test for systematic structuring of populations from lakes with and without sculpin. We chose mtDNA rather than whole genome sequences to minimize the effect of parallel evolution of shared standing variation, which can give a false signal of monophyly [14, 56]. Shared haplotypes among lake types would suggest a monophyletic origin, whereas haplotypes interspersed among populations would suggest multiple independent origins.

We Sanger sequenced a 352 bp stretch of the neutrally evolving mitochondrial control region (D-Loop) using the standard primers L-Pro-F [34] and TDK-D [35]. Sequences were aligned and visually checked in CodonCode Aligner (v6.0.2). A ML-phylogeny of haplotypes was inferred in RAxML (v8.0.0) [36] using the GTRCAT model of sequence evolution with rate heterogeneity among sites. The sequence alignment and phylogeny were then used to construct a haplotype genealogy with FITCHI [37]. The final analysis was based on 3–15 stickleback individuals per lake (112 sequences in total) and a total of 25 SNPs.

## Morphometrics

### Body Shape Measurements

Stickleback samples for body shape analysis were stained with alizarin red [57] and photographed on the left with a Nikon D300 camera. Twenty landmarks outlining the shape of the fish and the insertion points of spines and fins (Figure S2) [58] were placed using tpsDig 2.3 [38]. Landmarks were centered, scaled, and rotated using the *shapes* R package [39]. Shape differences among lakes were visualized using a linear discriminant function analysis (LDA), with population as the classification variable.

### Description of Body Shape Landmarks

1. Dorsal insertion of the caudal peduncle
2. Posterior midpoint of the caudal peduncle
3. Ventral insertion of the caudal peduncle
4. Posterior insertion of the anal fin at the first soft ray
5. Anterior insertion of the anal fin at the first soft ray
6. Hinge of the pelvic spine
7. Dorsal insertion of the pectoral fin
8. Ventral insertion of the pectoral fin
9. Anterior edge of the ectocoracoid bone
10. Anterior extent of the preopercle
11. Posteroventral extent of the maxilla
12. Anterior extent of the premaxilla
13. Naris
14. Anterior orbit in line with the midpoint of the eye
15. Posterior orbit in line with the midpoint of the eye
16. Posterior extent of the supraoccipital
17. Insertion point of the first dorsal spine
18. Insertion point of the second dorsal spine
19. Anterior insertion of the dorsal fin at the first soft ray
20. Posterior insertion of the dorsal fin at the first soft ray

### Armor Trait Measurements

The following traits were measured on stained stickleback specimens: length of the first dorsal spine, length of second dorsal spine, length of the pelvic spine, width of pelvic girdle, and number of lateral plates. All traits, with the exception of lateral plates, were size corrected following the methods in [11]. Armor differences among populations were visualized using a principal component analysis of log-transformed values.

### Whole Genome Re-sequencing

We chose a single representative fish from each lake by performing a PCA on size scaled and rotated morphological landmarks (see above) for up to 25 individuals per population. The fish nearest to the mean of PC1 and PC2 was selected for sequencing, whether male or female. Genomic DNA was extracted from fin clips using a standard phenol/chloroform method. Paired-end whole genome libraries were prepared for each fish using the Illumina TruSeq sample kit (Illumina, San Diego CA), with a target insert size of 500 bp and quantified using High-Sensitivity Bioanalyzer chips (Agilent Technologies, Inc). Libraries were sequenced using the Illumina HiSeq2000 (100 bp paired-end reads) at the University of British Columbia and at Génome Québec.

### Bioinformatics Pipeline

Raw sequence reads were aligned to the stickleback reference genome (gasAcu1 2006 assembly) using BWA (v0.7.13) [40]. Single Nucleotide Polymorphisms (SNPs) were identified using the HaplotypeCaller tool in GATK (v3.6) in conjunction with Picard (v2.8.2) (<http://broadinstitute.github.io/picard>) following the GATK 3.6 best practices recommendations [41]. Low quality SNPs and insertion/deletions were removed with VCFtools (v0.1.14) [42] using the filters `-minGQ 100`, `-min-meanDP 6`, `-remove-indels`, and `-max-missing-count 12` (for full dataset) or 8 (freshwater stickleback only). The seventeen freshwater genomes generated a final dataset, after filtering, of 6.3 million SNPs. Overall lower sequence coverage for marine samples (mean 5X) resulted in a dataset of 2.5 million SNPs for all 23 genomes, following filtering. This corresponds to approximately one SNP every 73 bp for the freshwater dataset, and one SNP every 184 bp for the full dataset.

Genome coordinates were converted to the improved stickleback reference genome assembly [15] using R scripts provided by the authors. The stickleback sex chromosome is chromosome XIX and was excluded from subsequent analyses.

### Genomic Divergence Among Lakes

The overall patterns of divergence among our samples were established using a PCA of SNP genotype values. We then determined if the presence of sculpin was correlated with the main axis of genetic differentiation among stickleback lake populations.

A VCF file was loaded into R using the *VariantAnnotation* package [43]. SNPs occurring in masked regions of the genome, as identified in the RepeatMasker track in the USCS stickleback genome table browser ([sticklebrowser.stanford.edu](http://sticklebrowser.stanford.edu)), were removed using custom R scripts. In each individual, SNPs were given a numerical value relative to the reference sequence (REF/REF = 0, ALT/ALT = 1; REF/ALT = 0.5). Missing values were filled in using the average value of that SNP across all samples. The PCA of the covariance matrix among all pairs of SNPs was calculated using the ‘svd’ function in the *pcaMethods* package [44].

We tested for spatial autocorrelation in the values of genome-wide principal components using Moran’s I test in R’s *Ape* package (v5) [45]. Distance-based Moran’s eigenvector map analysis (dbMEM) was then used to control for the effect of geographical autocorrelation [59]. This method was implemented using R’s *quanteco* package [46].

To measure genome-wide differentiation between stickleback from the two lake types, we calculated the raw variance between groups, in sliding windows across the genome using a modified version of *vcftools* ([https://github.com/dallogm/vcftools\\_custom/](https://github.com/dallogm/vcftools_custom/)). We refer to this value as  $FST_{NUM}$  because it is equivalent to the numerator of Weir-Cocherham  $FST$  [60].  $FST_{NUM}$  is a measure of absolute divergence, and has a maximum of 1 at bi-allelic SNPs and a value of 0 at invariant sites.  $FST_{NUM}$  in a window is the sum of its value over all nucleotide bases divided by the number of sequenced bases in each window. The statistical significance of  $FST_{NUM}$  values was assessed using a permutation test. Within each window, individual fish were randomly reassigned to the two groups, keeping the number of fish in each group the same. This process was repeated 10,000 times and a  $FST_{NUM}$  value between groups was calculated for each permutation. The p value is the proportion of times in which the permuted value exceeded  $FST_{NUM}$  calculated from the real data. Outlier windows were determined using a significance level corresponding to a genome-wide false discovery rate of 0.05 using the *qvalue* R package [47]. Outlier windows fulfilling this significance criterion are not only highly divergent but also show parallel allele frequency differences between stickleback in lakes with and without sculpin. A  $\chi^2$  goodness of fit test was performed to determine whether the number of outlier windows per chromosome was different from the number expected based on the number of nucleotides per chromosome.

### Candidate Genes

The number of genes within outlier windows were counted using the *biomaRt* R package [48]. Pseudogenes and ncRNAs were excluded by sorting genes based upon the “gene\_biotype” attribute. We classified “outlier genes” as those genes within 2000-bp upstream or downstream of outlier windows.

### Calculating Genomic Divergence Among Lakes using Other Metrics

#### FST

We also calculated  $FST$  in all genomic windows, a common measure of relative differentiation [60] that ignores invariant sites. Individuals from freshwater populations were grouped by whether their lakes contained sculpin. These two groups are not true subpopulations either spatially or via gene flow, which is absent between all lakes. We calculated  $FST$  between these groups in 10,000 bp sliding windows with a step size of 5,000 bp using *VCFtools* (v0.1.14) [42]. Outlier windows were identified using a permutation test, following the procedures described above for  $CS'$ .  $FST$  was highly correlated with  $FST_{NUM}$  and  $CS'$  (Figure S3).

#### CS'

To measure absolute genomic differentiation between stickleback from the two lake types, we used a modified version of the cluster separation score (CSS) from [14] in 10,000 bp sliding windows (step size 5,000 bp) across the genome. CSS is based on pairwise sequence divergence between individuals at variable sites within a window between individuals belonging to two different groups. Pairwise divergence is used to calculate the first two principal axes from a multi-dimensional scaling (MDS). The CSS score is the average pairwise Euclidean distance between individuals from different groups minus average distance between individuals within groups. CSS mainly describes divergence on axes of co-varying sequence variation among sites within a window. High CSS within a window indicates a comparatively large number of SNPs diverging in parallel between groups of lake.

We made two modifications of the method by Jones et al. [14]. First, we used the first two principal components of SNP variation within each window instead of MDS axes. Second, we scaled the score for each window by dividing it by the total number of called bases (variant and invariant) in the window. These modifications increased the speed of computation and converted CSS to a per-base metric of sequence divergence between groups, which we refer to as  $CS'$ .

A PCA was conducted within each 10,000 bp sliding window (step size 5,000 bp) on the numerical genotype scores (0, 0.5, and 1) at each SNP. After applying the data quality filters listed in the bioinformatics pipeline, a total of 69,215 windows containing 18,232 genes were available from the lake samples. We retained the first two principal components in each window and calculated the pairwise Euclidean distance  $D$  between individual fish from the two lake types.  $CS'$  is then calculated as:

$$CS' = \frac{\left[ \frac{\sum_{i=1}^s \sum_{j=1}^n D_{ij}}{(sn)} - \left( \frac{1}{s+n} \right) \left( \frac{\sum_{i=1}^{s-1} D_{i,i+1}}{(s-1)} + \frac{\sum_{j=1}^{n-1} D_{j,j+1}}{(n-1)} \right) \right]}{N}$$

The numerator of  $CS'$  corrects a typographical error in the formula from [14].  $D$  is the Euclidean distance between two fish,  $i$  and  $j$  are individual fish belonging to different groups, and  $s$  and  $n$  are the number of stickleback individuals from sculpin lakes and non-sculpin lakes. To convert to a per-base measure, we divided by  $N$ , the number of variant and invariant bases sequenced within a



window, yielding CS'. Windows containing fewer than 250 called bases (out of 10,000) or containing fewer SNPs than the total number of fish were dropped. CS' is occasionally negative, which occurs when the average pairwise distance between fish in different lake types is less than the average pairwise distance between fish of the same lake type.

Permutation tests were used to assess the statistical significance of CS' values. Within each window, individual fish were randomly reassigned to the two groups, keeping the number of fish in each group the same. This was repeated 10,000 times and a CS' score was calculated for each permutation. The p value is the proportion of times in which the permuted value exceeded the CS' score calculated from the real data. Outlier windows were determined using a significance level corresponding to a genome-wide false discovery rate of 0.05 using the *qvalue* R package [47]. Outlier windows fulfilling this significance criterion are not only highly divergent but also show parallel allele frequency differences between stickleback in lakes with and without sculpin. A  $\chi^2$  goodness of fit test was performed to determine whether the number of outlier windows per chromosome was different from the number expected based on the number of nucleotides per chromosome.

CS' was more sensitive at identifying outlier windows than the other metrics (Table S2). Using CS' we identified significant differences in 1,645 of 69,215 informative windows (hereafter outliers), representing ~2.4% of the genome. Median CS' was 0.0117 in outlier windows compared to 0.0001 in other windows. Outlier windows overlapped introns or exons of 684 of the 18,254 annotated genes in sampled windows (3.7%). There was an average of 0.84 genes (SD = 0.80) in each outlier window. Outlier windows were non-randomly distributed across the genome ( $\chi^2 = 1986$ , df = 20,  $p < 2.2e-16$ ). CS' and local recombination rate were negatively correlated ( $R = -0.206$ ,  $p = 3e-6$ ).

### Allele Frequency Difference

We also calculated Allele Frequency Difference (AFD) between groups within genomic windows. AFD is a commonly used metric of assessing genetic differentiation between populations [61]. Allele frequency was calculated between the two groups at each bi-allelic SNP using VCFtools [42]. The allele frequency difference was calculated for each SNP using the formula:

$$AFD = 0.5 * |SC - NS|$$

SC is the frequency of the reference allele in the sculpin-lake group and NS is the frequency of the reference allele in the non-sculpin lake group. The average AFD was calculated in 10,000 bp sliding windows with a step size of 5,000 bp using custom scripts in R. AFD was highly correlated with  $F_{ST_{NUM}}$  and CS' (Figure S4).

### Abiotic Lake Characteristics

Physical and chemical properties were compared among lakes to identify environmental differences that might be correlated with sculpin presence or absence. Information on the total surface area, perimeter, and mean depth of each study lake was obtained from Habitat Wizard (<http://www.env.gov.bc.ca/habwiz>). Lake elevation and shortest straight-line distance from the lake to the ocean was determined using Google maps (<http://maps.google.com>). To quantify the spatial distribution of study lakes, we calculated pairwise great circle distances among lakes (in km). Pairwise distances were then summarized using major axes from a multidimensional scaling analysis.

We collected water samples from the seventeen freshwater lakes to test for systematic differences in water chemistry between sculpin and non-sculpin lakes. Water samples were stored at 4°C for several months prior to testing. Sodium (Na) and Calcium (Ca) concentrations were assessed by averaging two independent measurements per lake water sample using flame atomic absorption spectroscopy (machine model: *Spectra AA-220FS*, Varian Inc.). In the same samples, we measured Soluble Reactive Phosphorus (SRP) with a spectrophotometer (machine model: *SpectraMAX 340pc*, Marshall Scientific) and pH. Readings from water samples taken at different time points from the same lake (i.e., in different years, or in different months between May and August within the same year) revealed little variation, indicating relatively high temporal stability in overall water chemistry within a lake. Whenever several water samples per lake were available from different dates, we report the mean of all measurements. All nine variables were log transformed to improve normality before performing a principal components analysis (PCA) of the correlation matrix using the *pcaMethods* package in R [44].

### Influence of Abiotic Environment on Genetic Divergence

Physical and chemical properties of lakes (Table S1) were weakly associated with sculpin presence or absence. To assess this association, nine measured abiotic environmental variables were log transformed then summarized using principal component analysis. The first major axis of environmental variation mainly reflected increased sodium and calcium concentration and decreased distance to the ocean. The second principle component axis (ePC2) mainly reflected increased distance of lakes to the ocean and decreased lake area and perimeter. Lakes with and without sculpin broadly overlapped along ePC1 and ePC2 (Figure S1) and their means were not detectably different (ePC1:  $U = 51$ ,  $p = 0.17$ ; ePC2:  $U = 40$ ,  $p = 0.74$ ).

Lakes with sculpin had higher pH and greater calcium concentrations than lakes without sculpin. We investigated the influence of variation in pH and calcium separately on the stickleback genome, focusing only on lakes without sculpin, whose range of values in pH and calcium was broad and encompassed the narrower range of values in sampled lakes with sculpin (Figure S1). We calculated  $F_{ST_{NUM}}$  between stickleback sequences from sculpin-absent lakes grouped by low ( $n = 5$ ) and high ( $n = 4$ ) calcium concentration (Figure S4). After false discovery rate correction, there were no statistically significant outlier windows, but this may be caused by the lower power from a reduced number of lakes in this analysis. Instead we used a relaxed significance threshold that considered the 1188 windows corresponding to the lowest 0.05% of uncorrected p values ( $p = 0.0082$ ) to be outliers. We repeated this analysis

between sequences from non-sculpin lakes grouped by low ( $n = 4$ ) and high ( $n = 5$ ) pH, which yielded 759 outlier windows under the relaxed criterion (Figure S4). Similar results were obtained when considering the top 1% of  $p$  values as outliers.

### Defining Boundaries of Divergent Regions

We used a two state Hidden Markov Model (HMM) implemented in the R package *depmixS4* [49], to estimate the number and size of contiguous blocks of highly divergent windows between stickleback from lakes with and without sculpin. The model assumed two underlying states of genomic divergence of windows, low and high, each with a distinct mean level of divergence and a normal probability distribution of emissions estimated from the data. Transitions between states, whose probabilities are also estimated from the data, were used to demarcate blocks of divergent windows. A dependent mixture model was fitted to  $\log FST_{NUM}$  values of contiguous (rather than sliding) 10,000 bp windows using the parameters 'nstates = 2' and 'instart = runif(2)'. The use of contiguous windows reduced the number of outlier windows in this analysis to 1,298. Other settings were program defaults. Each chromosome was analyzed separately.

### Correlation of $FST_{NUM}$ with Recombination rate

$FST_{NUM}$  values were averaged in non-overlapping 500 kb genomic windows, using the  $FST_{NUM}$  scores previously calculated between lakes with and without sculpin in 10,000-bp windows. Chromosomal positions were converted to the genome assembly positions in [17] using custom R scripts. Based on pedigree-derived crossover (recombination) rate estimates for the stickleback genome from [17], we calculated average recombination rates for the same genome-wide 500 kb sliding windows as above. For cases in which sliding windows overlapped regions with multiple recombination rate estimates, we weighted the estimates by the degree of overlap with a window, and then calculated the average of the recombination rates. A conventional correlation analysis was used to test the association between  $FST_{NUM}$  and recombination rate variation across the genome.

### Gene Ontology (GO) Enrichment in Outlier Windows

We searched for enriched Gene Ontology (GO) terms among genes identified in outlier windows. To be conservative, we restricted our analysis to the 456 genes identified as outlier by three different metrics,  $FST_{NUM}$ ,  $FST$ , and  $CS$ . Genes within outlier windows were identified using the *biomaRt* R package [48]. Pseudogenes and ncRNAs were excluded by sorting genes based upon the "gene\_biotype" attribute. We classified "outlier genes" as those genes within 2000-bp upstream or downstream of outlier windows. Outlier genes were searched for enriched GO terms using the 'weight01' algorithm in the *TopGO* R package [50], restricting the search to GO terms in the category of biological processes. *TopGO* takes the GO hierarchy into account when identifying enriched GO terms resulting in fewer false positives than other methods. Candidate enriched GO terms were identified using Fisher's exact test (Table S3).

## QUANTIFICATION AND STATISTICAL ANALYSIS

All analyses were conducted in R [51]. Statistical tests are described in Method Details (above).

## DATA AND SOFTWARE AVAILABILITY

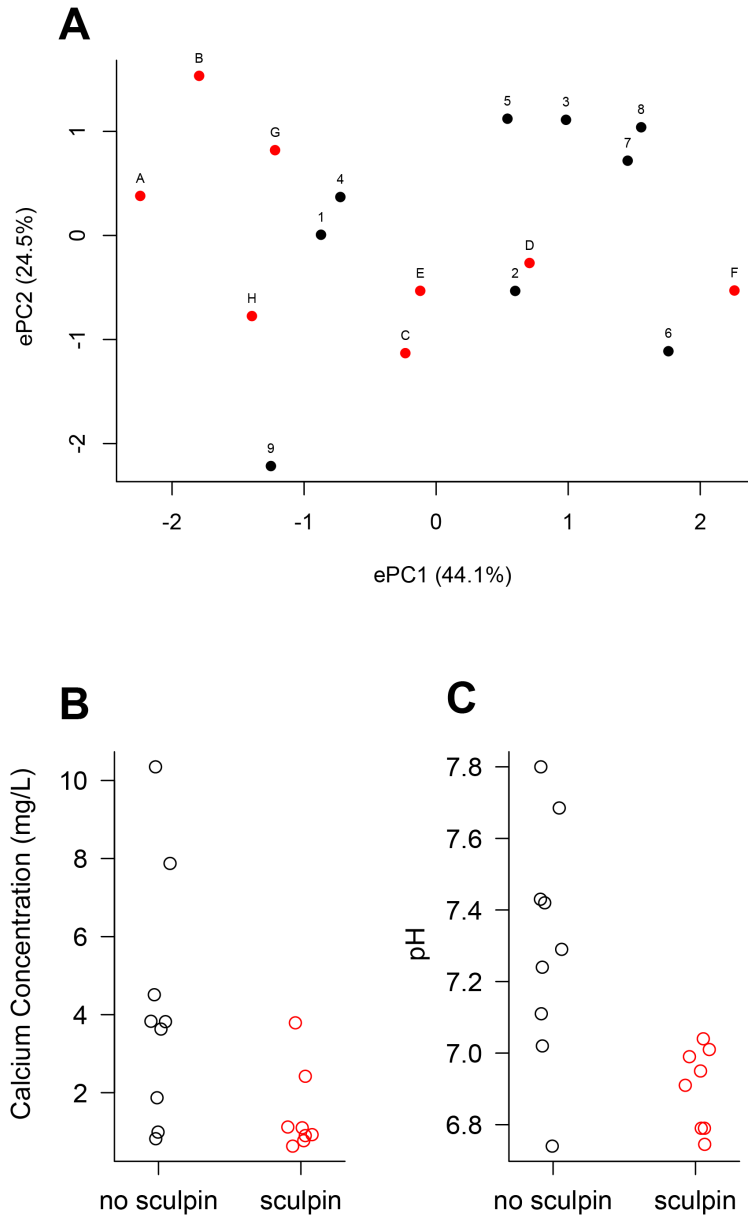
All data files and custom R scripts used to create these analyses are available at the Dryad data repository (<https://doi.org/10.5061/dryad.nb37pn3>). Sequence data used in this paper are available at the NCBI short read archive. The project accession number is PRJNA387728.

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**Supplemental Information**

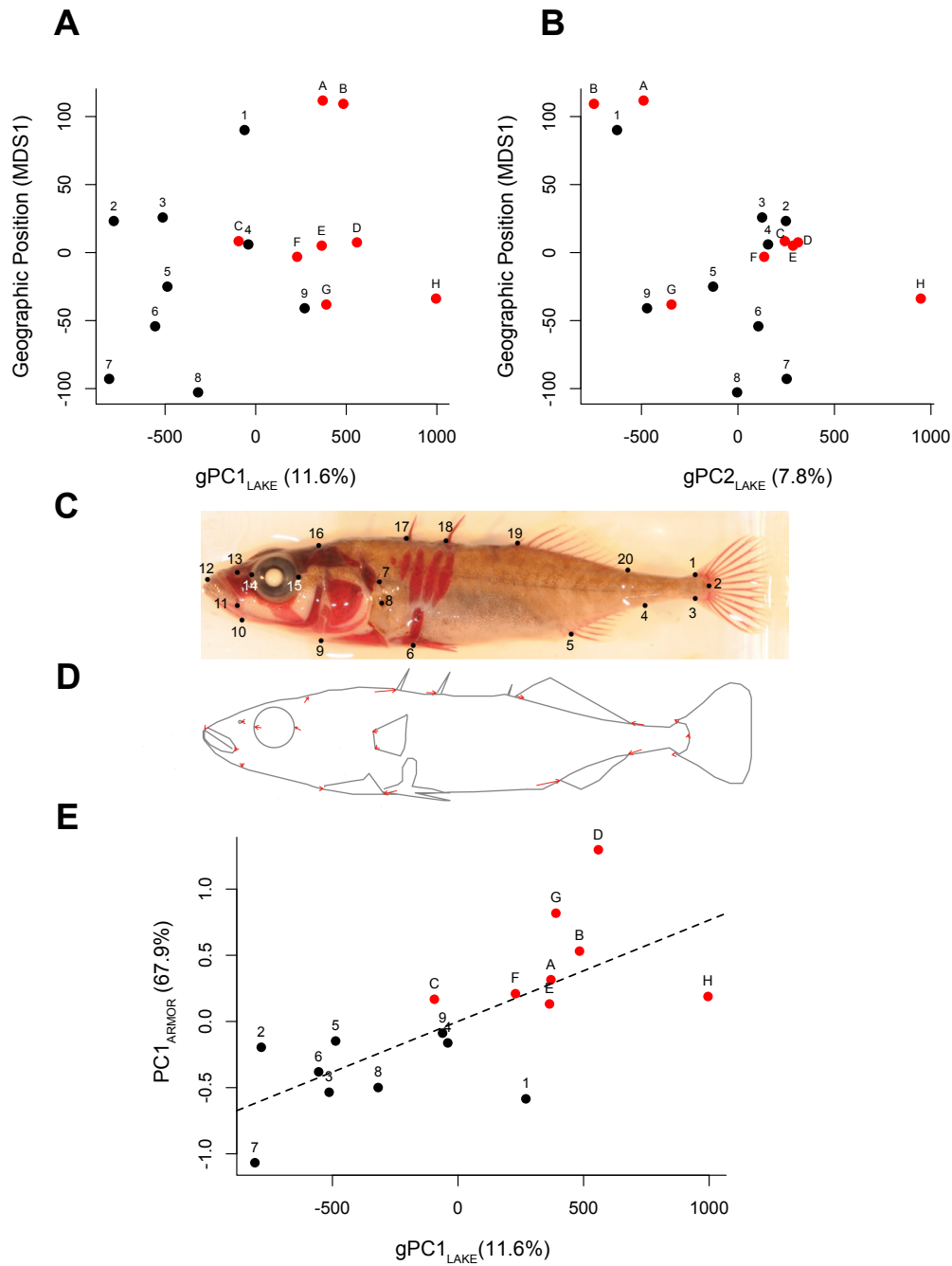
**A Single Interacting Species Leads to Widespread  
Parallel Evolution of the Stickleback Genome**

**Sara E. Miller, Marius Roesti, and Dolph Schluter**



**Figure S1: Little Environmental Difference Between Lakes with Sculpin and Lakes without Sculpin. Related to Figure 1B.**

(A) Population positions along the first two environmental principal components (ePC1 and ePC2) from an analysis of nine environmental variables measured on freshwater lakes with sculpin (red A-H) and without sculpin (black 1-9). The two lake types broadly overlap in ePC1 and ePC2. (B) Calcium concentration [Ca] and (C) pH in study populations. Lakes with sculpin have relatively low calcium concentration and pH, whereas lakes without sculpin range from low [Ca] and low pH to high [Ca] and high pH. Points have been jittered to show overlap. Raw data are given in Table S1.

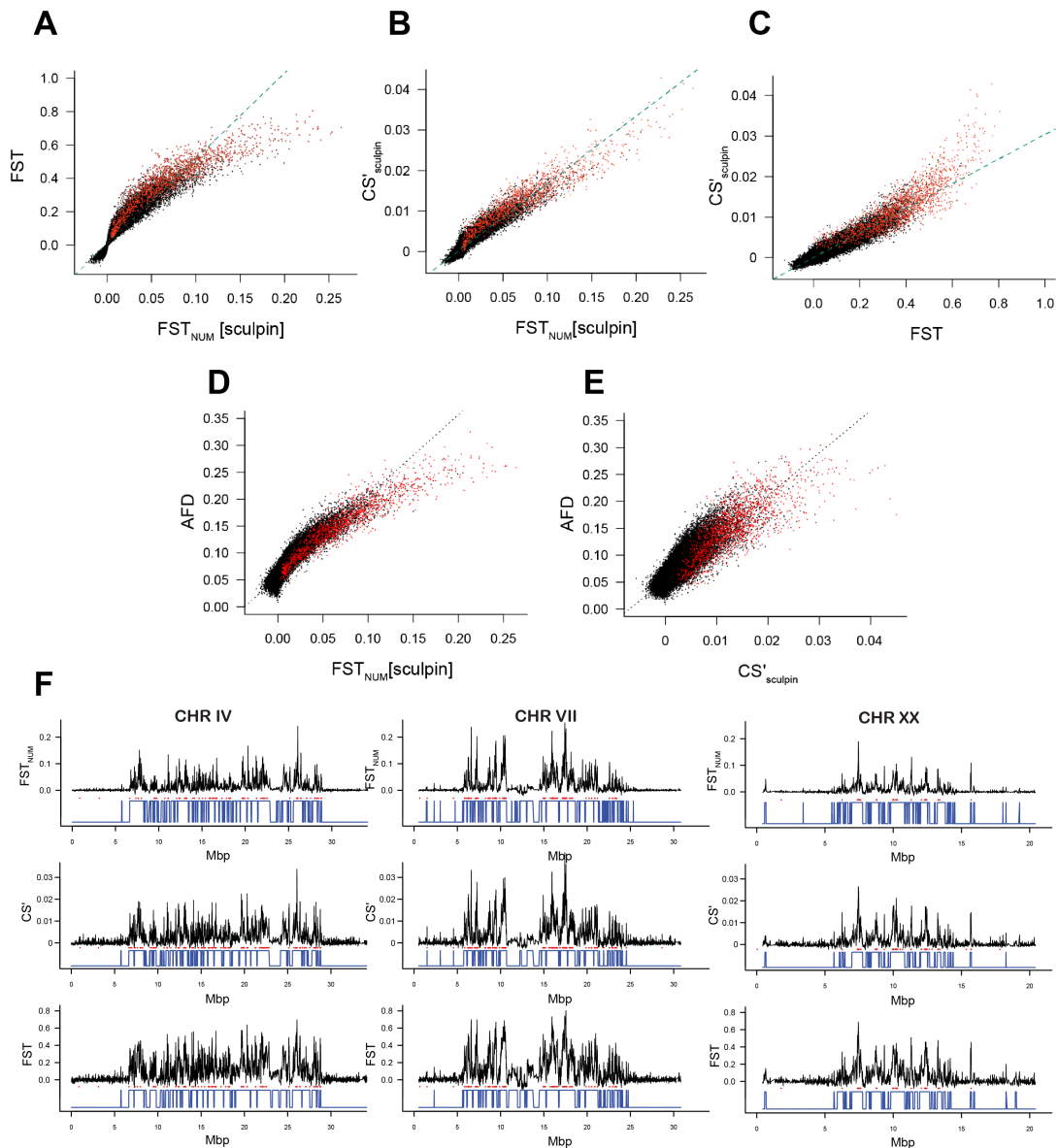


**Figure S2: Genomic Variation is Associated with Stickleback Morphology not Spatial Distribution of Study Lakes. Related to Figure 2.**

(A) Spatial distribution of lakes was summarized using the first multi-dimensional scaling axis of pairwise geographic distance among lakes, as in Figure 1B. The major axis of geographic position of populations (MDS1) does not correlate with the first principal component of genetic variation among all 17 freshwater lake populations of stickleback (gPC1<sub>LAKE</sub>, as in Figure 2A, see main text) ( $t=1.2$ ,  $df=15$ ,  $P=0.25$ ). (B) the second principal component of genetic variation (gPC2<sub>LAKE</sub>) is associated with geographic position of populations (MDS1) ( $t=-2.1$ ,  $df=15$ ,  $P=0.05$ ). (C) Twenty morphometric landmarks outlining body shape and fin insertion points. Landmarks were used to assess shape differences between stickleback populations from lakes with and without sculpin using stained stickleback samples. See supplementary methods for details. (D) Stickleback from lakes with and without sculpin differ predictably in body shape.

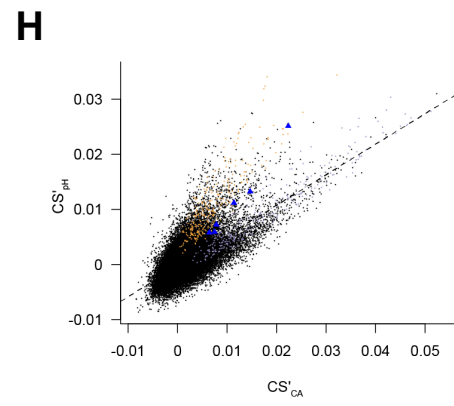
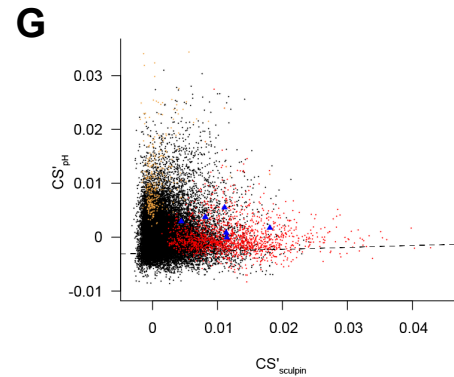
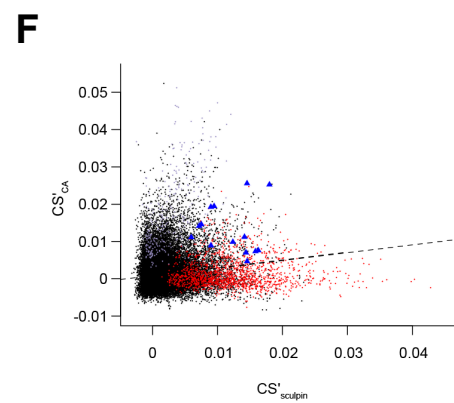
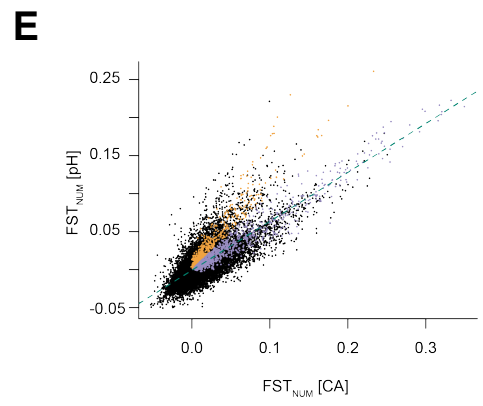
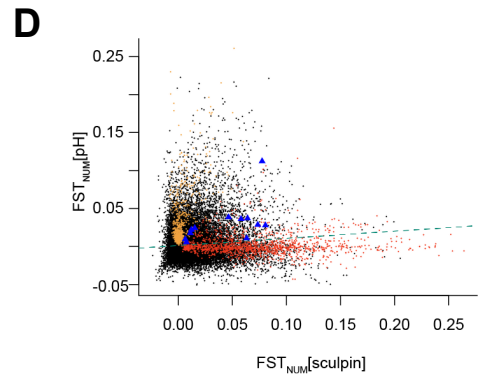
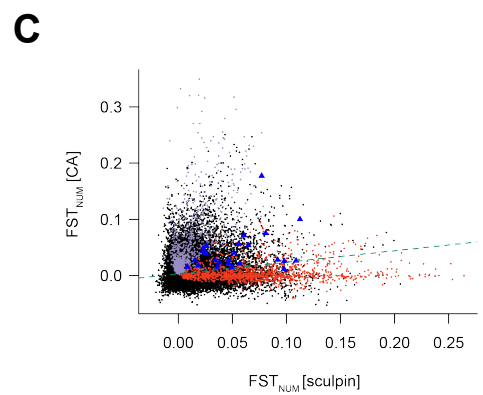
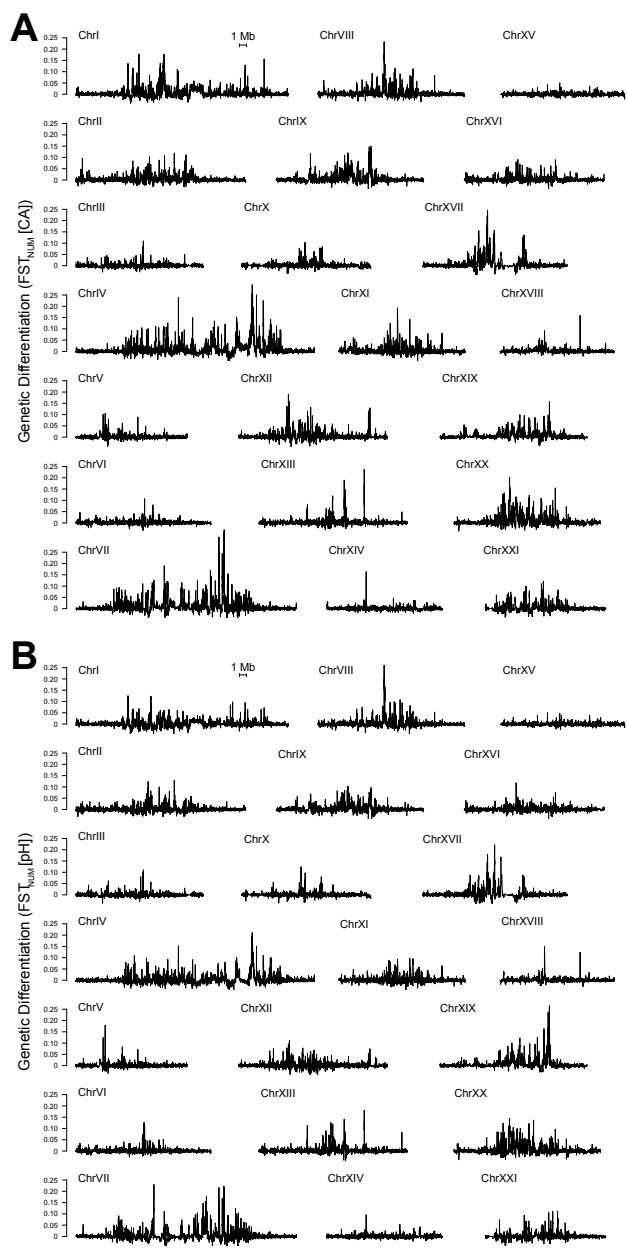


Arrows show the differences in the mean landmark positions for lakes without sculpin (arrow head) compared to the change in landmark positions for lakes with sculpin. Arrow lengths have been multiplied by four to increase visibility. **(E)** Comparison between the first genetic principal component (based on all SNPs in all lake populations, except those on the sex chromosome;  $gPC1_{LAKE}$ ) and the first principal component axis of armor traits ( $PC1_{ARMOR}$ ). A greater value of  $PC1_{ARMOR}$  indicates increased defensive armor. Greater armor is associated with more extreme genomic differentiation along  $gPC1_{LAKE}$  (overall:  $R=0.68$ ,  $N=17$ ,  $P=0.002$ ). Each point represents a single individual from a unique lake population. Lakes with sculpin are red (A-H) and lakes without sculpin are black (1-9).



**Figure S3: Correlation Between Different Metrics for Measuring Genetic Differentiation. Related to Figure 2 and STAR Methods.**

Comparison of differentiation metrics between stickleback from lakes with and without sculpin across the genome. All scores are measured in 10,000-bp windows (5,000-bp step size). **(A)** Plot of  $F_{ST}$  and  $F_{ST\_NUM}$ . Outlier windows for  $F_{ST\_NUM}$  are shown in red. These values are highly correlated ( $R^2 = 0.88$ ,  $P < 2e16$ ) **(B)** Plot of  $CS'_{sculpin}$  and  $F_{ST\_NUM}$ . Outlier windows for  $CS'_{sculpin}$  are shown in red. The two variables are highly correlated ( $R^2 = 0.91$ ,  $P < 2e16$ ). **(C)** Plot of  $CS'_{sculpin}$  and  $F_{ST}$ . Outlier windows for  $CS'_{sculpin}$  are shown in red. The two variables are highly correlated ( $R^2 = 0.86$ ,  $P < 2e16$ ). **(D)** Plot of  $F_{ST\_NUM}$  and Allele Frequency Differences (AFD). Outlier windows from the  $CS'_{sculpin}$  analysis are highlighted in red. The two variables are highly correlated ( $R^2 = 0.85$ ,  $P < 2e16$ ). **(E)** Plot of  $CS'_{sculpin}$  and Allele Frequency Differences (AFD). Outlier windows from the  $CS'_{sculpin}$  analysis are highlighted in red. The two variables are highly correlated ( $R^2 = 0.77$ ,  $P < 2e16$ ). **(F)**  $F_{ST\_NUM}$ ,  $CS'$ , and  $F_{ST}$  values between stickleback from lakes with and without sculpin plotted across three highly differentiated chromosomes. Outlier windows are indicated in red below each metric. State changes as predicted by a hidden markov model (HMM) are show for each metric with a blue line.



**Figure S4: Extensive Genomic Differentiation in Response to pH and Calcium Concentration is not Correlated with Genomic Differentiation Between Lakes with and Lakes Without Sculpin. Related to Figure 3.**

**(A)** Genome-wide distribution of genetic differentiation ( $F_{ST\_NUM}[CA]$ ) calculated between stickleback in low and high calcium concentration lakes lacking sculpin. **(B)** Genome-wide distribution of  $F_{ST\_NUM}[pH]$  calculated between stickleback in low and high pH lakes lacking sculpin. All chromosomes are plotted on the same scale. The sex chromosome (XIX) is included below but was not included in analyses reported in the manuscript. **(C)**  $F_{ST\_NUM}[CA]$  scores calculated between stickleback from lakes with low or high calcium concentrations plotted against  $F_{ST\_NUM}[sculpin]$  scores between lakes with and without sculpin. The two values were weakly correlated ( $R^2 = 0.039$ ). Outlier windows for the analysis between lakes with low and high calcium concentration are in purple, whereas outlier windows for  $F_{ST\_NUM}[sculpin]$  are in red. **(D)**  $F_{ST\_NUM}[pH]$  scores calculated between stickleback from lakes with low or high pH. The two scores are not correlated ( $R^2 = 0.013$ ). Outlier windows for  $F_{ST\_NUM}[pH]$  are in orange whereas outlier windows for  $F_{ST\_NUM}[sculpin]$  calculated between populations in lakes with and without sculpin are in red. **(E)** Scores for  $F_{ST\_NUM}[CA]$  and  $F_{ST\_NUM}[pH]$  were moderately correlated ( $R^2 = 0.64$ ). Outlier windows from the comparison between lakes with low and high pH are highlight in orange, whereas outlier windows for the comparison between lakes having low and high calcium concentration are highlighted with purple. There were no shared outlier windows. **(F)**  $CS'_{CA}$  scores calculated between stickleback from lakes with low or high calcium concentrations plotted against  $CS'_{sculpin}$  scores between lakes with and without sculpin. The two scores are only weakly correlated ( $R^2 = 0.024$ ). Outlier windows for the analysis between populations in sculpin-absent lakes with low and high calcium concentration are in purple, whereas outlier windows for  $CS'_{sculpin}$  between populations in lakes with and without sculpin are in red. The fourteen shared outlier windows for  $CS'_{CA}$  and  $CS'_{sculpin}$  are identified with blue triangles. **(G)**  $CS'_{pH}$  scores calculated between populations from sculpin-absent lakes having low and high pH plotted against  $CS'_{sculpin}$  scores between populations from lakes with and without sculpin across the same 10,000-bp windows (step size 5000-bp). The two scores are not correlated ( $R^2 = 0.002$ ). Outlier windows for  $CS'_{pH}$  are in orange whereas outlier windows for  $CS'_{sculpin}$  calculated between populations in lakes with and without sculpin are in red. The six shared outlier windows for  $CS'_{pH}$  and  $CS'_{sculpin}$  are identified with blue triangles. **(H)** Scores for  $CS'_{CA}$  and  $CS'_{pH}$  in 10,000-bp windows (step size 5,000-bp) calculated using only lakes without sculpin. These scores are moderately correlated ( $R^2 = 0.49$ ). Outlier windows from the comparison between lakes with low and high pH are highlight in orange, whereas outlier windows for the comparison between lakes having low and high calcium concentration are highlighted with purple. The six shared outlier windows for  $CS'_{CA}$  and  $CS'_{pH}$  are identified with blue triangles. All scores for CA and pH are calculated only between stickleback from sculpin-absent lakes.  $F_{ST\_NUM}$  and  $CS'$  scores were calculated using the same 10,000-bp sliding windows (5000-bp step size).

Population	Type	ID	Latitude	Longitude	Area (ha)	Perimeter (m)	Depth (m)	Elevation (m)	Distance to sea (m)	SRP (uM)	Na (uM)	Ca (mg/L)	pH	Env.PCI	Env.PC2
<b>Lake</b>															
Ambrose	sculpin	C	49.733195	-124.02437	29.8	3200	13.3	56	940	0.9	76.40	1.12	6.79	-0.2331	-1.1315
Black	no sculpin	9	48.773234	-125.096594	130.0	8400	11.1	111	1350	0	115.55	0.99	6.74	-1.2513	-2.2164
Brown	sculpin	E	49.741765	-123.915103	18.8	1796	3.5	49	1120	0.23	54.07	0.90	6.79	-0.1204	-0.5325
Bullocks	no sculpin	7	48.874112	-123.509031	9.4	1300	4	33	3310	0.97	424.75	3.82	7.43	1.4507	0.7180
Cedar	sculpin	A	50.204813	-125.565719	31.0	4900	3	204	11800	0.31	54.80	0.77	6.99	-2.2409	0.3795
Cranby	no sculpin	2	49.696084	-124.50654	44.6	3280	3.2	69	2530	0.34	159.76	7.88	7.69	0.5989	-0.5335
Hoggan	no sculpin	6	49.152667	-123.828194	19.7	2219	3	63	310	0	278.76	3.83	7.02	1.7593	-1.1130
Kirk	no sculpin	3	49.739441	-124.586289	8.3	1372	8.3	121	2543	0.73	126.94	10.35	7.80	0.9836	1.1108
Klein	no sculpin	4	49.730087	-123.968612	13.5	2650	12	135	2660	1.01	47.62	1.87	7.29	-0.7252	0.3689
North	sculpin	D	49.748275	-123.970265	12.8	1737	10.06	45	1040	0.29	100.66	2.42	7.01	0.7073	-0.2651
Ormond	sculpin	B	50.180256	-125.52577	8.0	1753	6.57	230	10140	0.07	59.74	0.63	6.95	-1.7948	1.5333
Pachena	sculpin	H	48.838338	-125.029038	58.7	4389	10.7	88	4730	0.295	87.76	1.10	6.91	-1.3951	-0.7751
Paq	sculpin	F	49.613543	-124.0229	12.1	1785	2.2	12	880	0.67	308.70	3.79	7.04	2.2593	-0.5296
Rosseau	sculpin	G	48.830345	-124.992733	13.1	2120	5.8	137	6405	0.28	74.10	0.93	6.75	-1.2199	0.8192
Stowell	no sculpin	8	48.781861	-123.444086	5.6	983	4.6	77	1400	2.03	218.87	4.51	7.42	1.5530	1.0392
Tom	no sculpin	1	50.256375	-124.923292	17.0	2600	1.7	198	1689	0.03	60.95	0.82	7.11	-0.8719	0.0063
Trout	no sculpin	5	49.50843	-123.876057	7.6	1308	5.8	157	2420	0.92	162.34	3.63	7.24	0.5403	1.1214
<b>Mean</b>	no sculpin				28.4 ± 13.3	2679 ± 761	6.0 ± 1.2	107 ± 17	2023 ± 304	0.67 ± 0.22	177.3 ± 39	4.19 ± 1.05	7.3 ± 0.11	0.45 ± 0.38	0.06 ± 0.38
	sculpin				23 ± 5.9	2710 ± 458	6.9 ± 1.4	102 ± 28	4632 ± 1569	0.38 ± 0.09	102 ± 30	1.45 ± .04	6.9 ± .01	-0.51 ± 0.06	-0.06 ± .04
<b>Mann-Whitney Test</b>					U=30; P=0.61	U=29; P=0.54	U=32; P=0.74	U=40; P=0.74	U=32; P=0.74	U=45; P=0.41	U=53; P=0.11	U=59; <b>P=0.03</b>	U=63; <b>P=0.01</b>	U=51; P=0.17	U=40; P=0.74
<b>Marine</b>															
Bamfield Inlet	marine	M6	48.824876	-125.136342											
Little Cambell River	marine	M3	49.014982	-122.772487											
Oyster Lagoon	marine	M2	49.613611	-124.029722											
Salmon River	marine	M5	49.175163	-122.59434											
Seyward Estuary	marine	M1	50.373634	-125.929494											
West Creek	marine	M4	49.191545	-122.655709											

**Table S1: Abiotic Characteristics of Study Populations. Related to Figure 1B.**

Area, perimeter, and mean depth were obtained from HabitatWizard (<http://www.env.gov.bc.ca/habwiz/>). Elevation and the distance from the lake to the nearest ocean were calculated using Google Maps. Soluble reactive phosphorus (SRP), sodium concentration (Na), calcium concentration (Ca), and pH were directly measured from water samples. Mann-Whitney U tests were used to test for differences in abiotic variables between lakes with sculpin and without sculpin. Statistically significant differences are shown in bold.



	FSTNUM		FST		CS'	
Chromosome	Number of outlier windows	Number of regions of high divergence	Number of outlier windows	Number of regions of high divergence	Number of outlier windows	Number of regions of high divergence
I	101	17	102	14	120	16
II	49	8	49	6	50	9
III	9	0	9	0	6	3
IV	265	28	265	23	331	32
V	2	0	2	0	4	0
VI	11	2	11	2	8	4
VII	432	20	433	16	493	23
VIII	70	10	70	10	86	12
IX	62	10	62	9	71	12
X	8	2	8	2	9	2
XI	27	8	27	9	31	10
XII	142	9	141	8	164	11
XIII	6	2	6	1	11	3
XIV	5	1	5	0	5	1
XV	6	0	6	1	7	4
XVI	11	3	11	3	14	4
XVII	38	6	38	4	48	7
XVIII	17	3	17	2	16	2
XIX (Sex)	Not included	Not included	Not included	Not included	Not included	Not included
XX	110	8	111	8	120	9
XXI	21	4	21	4	45	7
unassembled	3	Not included	3	Not included	6	Not included
<b>Total</b>	1395	141	1397	122	1645	171

**Table S2: Outlier Windows Between Stickleback from Lakes with and without Sculpin are Non-Randomly Distributed. Related to Figure 3.**

The number of 10,000 bp outlier windows for CS'<sub>sculpin</sub> and the number of regions of high divergence identified by the hidden markov model (HMM). Chromosome refer to the genome alignment in [S1].

GO.ID	GO Term Description	Annotated	Significant	Expected	P-value
GO:0006811	ion transport	486	19	11.61	0.00016
GO:0007271	synaptic transmission, cholinergic	4	2	0.1	0.0033
GO:0006897	endocytosis	15	3	0.36	0.00494
GO:0001878	response to yeast	19	3	0.45	0.00981
GO:0006506	GPI anchor biosynthetic process	8	3	0.19	0.01091
GO:0051260	protein homooligomerization	64	5	1.53	0.01806
GO:0016311	dephosphorylation	135	5	3.22	0.0183
GO:0010842	retina layer formation	24	3	0.57	0.01878
GO:0007423	sensory organ development	307	9	7.33	0.0238
GO:0003403	optic vesicle formation	1	1	0.02	0.02389
GO:0006809	nitric oxide biosynthetic process	1	1	0.02	0.02389
GO:0045602	negative regulation of endothelial cell differentiation	1	1	0.02	0.02389
GO:0045737	positive regulation of cyclin-dependent protein kinase activity	1	1	0.02	0.02389
GO:0001886	endothelial cell morphogenesis	1	1	0.02	0.02389
GO:0016444	somatic cell DNA recombination	1	1	0.02	0.02389
GO:0006622	protein targeting to lysosome	1	1	0.02	0.02389
GO:0006047	UDP-N-acetylglucosamine metabolism	1	1	0.02	0.02389
GO:0010749	regulation of nitric oxide mediated signal transduction	1	1	0.02	0.02389
GO:0051570	regulation of histone H3-K9 methylation	1	1	0.02	0.02389
GO:0000045	autophagosome assembly	1	1	0.02	0.02389
GO:0021905	forebrain-midbrain boundary formation	1	1	0.02	0.02389
GO:0016255	attachment of GPI anchor to protein	1	1	0.02	0.02389
GO:0016055	Wnt signaling pathway	83	5	1.98	0.02554
GO:0003007	heart morphogenesis	126	7	3.01	0.03042
GO:0060037	pharyngeal system development	14	2	0.33	0.04278
GO:0006189	'de novo' IMP biosynthetic process	2	1	0.05	0.0472
GO:0050890	cognition	2	1	0.05	0.0472
GO:0008057	eye pigment granule organization	2	1	0.05	0.0472
GO:0021884	forebrain neuron development	2	1	0.05	0.0472
GO:0019731	antibacterial humoral response	2	1	0.05	0.0472
GO:0048755	branching morphogenesis of a nerve	2	1	0.05	0.0472
GO:0003308	negative regulation of Wnt signaling pathway involved in heart development	2	1	0.05	0.0472
GO:0046654	tetrahydrofolate biosynthetic process	2	1	0.05	0.0472
GO:0048814	regulation of dendrite morphogenesis	2	1	0.05	0.0472

**Table S3: Gene Ontology (GO) Terms Enriched in Outlier Windows. Related to Figure 3.**

Enriched Biological Processes Gene Ontology (GO) terms calculated using topGO for windows that were identified as outliers using  $FST_{NUM}$ ,  $CS'_{sculpin}$ , and FST. The term annotated refers to the number of genes in the dataset that were annotated with that GO term. Significant refers to the number of genes with the given GO term in outlier windows. Expected is the number of genes with the GO term expected by chance. P-values were calculated from Fisher exact tests.

## **Supplemental Reference**

S1. Glazer A.M., Killingbeck E.E., Mitros T., Rokhsar D.S., and Miller C.T. (2015). Genome Assembly Improvement and Mapping Convergently Evolved Skeletal Traits in Sticklebacks with Genotyping-by-Sequencing. *G3-Genes Genom. Genet.* 5, 1463–72.