

**Western Alaska Salmon Stock Identification Program  
Technical Document 6: Selection of the 96 SNP  
Marker Set for Sockeye Salmon**

by

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Alaska Department of Fish and Game

Division of Commercial Fisheries



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Weights and measures (metric)		General		Mathematics, statistics	
centimeter	cm	Alaska Administrative		<i>all standard mathematical</i>	
deciliter	dL	Code	AAC	<i>signs, symbols and</i>	
gram	g	all commonly accepted		<i>abbreviations</i>	
hectare	ha	abbreviations	e.g., Mr., Mrs., AM, PM, etc.	alternate hypothesis	H <sub>A</sub>
kilogram	kg			base of natural logarithm	<i>e</i>
kilometer	km	all commonly accepted		catch per unit effort	CPUE
liter	L	professional titles	e.g., Dr., Ph.D., R.N., etc.	coefficient of variation	CV
meter	m		@	common test statistics	(F, t, $\chi^2$ , etc.)
milliliter	mL	at		confidence interval	CI
millimeter	mm	compass directions:		correlation coefficient	
		east	E	(multiple)	R
		north	N	correlation coefficient	
		south	S	(simple)	r
		west	W	covariance	cov
		copyright	©	degree (angular )	°
		corporate suffixes:		degrees of freedom	df
		Company	Co.	expected value	<i>E</i>
		Corporation	Corp.	greater than	>
		Incorporated	Inc.	greater than or equal to	≥
		Limited	Ltd.	harvest per unit effort	HPUE
		District of Columbia	D.C.	less than	<
		et alii (and others)	et al.	less than or equal to	≤
		et cetera (and so forth)	etc.	logarithm (natural)	ln
		exempli gratia		logarithm (base 10)	log
		(for example)	e.g.	logarithm (specify base)	log <sub>2</sub> , etc.
		Federal Information		minute (angular)	'
		Code	FIC	not significant	NS
		id est (that is)	i.e.	null hypothesis	H <sub>O</sub>
		latitude or longitude	lat. or long.	percent	%
		monetary symbols		probability	P
		(U.S.)	\$, ¢	probability of a type I error	
		months (tables and		(rejection of the null	
		figures): first three		hypothesis when true)	α
		letters	Jan,....,Dec	probability of a type II error	
		registered trademark	®	(acceptance of the null	
		trademark	™	hypothesis when false)	β
		United States		second (angular)	"
		(adjective)	U.S.	standard deviation	SD
		United States of		standard error	SE
		America (noun)	USA	variance	
		U.S.C.	United States	population	Var
			Code	sample	var
		U.S. state	use two-letter		
			abbreviations		
			(e.g., AK, WA)		
Weights and measures (English)					
cubic feet per second	ft <sup>3</sup> /s				
foot	ft				
gallon	gal				
inch	in				
mile	mi				
nautical mile	nmi				
ounce	oz				
pound	lb				
quart	qt				
yard	yd				
Time and temperature					
day	d				
degrees Celsius	°C				
degrees Fahrenheit	°F				
degrees kelvin	K				
hour	h				
minute	min				
second	s				
Physics and chemistry					
all atomic symbols					
alternating current	AC				
ampere	A				
calorie	cal				
direct current	DC				
hertz	Hz				
horsepower	hp				
hydrogen ion activity	pH				
(negative log of)					
parts per million	ppm				
parts per thousand	ppt, ‰				
volts	V				
watts	W				

***REGIONAL INFORMATION REPORT NO. 5J12-11***

**WESTERN ALASKA SALMON STOCK IDENTIFICATION PROGRAM  
TECHNICAL DOCUMENT 6: SELECTION OF THE 96 SNP MARKER  
SET FOR SOCKEYE SALMON**

by

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# ABSTRACT

Uncertainty about the magnitude, frequency, location, and timing of the non-local harvest of sockeye and chum salmon in Western Alaska fisheries was the impetus for the Western Alaska Salmon Stock Identification Project (WASSIP). The project was designed to use genetic data in mixed stock analysis (MSA) to reduce this uncertainty. A baseline of allele frequencies is required for use in mixed stock analysis to estimate the stock of origin of harvested fish. In 2008 the WASSIP Advisory Panel requested new baselines based upon 96 single nucleotide polymorphisms (SNPs) for both sockeye salmon *Oncorhynchus nerka* and chum salmon *O. keta* to improve the precision and accuracy of stock composition estimates. We investigated 124 SNPs previously developed by the Alaska Department of Fish and Game (ADFG), contractors, or other laboratories to identify the best-performing SNPs for sockeye salmon MSA. We screened these SNPs for a total of 3,447 fish from 36 test populations, representing regions producing large numbers of sockeye salmon and genetic diversity on different geographic scales, to choose the 96 SNPs for the WASSIP baseline. Our primary factors in marker selection were laboratory performance, conformance to population genetic assumptions, Principal Component Analysis (PCA),  $F_{st}$ 's, log-likelihood ratios, and  $f_{ORCA}$ . We selected 96 of the 124 SNPs that perform better in the laboratory and distinguish among populations and regions more clearly than the previous set of 45 SNPs. This marker set should create consistent, reliable data on which to base further analyses for WASSIP. The development and application of methods used to select SNPs in sockeye salmon will provide insight into SNP selection for chum salmon.

Key words: Western Alaska Salmon Stock Identification Project, WASSIP, sockeye salmon, *Oncorhynchus nerka*, mixed stock analysis, genetic baseline, marker selection, single nucleotide polymorphism, SNP

# INTRODUCTION

The WASSIP Advisory Panel requested that 96 single nucleotide polymorphism (SNP) markers be incorporated into the baselines for both sockeye salmon (*Oncorhynchus nerka*) and chum salmon (*O. keta*) to improve the precision and accuracy of stock composition estimates (March 24, 2008). To meet this request for sockeye salmon, we contracted the development (Everett et al. 2011) and validation (J. E. Seeb, Research Professor, School of Aquatic & Fishery Sciences, University of Washington, personal communication) of at least 55 SNP markers that were targeted to differentiate among populations spawning within western Alaska and the Alaska Peninsula drainages and we requested novel SNPs developed by other laboratories. Through these sources, we received a total of 79 novel SNPs. Here we present the methodology for how the best-performing SNP markers for sockeye salmon were selected for WASSIP. These selected markers are being screened in baseline collections at this time and the new baseline genotypes should be complete by the end of July, 2010.

The purpose of this technical document is to describe the methods that the Gene Conservation Laboratory (GCL) used to choose the set of SNPs to be assayed in sockeye salmon for the WASSIP project. We intend to gather feedback from the Technical Committee on the methodology for marker selection in chum salmon. We anticipate having more markers to select from in chum salmon (i.e., 240 SNPs) than we did for sockeye salmon (i.e., 124 SNPs). For sockeye salmon, laboratory performance, conformance to Hardy-Weinberg expectations, linkage among loci, and discrimination among pairs of populations of interest were the primary judges used in marker selection. For chum salmon, we anticipate that population-discrimination factors will contribute to the marker selection to a greater degree because we will be starting from a larger pool of markers and the genetic divergence among chum salmon populations within the WASSIP study area is less than that for sockeye salmon in the current baselines.

# METHODS

## CHOICE OF TEST POPULATIONS

We chose 24 populations from across the species range to represent the regions that produce the majority of sockeye salmon as well as the geographic and genetic diversity observed in previous analyses (Habicht et al. 2010). In addition to these production and diversity criteria, we included populations where collections met the following criteria: 1) fin, heart or liver tissue was available, 2) 8-10 DNA extractions worth of tissue was available for future analyses, and 3) 95 individuals were available for adequate estimates of allele frequencies. We intended this set of 24 populations to serve as a set of test populations for all laboratories interested in the population genetics of Pacific Rim sockeye salmon. In addition to one pair of populations in the set of 24 test populations, we included an additional six pairs of populations that were of interest to ADF&G for a total of 36 populations. Each pair of populations represented two regions which the department desired greater genetic divergence between to aid in mixed stock analyses (MSA) for management purposes. Populations were assigned to fine- and broad-scale regions for use in regional measures of diversity (Figure 1).

## LABORATORY MEASURES

### Assaying genotypes

Genomic DNA for more recent collections was extracted using a DNeasy® 96 Tissue Kit by QIAGEN® (Valencia, CA), while an inorganic method was used for some older collections. We screened the 45 current sockeye salmon SNP markers as well as 77 new SNP markers performed using Fluidigm® 96.96 Dynamic Arrays (<http://www.fluidigm.com>) and 1 current and 1 new marker on the Applied Biosystems platform. The Fluidigm® 96.96 Dynamic Array contains a matrix of integrated channels and valves housed in an input frame. On one side of the frame are 96 inlets to accept the sample DNA from each individual fish and on the other are 96 inlets to accept the assays for each SNP marker. Once in the wells, the components are pressurized into the chip using the IFC Controller HX (Fluidigm). The 96 samples and 96 assays are then systematically combined into 9,216 parallel reactions. In this study, 122 assays were loaded on two separate arrays. Each reaction is a mixture of 4µl of assay mix [1×DA Assay Loading Buffer (Fluidigm), 10×TaqMan SNP Genotyping Assay (Applied Biosystems), and 2.5×ROX (Invitrogen)] and 5µl of sample mix (1×TaqMan® Universal Buffer (Applied Biosystems), 0.05×AmpliTaq® Gold DNA Polymerase (Applied Biosystems), 1×GT Sample Loading Reagent (Fluidigm) and 60–400ng/µl DNA) combined in a 7.2nL chamber. Thermal cycling was performed on an Eppendorf IFC Thermal Cycler as follows: 70°C for 30 min for “Hot-Mix” step, initial denaturation of 10 min at 96°C followed by 40 cycles of 96° for 15 s and 60° for 1 min. The Dynamic Arrays were read on a Fluidigm® EP1™ System after amplification and scored using Fluidigm® SNP Genotyping Analysis software.

The 2 assays genotyped on the Applied Biosystems platform was performed in 384-well reaction plates. Each reaction was conducted in a 5µL volume consisting of 5–40ng/µl of template DNA, 1×TaqMan® Universal PCR Master Mix (Applied Biosystems), and 1×TaqMan® SNP Genotyping Assay (Applied Biosystems). Thermal cycling was performed on a Dual 384-Well GeneAmp® PCR System 9700 (Applied Biosystems) as follows: an initial denaturation of 10 min at 95°C followed by 50 cycles of 92°C for 1 s and annealing/extension temperature for 1 min. The plates were scanned on an Applied Biosystems Prism 7900HT Sequence Detection

System after amplification and scored using Applied Biosystems' Sequence Detection Software (SDS) version 2.2.

### **Scoring genotypes and rating assays**

The most important measure of an assay's utility was the genotyping performance on our platform. We assessed the performance of these assays in our laboratory to determine their utility as markers used in the future for MSA.

Dynamic Arrays that experienced the same DNA and assay loading and PCR process were combined for scoring purposes, and are referred to as combined chip runs. Two Dynamic Arrays always experienced these same conditions on our platform, and so we scored 18 combined chip runs independently that serve as repeated measures for further laboratory performance judges. We entered collected genotypes into the GCL Oracle database, LOKI, for access for further analyses.

During the scoring process, we rated each combined chip run for three measures to judge laboratory performance. These measures were: 1) the spread of, 2) the space between, and 3) the alignment of genotype clusters. The spread of a genotype cluster was a measure of the variation among individuals of a genotype; an assay with very little variability in genotype clusters spread had tight genotype clusters that were easier to score and produced more reliable data (Figure 2). The space between genotype clusters was a measure of the distance between the edges of genotype clusters; an assay with large spacing between clusters had distinct clusters that were easier to score and produced more reliable data (Figure 3). The alignment of genotype clusters was a measure of the alignment of genotype clusters relative to the origin; an assay with separated cluster alignment from the origin had distinct genotype clusters. This measure is correlated with the space between clusters but can differ depending upon the distance from the origin (Figure 4).

For each of these measures, we gave each combined chip run a subjective score ranging between 1 (worst) and 5 (best). To avoid scoring and rating bias, each combined chip run was scored and rated by two people, one of which was always our most senior laboratory staff member for consistency and the other was one of the three other laboratory staff members involved with this project. In addition to these measures, we also calculated the success rate of an assay as the number of successfully genotyped individuals divided by the total number amplified. We tabulated the mean, variance and coefficient of variation (CV) for each of these measures of each assay and ranked each assay by its mean and CV for each measure. Rankings for all judges in this study were corrected for ties when necessary.

## **POPULATION GENETICS MEASURES**

### **Conformance to Hardy-Weinberg expectations**

We tested population genotype frequencies at each marker for conformance to Hardy-Weinberg expectations (HWE) using genetic data analysis (Lewis and Zaykin 2001). We tabulated the number of populations that failed to conform to HWE for three levels of significance ( $\alpha = 0.05$ , 0.01, and 0.001), and ranked each marker based upon the number of populations that failed to conform to HWE for  $\alpha = 0.05$ . The number of the 36 test populations expected to fail to conform to HWE by chance at these three criteria are approximately 2, 0 and 0, respectively.

## Linkage disequilibrium

We tested all pairs of nuclear markers for gametic disequilibrium within each collection using genetic data analysis (Lewis and Zaykin 2001). We defined a pair of markers to be significantly out of gametic equilibrium if tests for gametic disequilibrium were significant ( $P < 0.01$ ) for greater than half of all collections. When gametic linkage was significant, we examined the distribution of the linkage across the range of the test populations to determine if there was a pattern to the phase of linkage that might be useful for MSA. We used the measure of linkage  $r$  (Hedrick 2005), which is  $D$  as described by Hill (1974) but corrected for allele frequency, to assess the phasing of linkage. We also estimated the haplotype frequencies as described by Hill (1974), and, if a haplotype was missing, we interpreted this as indicating the loci were in phase, in which case one locus provides the same amount of information as both linked loci. When a pair of loci were linked, and no pattern in the phase of linkage was observed, we chose which locus to keep for further analyses based upon ratings for the laboratory measures described above and observed heterozygosities. We did not rank markers based upon linkage; rather, we treated this measure as a “gating measure” beyond which only unlinked or usefully linked markers were allowed to continue on for ranking in further analyses.

## Principal component analysis

We conducted a principal component analysis as a measure of how much of the overall genetic variation each marker explained. We calculated the contribution of each marker to the first and second principal components (PC1 and PC2, respectively), as well as the average contribution to the principal components that explained 80% of total variation. We ranked each marker based upon these three judges of informational content.

## Measures of population divergence based upon $F_{ST}$

We calculated three measures of  $F_{ST}$  to assess how each marker described differentiation among populations and regions using the Weir and Cockerham measure of  $F_{ST}$  ( $\theta$ ) calculated in genetic data analysis (Lewis and Zaykin 2001). These measured variation partitioned among populations within fine-scale regions ( $\theta_S$ ), among fine-scale regions within broad-scale regions ( $\theta_P$ ), and between populations within pairs of populations of interest ( $\theta_{Pairs}$ ). We ranked each marker based upon each of these three  $F_{ST}$  measures; the marker with the highest  $F_{ST}$  value received the top rank.

## Measures of population divergence among pairs of populations of interest

We used two measures to examine the divergence among pairs of populations of interest to assess how each marker differentiated populations of interest to ADF&G. These measures were the log likelihood ratio  $G$  statistic (Sokal and Rohlf 1995) from a test of the homogeneity of allele frequencies and  $f_{ORCA}$ .  $f_{ORCA}$  is a measure of informativeness described by Rosenberg (2005; Rosenberg et al. 2003) that indicates how well each marker assigns individuals back to their population of origin. The two measures differ in that the  $G$  test is based solely upon differences in population allele frequencies of a marker while  $f_{ORCA}$  measures a markers utility for population assignment.

## Overall $f_{ORCA}$

We implemented the univariate accumulation algorithm described by Rosenberg (2005), using the Optimal Rate of Correct Assignment as a performance function ( $f_{ORCA}$ ; Rosenberg et al. 2003; Rosenberg 2005). While the univariate accumulation method does not consider synergies

between markers, Rosenberg (2005) demonstrated that it performs as well as methods that do consider synergies.<sup>a</sup>

### **Sum of rankings and selection of final marker set**

We investigated a nonparametric approach to determine how concordant rankings among differing measures of marker performance were. Specifically, we examined Kendall's coefficient of concordance ( $W$ ), which measures the agreement among rankings of different judges (Sokal and Rohlf 1995). It is calculated as:

$$W = \frac{X^2}{k(n-1)} \quad (1)$$

where  $k$  is the number of variables or judges,  $n$  is the number of items per variable or items being ranked, and  $X^2$  is a component of Friedman's method for randomized blocks, the nonparametric analog of a randomized block ANOVA (Sokal and Rohlf 1995). The statistic  $X^2$  is calculated as:

$$X^2 = \left[ \frac{12}{ab(a+1)} \sum^a (\sum^b R_{ij})^2 \right] - 3b(a+1) \quad (2)$$

where  $a$  is the number of treatments and is equal to  $n$  in Equation 1,  $b$  is the number of blocks and is equal to  $k$  in Equation 1, and  $R_{ij}$  is the sum of ranks across  $b$  blocks.

We were more interested in an overall measure of performance across many different judges, so we chose the  $R_{ij}$  of Equation 2 as our overall measure of marker performance. Markers were ranked based upon their  $R_{ij}$  values with the lowest value receiving the highest ranking. Some Markers lacked scores for some judges for various reasons (e.g.,  $G$  tests when the marker was fixed between the populations being tested), and these markers were given rankings equal to the worst ranking for these judges. In contrast, mitochondrial SNPs (*One\_CO1*, *One\_Cytb\_17*, and *One\_Cytb\_26*) were given rankings equal to the highest ranking for our Hardy-Weinberg judge, and both the mitochondrial SNPs and the two linked MHC SNPs (*One\_MHC2\_190* and *One\_MHC2\_251*) were given the rankings that their combined haplotype/phenotype marker received.

We examined the top-ranked 96 markers from this final list with senior laboratory staff to ensure that their performance in the lab would produce reliable data. When it was deemed necessary, we replaced SNPs that performed very poorly with the next highest ranked SNP that we believed would produce reliable data.

The final consideration in choosing 96 markers was correspondence with stakeholder laboratories. We distributed our methodology and final list of 96 SNPs to these laboratories to see if any of the SNPs we chose to remove were highly valuable to other laboratories and could be easily replaced to maximize the efficiency of SNP data collection and standardize data sets across the Pacific Rim.

### **POPULATION GENETICS MEASURES NOT INCLUDED IN THE RANKING OF SNPs**

We conducted three other analyses of genetic variation among test populations that we ultimately did not include as judges for final rankings and marker selection. We present those analyses here

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<sup>a</sup> This sentence is commented on in the section entitled "Technical Committee Review and Comments."

for completeness and to help guide the methodology to be employed for chum salmon marker selection.

### **Backward elimination locus selection**

We examined the marker contribution to correct individual assignment with the backward elimination locus selection algorithm incorporated in the program BELS (Bromaghin 2008). We did not define reporting groups, as our regional groupings and numbers of populations within regions for this project were not representative of the reporting groups we have used in the past—or that we intend to use with the sockeye baseline currently in production. Instead, we chose to maximize the mean individual assignment accuracy as our measure of performance and set a minimum performance of 0. We chose to resample baseline data, with equal baseline collection sample sizes of 95 individuals, to accurately represent our baseline for sockeye salmon. We chose to simulate genotypes for individual assignment from baseline allele frequencies with a fixed number of individuals per population. We set this number of individuals to be 190, which is representative of our desired minimum sample size for mixtures, and conducted 250 replications. Since the analysis took 54 days to complete with these parameters, and was not nearing completion when we chose our marker set, we did not include BELS rankings as a judge but report them here for completeness.

### **Hierarchical log-likelihood analysis**

We examined the homogeneity of allele frequencies among populations within nine fine-scale regions (Table 1) using a hierarchical log-likelihood ratio test ( $G$  test; Sokal and Rohlf, 1995). We included data from all independent nuclear markers and haplotype data for the mitochondrial SNPs and the two linked MHC SNPs. As the number of populations within regions differed greatly (i.e., 3 populations in the North Peninsula region, 7 populations in the Cook Inlet region), we divided  $G$  statistics by degrees of freedom to examine a measure of regional diversity less biased by sampling effort.

### **Nei's gene diversity analysis**

Finally, we examined how diversity was distributed among the different hierarchical levels described above with Nei's Gene Diversity analysis (Nei 1987). We tabulated the percentage of variation attributable to allele frequency differences within populations ( $G_{WP}$ ), among populations within fine-scale regions ( $G_{PF}$ ), among fine-scale regions within broad-scale regions ( $G_{FB}$ ), and among broad-scale regions within the total ( $G_{BT}$ ) for each of the 124 SNPs (Tables 2 and 3) screened for 36 test populations (Table 1) of sockeye salmon.

## **RESULTS**

### **CHOICE OF TEST POPULATIONS**

The 36 populations chosen as test populations represent regions producing the majority of sockeye salmon (Bristol Bay, 8 populations; Kamchatka Peninsula, 4 populations; Eggers and Irvine 2007; Bugaev et al. 2008), much of the genetic diversity that has been observed (6 populations from the Alaska Peninsula and Kodiak; 8 populations from Southeast Alaska and British Columbia; Habicht et al. 2010), and a broad expanse of the species' range (Table 1; Figure 1). The pairs of populations of interest include 3 pairs of populations from Bristol Bay where the GCL has noted a lack of genetic divergence between reporting groups in the past (i.e., Igushik and Wood rivers, Egegik and Ugashik rivers, and Ugashik and Meshik rivers; Habicht et

al. 2007; Dann et al. 2009), and where we hoped to find new discriminating markers. Similar reporting group overlap was identified in the Chignik River system (i.e., Black and Chignik lakes; Creelman et al. *In prep*), in Cook Inlet (i.e., Yentna and Susitna river sloughs; Larson and Mama and Papa Bear lakes; Barclay et al. 2010), and in southern Southeast Alaska (i.e., McDonald and Hugh Smith lakes; Gilk-Baumer et al. *In prep*). With the exception of Glacial Lake in the Norton Sound region, all of our regional groupings are represented by multiple populations; Norton Sound was subsequently included with the western Bristol Bay region.

## LABORATORY MEASURES

### Assaying genotypes

A total of 3,447 fish from the 36 test populations (Table 1) were genotyped for the 124 markers included in this study (Table 2). When all markers were included, individuals were genotyped with a failure rate of 4.11%,<sup>b</sup> but this is inflated artificially high because some markers failed in the laboratory and their genotypes for all individuals were zeroed out during the scoring process (see Scoring Genotypes and Rating Assays below). After correcting for these two markers, individuals were genotyped with a failure rate of 2.52%, which is common for our laboratory. A comparison of genotypes for the 45 SNPs previously genotyped for these collections and genotypes produced in this project revealed a low discrepancy rate of 0.30%. Assuming an equal error rate in the original and current genotyping process, and that this project accurately represents our genotyping process, these collections were genotyped with a process that produced genotypes with an error rate of 0.15%.<sup>c</sup>

### Scoring genotypes and rating assays

A majority of assays performed well in our laboratory. The average cluster tightness was 3.7, and in general, ratings were normally distributed. Five assays were always rated 5 (best), few were rated very poorly (i.e., 5 assays rated < 3), and many assays were rated intermediate producing a ‘broad shoulder’ of intermediate ratings (Table 4; Figure 5). A more left-skewed distribution was observed for the measure of space between genotype clusters. (i.e. 26 assays rated < 3), with an average rating of 3.3 (Table 4; Figure 6). In contrast, a majority of assays produced genotype clusters with good alignment from the origin, with an average rating of 4.3, 34 with a rating of 5 and only 8 with a rating below 3 (Table 4; Figure 7). Three assays failed in the laboratory; two that exhibited multiple clusters, indicative of multiple SNPs (*One\_PPM1K-118* and *One\_U1207-231*) and one due to massive drop-outs (*One\_UCA-24*). These assays were removed from further analyses. A majority of remaining assays produced quality genotypes for the 36 test populations, with an average success rate of 98% (Figure 8).

## POPULATION GENETICS MEASURES

### Conformance to Hardy-Weinberg expectations<sup>d</sup>

Forty-one markers conformed to HWE in all populations, 38 for all but 1 population, and 26 for all but 2 populations at  $\alpha = 0.05$  (Table 5; Figure 9). The few variates observed for this measure resulted in many tied ranks (e.g., markers out of HWE for 1 population were ranked 60.5). While 12 markers failed to conform to HWE at more than 2 populations ( $\alpha = 0.05$ ), only one showed a

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<sup>b</sup> This sentence is commented on in the section entitled “Technical Committee Review and Comments.”

<sup>c</sup> This sentence is commented on in the section entitled “Technical Committee Review and Comments.”

<sup>d</sup> This section is commented on in the section entitled “Technical Committee Review and Comments.”

considerable departure indicative of major problems (i.e., *One\_U1021-57*; 14 populations). As a result, we removed this SNP from further analyses.

### Linkage disequilibrium

Three pairs of SNPs exhibited significant linkage disequilibrium in a majority of populations. These were *One\_aldB-152/One\_ALDOB-135* (36 populations at  $\alpha = 0.05$ ), *One\_GPH-414/One\_GTHa* (36 populations at  $\alpha = 0.05$ ), and *One\_MHC2\_190/One\_MHC2\_251* (26 populations at  $\alpha = 0.05$ ; Table 6). The linked MHC SNPs exhibit a pattern of linkage that is useful for MSA (i.e., a different phasing of linkage across regions) whereas the other two pairs do not (Figure 10; Appendix A.113). For the two pairs of SNPs without a useful pattern of linkage, we chose one SNP to retain for further analyses based upon laboratory ratings and observed heterozygosity (*One\_aldB-152* and *One\_GTHa*). For the remaining pair of linked nuclear SNPs and the triplet of mitochondrial SNPs (*One\_CO1*, *One\_Cytb\_17*, and *One\_Cytb\_26*), genotypes from each locus were pooled to form phenotype/haplotype markers: *One\_MHC2\_190\_251* and *One\_CO1\_Cytb17\_26*, respectively.

### Principal component analysis

The first 2 principal components, PC1 and PC2, explained 25% and 10% of the overall variation, respectively, while the first 12 principal components explained 80% of total variation. Marker contributions to PC1 and PC2 exhibited similar distributions, with few markers contributing greater than 2% to each PC (9 and 10 markers, respectively) and a sharply dropping tail of marker contribution (Table 7; Figures 11 and 12). In contrast, most markers had a similar contribution to the 12 PC's that explained 80% of the overall variation (64 markers 0.5-1%), with one exception: the combined MHC marker contributed 3.9% (Table 7; Figure 13).

### Measures of population divergence based upon $F_{ST}$

The distribution of marker  $F_{ST}$  values among populations within fine-scale region and among fine-scale regions within broad-scale regions were very similar, with few markers having relatively high values, a rapidly declining tail to many intermediate values, and some very small values (Table 8; Figures 14 and 15).  $F_{ST}$  values were greater among populations than among regions, and the distribution of  $F_{ST}$  values reflect marked differences of allele frequencies among populations and regions for a few markers and small differences for many markers. For example, *One\_U1103* exhibited nearly fixed allele frequencies west of the Copper River but highly variable frequencies among populations in the Eastern Gulf region ( $\theta_p = 0.261$ ; rank = 1; Appendix A.82).

The distribution of marker  $F_{ST}$  values between populations within pairs of populations was different from the other two measures in that one marker had a high  $F_{ST}$  while most were of intermediate values (Table 8; Figure 16). *One\_U1004-184* exhibited substantial variation between these pairs of populations and had an  $F_{ST}$  of 0.49 (Rank = 1; Appendix A.71), while 87 markers had an  $F_{ST}$  between 0 and 0.1. However, some markers (e.g., *One\_U404-229*) showed highly divergent allele frequencies between some pairs of populations (McDonald Lake minor allele frequency (MAF) = 0.214, Hugh Smith Lake MAF = 0.096), but were either fixed (Ualik – Pick) or showed very small differences in MAF for other pairs of populations (maximum MAF difference for other 5 pairs = 0.035; Appendix A.101).



## Measures of population divergence among pairs of populations of interest

Log-likelihood ratio ( $G$ ) statistics and  $f_{ORCA}$  generally gave very similar marker rankings (Tables 9 and 10). Markers were generally either good at discriminating between pairs of populations or they were not; that is, markers were either ranked in the top 96 for many population discriminating judges or they were ranked in the top 96 for very few (Figure 17). Notably, *One\_U1004-184* exhibited substantial allele frequency differences between populations in 5 of the 7 pairs, and was the top ranked marker in 6 judges (both  $G$ -statistic and  $f_{ORCA}$  for Ualik-Pick, Deer-Cinder, and Broadway-Hatchery comparisons; Appendix A.71).

## Overall $f_{ORCA}$

The combined MHC and mtDNA markers were the top and second-ranked markers as measured by overall  $f_{ORCA}$ , and *One\_U1004-183* ranked third (Table 11). This may be explained by the fact that MHC and mtDNA, the top and second-ranked markers, were the only two with more than two alleles.

## Sum of rankings and selection of final marker set

Some of our judges produced very different rankings, and so the nonparametric measure of concordance suggested little agreement among judges (e.g., Figure 18). For example, measures of laboratory performance were often not highly correlated with measures of diversity across broad-scale regions.

One hundred fifteen markers passed each of the gating judges and were ranked for each of the 30 judges. The distribution of the summed rankings was approximately normal with an average of 1,704 (SD = 363), and ranged from a low of 704.5 (*One\_MHC2\_251*) to a high of 2,697 (*One\_serpin*) (Table 12; Figure 19). The top-ranked markers (i.e., those with the lowest sum of ranks) included 5 markers with sum of ranks lower than 1,000, 10 markers with sum of ranks between 1,000 and 1,500, and 81 markers with sum of ranks greater than 1,500.

The final examination of the 96 markers with the lowest sum of ranks revealed 2 that performed poorly in the laboratory. These were: *One\_dds-529* (original rank 90; loose, poorly separated genotype clusters) and *One\_psme2-354* (original rank 91; indistinct separation between heterozygote and minor allele homozygote clusters). We replaced these with markers originally ranked 97 and 99 (*One\_U1205-57* and *One\_c3-98*; sum of ranks 2,064 and 2,083, respectively), which performed much better in the laboratory and will likely produce much more accurate and repeatable genotypic data.

Following our correspondence with stakeholder laboratories, we exchanged 1 SNP with its linked complementary SNP to maximize marker set alignment with other laboratories. This exchange was of *One\_GPH-414* for *One\_GTHa*, which was a relatively benign transition, as the 2 SNPs had very similar observed heterozygosities (0.38 for both) and average cluster tightness (3.2 and 3.6, respectively), space between (3.9 and 4.2, respectively), alignment (4.4 and 4.7, respectively), and success rate (98% for both) ratings.

## POPULATION GENETICS MEASURES NOT INCLUDED IN THE RANKING OF SNPs

### Backward elimination locus selection

The BELS analysis indicated that the average individual assignment accuracy was 89% with all 114 markers included in the analysis (Table 13; Figure 14).<sup>e</sup> The first markers dropped contributed little to individual assignment accuracy, with 5 contributing less than 1% individually and 87 contributing 1%. Only 2 markers contributed greater than 7% (combined *One\_MHC2-190\_251* = 14%) and the mitochondrial marker contributed 7% (*One\_CO1\_Cytb17\_26*).

### Hierarchical log-likelihood analysis

The *G* statistics scaled by degrees of freedom varied considerably among markers within a region and among regions (Table 14). As measured by this scaled *G*, the greatest diversity was observed in the British Columbia – Washington region (average *G*/df across markers of 23.72), while the least diversity was observed in the Eastern Bristol Bay region (4.31; average across all markers and all regions = 10.94). When averaged across regions for individual markers, the distribution of *G*/df was approximately normal and *One\_U1004-183* exhibited the greatest intra-regional allele frequency diversity (28.9; Appendix A.71).

### Nei's gene diversity analysis

Overall, 88% of total allele frequency variation was attributed to within populations, with 6% among populations within fine-scale regions, 2% among fine-scale regions within broad-scale regions and 5% among broad-scale regions within the total. *One\_ppie-74* had the largest percent of variation attributable to among populations within regions (25%), although this was largely due to a drastically different allele frequency for the Issaquah population relative to other British Columbia/Washington populations (Table 15; Appendix A.44). The combined *One\_MHC2-190\_251* (18%), *One\_U1214-107* (15%), the combined *One\_CO1\_Cytb17\_26* (14%), *One\_metA-253* (11%), and *One\_U1004-183* (11%) were the other markers with greater than 10% attributable to frequency variation among populations within regions. *One\_HpaI-99* (11%), *One\_metA-253* (6%), *One\_STC-410* (5%) and *One\_STR07* (5%) were the markers that varied the most among fine-scale regions within broad-scale regions.

## DISCUSSION

Laboratory performance, conformance to Hardy-Weinberg expectations, and linkage among SNPs were the primary factors in marker selection for sockeye salmon. These factors played the dominant role in marker selection because they were gating factors and we were selecting from a small set of markers (124 SNPs) that was only 29% larger than our targeted number of 96 SNPs to run for the full baseline. In other words, after exclusion based on these factors, there were few loci to exclude based on other factors to achieve our target of 96 SNPs. Of the 124 SNPs we screened, 6 were excluded outright based on these 3 factors and 3 SNPs were excluded due to fixation. An additional 10 markers were excluded primarily due to low rankings heavily influenced by poor laboratory performance (Figures 18 and 19).

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<sup>e</sup> This sentence is commented on in the section entitled "Technical Committee Review and Comments."

Other factors that influenced marker selection included both positive and negative factors. The positive factors included large allele frequency differences at both wide and narrow geographic scales and a premium on selecting markers that would allow standardization among laboratories. The two MHC SNPs (*One\_MHC2-190* and *One\_MHC2-251*), for example, had relatively poor but acceptable laboratory performance, yet rated at the top based on high measures of genetic variation both on the wide ( $F_{ST}$  and overall  $f_{ORCA}$ ) and narrow (pairwise  $G$  and  $f_{ORCA}$ ) geographic scales (Figure 18). We also selected *One\_GPH-414*, which was the lower laboratory-ranked locus of the linked locus pair that included *One\_GTHa*. We selected *One\_GPH-414* because it is part of the standard set used by Canadian Department of Fisheries and Oceans (CDFO). CDFO is one of the stakeholder laboratories that analyze SNPs in sockeye salmon. Both of these two loci had acceptable laboratory performance, and this change would allow for data sharing across laboratories.

Aside from the three primary factors, the largest negative factor that affected marker selection was low levels of genetic variation. Low variation at all levels was the main factor in the exclusion of an additional 8 markers (Tables 7, 8, 14, and 15). These excluded markers either had very low levels of variability within populations, among populations within fine-scale regions, among fine-scale regions within broad-scale regions, and/or among broad-scale regions within the total. Little variation also heavily influenced some measures as divergent allele frequencies for ‘outlier’ populations provided great contrast from most populations. For example, *One\_metA-253* was the highest ranking SNP as measured by PC1 due to near fixation for all populations west and north of the Copper River, but a frequency of approximately 0.2 for Meziadin Beach in British Columbia (Appendix A.36).

The choice of populations to test SNPs on could impact the ranking of loci due to differences in genetic variation at each of these levels based on different test populations. We chose populations that represented large sockeye salmon producing regions (e.g., Bristol Bay and the Kamchatka Peninsula) and genetic diversity on different geographic scales (e.g., North Alaska Peninsula and the Eastern Gulf of Alaska). Some populations were chosen to ensure that the full Pacific Rim was represented so that we could identify loci that might differentiate among populations from throughout the range of the species. Other collections were included to identify loci that were likely to provide discriminating power between specific populations. How these populations are chosen and how they are grouped into regional groups clearly has substantial influence on marker selection based upon differing population and regional allele frequencies. For example, we included the Glacial Lake population from the Seward Peninsula for geographic representation but did not include other Norton Sound area populations. So we grouped that population with others from the nearest regional group (western Bristol Bay) for analyses of measures based upon regional diversity. This may have artificially inflated the relative intra-regional diversity for that region, and corresponding rankings for measures based upon this diversity, due to this necessary inclusion of a distantly related population.

Similarly, the methodology used for the final selection of loci can introduce potential bias. Test statistics and scores for the 30 judges we chose to include were on different scales, and instead of ranking each independently and summing ranks we could have unitized all scores to a common range. Some of our judges exhibited very little variability (e.g., the number of populations failing to conform to HWE at  $\alpha = 0.05$ ), so a difference of one variate had profound consequences on

rankings when corrections for ties were made. Such a unitizing across judges would have allowed us to include a weighting scheme if we wanted.

After gating judges were considered, we weighted our sockeye salmon SNP selection methodology based upon the number of judges from a category of analyses. For example, because the genetic diversity among populations in Western Alaska is greater for sockeye salmon (e.g., Dann et al. 2012 ) than that for chum salmon (e.g., Jasper et al. *In prep*), we focused our sockeye salmon SNP selection on discriminating between populations from the few reporting groups that still misallocate to one another. By defining half of our judges (i.e., 15 of 30) to be measures of discrimination between these pairs of populations, we weighted our methodology heavily towards this effort (Figures 18 and 20). Our next most important focus was laboratory performance (8 judges) and, lastly, information content of genetic variation among populations, regional groupings and overall (6 judges). Our thinking was that if we selected the loci with the most genetic variation between populations within fine-scale regions, we would select loci that would also be useful at the larger geographic scales. This hypothesis seems to be graphically borne out in Figure 18, where the distribution of good (green) and poor (red) performing loci appear to be correlated between the PCA and  $F_{ST}$  columns and the “Pairs of population measures” columns. Cases where these measures scored high but the loci scored low overall were driven by poor laboratory performance scores.

The set of 96 SNPs we have chosen appear very useful for our needs. The new marker set performs much better in our laboratory than our previous set of 45 SNPs, which will create more consistent, reliable data to base analyses on. Similarly, the new marker set distinguishes among populations and regions better than the old set, and improves the correct assignment as measured by  $f_{ORCA}$  (Figure 20). The rate of improvement to correct assignment with each additional SNP was slightly greater for our chosen 96 SNPs than it was for the original 45 SNPs, indicating that the intermediate SNPs in the 96 SNP set contributed to correct assignment more than the intermediate SNPs in the 45 SNP set. Furthermore, the correct-assignment curve asymptotes at a much higher correct assignment (i.e., 91% compared to ~72%; Figure 20). In general, it appears as though this set of SNPs should provide adequate power to correctly identify reporting groups in mixed stock analyses of WASSIP area fisheries.

## **INCORPORATING LESSONS ON SNP SELECTION FROM SOCKEYE SALMON TO CHUM SALMON**

The development and application of methods to select SNP loci in sockeye salmon provides insights into the most appropriate methods to select SNP loci for chum salmon. Higher numbers of loci available for selection for chum salmon, differences among life histories between the two species that lead to different population structure, and different management needs will all factor into the best methodology. One of the likely parallels between these two methods will be the incorporation of first gating judges (including Hardy-Weinberg equilibrium expectations, linkage disequilibrium, acceptable laboratory performance) and then ranking judges (including laboratory performance, principal component analysis, measures of population divergence based upon  $F_{ST}$ , measures of population divergence among pairs of populations of interest, overall  $f_{ORCA}$ , backward elimination locus selection, hierarchical log-likelihood analysis, and Nei’s gene diversity analysis). The likely differences will be in the focus and weighting of the ranking variables. These ranking judges will be more important for chum salmon than sockeye salmon SNP selection because of the larger number of loci likely available to chose from in chum

salmon. Retaining markers to increase correspondence of marker sets among stakeholder laboratories will affect the final marker selection process but only for markers with intermediate scores (all critical top-scoring markers will be retained and markers that do not pass gating judges will be excluded).

We anticipate focusing the selection of test populations based on a hierarchy of discrimination. At the highest level of the hierarchy, we will score loci based on their among-continent variation (Asia, North America), then move to among coast-wide areas (Japan, Russia, Western Alaska, Alaska Peninsula, Cook Inlet and Prince William Sound, Southeast Alaska, British Columbia, Washington), then among Western Alaska/Alaska Peninsula regions (Alaska Peninsula, eastern Bristol Bay, western Bristol Bay, Kuskokwim River summer, Kuskokwim River fall, Yukon River summer, Yukon River fall, Norton Sound, Kotzebue Sound) and finally within Western Alaska/Alaska Peninsula regions (two populations within each region). Based on known population structure of chum salmon, we expect to find adequate genetic differentiation for the two levels of this hierarchy and to distinguish among the Yukon River fall, Kuskokwim River fall and Alaska Peninsula regions and coastal Western Alaska combined regions. Therefore we propose to emphasize loci that are particularly powerful at discriminating among coastal Western Alaska regions (eastern Bristol Bay, western Bristol Bay, Kuskokwim River summer, Yukon River summer, Norton Sound and Kotzebue Sound). The proposed list of test populations along with our objectives will be shared with the Advisory Panel for comment (Table 16).

One other change we might propose for chum salmon is to change the methods for the scored variables from the ranking method used in sockeye salmon (1 for the best to 124 for the worst) to a rating based on the score for the variable scaled to 1. Let us take the measures of population divergence based upon  $F_{ST}$  among populations within fine-scale regions (Table 8; Figure 14) to demonstrate the difference in methods. For sockeye, we simply ranked each marker and gave a rank value to each marker. However, in looking at the distribution of  $F_{ST}$  across ranked markers, we do not see a linear increase in  $F_{ST}$  from the poorest to best-performing loci. Rather the curve is S-shaped with an initial steep increase in  $F_{ST}$ , followed by a much flatter increase through the middle-ranked markers followed by a steep increase in  $F_{ST}$  for the highest-ranked markers. The ranking method used for sockeye salmon does not take this non-linear information into account. For example the 19<sup>th</sup> ranked marker is ranked 92 rankings higher than the 111<sup>th</sup> marker (111-19), even though it only had a 32% higher scaled  $F_{ST}$  value  $((0.163-0.036)/0.398)$ . On the other hand, the highest ranked marker was ranked only slightly higher (110 rankings higher) than the 19<sup>th</sup> marker even though it has a 91% higher scaled  $F_{ST}$  value  $((0.398-0.036)/0.398)$ . Using the scaled  $F_{ST}$  values will add this information into the marker rating methods. Most of the scored variables did not have linear relationships (see Figures 5-8 and 11-17) and should benefit from the scaled rating method.

## **ACKNOWLEDGEMENTS**

The Technical Document series served as a record of communication between the Alaska Department of Fish and Game Commercial Fisheries Division and the Western Alaska Salmon Stock Identification Program Technical Committee during the implementation of the program.

## **FUTURE ANALYSES**

1. Finish running the baseline. We have currently genotyped approximately 30,500 individuals from 324 populations, and have approximately another 5,700 from 60 populations to genotype.
2. Identify changes to the marker selection methodology used for sockeye salmon to be employed for chum salmon.

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## TECHNICAL COMMITTEE REVIEW AND COMMENTS

### Document 6: Selection of the 96 SNP marker set for sockeye salmon

*Unedited comments by the WASSIP Technical Committee on documents discussed at 23 September 2009 meeting of the WASSIP Advisory Panel.*

Comments on this document appear to be somewhat moot, as the 96-locus panel has already been selected and implemented for sockeye. However, many of the issues are relevant for the parallel exercise for chum.

In general, the approach seems logical and well thought out. Each of the ‘judges’ has some potential value in helping to screen candidate loci, but the criteria differ in the types of information they provide. Some criteria are directly relevant only to local WASSIP issues, while others address issues of broader coastwide relevance. It should be recognized that the number of criteria used for each category produces a *de facto* weighting scheme, and those involved should carefully examine this to ensure that the results adequately reflect the relative importance of different overlapping goals. We believe that substantial coastwide benefits, at little cost to local resolution within WASSIP, can be achieved by giving some consideration to loci that are strongly divergent around the Pacific Rim but relatively homogeneous within WASSIP.

We agree with the general idea that an important initial screening should eliminate from consideration loci that cannot reliably be resolved in the laboratory. From the information presented, however, it is difficult to determine exactly how the tolerance level for error was set.

We also agree with the idea to create a common scale for each of the criteria (e.g., each locus gets a score from 0 to 1). This should facilitate a quantitative rating scale that summarizes information across criteria.

*Comments keyed to specific sentences:*

page 5, 1<sup>st</sup> ¶, last sentence: It is not clear why this would be a general result

page 7, 1<sup>st</sup> full ¶, second sentence: failure rate of ~2.5% in getting genotypes. The key is whether this is random or whether the probability of failure depends on the genotype. If, for example, heterozygotes are more likely to be scored as “failed,” this would bias genotypic and allele frequencies.

page 7, 1<sup>st</sup> full ¶, last sentence: estimated error rate of 0.15%. Actually, this is not a measure of absolute error rate, which would require knowing what the true genotype is. What has been quantified is a measure of *consistency*. For example, independent runs or scorers could get the same answer but both be wrong.

page, 1<sup>st</sup> full ¶, and Table 5: How many total HWE departures were there? How does this compare with the number expected by chance?

page 10, 1<sup>st</sup> full ¶, first sentence: 423: BELS appears to be an appropriate method to select informative loci from a larger set of candidate loci. However, because they do not implement proper cross-validation, BELS and other commonly used programs provide an overly optimistic assessment of assignment success of the selected loci; see Anderson (2010).



## TABLES

Table 1.—Populations of sockeye salmon screened for 124 SNPs (Tables 2 and 3) and their associated map number, pair number, and fine- and broad-scale regional groupings.

Population	Map number	Pair number	Fine-scale region	Broad-scale region
Palana River	1		Russia	Russia
Ozernaya River - Kuril Lake	2		Russia	Russia
Kamchatka River	3		Russia	Russia
Severnaya Lagoon	4		Russia	Russia
Glacial Lake	5		Norton Sound and western Bristol Bay	Western Alaska
Ualik Lake	6	1	Norton Sound and western Bristol Bay	Western Alaska
Pick Creek	7	1	Norton Sound and western Bristol Bay	Western Alaska
Upper Nushagak-Klutapuk Creek	8		Norton Sound and western Bristol Bay	Western Alaska
Tikchik River	9		Norton Sound and western Bristol Bay	Western Alaska
Upper Talarik Creek	10		Eastern Bristol Bay	Western Alaska
Margot Creek	11		Eastern Bristol Bay	Western Alaska
Becharof Creek	12	2	Eastern Bristol Bay	Western Alaska
Deer Creek	13	2, 3	Eastern Bristol Bay	Western Alaska
Mainstem - Cinder River	14	3	North Peninsula	Alaska Peninsula and Kodiak
Bear Lake	15		North Peninsula	Alaska Peninsula and Kodiak
Outer Marker Lake	16		North Peninsula	Alaska Peninsula and Kodiak
Broadway Creek, Black Lake	17	4	Chignik and Kodiak	Alaska Peninsula and Kodiak
Hatchery Beach, Chignik Lake	18	4	Chignik and Kodiak	Alaska Peninsula and Kodiak
Lower Thumb River, Karluk Lake	19		Chignik and Kodiak	Alaska Peninsula and Kodiak
Pyramid Creek - Crescent Lake	20		Cook Inlet	Cook Inlet
West Fork Yentna River	21	5	Cook Inlet	Cook Inlet
Susitna River Slough	22	5	Cook Inlet	Cook Inlet
Larson Lake	23	6	Cook Inlet	Cook Inlet

-continued-

Table 1. Page 2 of 2.

Population	Map number	Pair number	Fine-scale region	Broad-scale region
Mama and Papa Bear Lakes	24	6	Cook Inlet	Cook Inlet
Kenai River	25		Cook Inlet	Cook Inlet
Moose Creek	26		Cook Inlet	Cook Inlet
Mahlo River	27		Copper River and northern Southeast Alaska	Eastern Gulf of Alaska
Klukshu River, Alsek	28		Copper River and northern Southeast Alaska	Eastern Gulf of Alaska
Taku River	29		Copper River and northern Southeast Alaska	Eastern Gulf of Alaska
Scud River, Stikine	30		Southern Southeast Alaska	Eastern Gulf of Alaska
McDonald Lake	31	7	Southern Southeast Alaska	Eastern Gulf of Alaska
Hugh Smith Lake	32	7	Southern Southeast Alaska	Eastern Gulf of Alaska
Meziadin Beach, Nass	33		British Columbia and Washington	Eastern Gulf of Alaska
Slamgeesh River, Skeena	34		British Columbia and Washington	Eastern Gulf of Alaska
Kitlope Lake, Central	35		British Columbia and Washington	Eastern Gulf of Alaska
Issaquah Creek	36		British Columbia and Washington	Eastern Gulf of Alaska

Table 2.–Forward and reverse primer sequences for 124 SNP assays screened for 36 test populations (Table 1).

Assay	Forward sequence	Reverse sequence
<i>One_ACBP-79<sup>a</sup></i>	GAGGTGTGGGCTGACCA	TCGACCGCTGGCAGTG
<i>One_agt-132<sup>b</sup></i>	GACCCAGATCAACAACCTTCATCCA	TGGTTGAGCTAAGGTCCTTGAAC
<i>One_aldB-152<sup>c</sup></i>	CGATCAGGTGACGCTAAAATTAAGTC	GTGGCTTCCTCTTCACTCTGA
<i>One_ALDOB-135<sup>a</sup></i>	CCCGTGCCGGACTTGTT	TCAGCCATGTCAATTGGAATGTGA
<i>One_apoe-83<sup>b</sup></i>	CGCCATGGACAAGGTCAAG	GGCACAGTGCTTCCAAACC
<i>One_bckB-137<sup>c</sup></i>	TCATCTCTCCCTCTCACCAATATCTC	CATTGGGCGGAGTGTATTTCC
<i>One_c3-98<sup>b</sup></i>	GAGTGTGGAAGTGGTTCTTGTTG	GCCGGCAGGGCATCA
<i>One_ccd16-131<sup>b</sup></i>	CCGTGACCTGTTGAACTTTGTTTAG	TCACGTTCTTGGAACACAGC
<i>One_CD9-269<sup>b</sup></i>	ACGCTCTGAGGTGATATGAAACAC	CATCCGACGTCAACATCCAAAC
<i>One_cetn1-167<sup>b</sup></i>	CAGAAATCCTGACTGTAAAACAATGCA	CTGCTCGTTGATCTCTCCATCTC
<i>One_CFP1<sup>d</sup></i>	CGCAGGTCAAAGTAGTACTTAGCAT	GAGCGTCACTTCCTGGAAGTT
<i>One_cin-177<sup>c</sup></i>	CCTCAGACTAGTGACCGTACCTA	CGCTCACCGTGGTTACGT
<i>One_CO1<sup>a</sup></i>	CATAGTAATGCCTGCTGCTAGGA	CCACTTTTTGTTTGAGCTGTGCTAA
<i>One_ctgf-301<sup>a</sup></i>	AAGGACAGAAACATATATGCGTATATTCAATGT	CTGTCTTTCGTCCCCTCTTTAGG
<i>One_Cytb_17<sup>a</sup></i>	CCTGGGAGATCCAGACAATTTTA	CGTAAGCGAAAAGGAAGTATCACTCT
<i>One_Cytb_26<sup>a</sup></i>	CCTGGGAGATCCAGACAATTTTA	CGTAAGCGAAAAGGAAGTATCACTCT
<i>One_dds-529<sup>c</sup></i>	CATAATGCTCCCCATCTTGAATTGG	CACTCAGCCCTTTAGGGAAGA
<i>One_DDX5-86<sup>b</sup></i>	CTCCCACATTGATCTGGACGTA	TGCCACTTGGCCCAAAGAG
<i>One_E2-65<sup>a</sup></i>	GTGGCACCCCTTTCTCT	TGCAAACCTCAGTGGAGAACC
<i>One_gadd45-269<sup>c</sup></i>	AGTTCTCATCCTCTGCGGAAAG	CCAAAATGGCTGGGCAAACAG
<i>One_gdh-212<sup>c</sup></i>	CCTGTGTTGAAGTGGAGTAGGTAA	GCTTTATACTGTAAGTGGACTGACCTT
<i>One_GHII-2165<sup>a</sup></i>	GGCATCAACCTGCTCATCGA	TGCACAAAGTGCGGCAC

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Table 2. Page 2 of 6.

Assay	Forward sequence	Reverse sequence
<i>One_ghsR-66<sup>c</sup></i>	TGTAACAATACAAGGATAATGCAAATAATGTAGGT	GGTTATTAGGTTACTGTGCTGACTGT
<i>One_GPDH-201<sup>a</sup></i>	GAAGCTGATCCTAGACCTGTACCTA	TGGTATGATGGTGCTACTGGAAGT
<i>One_GPDH2-187<sup>a</sup></i>	TCACATCCTTGAGTCGTGTTTGTGTC	GGGCGTAACCGCAAGGT
<i>One_GPH-414<sup>a</sup></i>	CAAGAAGAATCAAGAGAAAGAGAGATGGT	CCTAGTGTCATGCACATAACGTGTA
<i>One_GTHa<sup>d</sup></i>	CAAGAAGAATCAAGAGAAAGAGAGATGGT	CCTAGTGTCATGCACATAACGTGTA
<i>One_HGFA-49<sup>a</sup></i>	ACTTGCTACTTCAGGGTTTTTGTGA	TGGCAGAACAATTCCTCAATGCATA
<i>One_HpaI-71<sup>a</sup></i>	TGTTGTTCCCTAGGCTGTCATTGAAA	CCCTGCGTATTACTAAGGCCATATTTATT
<i>One_HpaI-99<sup>a</sup></i>	CCTGAGTTGTGTTCAATGGGCATAA	TGGGTCATGTTTCATTAGAGCACAAA
<i>One_hsc71-220<sup>a</sup></i>	ACAGCGAAACTATTGATTTAAGGCTCAT	CGCAGGTAAATCACTGATCATGTTT
<i>One_Hsp47<sup>d</sup></i>	CGTTCAAATAAAATGCTGTTTGGCCTTT	GTGGTGTTCGGATTTTTTCCTGAAA
<i>One_Ig-90<sup>b</sup></i>	GGATTGTGGTAACTCTGACAGTAGT	CATCTAAATTCAGTGGCAGTGGGTTA
<i>One_IL8r-362<sup>a</sup></i>	TTGCTAGAAGCGTTGGTTATGATGA	CAGCAAAATTGAGAAGTCACTAGGAAAA
<i>One_ins-107<sup>a</sup></i>	GGAACCCCTGCAAGAGGAGAAAA	GAAATGAATGTGAAGGCAATGATGAGA
<i>One_KCT1-453<sup>b</sup></i>	GGGAAAGTATGCTGTGGGATCAG	GGTTCCTCAGTGAGTGTTCTCTATG
<i>One_KPNA-422<sup>a</sup></i>	TGGGCCCTGGGAAACATC	CCATAGCCACTTTCGATACAGGTAA
<i>One_LEI-87<sup>a</sup></i>	ACAGCGCATCCCCATAATGG	GCCTTTGTGGAGGTCAACGA
<i>One_leptin-92<sup>c</sup></i>	CAGTTGCGCTAAACAGACTCAAG	CAGTTGCTCAGTGATTGTCAACATT
<i>One_lpp1-44<sup>b</sup></i>	GGTCCAATAGGGAGCTCAGACA	GGGAATGAACCAGACATGTGAATG
<i>One_MARCKS-241<sup>a</sup></i>	CCTATCACAGCTTGGTTGAGTTCAA	TCCACCCGCTCATTTTTTGTAAAGAT
<i>One_metA-253<sup>c</sup></i>	TTCTTATCGCTGGTGGCACTTT	GACCAAAGACTATTTAGTTGCCACCTA
<i>One_MHC2_190<sup>a</sup></i>	GTATGGTGTGAAGAATGCA	GCTCACCTGTCTTGTCCAGTA
<i>One_MHC2_251<sup>a</sup></i>	CTGGACAAGACAGGTGAGCA	AAAGTAATGGTCTTGACTTGATCA

-continued-

Table 2. Page 3 of 6.

Assay	Forward sequence	Reverse sequence
<i>One_Mkpro-129<sup>c</sup></i>	TGACGTATGTGCAATGCATGTCTAT	AGATGAAGGACATGGCTGAAAACAT
<i>One_ODC1-196<sup>b</sup></i>	CCGAGGTGGGATTCAACATGAC	TGTCCTCAGACCCAGGGAAA
<i>One_Ots208-234<sup>c</sup></i>	CAGCCGACATGCATCAGTTA	TGACCCCATGTTTCATGCT
<i>One_Ots213-181<sup>a</sup></i>	CCATAGTGTATCACACAATACTCATGTCT	TCTATCATCTGCAAATCTGTGTACTAGACT
<i>One_p53-534<sup>a</sup></i>	GACAATCTTAAAGCGGTGGTCTTG	AACCTTTATCAGCCATCATCCAAC
<i>One_parp3-170<sup>c</sup></i>	TGTGCACCGTTGCCTTTCT	ACAGTAACAAACCAGAGTTACAAGTGG
<i>One_pax7-248<sup>c</sup></i>	AGTAAAGGTAGTGATGCAATGATGCA	AACCGCATAGGACGTAAAGCA
<i>One_PIP<sup>d</sup></i>	ACAGAGTCAGGACTTGATATGTACAGA	CCTGACGAGGGTCTACTACACT
<i>One_ppie-74<sup>c</sup></i>	GTTGATTCCACCTTCTCTGTGATGT	GTGAAATTGACACAGAAGCTGTTCA
<i>One_PPM1K-118<sup>b</sup></i>	GGGATCCAAGCTAACCACAACTT	CACATCAACGCAGGGTTACATTATT
<i>One_Prl2<sup>a</sup></i>	ACCTCTCTCTCTCTCAGGACTCTCA	GAGGAGGTGTGACACATAGATGGA
<i>One_psme2-354<sup>b</sup></i>	TGGTCCTTCAGGTACTTTTCAGAGA	CAAATGCCAATTCTCACCACATGA
<i>One_rab1a-76<sup>b</sup></i>	TCGCCATATTCTCTCTCCCTATCC	ATCCACTCAGACCCATATCTACCAA
<i>One_RAG1-103<sup>a</sup></i>	AGCTCACACATACAACAAATATGATCTAATGT	GTGAACTGCATCTTTGAACAAATGC
<i>One_RAG3-93<sup>a</sup></i>	AGATAAAGATGGTTTCAAAGTCACCCA	GGGCTGCCATCTAAAAAATATTGCT
<i>One_redd1-414<sup>c</sup></i>	GTTGGCTACATCCTAAAACACAATGG	CAGCCCTGGAGTACTGAATCAG
<i>One_RFC2-102<sup>a</sup></i>	TCCAGGAGCTGCATTTTGAGTTAAA	AAGGTGGATGACAATGTGTTAGTGT
<i>One_RFC2-285<sup>a</sup></i>	GGATGAGGCTGACAGGTAAGTC	ACAGTCGTTATAGGTACAGGTACACT
<i>One_RH2op-395<sup>a</sup></i>	GCTGCTAGGTCAAACCTCGAAGAG	CAGCCTTGTTCAACCCCATATCTA
<i>One_rpo2j-261<sup>c</sup></i>	GATTCTGAGATCATACAGTGGATTGGT	GCTTGTCATCTTTCAGCACATACCTA
<i>One_sast-211<sup>c</sup></i>	TGTACTTAGTCCAATAAGCATTTCACAGT	TGGCTAGATTACATGGTCAACAAA
<i>One_serpin-75<sup>a</sup></i>	ACACCTGCAACCAAATTATCATTGC	AACAGGCCTTAACCAATTTCCATCT

-continued-



Table 2. Page 4 of 6.

Assay	Forward sequence	Reverse sequence
<i>One_spf30-207<sup>c</sup></i>	AGCATTTTCAGTTTTGTACATTTACAGTAAAACA	ACCTACTCGTAATTTTCAGGGCAAAA
<i>One_srp09-127<sup>c</sup></i>	CGGAGCTGGAATGACGACAT	AGG TTCAGCAAATCCCTCTTTAGAG
<i>One_ssrd-135<sup>c</sup></i>	TGGAAACTCCTAGTGTACTTCATTCTCA	CGTTCACGCTCCCTAGAATAGA
<i>One_STC-410<sup>a</sup></i>	CAACACAACATCAACATCATTAATAAACATTCTG	AACATCCCCGTTTTGACCACTTAT
<i>One_STR07<sup>a</sup></i>	CACACCTGAGGCACAAGCT	GTATGTCTACCAGAGAGGTCAAGGA
<i>One_SUMO1-6<sup>c</sup></i>	GCACAAGCCAAAAAGTTTTCTCCAT	GGACATAGTTGGAGGCAGACAAAA
<i>One_sys1-230<sup>c</sup></i>	CTACCTGTCTAACAGTGAATGCTAACTT	TGAAACCATTAAGCTCTTTGTAGGACAA
<i>One_taf12-248<sup>c</sup></i>	ACCTTCAATATGGTGGTGGTTACC	ACTAAACGCACAACAGCAAACG
<i>One_Tf_ex11-750<sup>a</sup></i>	AGCAGGTGTAAGCATGTGTACTT	CCTGCTCTGCCTCAACAATGTAA
<i>One_Tf_in3-182<sup>a</sup></i>	GCCCTTAGCACTTCAGTTGCA	CAGACAGAAACCATTGATCCGATTC
<i>One_tshB-92<sup>c</sup></i>	GCATTGTCGTACTCGTGTGTTTG	CACAACAGCAACAATACATGTCACA
<i>One_txnlp-401<sup>c</sup></i>	GCCAGATCCCTTCAGTTGGA	GGCCATTTCAAAAGGCTGCAT
<i>One_U1002-101<sup>b</sup></i>	GCCAACCCTATACTGTACGGATTTTT	TCCGTTGCATTGTCCATCCA
<i>One_U1003-75<sup>b</sup></i>	TCACGAGCCCCAGTCAGA	CGGGTTTCGGTGGTTTAGTATTCTA
<i>One_U1004-183<sup>b</sup></i>	GGTGTGACTGCTGTGTTTAATTGC	ACCATCATTACACAGCAATTCTGAGT
<i>One_U1009-91<sup>b</sup></i>	CTCTGTCCTTGAAGTGTGTCTGTT	GCCGCTGCTACTCTTCCT
<i>One_U1010-81<sup>b</sup></i>	CAGCCCCCTCGAGGTAAGT	GTTGAGACAACAAAACGTCTACTGT
<i>One_U1012-68<sup>b</sup></i>	TCTATTACCATACAGGCCAGTACA	CCTTTTGTGTCTTCCAGTCATGTGA
<i>One_U1013-108<sup>b</sup></i>	TCTGTGCTCTCCTCCAGGAT	CGAAACTGAGGAGTGCTCTGA
<i>One_U1014-74<sup>b</sup></i>	TCCCCTGCAGCAACTGTTTT	GGCAGAGACGGCATCCT
<i>One_U1016-115<sup>b</sup></i>	GGATTTTTGACTTGACCGTTTTGTGT	ATTAACATGTGCAAAGGGAGAATGC
<i>One_U1017-62<sup>b</sup></i>	CAGAGAAGGACGTACCATTGATACAT	CCGGTAGATTGGCGTTGCT

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Table 2. Page 5 of 6.

Assay	Forward sequence	Reverse sequence
<i>One_U1021-57<sup>b</sup></i>	ACAGTGCTACAGGGAGAGAGATTT	GATGGTCAGCGTAGAGAAGCAA
<i>One_U1024-197<sup>b</sup></i>	CTGAACTGATCTACCGCTCTGT	GGAACAGATACTCCAGGAGAGATGA
<i>One_U1101<sup>b</sup></i>	CTATGACATGTTTATTTTAATTAGCCACCAACT	AGTATAGCTAGGGAACCTTTTCGATCTT
<i>One_U1102-220<sup>b</sup></i>	TCCCTCTGCTGGAGAACTACAG	GGAACAGCAGTCCTGAGTACAG
<i>One_U1103<sup>b</sup></i>	CCCAGCCGCCATGTGTA	TGTAGTTCAGCCACCATCTTTGG
<i>One_U1104-138<sup>b</sup></i>	GGAACAGAACACTGAGAATGAATGC	GGGAATATGTCGACTGCTCACT
<i>One_U1105<sup>b</sup></i>	GCCTTAATAGTGTCTTCTGATCCCTTT	CCCTCTGTTGTCCAGACTCTTAG
<i>One_U1201-492<sup>b</sup></i>	GCTTATGACGGAGAAGAGATGCA	AGGATACTGAAGCCCAGAGACA
<i>One_U1202-1052<sup>b</sup></i>	CGATTTGAGTCTCCAATGGTCTCT	ATTCCTATGGTTAACATCAATTCTATAAAGTCAT
<i>One_U1203-175<sup>b</sup></i>	CCCGGAGACATACTTGATGCA	GGAGGACCTGCAGGATCAC
<i>One_U1204-53<sup>b</sup></i>	GTAAAACCCTTCATGTTGGCCATT	CTCCATGTCTGAATGTCCCATCA
<i>One_U1205-57<sup>b</sup></i>	AGTAAATGGTTATTCACGTAACGGATAAG	CAGGACAGTTCACATTCTAACAGA
<i>One_U1206-108<sup>b</sup></i>	CTGAGATGGTGCTTTCTGAGGATA	TGGATGAAAGGGAAATTCTGTCAACA
<i>One_U1207-231<sup>b</sup></i>	GGCCAAACTGACAGGGATCTATTAA	GGGTCCAGTCTGTACACCATCTAT
<i>One_U1208-67<sup>b</sup></i>	ACTTGAATGTCTGTTTCGTAGGTGAT	ACACAGTTGACAGTGGAGCAA
<i>One_U1209-111<sup>b</sup></i>	GTCACGTAATCACGAGAAAGATACTAAATGT	TCTGCGTCTCCAGAGAGGTT
<i>One_U1210-173<sup>b</sup></i>	ACAAAGTCTCTCTCTGAGTAGGAGTAC	CAAAGTATCTCAGAGTGCTGATCTAGGA
<i>One_U1211-97<sup>b</sup></i>	GCGTGTCTCTCCCATTAGAAGA	CTGCAGAAGTACAGCATCTATCTGA
<i>One_U1212-106<sup>b</sup></i>	CGTAATGACCTACCACCATATCAGT	TGGCATGACTTTTAAACAATTCCCAAAAAA
<i>One_U1214-107<sup>b</sup></i>	CCAAATGTACTCCATGTTGGTTAGC	TGCCTGAGTATTAAGCTATATCATTGAAGTTTT
<i>One_U1215-82<sup>b</sup></i>	GTTGCTTGGTTCGTTTGGAGTAG	CTCCAGAAGAGGAATACCACAGTTC
<i>One_U1216-230<sup>b</sup></i>	TGGGATCGGACGTCAATAGATTTC	GTAATACAGAGTGAGCGTGATACATTGT

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Table 2. Page 6 of 6.

Assay	Forward sequence	Reverse sequence
<i>One_U301-92<sup>a</sup></i>	AGCCAGTAGCCGATAATGTTTGTC	CCCCTCCCAAATTGCTAGCT
<i>One_U401-224<sup>a</sup></i>	GGGTGGAGACGAACGGATTC	GTACGATTTTTTTGTAGCCCCAAGT
<i>One_U404-229<sup>a</sup></i>	GTTTGTGTGTTGGTGTTCCTT	CATTTATCTTGGTGGACGTGTGAGT
<i>One_U502-167<sup>a</sup></i>	GCTTTTGTGCAATAGCTATGTTGCT	GCAAAGGTAGGCAGCAGATTG
<i>One_U503-170<sup>a</sup></i>	GATTCAGAATTGCCACGACAAAGAA	GTGATTGGTACATGTCTGTCGAGTT
<i>One_U504-141<sup>a</sup></i>	GCTATAGCTCACAGAGGATCCCA	TATTGGCGGGTGAGGGATG
<i>One_U508-533<sup>a</sup></i>	AGGCACAACCTCACATTTGGAA	CTCAAAGGGTCTGAATACTTATGTAAATAAGGT
<i>One_UCA-24<sup>b</sup></i>	AACTCTGCGTCTGTCTGCTT	TCAGATGGTTCATTATGACAGCAACA
<i>One_vamp5-255<sup>c</sup></i>	GGTTGACTTTTCTTAACCTTTTAATCTGTGATATTGT	GCTGAGCTAGTGATGGTACCATT
<i>One_vatf-214<sup>c</sup></i>	TCATTCCTTTGCCTGGAGCATT	GGCATAACAGCAAAACAATTCAACCA
<i>One_VIM-569<sup>a</sup></i>	TTCTGGGTGGACTCATTGATCAC	ATGCGTTATACCTGTAATCTGCAAGT
<i>One_zn706-68<sup>c</sup></i>	CCACTCTACGTACATCCCATATTCC	GCAGTATACAGATGAGAAAAAGTAGCAAAAAAA
<i>One_ZNF-61<sup>a</sup></i>	CCATTCATGTTCTATTTCAGATATATTTTGTGCA	CCTAGCTAGAGCTCAACAATATGCA
<i>One_Zp3b-49<sup>a</sup></i>	TCCTCGTGGTTATAGTTATAAAGATGTCAGT	TTGGCTCTGCACTCGGTTTA

<sup>a</sup> Assay developed by the Gene Conservation Laboratory of the Alaska Department of Fish and Game (Elfstrom et al. 2006; Smith et al. 2005).

<sup>b</sup> Assay developed by the International Program for Salmon Ecological Genetics at the University of Washington.

<sup>c</sup> Assay developed by the Hagerman Genetics Laboratory of the Columbia River Inter-Tribal Fish Commission.

<sup>d</sup> Assay developed by the Molecular Genetics Laboratory at the Canadian Department of Fisheries and Oceans.

Table 3.–VIC and FAM definitions and sequences for 124 SNP assays (Table 2) screened for 36 test populations (Table 1) and observed heterozygosities ( $H_o$ ).

Assay	VIC	FAM	VIC sequence	FAM sequence	$H_o$
<i>One_ACBP-79<sup>a</sup></i>	G	A	CAGAGGTCATGGTTCTA	CAGAGGTCATAGTTCTA	0.433
<i>One_agt-132<sup>b</sup></i>	A	C	ACAGGAAAATCACGAGCCT	CAGGAAAATCCCGAGCCT	0.425
<i>One_aldB-152<sup>c</sup></i>	A	G	CTCAGGCATTACCTTC	CAGGCATCACCTTC	0.369
<i>One_ALDOB-135<sup>a</sup></i>	G	A	ACAGCACGAAATTA	ACAGCACAAAATTA	0.300
<i>One_apoe-83<sup>b</sup></i>	C	T	TTTAGACGGCGGTCTC	ATTTAGACAGCGGTCTC	0.328
<i>One_bckB-137<sup>c</sup></i>	T	G	TTGATGTAGTTAAGATTATTG	TGATGTAGTTAAGCTTATTG	0.000
<i>One_c3-98<sup>b</sup></i>	C	T	GTTGATGGACCACCTGGT	TTGATGGACCACTTGGT	0.135
<i>One_ccd16-131<sup>b</sup></i>	C	T	AAGGAGAAAGTTGCCGAGCT	ATAAGGAGAAAGTTACCGAGCT	0.002
<i>One_CD9-269<sup>b</sup></i>	C	T	TGGAATGGAGAAATC	ATGGAATGAAGAAATC	0.353
<i>One_cetm1-167<sup>b</sup></i>	A	C	TTGACGAAGCAGACCGA	TTGACGAAGCCGACCGA	0.442
<i>One_CFP1<sup>d</sup></i>	C	T	TGCAGTTCAACATCAA	CTGCAGTTCAATATCAA	0.235
<i>One_cin-177<sup>c</sup></i>	C	T	TCACGCACGGGACAG	CACGCACGGAACAG	0.466
<i>One_CO1<sup>a</sup></i>	T	C	ACTTCTACTACTTTCCC	ACTTCTACTACTCTCCC	N/A <sup>e</sup>
<i>One_ctgf-301<sup>a</sup></i>	G	T	TGATGGATGTGTAGGGC	TGATGGATGTTTAGGGC	0.047
<i>One_Cytb_17<sup>a</sup></i>	A	G	CAACCCGCTAGTTAC	AACCCGCTGGTTAC	N/A <sup>e</sup>
<i>One_Cytb_26<sup>a</sup></i>	A	G	TTTGATATGAGGTGGAGTAA	TGATATGAGGTGGGGTAA	N/A <sup>e</sup>
<i>One_dds-529<sup>c</sup></i>	A	G	AGCAATCCCATCTCTC	AGCAACCCCATCTCTC	0.405
<i>One_DDX5-86<sup>b</sup></i>	C	T	AGGACTTCCTGAAGGAC	AGGACTTCCTAAAGGAC	0.441
<i>One_E2-65<sup>a</sup></i>	C	T	CATTGTCCCTAGGAAAG	ATTGTCCCTAGAAAAG	0.280
<i>One_gadd45-269<sup>c</sup></i>	G	C	CTCCAGCCGATACTT	TCCAGCGGATACTT	0.001
<i>One_gdh-212<sup>c</sup></i>	C	A	ATCTGTTACCAGAATGTTT	ATCTGTTACCATAATGTTT	0.451

-continued-

Table 3. Page 2 of 6.

Assay	VIC	FAM	VIC sequence	FAM sequence	H <sub>0</sub>
<i>One_GHII-2165<sup>a</sup></i>	T	A	CACAAATGGAAATTGA	CACAAATGGTAATTGA	0.213
<i>One_ghsR-66<sup>c</sup></i>	A	T	AGGTTAAGCTGTGTATAAGT	TTAAGCTGTGAATAAGT	0.389
<i>One_GPDH-201<sup>a</sup></i>	T	C	CTTCACCCCTGGAGCC	CACCCCGGAGCC	0.438
<i>One_GPDH2-187<sup>a</sup></i>	G	C	CCTTGGAGGTCTTG	ACCTTGGACGTCTTG	0.187
<i>One_GPH-414<sup>a</sup></i>	C	T	AAGAACTAGAATGGAACAGA	AAGAACTAGAATGAAACAGA	0.381
<i>One_GTHa<sup>d</sup></i>	A	G	CAAGAACTAGAATGAAACAGA	AAGAACTAGAATGGAACAGA	0.381
<i>One_HGFA-49<sup>a</sup></i>	A	T	CTAAAGCACCATGTTGC	ACTAAAGCACCTTGTTGC	0.275
<i>One_HpaI-71<sup>a</sup></i>	A	T	TCAGTTAAGAACTAATTCT	AGTTAAGAACAAATTCT	0.392
<i>One_HpaI-99<sup>a</sup></i>	C	T	AACGGAAGAAACCCCTCAA	AACGGAAGAACTCCTCAA	0.157
<i>One_hsc71-220<sup>a</sup></i>	C	A	ATTGGCCACAGCGC	ATTGGCAACAGCGC	0.352
<i>One_Hsp47<sup>d</sup></i>	A	G	TTATTGACTATGGCACATTG	TTGACTATGGCGCATTG	0.307
<i>One_Ig-90<sup>b</sup></i>	C	G	CTCCTGCATCTTCAGCC	CCTGCATGTTTCAGCC	0.056
<i>One_IL8r-362<sup>a</sup></i>	C	T	CAGCCAAAGAAGAGTC	AGCCAAAAAAGAGTC	0.150
<i>One_ins-107<sup>a</sup></i>	C	T	ATATGTTGTATGGACTACTG	ATATGTTGTATGAACTACTG	0.435
<i>One_KCT1-453<sup>b</sup></i>	G	T	TGGTCAGGGTATCGCCATA	TGGTCAGGGTATCTCCATA	0.198
<i>One_KPNA-422<sup>a</sup></i>	A	G	CTGGTATGAGAAGGCACA	TGGTATGAGGAGGCACA	0.350
<i>One_LEI-87<sup>a</sup></i>	A	G	ACTCGCCACCTCTGT	TCGCCGCCTCTGT	0.430
<i>One_leptin-92<sup>c</sup></i>	T	A	CTGATCCAGGTTCTGTAGTA	CTGATCCAGGTTCTGTAGTA	0.000
<i>One_lpp1-44<sup>b</sup></i>	C	T	TTGTGCTTTCCTGACCTAT	TTGTGCTTTCCTAACCTAT	0.371
<i>One_MARCKS-241<sup>a</sup></i>	T	A	TTGCTTAAAAGGTCTTCC	TTGCTTAAAAGGTCATCC	0.035
<i>One_metA-253<sup>c</sup></i>	C	G	AGGCAATTGAGGTTAAT	AGGCAATTGACGTTAAT	0.094
<i>One_MHC2_190<sup>a</sup></i>	G	T	CTGCTATCGACTACAGC	CGCTGCTATCTACTACAG	0.298

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Table 3. Page 3 of 6.

Assay	VIC	FAM	VIC sequence	FAM sequence	H <sub>0</sub>
<i>One_MHC2_251<sup>a</sup></i>	C	T	CACTTACAGGCCCTG	CACTTACAGGCCTCTG	0.322
<i>One_Mkpro-129<sup>c</sup></i>	A	G	ATGCATATACATGTAATATAT	TGCATATACATGTAACATAT	0.432
<i>One_ODC1-196<sup>b</sup></i>	C	T	CCACCTCCGATGTCC	CACCTCCAATGTCC	0.416
<i>One_Ots208-234<sup>c</sup></i>	-	A	CACACGTTACATCAGATAACT	CACACAATGTTACATCAGATAAC	0.192
<i>One_Ots213-181<sup>a</sup></i>	T	A	CTTTGAATTAAAAACATTTTT	CTTTGAATTAAAAACTTTTTT	0.267
<i>One_p53-534<sup>a</sup></i>	C	A	ATGTCCAAAGATCTGG	AATGTCCAAATATCTGG	0.059
<i>One_parp3-170<sup>c</sup></i>	T	A	ACACAGGAAAAGTTG	ACACAGGTAAAGTTG	0.000
<i>One_pax7-248<sup>c</sup></i>	C	A	AATTCAAAACGAAATGTG	TGAATTCAAAACTAAATGTG	0.212
<i>One_PIP<sup>d</sup></i>	C	T	AACACACATTTCTCAACACA	ACACACATTTTTCAACACA	0.448
<i>One_ppie-74<sup>c</sup></i>	-	A	TGCAAACACTTTTTTTTATAATG	TGCAAACACTTTTTTTTATAATG	0.033
<i>One_PPM1K-118<sup>b</sup></i>	G	T	ATCTCACTTATGGTGCTTC	ATATCTCACTTATTGTGCTTC	N/A <sup>f</sup>
<i>One_Prl2<sup>a</sup></i>	G	T	ACCAATGGGACGAGTG	CCACCAATTGGACGAG	0.448
<i>One_psme2-354<sup>b</sup></i>	A	G	TGATGCAGTAGCTAAAG	ATGCAGTGGCTAAAG	0.373
<i>One_rab1a-76<sup>b</sup></i>	G	T	TGTGGAGCAAGGTAAC	TGTGGAGCAATGTAAC	0.193
<i>One_RAG1-103<sup>a</sup></i>	T	A	CGAATCTCAACAATAAGT	CTCGAATCTCAACTATAAGT	0.079
<i>One_RAG3-93<sup>a</sup></i>	C	T	CATTTTGGACTTCGGGACC	CATTTTGGACTTTGGGACC	0.148
<i>One_redd1-414<sup>c</sup></i>	T	C	CCTAAGTCAGTCACTGTAG	CCCAAGTCAGTCACTGTA	0.410
<i>One_RFC2-102<sup>a</sup></i>	A	G	ATCACGTTGTATTTCTTT	CACGTTGTGTTTCTTT	0.290
<i>One_RFC2-285<sup>a</sup></i>	A	T	CACGACATCTAAGCTGAA	CACGACATCTATGCTGAA	0.064
<i>One_RH2op-395<sup>a</sup></i>	G	T	TGGGAACATCATTTTTTAA	TTGGGAACATAATTTTTTAA	0.016
<i>One_rpo2j-261<sup>c</sup></i>	G	T	CACATGTTTTACTCATTTGA	CACATGTTTTACTAATTTGA	0.312
<i>One_sast-211<sup>c</sup></i>	G	T	CATCATTTGCATTATTG	CATCATTTGAATTATTG	0.073

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Assay	VIC	FAM	VIC sequence	FAM sequence	H <sub>0</sub>
<i>One_serpin-75<sup>a</sup></i>	G	T	CAGTGTGTAATTTAATGATATAT	ACAGTGTGTAATTTAATTATATAT	0.039
<i>One_spf30-207<sup>c</sup></i>	G	T	AGGGACATCTTACCTCAAAA	AGGGACATCTTACCTAAAAA	0.295
<i>One_srp09-127<sup>c</sup></i>	T	A	CAGCGAAGGATATGCT	CAGCGAAGGTTATGCT	0.082
<i>One_ssrd-135<sup>c</sup></i>	-	T	CTGCGGCTTTTGTCTTG	TGCGGCTTTTGTCTTG	0.475
<i>One_STC-410<sup>a</sup></i>	T	C	CCGATGGGTATATTATTATA	CCGATGGGTATATTGTTATA	0.336
<i>One_STR07<sup>a</sup></i>	G	C	ACGCACACTGTCCTT	ACGCACACTCTCCTT	0.399
<i>One_SUMO1-6<sup>c</sup></i>	C	A	CAAGATTGAAATTGGTTTGC	CAAGATTGAAATTTGTTTGC	0.297
<i>One_sys1-230<sup>c</sup></i>	T	G	CAAAGCAAGTGATATATTAGTG	AAAGCAAGTGATATCTTAGTG	0.413
<i>One_taf12-248<sup>c</sup></i>	C	T	CCAGACAAAATCAAATTA	CCAGACAAAATAAAATTA	0.047
<i>One_Tf_ex11-750<sup>a</sup></i>	G	A	CAGGGTCGCTGCAC	CCAGGGTCACTGCAC	0.380
<i>One_Tf_in3-182<sup>a</sup></i>	A	G	AACAGAAAGTCTACACTTT	ACAGAAAGTCTGCACTTT	0.109
<i>One_tshB-92<sup>c</sup></i>	A	C	ACCACCCTGTAGCTCA	CACCCTGGAGCTCA	0.111
<i>One_txnip-401<sup>c</sup></i>	C	T	TGACTGCACTAGTTTAGAC	TGACTGCACTAATTTAGAC	0.047
<i>One_U1002-101<sup>b</sup></i>	G	T	TCGTTCCAAAGAATGTTGTG	CGTTCCAAAGAATTTTGTG	0.009
<i>One_U1003-75<sup>b</sup></i>	C	T	AGAGACTACTTCCTTTTTG	AGAGACTACTTCTTTTTTG	0.294
<i>One_U1004-183<sup>b</sup></i>	A	G	AAGTTCCTGTATTTCTT	TCCCTGCATTTCTT	0.345
<i>One_U1009-91<sup>b</sup></i>	A	G	CATGTTCTGTATGGACCC	TGTTCTGTGTGGACCC	0.283
<i>One_U1010-81<sup>b</sup></i>	A	G	CACACCAACGTTATGTAGAG	CACCAACGTTGTGTAGAG	0.064
<i>One_U1012-68<sup>b</sup></i>	C	T	TGACGGGTGTTCTTGATAA	TGACGGGTGTTCTTGATAA	0.251
<i>One_U1013-108<sup>b</sup></i>	G	T	ACGGAATTCCTGTTGCCCT	ACGGAATTCCTTTTGCCCT	0.250
<i>One_U1014-74<sup>b</sup></i>	C	T	TTGACCTGCGCCAGTAT	TTTTGACCTGCACCAGTAT	0.212
<i>One_U1016-115<sup>b</sup></i>	-	T	AATGGCAGTTTTTTATTGA	ATGGCAGTTTTTTTATTGA	0.402

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Assay	VIC	FAM	VIC sequence	FAM sequence	H <sub>0</sub>
<i>One_U1017-62<sup>b</sup></i>	A	T	CAGAAAAACTGGTACTTGTT	CAGAAAAACTGGTTCTTGTT	0.031
<i>One_U1021-57<sup>b</sup></i>	A	G	AGTTGAACGTTTGGTTTGA	GTTGAACGTTTCGGTTTGA	0.414
<i>One_U1024-197<sup>b</sup></i>	G	T	ACCTGACCCAACAAA	ACCTGACACAACAAA	0.199
<i>One_U1101<sup>b</sup></i>	C	A	TGGACGTATGTCATATTT	TGGACGTATGTAATATTT	0.303
<i>One_U1102-220<sup>b</sup></i>	C	T	CCAGTAGTGTTTTCTG	CAGTAGTGCTTTCTG	0.168
<i>One_U1103<sup>b</sup></i>	G	A	TCGGCGAAAACT	TCGGCAAAAACT	0.050
<i>One_U1104-138<sup>b</sup></i>	G	T	CCTTCTCAGAGGGTAGAGA	CCTTCTCAGAGGTTAGAGA	0.009
<i>One_U1105<sup>b</sup></i>	T	A	CCTGTTTTTTTTTAAAGAC	TCCTGTTTTTTTTTAAAGAC	0.332
<i>One_U1201-492<sup>b</sup></i>	A	G	AAGACTTCCTCCAGGCTC	ACTTCCCCCAGGCTC	0.445
<i>One_U1202-1052<sup>b</sup></i>	T	C	CAAACCTTTTTCATCTACATTTA	ACTTTTTCATCCACATTTA	0.370
<i>One_U1203-175<sup>b</sup></i>	G	A	CCATAGTTGCTGGGCTT	CTCCATAGTTACTGGGCTT	0.397
<i>One_U1204-53<sup>b</sup></i>	C	T	ATGCATACACGCTGATGC	ATGCATACACACTGATGC	0.318
<i>One_U1205-57<sup>b</sup></i>	A	G	AGTTATCATGGTCATCTCT	AGTTATCATGGTCGTCTCT	0.049
<i>One_U1206-108<sup>b</sup></i>	G	T	AACATTGAGCTTCCC	ATAACATTGATCTTCCC	0.300
<i>One_U1207-231<sup>b</sup></i>	C	T	ACATTCCTTGGCATTGC	CATTTCCTTGACATTGC	N/A <sup>f</sup>
<i>One_U1208-67<sup>b</sup></i>	A	C	CCAATGTGATTGTCAC	CCAATGTGCTTGTCAC	0.398
<i>One_U1209-111<sup>b</sup></i>	C	T	CTCACATCGAGATGATC	TCACATCGAAATGATC	0.170
<i>One_U1210-173<sup>b</sup></i>	A	G	CCCTCCTATTCAATTATGATTGT	CCTCCTATTCAATTACGATTGT	0.149
<i>One_U1211-97<sup>b</sup></i>	C	T	CTGTTTCAGTGTGCTTG	CTGTTTCAGTATGCTTG	0.165
<i>One_U1212-106<sup>b</sup></i>	A	G	TTTTGACATACAAAAAATA	TTTGACATACAGAAAATA	0.445
<i>One_U1214-107<sup>b</sup></i>	A	C	TAGTGACCTATTAAATTGC	TGACCTATTCAATTGC	0.137
<i>One_U1215-82<sup>b</sup></i>	A	C	AATGAGACAAAGTATTTGGT	AATGAGACAAAGTCTTTGGT	0.469

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Assay	VIC	FAM	VIC sequence	FAM sequence	H <sub>0</sub>
<i>One_U1216-230<sup>b</sup></i>	A	T	CCTGGCTACTAAGTAAC	CTGGCTACAAAGTAAC	0.452
<i>One_U301-92<sup>a</sup></i>	T	G	CCATGGATTAAAATATTT	CCATGGATTAAACTATTT	0.230
<i>One_U401-224<sup>a</sup></i>	C	A	CACCTGGAAAGGACTGA	ACACCTGGAAATGACTGA	0.439
<i>One_U404-229<sup>a</sup></i>	C	T	CATGTTCTTCAGTGAACC	ATGTTCTTCAATGAACC	0.108
<i>One_U502-167<sup>a</sup></i>	A	G	CTTCTTGATCAATAACG	CTTCTTGATCGATAACG	0.034
<i>One_U503-170<sup>a</sup></i>	T	G	AAGTACTAAAATCAGTTTTACATTG	TACTAAAATCAGTTGTACATTG	0.231
<i>One_U504-141<sup>a</sup></i>	C	A	TCAAGGACACAAACAA	TCAAGGACAAAAACAA	0.362
<i>One_U508-533<sup>a</sup></i>	C	T	ACACTACAGCCTTATTC	ACACTACAGCTTTATTC	0.090
<i>One_UCA-24<sup>b</sup></i>	C	T	CGAACAGGGCTGGATG	CGAACAGGACTGGATG	N/A <sup>f</sup>
<i>One_vamp5-255<sup>c</sup></i>	C	T	TAGGCTCCGTGCTCAGT	TAGGCTCCGTACTCAGT	0.309
<i>One_vatf-214<sup>c</sup></i>	C	A	TGGTATTACTGTGCATTGAC	ATGGTATTACTGTTCAATTGAC	0.089
<i>One_VIM-569<sup>a</sup></i>	G	A	AAGTGTTTCCATACTCACTATA	AAGTGTTTCCATATTCCTATA	0.207
<i>One_zn706-68<sup>c</sup></i>	C	T	ATTAAGTGAAGGGAGCAGC	AAGTGAAGGAAGCAGC	0.002
<i>One_ZNF-61<sup>a</sup></i>	C	A	CTATGGACATGATCTTT	TTCTATGGACATTATCTTT	0.342
<i>One_Zp3b-49<sup>a</sup></i>	C	A	AGGCCCAATCCTT	AGGCCAAATCCTT	0.182

<sup>a</sup> Assay developed by the Gene Conservation Laboratory of the Alaska Department of Fish and Game (Elfstrom et al. 2006; Smith et al. 2005).

<sup>b</sup> Assay developed by the International Program for Salmon Ecological Genetics at the University of Washington.

<sup>c</sup> Assay developed by the Hagerman Genetics Laboratory of the Columbia River Inter-Tribal Fish Commission.

<sup>d</sup> Assay developed by the Molecular Genetics Laboratory at the Canadian Department of Fisheries and Oceans.

<sup>e</sup> These assays are mitochondrial SNPs and were not measured for heterozygosity.

<sup>f</sup> These assays failed in the laboratory and were not measured for heterozygosity.

Table 4.–Summary of the average, coefficient of variation (CV) and associated ranks for four measures of laboratory performance of 124 assays (Tables 2 and 3) screened for 36 test populations (Table 1). See footnotes for explanations of the lack of scores or rankings.

Assay	Cluster tightness				Space between clusters				Cluster alignment				Success rate			
	Average		CV		Average		CV		Average		CV		Average		CV	
	Score	Rank	Score	Rank	Score	Rank	Score	Rank	Score	Rank	Score	Rank	Score	Rank	Score	Rank
<i>One_ACBP-79</i>	3.28	90	0.14	23	3.67	55	0.23	51	4.56	58	0.11	41.5	97.69	79	0.02	73
<i>One_agt-132</i>	3.39	77.5	0.18	58	3.72	53	0.20	33	4.28	81	0.19	90	98.04	56	0.02	74
<i>One_aldB-152</i>	3.39	77.5	0.21	85.5	3.89	47	0.25	60	4.44	71	0.12	47.5	98.08	52	0.02	59
<i>One_ALDOB-135</i>	2.89	111.5	0.16	42	2.83	91.5	0.33	78	3.89	99	0.21	96	96.01	107	0.09	109
<i>One_apoe-83</i>	3.78	44.5	0.21	89	4.44	14.5	0.19	23.5	4.78	31	0.09	26	98.26	48	0.01	16
<i>One_bckB-137<sup>a</sup></i>	4.89	10	0.10	11.5	1.00	117.5	N/A	N/A	4.89	20	0.10	30	99.50	3	0.01	2
<i>One_c3-98</i>	3.89	37.5	0.20	68.5	3.56	63	0.34	80	4.17	87	0.22	101	99.25	9	0.01	12
<i>One_ccd16-131</i>	4.94	7.5	0.05	7.5	0.28	115.5	4.24	115.5	4.94	12	0.05	12	99.54	1	0.01	6
<i>One_CD9-269</i>	3.39	77.5	0.21	85.5	4.28	19.5	0.19	25	4.78	31	0.09	26	97.97	59	0.01	33
<i>One_cetn1-167</i>	3.61	56.5	0.17	43.5	3.56	63	0.24	56.5	4.00	94	0.19	89	97.84	70	0.02	76
<i>One_CFP1</i>	3.67	52	0.23	101	4.17	24	0.24	54	4.56	58	0.15	75.5	98.26	47	0.01	48
<i>One_cin-177</i>	3.56	61.5	0.20	72.5	3.28	78.5	0.29	77	3.83	101.5	0.22	102	97.37	92	0.02	64
<i>One_CO1</i>	5.00	3.5	0.00	3.5	5.00	1.5	0.00	1.5	4.94	12	0.05	12	99.13	15	0.01	24
<i>One_CTGF-301</i>	4.00	32.5	0.23	98	3.33	75	0.58	97	4.67	45	0.13	53	97.01	98	0.09	110
<i>One_Cytb_17</i>	5.00	3.5	0.00	3.5	5.00	1.5	0.00	1.5	5.00	4.5	0.00	4.5	99.54	2	0.01	1
<i>One_Cytb_26</i>	5.00	3.5	0.00	3.5	4.94	3	0.05	3	5.00	4.5	0.00	4.5	99.10	16	0.01	15
<i>One_dds-529</i>	2.50	119	0.25	111	2.00	105.5	0.34	82	2.11	118.5	0.32	110	97.62	83	0.01	29
<i>One_DDX5-86</i>	3.33	84.5	0.25	112	2.78	95	0.40	86	3.67	104	0.25	106	96.91	101	0.02	90
<i>One_E2-65</i>	3.28	90	0.14	23	3.50	67.5	0.20	34	4.44	71	0.14	69	95.23	112	0.11	114
<i>One_gadd45-269</i>	5.00	3.5	0.00	3.5	1.00	118.5	N/A	115.5	4.89	20	0.10	30	99.20	13	0.01	14
<i>One_gdh-212</i>	3.50	67.5	0.18	52	4.00	35.5	0.27	69.5	4.61	51	0.13	56.5	97.81	72	0.01	20
<i>One_GHII-2165</i>	3.72	47.5	0.24	107	4.56	8	0.17	15	4.78	31	0.09	26	98.62	34	0.02	58
<i>One_ghsR-66</i>	3.39	77.5	0.23	104	4.00	35.5	0.27	69.5	4.56	58	0.14	62	97.94	63	0.01	21

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Table 4. Page 2 of 6.

Assay	Cluster tightness				Space between clusters				Cluster alignment				Success rate			
	Average		CV		Average		CV		Average		CV		Average		CV	
	Score	Rank	Score	Rank	Score	Rank	Score	Rank	Score	Rank	Score	Rank	Score	Rank	Score	Rank
<i>One_GPDH-201</i>	3.72	47.5	0.22	93	3.94	40.5	0.24	55	4.44	71	0.14	69	98.41	43	0.01	46
<i>One_GPDH2-187</i>	3.11	99.5	0.24	109	2.78	95	0.42	89	4.11	88.5	0.20	93	98.74	32	0.01	8
<i>One_GPH-414</i>	3.22	94	0.13	19	3.89	47	0.20	27	4.39	75.5	0.16	79	97.95	61	0.01	38
<i>One_GTHa</i>	3.61	56.5	0.22	91	4.17	24	0.17	13	4.72	38.5	0.10	34	98.06	54	0.01	41
<i>One_HGFA-49</i>	2.94	109	0.14	26	3.28	78.5	0.23	52	4.56	58	0.14	62	97.60	86	0.05	100
<i>One_HpaI-71</i>	2.89	112.5	0.11	14	2.89	90	0.20	31.5	3.83	101.5	0.16	80	96.11	106	0.04	96
<i>One_HpaI-99</i>	3.39	77.5	0.21	85.5	3.56	63	0.20	29	3.94	98	0.22	100	95.71	111	0.10	111
<i>One_hsc71-220</i>	3.61	56.5	0.17	43.5	2.33	103	0.36	83	3.28	111	0.34	112	98.24	49	0.02	52
<i>One_Hsp47</i>	3.83	41.5	0.22	95.5	4.06	32	0.25	59	4.33	78	0.19	91	98.46	41	0.01	42
<i>One_Ig-90</i>	4.50	16.5	0.21	82	1.78	107	1.23	111	4.67	45	0.15	72	99.34	7	0.01	7
<i>One_IL8r-362</i>	3.83	41.5	0.22	95.5	4.44	14.5	0.18	17	4.72	38.5	0.10	34	99.02	18	0.01	27
<i>One_ins-107</i>	3.00	105	0.16	41	3.28	78.5	0.27	71.5	4.00	94	0.17	82	95.12	113	0.06	104
<i>One_KCT1-453</i>	3.89	37.5	0.15	31	3.89	47	0.21	42.5	4.28	81	0.22	103	98.48	40	0.02	71
<i>One_KPNA-422</i>	3.22	94	0.17	45	3.94	40.5	0.20	35	4.67	45	0.13	53	97.96	60	0.02	57
<i>One_LEI-87</i>	3.89	37.5	0.20	68.5	3.61	58	0.32	78	4.50	64.5	0.17	84	98.29	45	0.02	53
<i>One_leptin-92<sup>a</sup></i>	5.00	3.5	0.00	3.5	1.00	117.5	N/A	N/A	5.00	4.5	0.00	4.5	99.49	4	0.01	9
<i>One_lpp1-44</i>	3.67	52	0.23	101	4.56	8	0.15	9	4.83	26	0.08	22.5	98.01	57	0.01	32
<i>One_MARCKS-241</i>	3.22	94	0.27	115	2.50	101	0.48	92	3.00	113	0.36	114	95.77	110	0.12	115
<i>One_metA-253</i>	4.56	14.5	0.17	47	3.17	85	0.74	105	4.94	12	0.05	12	98.84	23	0.01	44
<i>One_MHC2_190</i>	2.78	116.5	0.20	70	2.78	95	0.36	84	3.56	106	0.22	99	94.95	115	0.07	106
<i>One_MHC2_251</i>	3.94	34.5	0.18	59.5	3.89	47	0.21	42.5	4.22	84.5	0.19	87.5	97.93	64	0.02	85
<i>One_Mkpro-129</i>	3.33	84.5	0.23	105	3.67	55	0.26	66	4.50	64.5	0.14	65	96.75	102	0.04	99
<i>One_ODC1-196</i>	3.06	103	0.08	10	3.22	83	0.23	49	4.06	90.5	0.13	59.5	97.89	67	0.02	79
<i>One_Ots208-234<sup>b</sup></i>	N/A	61.5	N/A	60.5	N/A	63	N/A	58.5	N/A	58	N/A	60.75	N/A	60.5	N/A	60.5

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Table 4. Page 3 of 6.

Assay	Cluster tightness				Space between clusters				Cluster alignment				Success rate			
	Average		CV		Average		CV		Average		CV		Average		CV	
	Score	Rank	Score	Rank	Score	Rank	Score	Rank	Score	Rank	Score	Rank	Score	Rank	Score	Rank
<i>One_Ots213-181</i>	3.50	67.5	0.18	52	3.89	47	0.21	42.5	4.61	51	0.13	56.5	96.35	104	0.08	107
<i>One_p53-534</i>	4.22	25.5	0.19	64	2.83	92.5	0.84	107	4.89	20	0.07	18.5	95.99	108	0.10	113
<i>One_parp3-170<sup>a</sup></i>	5.00	3.5	0.00	3.5	1.00	117.5	N/A	N/A	5.00	4.5	0.00	4.5	98.82	25	0.02	66
<i>One_pax7-248</i>	4.22	25.5	0.17	48.5	4.50	11	0.27	67	4.89	20	0.07	18.5	98.74	31	0.02	65
<i>One_PIP</i>	3.11	99.5	0.15	33	3.50	67.5	0.18	18	4.06	90.5	0.13	59.5	97.04	96	0.03	91
<i>One_ppie-74</i>	4.06	30.5	0.27	116	1.22	111	1.21	110	3.61	105	0.40	117	93.63	117	0.25	118
<i>One_PPM1K-118<sup>c</sup></i>	1.72	N/A	0.65	N/A	1.33	N/A	0.68	N/A	2.11	N/A	0.58	N/A	5.44	N/A	4.24	N/A
<i>One_Prl2</i>	3.11	99.5	0.15	33	3.56	63	0.24	56.5	4.56	58	0.14	62	96.93	100	0.02	82
<i>One_psme2-354</i>	2.94	109	0.27	114	2.56	99.5	0.52	95	3.28	111	0.33	111	97.01	97	0.04	97
<i>One_rab1a-76</i>	3.28	90	0.23	103	4.06	32	0.20	28	4.44	71	0.12	47.5	97.62	84	0.02	51
<i>One_RAG1-103</i>	4.33	22	0.18	54	3.94	40.5	0.47	91	4.89	20	0.07	18.5	98.08	53	0.04	98
<i>One_RAG3-93</i>	3.50	67.5	0.22	97	3.39	73	0.34	81	4.00	94	0.24	105	96.13	105	0.10	112
<i>One_redd1-414</i>	3.33	84.5	0.18	55.5	2.89	90	0.26	65	3.28	111	0.31	109	97.75	75	0.02	61
<i>One_RFC2-102</i>	4.00	32.5	0.17	46	4.11	28	0.22	45.5	4.50	64.5	0.16	77	98.61	35	0.01	28
<i>One_RFC2-285</i>	4.39	20	0.16	40	3.94	40.5	0.40	87	4.61	51	0.15	73	99.36	6	0.01	10
<i>One_RH2op-395</i>	4.50	16.5	0.16	38	3.11	86	0.74	104	4.78	31	0.11	44.5	97.81	71	0.06	103
<i>One_rpo2j-261</i>	3.56	61.5	0.20	72.5	4.11	28	0.16	12	4.72	38.5	0.10	34	97.61	85	0.01	36
<i>One_sast-211</i>	3.56	61.5	0.28	117	4.11	28	0.29	76	4.72	38.5	0.12	50.5	98.87	22	0.01	25
<i>One_serpin-75</i>	2.83	114.5	0.39	119	1.72	108	0.77	106	2.50	116.5	0.42	118	85.94	120	0.31	120
<i>One_spf30-207</i>	3.56	61.5	0.20	72.5	4.22	21.5	0.19	22	4.61	51	0.11	39.5	98.52	39	0.01	40
<i>One_srp09-127</i>	4.22	25.5	0.17	48.5	3.67	55	0.56	96	4.94	12	0.05	12	98.91	21	0.01	49
<i>One_ssrd-135</i>	3.44	72	0.20	80.5	4.11	28	0.18	21	4.61	51	0.11	39.5	97.56	88	0.02	78
<i>One_STC-410</i>	2.94	109	0.25	110	1.67	109	0.41	88	2.61	115	0.37	115	90.47	119	0.13	116
<i>One_STR07</i>	2.78	116.5	0.15	36	3.28	78.5	0.27	71.5	3.94	98	0.16	81	97.20	93	0.02	80

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Table 4. Page 4 of 6.

Assay	Cluster tightness				Space between clusters				Cluster alignment				Success rate			
	Average		CV		Average		CV		Average		CV		Average		CV	
	Score	Rank	Score	Rank	Score	Rank	Score	Rank	Score	Rank	Score	Rank	Score	Rank	Score	Rank
<i>One_SUMO1-6</i>	3.17	96	0.22	94	3.61	58	0.25	62	4.56	58	0.11	41.5	97.64	82	0.02	70
<i>One_sys1-230</i>	3.39	77.5	0.21	85.5	3.83	51.5	0.22	48	4.67	45	0.10	37	97.89	66	0.01	35
<i>One_taf12-248</i>	4.28	23	0.18	50	2.22	104	1.05	109	4.83	26	0.08	22.5	99.09	17	0.01	13
<i>One_Tf_ex11-750</i>	3.39	77.5	0.15	29.5	4.00	35.5	0.21	40	4.44	71	0.12	47.5	95.89	109	0.07	105
<i>One_Tf_in3-182</i>	4.22	25.5	0.15	35	4.44	14.5	0.28	74	4.83	26	0.11	38	99.15	14	0.01	19
<i>One_tshB-92</i>	3.67	52	0.19	63	4.06	32	0.18	19	4.44	71	0.14	69	98.77	27	0.01	11
<i>One_txnlp-401</i>	4.72	13	0.12	18	1.61	110	1.46	112	5.00	4.5	0.00	4.5	99.40	5	0.01	5
<i>One_U1002-101</i>	4.89	10	0.07	9	0.83	113	2.30	113	5.00	4.5	0.00	4.5	99.28	8	0.01	23
<i>One_U1003-75</i>	3.33	84.5	0.18	55.5	3.94	40.5	0.18	20	4.44	71	0.12	47.5	97.92	65	0.02	83
<i>One_U1004-183</i>	3.50	67.5	0.15	28	3.56	63	0.28	73	4.28	81	0.18	85	98.20	50	0.02	63
<i>One_U1009-91</i>	3.72	47.5	0.15	37	4.67	5	0.13	6	4.94	12	0.05	12	98.28	46	0.02	55
<i>One_U1010-81</i>	4.56	14.5	0.11	15	4.33	18	0.37	85	4.94	12	0.05	12	98.83	24	0.01	37
<i>One_U1012-68</i>	4.06	30.5	0.16	39	4.50	11	0.11	5	4.78	31	0.09	26	98.74	33	0.01	39
<i>One_U1013-108</i>	3.67	52	0.23	101	4.00	35.5	0.26	63	4.72	38.5	0.10	34	97.88	68	0.02	67
<i>One_U1014-74</i>	3.78	44.5	0.19	67	4.39	17	0.19	26	4.78	31	0.11	44.5	98.04	55	0.02	69
<i>One_U1016-115</i>	3.11	99.5	0.10	13	2.00	105.5	0.17	14	2.11	118.5	0.15	74	97.86	69	0.01	45
<i>One_U1017-62</i>	4.44	18	0.19	65	3.22	83	0.73	103	5.00	4.5	0.00	4.5	98.75	29	0.02	54
<i>One_U1021-57</i>	2.56	117	0.24	107	2.44	101	0.25	60	3.33	108	0.21	93	97.17	94	0.02	75
<i>One_U1024-197</i>	3.44	72	0.20	80.5	3.89	47	0.17	16	4.22	84.5	0.19	87.5	98.57	36	0.01	34
<i>One_U1101</i>	3.11	99.5	0.22	92	3.89	47	0.21	42.5	4.50	64.5	0.11	43	97.94	62	0.02	88
<i>One_U1102-220</i>	3.00	105	0.23	99	2.56	99.5	0.63	101	2.50	116.5	0.68	120	98.46	42	0.01	18
<i>One_U1103</i>	4.39	20	0.11	16.5	3.22	83	0.66	102	4.72	38.5	0.12	50.5	98.97	19	0.01	4
<i>One_U1104-138</i>	4.78	12	0.14	21	0.78	114	2.31	114	4.89	20	0.10	30	99.23	10	0.01	17
<i>One_U1105</i>	3.11	99.5	0.15	33	3.28	78.5	0.20	36	3.94	98	0.18	86	97.74	77	0.02	56

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Table 4. Page 5 of 6.

Assay	Cluster tightness				Space between clusters				Cluster alignment				Success rate			
	Average		CV		Average		CV		Average		CV		Average		CV	
	Score	Rank	Score	Rank	Score	Rank	Score	Rank	Score	Rank	Score	Rank	Score	Rank	Score	Rank
<i>One_U1201-492</i>	3.33	84.5	0.21	83	3.44	70	0.27	68	4.22	84.5	0.17	83	97.67	80	0.02	81
<i>One_U1202-1052</i>	2.83	114.5	0.14	20	2.89	90	0.20	31.5	3.50	107	0.26	108	97.55	89	0.02	86
<i>One_U1203-175</i>	3.44	72	0.18	57	4.17	24	0.21	37	4.89	20	0.07	18.5	98.36	44	0.02	68
<i>One_U1204-53</i>	3.50	67.5	0.20	77	4.22	21.5	0.21	39	4.78	31	0.09	26	97.77	74	0.02	84
<i>One_U1205-57</i>	4.11	28.5	0.23	106	3.28	78.5	0.59	99	4.33	78	0.21	96	98.94	20	0.01	22
<i>One_U1206-108</i>	2.94	109	0.14	26	3.39	73	0.21	38	3.78	103	0.25	107	92.54	118	0.25	119
<i>One_U1207-231<sup>c</sup></i>	1.56	N/A	0.63	N/A	1.72	N/A	0.74	N/A	2.67	N/A	0.66	N/A	5.32	N/A	4.24	N/A
<i>One_U1208-67</i>	3.67	52	0.21	88	4.56	8	0.14	8	4.89	20	0.07	18.5	97.38	91	0.02	62
<i>One_U1209-111</i>	3.83	41.5	0.18	61.5	4.72	4	0.10	4	4.72	38.5	0.10	34	98.77	26	0.01	43
<i>One_U1210-173</i>	3.56	61.5	0.20	72.5	3.61	58	0.24	53	4.00	94	0.23	104	98.76	28	0.01	31
<i>One_U1211-97</i>	3.33	84.5	0.29	118	2.67	98	0.50	93	3.44	108	0.35	113	97.74	76	0.02	89
<i>One_U1212-106</i>	3.28	90	0.14	23	3.56	63	0.22	47	4.39	75.5	0.14	67	96.97	99	0.03	94
<i>One_U1214-107</i>	3.61	56.5	0.19	66	4.28	19.5	0.29	75	4.50	64.5	0.14	65	98.56	38	0.01	47
<i>One_U1215-82</i>	2.11	120	0.46	120	1.17	112	0.61	100	1.94	120	0.54	119	95.12	114	0.03	92
<i>One_U1216-230</i>	3.28	90	0.20	79	4.11	28	0.22	45.5	4.67	45	0.13	53	97.44	90	0.02	87
<i>One_U301-92</i>	3.94	34.5	0.18	59.5	4.44	14.5	0.19	23.5	4.72	38.5	0.14	71	96.64	103	0.08	108
<i>One_U401-224</i>	3.39	77.5	0.15	29.5	3.83	51.5	0.24	58	4.50	64.5	0.14	65	97.57	87	0.01	50
<i>One_U404-229</i>	4.11	28.5	0.20	78	3.44	70	0.51	94	4.56	58	0.15	75.5	98.75	30	0.02	77
<i>One_U502-167</i>	4.94	7.5	0.05	7.5	2.72	97	0.92	108	4.94	12	0.05	12	99.21	11	0.01	30
<i>One_U503-170</i>	4.39	20	0.11	16.5	4.61	6	0.13	7	4.61	51	0.13	56.5	97.73	78	0.05	101
<i>One_U504-141</i>	2.94	109	0.14	26	3.44	70	0.23	50	4.33	78	0.16	78	97.66	81	0.02	60
<i>One_U508-533<sup>d</sup></i>	3.00	105	0.2	75	3.00	87	0.2	30	4.00	94	0.2	92	93.7	116	0.2	117
<i>One_UCA-24<sup>c</sup></i>	3.39	N/A	0.27	N/A	3.28	N/A	0.29	N/A	3.06	N/A	0.38	N/A	67.08	N/A	0.18	N/A
<i>One_vamp5-255</i>	3.50	67.5	0.18	52	3.56	63	0.26	64	4.11	88.5	0.22	98	97.79	73	0.02	72

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Table 4. Page 6 of 6.

Assay	Cluster tightness				Space between clusters				Cluster alignment				Success rate			
	Average		CV		Average		CV		Average		CV		Average		CV	
	Score	Rank	Score	Rank	Score	Rank	Score	Rank	Score	Rank	Score	Rank	Score	Rank	Score	Rank
<i>One_vatf-214</i>	3.89	37.5	0.21	90	3.39	73	0.58	98	4.61	51	0.13	56.5	99.21	12	0.01	3
<i>One_VIM-569</i>	3.83	41.5	0.18	61.5	3.94	40.5	0.16	11	4.22	84.5	0.21	95	98.14	51	0.03	93
<i>One_zn706-68</i>	4.89	10	0.10	11.5	0.28	115.5	4.24	115.5	5.00	4.5	0.00	4.5	98.00	58	0.03	95
<i>One_ZNF-61</i>	3.72	47.5	0.20	76	4.50	11	0.16	10	4.89	20	0.07	18.5	97.16	95	0.05	102
<i>One_Zp3b-49</i>	3.56	61.5	0.26	113	2.94	88	0.46	90	2.78	114	0.40	116	98.56	37	0.01	26

<sup>a</sup> This marker was excluded from subsequent analyses due to fixation in 36 test populations.

<sup>b</sup> This marker was screened on the ABI platform for the 36 test populations and was given median rankings for each laboratory measure.

<sup>c</sup> These markers were not given ratings or included in subsequent analyses due to massive laboratory failure or the appearance of excessive clusters.

<sup>d</sup> This marker was screened on the ABI platform for the 36 test populations and was given ratings based upon previous analyses of other populations on the Biomark platform.

Table 5.–The number of populations that failed to conform to Hardy-Weinberg expectations for 124 SNP assays (Tables 2 and 3) screened for 36 test populations (Table 1) at three levels of significance ( $\alpha = 0.001$ , 0.01, and 0.05), and rankings based upon the number of populations at  $\alpha = 0.05$ . *This table is commented on in the section entitled “Technical Committee Review and Comments.”*

Assay	$\alpha = 0.001$	$\alpha = 0.01$	$\alpha = 0.05$	
	Number	Number	Number	Rank
<i>One_ACBP-79</i>	0	1	1	60.5
<i>One_agt-132</i>	0	2	3	109.5
<i>One_aldB-152</i>	0	0	1	60.5
<i>One_ALDOB-135</i>	0	1	1	60.5
<i>One_apoe-83</i>	1	1	1	60.5
<i>One_bckB-137</i>	0	0	0	21
<i>One_c3-98</i>	1	1	2	92.5
<i>One_ccd16-131</i>	0	0	0	21
<i>One_CD9-269</i>	0	0	2	92.5
<i>One_cetn1-167</i>	0	0	3	109.5
<i>One_CFP1</i>	0	1	2	92.5
<i>One_cin-177</i>	0	1	2	92.5
<i>One_COI<sup>a</sup></i>	N/A	N/A	N/A	21
<i>One_CTGF-301</i>	0	0	0	21
<i>One_Cytb_17<sup>a</sup></i>	N/A	N/A	N/A	21
<i>One_Cytb_26<sup>a</sup></i>	N/A	N/A	N/A	21
<i>One_dds-529</i>	0	0	2	92.5
<i>One_DDX5-86</i>	1	1	4	114.5
<i>One_E2-65</i>	0	0	0	21
<i>One_gadd45-269</i>	0	0	0	21
<i>One_gdh-212</i>	0	0	1	60.5
<i>One_GHII-2165</i>	1	1	3	109.5
<i>One_ghsR-66</i>	0	0	0	21
<i>One_GPDH-201</i>	0	0	0	21
<i>One_GPDH2-187</i>	0	0	1	60.5
<i>One_GPH-414</i>	0	0	1	60.5
<i>One_GTHa</i>	0	0	1	60.5
<i>One_HGFA-49</i>	0	0	1	60.5
<i>One_HpaI-71</i>	0	0	0	21
<i>One_HpaI-99</i>	0	1	1	60.5
<i>One_hsc71-220</i>	0	0	1	60.5
<i>One_Hsp47</i>	0	0	0	21
<i>One_Ig-90</i>	1	1	1	60.5

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Table 5. Page 2 of 4.

Assay	$\alpha = 0.001$	$\alpha = 0.01$	$\alpha = 0.05$	
	Number	Number	Number	Rank
<i>One_IL8r-362</i>	0	0	0	21
<i>One_ins-107</i>	0	2	5	116
<i>One_KCT1-453</i>	0	0	2	92.5
<i>One_KPNA-422</i>	0	0	3	109.5
<i>One_LEI-87</i>	0	0	2	92.5
<i>One_leptin-92</i>	0	0	0	21
<i>One_lpp1-44</i>	0	0	1	60.5
<i>One_MARCKS-241</i>	0	0	0	21
<i>One_metA-253</i>	0	0	0	21
<i>One_MHC2_190</i>	0	1	2	92.5
<i>One_MHC2_251</i>	0	0	2	92.5
<i>One_Mkpro-129</i>	0	1	3	109.5
<i>One_ODC1-196</i>	0	0	0	21
<i>One_Ots208-234</i>	0	0	2	92.5
<i>One_Ots213-181</i>	0	0	0	21
<i>One_p53-534</i>	0	0	0	21
<i>One_parp3-170<sup>b</sup></i>	N/A	N/A	N/A	N/A
<i>One_pax7-248</i>	0	0	0	21
<i>One_PIP</i>	0	0	1	60.5
<i>One_ppie-74</i>	0	0	0	21
<i>One_PPM1K-118<sup>b</sup></i>	N/A	N/A	N/A	N/A
<i>One_Prl2</i>	0	2	2	92.5
<i>One_psme2-354</i>	0	0	2	92.5
<i>One_rab1a-76</i>	0	0	2	92.5
<i>One_RAG1-103</i>	0	0	0	21
<i>One_RAG3-93</i>	0	0	0	21
<i>One_redd1-414</i>	0	0	0	21
<i>One_RFC2-102</i>	0	0	0	21
<i>One_RFC2-285</i>	0	0	0	21
<i>One_RH2op-395</i>	0	0	1	60.5
<i>One_rpo2j-261</i>	0	0	2	92.5
<i>One_sast-211</i>	0	0	0	21
<i>One_serpin-75</i>	0	0	0	21
<i>One_spf30-207</i>	0	0	0	21
<i>One_srp09-127</i>	0	0	1	60.5
<i>One_ssrd-135</i>	0	0	1	60.5

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Table 5. Page 3 of 4.

Assay	$\alpha = 0.001$	$\alpha = 0.01$	$\alpha = 0.05$	
	Number	Number	Number	Rank
<i>One_STC-410</i>	0	1	2	92.5
<i>One_STR07</i>	0	0	1	60.5
<i>One_SUMO1-6</i>	0	0	1	92.5
<i>One_sys1-230</i>	0	1	2	92.5
<i>One_taf12-248</i>	0	0	2	60.5
<i>One_Tf_ex11-750</i>	0	0	0	21
<i>One_Tf_in3-182</i>	0	0	0	21
<i>One_tshB-92</i>	0	0	2	92.5
<i>One_txnip-401</i>	1	1	1	60.5
<i>One_U1002-101</i>	0	0	0	21
<i>One_U1003-75</i>	0	0	0	21
<i>One_U1004-183</i>	0	0	4	114.5
<i>One_U1009-91</i>	0	0	0	21
<i>One_U1010-81</i>	0	0	0	21
<i>One_U1012-68</i>	0	0	1	60.5
<i>One_U1013-108</i>	1	1	2	92.5
<i>One_U1014-74</i>	1	1	1	60.5
<i>One_U1016-115</i>	0	0	1	60.5
<i>One_U1017-62</i>	0	0	0	21
<i>One_U1021-57</i>	8	12	14	117
<i>One_U1024-197</i>	0	0	2	92.5
<i>One_U1101</i>	0	0	2	92.5
<i>One_U1102-220</i>	2	2	3	109.5
<i>One_U1103</i>	0	0	0	21
<i>One_U1104-138</i>	0	0	0	21
<i>One_U1105</i>	1	1	1	60.5
<i>One_U1201-492</i>	0	0	2	92.5
<i>One_U1202-1052</i>	1	1	1	60.5
<i>One_U1203-175</i>	1	1	1	60.5
<i>One_U1204-53</i>	0	0	2	92.5
<i>One_U1205-57</i>	0	0	2	92.5
<i>One_U1206-108</i>	0	0	1	60.5
<i>One_U1207-231<sup>b</sup></i>	N/A	N/A	N/A	N/A
<i>One_U1208-67</i>	0	0	2	92.5
<i>One_U1209-111</i>	0	0	0	21
<i>One_U1210-173</i>	0	1	1	60.5
<i>One_U1211-97</i>	0	0	1	60.5

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Table 5. Page 4 of 4.

Assay	$\alpha = 0.001$	$\alpha = 0.01$	$\alpha = 0.05$	
	Number	Number	Number	Rank
<i>One_U1212-106</i>	0	1	1	60.5
<i>One_U1214-107</i>	0	0	2	92.5
<i>One_U1215-82</i>	0	0	1	60.5
<i>One_U1216-230</i>	0	0	3	109.5
<i>One_U301-92</i>	0	1	1	60.5
<i>One_U401-224</i>	0	0	1	60.5
<i>One_U404-229</i>	0	1	1	60.5
<i>One_U502-167</i>	0	0	0	21
<i>One_U503-170</i>	0	0	0	21
<i>One_U504-141</i>	0	0	1	60.5
<i>One_U508-533</i>	0	0	1	60.5
<i>One_UCA-24<sup>b</sup></i>	N/A	N/A	N/A	N/A
<i>One_vamp5-255</i>	0	0	1	60.5
<i>One_vatf-214</i>	0	0	0	21
<i>One_VIM-569</i>	0	0	1	60.5
<i>One_zn706-68</i>	0	0	0	21
<i>One_ZNF-61</i>	0	0	3	109.5
<i>One_Zp3b-49</i>	0	0	0	21

<sup>a</sup> These mitochondrial assays were not included in tests of conformance to Hardy-Weinberg expectations, and were given rankings equal to the highest rank for this judge.

<sup>b</sup> These assays were not included in tests of conformance to Hardy-Weinberg expectations due to poor laboratory performance.

Table 6.—Percent of total test populations of sockeye salmon (Table 1) exhibiting significant ( $\alpha = 0.001, 0.01$ , and  $0.05$ ) gametic disequilibrium for the pairs of loci for which disequilibrium was most commonly observed.

Pair of loci		$\alpha = 0.001$		$\alpha = 0.01$		$\alpha = 0.05$	
		Number of populations	Percentage of total	Number of populations	Percentage of total	Number of populations	Percentage of total
<i>One_aldB-152</i>	<i>One_ALDOB-135</i>	34	94%	34	94%	36	100%
<i>One_GPH-414</i>	<i>One_GTHa</i>	35	97%	36	100%	36	100%
<i>One_MHC2_190</i>	<i>One_MHC2_251</i>	16	44%	23	64%	26	72%

Table 7.—The contribution (%) to the first (PC1) and second (PC2) principal components, the average contribution to the first 12 principal components that explained 80% of total variation, and associated rankings for 124 sockeye salmon SNPs (Tables 2 and 3) screened for 36 test populations (Table 1).

Assay	PC1		PC2		Average contribution	
	%	Rank	%	Rank	%	Rank
<i>One_ACBP-79</i>	0.8	48	1.5	25.5	0.9	21
<i>One_agt-132</i>	0.5	72	1.3	29	0.7	49.5
<i>One_aldB-152</i>	0.6	62.5	0.2	91	0.5	89
<i>One_ALDOB-135<sup>a</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_apoe-83</i>	2.1	9	0.9	44.5	1.1	11
<i>One_bckB-137<sup>b</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_c3-98</i>	0.4	84	0.8	50.5	0.5	89
<i>One_ccd16-131</i>	0.2	100	0.1	101	0.2	109.5
<i>One_CD9-269</i>	0.2	100	0	109	0.3	107.5
<i>One_cetn1-167</i>	0.5	72	0	109	0.5	89
<i>One_CFP1</i>	1.5	20.5	2.4	5	1	14.5
<i>One_cin-177</i>	0.4	84	1.2	32	0.6	69
<i>One_COI<sup>c</sup></i>	2.3	7	1.8	17.5	1.8	2
<i>One_CTGF-301</i>	0.4	84	0.6	60.5	0.4	102
<i>One_Cytb_17c</i>	2.3	7	1.8	17.5	1.8	2
<i>One_Cytb_26c</i>	2.3	7	1.8	17.5	1.8	2
<i>One_dds-529</i>	0.7	55	1.1	35.5	0.6	69
<i>One_DDX5-86</i>	0	110	0.2	91	0.4	102
<i>One_E2-65</i>	0.2	100	0.9	44.5	0.5	89
<i>One_gadd45-269</i>	0	110	0	109	0.1	111.5
<i>One_gdh-212</i>	0.6	62.5	1.9	14	0.7	49.5
<i>One_GHII-2165</i>	2.9	3	1.2	32	1.4	3.5
<i>One_ghsR-66</i>	0.7	55	1.3	29	0.6	69
<i>One_GPDH-201</i>	1.4	26	0.6	60.5	0.7	49.5
<i>One_GPDH2-187</i>	1.8	12.5	0.1	101	0.9	21
<i>One_GPH-414<sup>a</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_GTHa</i>	1.1	35	0.8	50.5	0.7	49.5
<i>One_HGFA-49</i>	0.3	93.5	0.4	71	0.4	102
<i>One_HpaI-71</i>	0.8	48	1	39	0.9	21
<i>One_HpaI-99</i>	2.4	5.5	1.5	25.5	1.3	5.5
<i>One_hsc71-220</i>	0.7	55	0.3	81	0.7	49.5
<i>One_Hsp47</i>	0.4	84	2.1	9	0.7	49.5
<i>One_Ig-90</i>	1.6	16.5	0.9	44.5	0.8	33.5

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Table 7. Page 2 of 4.

Assay	PC1		PC2		Average contribution	
	%	Rank	%	Rank	%	Rank
<i>One_IL8r-362</i>	0.4	84	0.6	60.5	0.6	69
<i>One_ins-107</i>	0.7	55	0.6	60.5	0.5	89
<i>One_KCT1-453</i>	0.6	62.5	0.7	55	0.6	69
<i>One_KPNA-422</i>	0.3	93.5	0.8	50.5	0.4	102
<i>One_LEI-87</i>	1.3	30	0.3	81	0.9	21
<i>One_leptin-92<sup>b</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_lpp1-44</i>	0.7	55	0.4	71	0.8	33.5
<i>One_MARCKS-241</i>	0.4	84	0.6	60.5	0.4	102
<i>One_metA-253</i>	3.3	1	2.7	4	1.4	3.5
<i>One_MHC2_190<sup>d</sup></i>	3.1	2	3.7	1.5	3.9	1
<i>One_MHC2_251<sup>d</sup></i>	3.1	2	3.7	1.5	3.9	1
<i>One_Mkpro-129</i>	1.0	37	1.7	20.5	1	14.5
<i>One_ODC1-196</i>	0.3	93.5	1.8	17.5	0.6	69
<i>One_Ots208-234</i>	2.4	5.5	0.4	71	0.9	21
<i>One_Ots213-181</i>	1.6	16.5	0.2	91	0.7	49.5
<i>One_p53-534</i>	1.4	26	2.1	9	0.8	33.5
<i>One_parp3-170<sup>b</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_pax7-248</i>	0.7	55	0.1	101	0.6	69
<i>One_PIP</i>	0	110	0.7	55	0.5	89
<i>One_ppie-74</i>	1.4	26	3.4	3	1.3	5.5
<i>One_PPM1K-118<sup>e</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_Prl2</i>	0.1	105.5	2.0	11.5	0.7	49.5
<i>One_psme2-354</i>	0.4	84	0.3	81	0.6	69
<i>One_rab1a-76</i>	0.4	84	0.9	44.5	0.6	69
<i>One_RAG1-103</i>	0.1	105.5	1.5	25.5	0.7	49.5
<i>One_RAG3-93</i>	0.5	72	0.6	60.5	0.6	69
<i>One_redd1-414</i>	1.6	16.5	0.4	71	1	14.5
<i>One_RFC2-102</i>	0.5	72	0.5	64.5	0.5	89
<i>One_RFC2-285</i>	0.6	62.5	0.9	44.5	0.5	89
<i>One_RH2op-395</i>	0	110	0	109	0.2	109.5
<i>One_rpo2j-261</i>	0.4	84	1.8	17.5	0.6	69
<i>One_sast-211</i>	0.5	72	0.2	91	0.5	89
<i>One_serpin-75</i>	0.2	100	0	109	0.3	107.5
<i>One_spf30-207</i>	0.7	55	1.1	35.5	0.6	69
<i>One_srp09-127</i>	1.7	14	0.5	64.5	0.7	49.5

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Table 7. Page 3 of 4.

Assay	PC1		PC2		Average contribution	
	%	Rank	%	Rank	%	Rank
<i>One_ssr1-135</i>	0.6	62.5	1	39	0.6	69
<i>One_STC-410</i>	1.4	26	0.8	50.5	1.2	8
<i>One_STR07</i>	0.8	48	1.7	20.5	0.9	21
<i>One_SUMO1-6</i>	1	37	0.4	71	0.7	49.5
<i>One_sys1-230</i>	0	110	0.2	91	0.5	89
<i>One_taf12-248</i>	1.5	20.5	1.5	25.5	0.8	33.5
<i>One_Tf_ex11-750</i>	1.9	10.5	0.4	71	1.2	8
<i>One_Tf_in3-182</i>	0.2	100	1.3	29	0.9	21
<i>One_tshB-92</i>	0.2	100	0.3	81	0.4	102
<i>One_txn1p-401</i>	1.9	10.5	0.1	101	0.8	33.5
<i>One_U1002-101</i>	0.5	72	0.4	71	0.5	89
<i>One_U1003-75</i>	2.2	8	0.3	81	1.1	11
<i>One_U1004-183</i>	0.6	62.5	2.3	6	1.1	11
<i>One_U1009-91</i>	0.5	72	1.1	35.5	0.7	49.5
<i>One_U1010-81</i>	0.6	62.5	1.8	17.5	0.6	69
<i>One_U1012-68</i>	0.6	62.5	0.7	55	0.8	33.5
<i>One_U1013-108</i>	0.5	72	0.9	44.5	0.5	89
<i>One_U1014-74</i>	0.5	72	0.9	44.5	0.5	89
<i>One_U1016-115</i>	1.2	32.5	0.3	81	0.9	21
<i>One_U1017-62</i>	0.9	41.5	1	39	0.5	89
<i>One_U1021-57<sup>f</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_U1024-197</i>	1.2	32.5	0.9	44.5	0.8	33.5
<i>One_U1101</i>	0.2	100	2	11.5	0.6	69
<i>One_U1102-220</i>	1.5	20.5	0.4	71	0.8	33.5
<i>One_U1103</i>	0.9	41.5	0.2	91	0.7	49.5
<i>One_U1104-138</i>	0.3	93.5	0	109	0.4	102
<i>One_U1105</i>	1.5	20.5	0.7	55	0.8	33.5
<i>One_U1201-492</i>	0.3	93.5	1.6	22.5	0.6	69
<i>One_U1202-1052</i>	1.4	26	0.4	71	0.7	49.5
<i>One_U1203-175</i>	0.8	48	0.1	101	0.5	89
<i>One_U1204-53</i>	0.5	72	1.9	14	0.7	49.5
<i>One_U1205-57</i>	0.4	84	0.2	91	0.4	102
<i>One_U1206-108</i>	0.9	41.5	0.1	101	0.6	69
<i>One_U1207-231<sup>e</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_U1208-67</i>	1.4	26	0.3	81	0.8	33.5
<i>One_U1209-111</i>	0.8	48	0.7	55	0.8	33.5

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Table 7. Page 4 of 4.

Assay	PC1		PC2		Average contribution	
	%	Rank	%	Rank	%	Rank
<i>One_U1210-173</i>	0.9	41.5	0.2	91	0.6	69
<i>One_U1211-97</i>	1.2	32.5	0.3	81	0.6	69
<i>One_U1212-106</i>	0.4	84	2.2	7	0.6	69
<i>One_U1214-107</i>	1.4	26	3.7	1.5	1.2	8
<i>One_U1215-82</i>	0.3	93.5	0	109	0.4	102
<i>One_U1216-230</i>	0.5	72	0.1	101	0.5	89
<i>One_U301-92</i>	0.1	105.5	0.3	81	0.5	89
<i>One_U401-224</i>	1.6	16.5	0.2	91	0.8	33.5
<i>One_U404-229</i>	1.8	12.5	1.2	32	0.9	21
<i>One_U502-167</i>	0.9	41.5	1.9	14	0.8	33.5
<i>One_U503-170</i>	0.4	84	0.1	101	0.6	69
<i>One_U504-141</i>	0.4	84	2.1	9	0.7	49.5
<i>One_U508-533</i>	0.8	48	0.2	91	0.6	69
<i>One_UCA-24<sup>e</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_vamp5-255</i>	0.8	48	1.1	35.5	0.8	33.5
<i>One_vatf-214</i>	0.9	41.5	1.6	22.5	0.8	33.5
<i>One_VIM-569</i>	1	37	0.4	71	0.6	69
<i>One_zn706-68</i>	0.1	105.5	0.1	101	0.1	111.5
<i>One_ZNF-61</i>	1.2	32.5	0.2	91	0.8	33.5
<i>One_Zp3b-49</i>	2.6	4	0.4	71	1	14.5

<sup>a</sup> These assays were dropped due to significant linkage and were not included in this analysis.

<sup>b</sup> These assays were dropped due to fixation in the 36 test populations and were not included in this analysis.

<sup>c</sup> These assays are linked and were included as a haplotype marker in this analysis.

<sup>d</sup> These assays are linked and were included as a haplotype marker in this analysis.

<sup>e</sup> These assays were dropped due to laboratory failure and were not included in this analysis.

<sup>f</sup> This assay was dropped due to failure to conform to Hardy-Weinberg expectations and was not included in this analysis.



Table 8.— $F_{ST}$  values and associated rankings among populations within fine-scale regions ( $\theta_S$ ), among fine-scale regions within broad-scale regions ( $\theta_P$ ), and between populations within pairs of populations of interest ( $\theta_{Pairs}$ ) for 124 assays (Tables 2 and 3) screened for 36 test populations (Table 1) using the Weir and Cockerham method (1984).

Assay	$\theta_S$		$\theta_P$		$\theta_{Pairs}$	
	Statistic	Rank	Statistic	Rank	Statistic	Rank
<i>One_ACBP-79</i>	0.125	31	0.012	90	0.166	9
<i>One_agt-132</i>	0.112	43	0.016	81	0.049	78
<i>One_aldB-152</i>	0.090	64	0.018	76	0.065	56
<i>One_ALDOB-135<sup>a</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_apoe-83</i>	0.225	11	0.116	12	0.062	60
<i>One_bckB-137<sup>b</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_c3-98</i>	0.066	86	-0.014	111	0.037	86
<i>One_ccd16-131</i>	0.014	109	0.017	77	N/A <sup>g</sup>	112
<i>One_CD9-269</i>	0.048	99	0.078	24	0.061	63
<i>One_cetn1-167</i>	0.096	58	0.002	105	0.082	37
<i>One_CFP1</i>	0.152	18	0.013	88	0.091	30
<i>One_cin-177</i>	0.079	77	0.136	10	0.032	95
<i>One_CO1<sup>c</sup></i>	0.243	8	0.016	79	0.203	4
<i>One_CTGF-301</i>	0.036	108	0.020	69	0.026	99
<i>One_Cytb_17c</i>	0.243	8	0.016	79	0.203	4
<i>One_Cytb_26c</i>	0.243	8	0.016	79	0.203	4
<i>One_dds-529</i>	0.052	98	0.017	78	0.036	89
<i>One_DDX5-86</i>	0.058	94	0.006	96	0.071	49
<i>One_E2-65</i>	0.080	76	0.188	4	0.085	36
<i>One_gadd45-269</i>	0.000	112	0.087	19	0.000	111
<i>One_gdh-212</i>	0.081	72	0.024	63	0.019	101
<i>One_GHII-2165</i>	0.314	3	0.161	7	0.156	12
<i>One_ghsR-66</i>	0.120	36	0.034	50	0.074	48
<i>One_GPDH-201</i>	0.093	60	0.035	49	0.063	57
<i>One_GPDH2-187</i>	0.141	25	0.009	95	0.200	6
<i>One_GPH-414<sup>a</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_GTHa</i>	0.109	46	0.097	14	0.065	55
<i>One_HGFA-49</i>	0.038	106	0.054	40	0.026	98
<i>One_HpaI-71</i>	0.163	16	0.092	15	0.115	22
<i>One_HpaI-99</i>	0.277	5	-0.001	108	0.055	71
<i>One_hsc71-220</i>	0.104	51	0.079	23	0.122	19
<i>One_Hsp47</i>	0.111	44	0.005	98	0.126	18

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Table 8. Page 2 of 4.

Assay	$\theta_S$		$\theta_P$		$\theta_{\text{Pairs}}$	
	Statistic	Rank	Statistic	Rank	Statistic	Rank
<i>One_Ig-90</i>	0.124	32	0.030	57	0.081	38
<i>One_IL8r-362</i>	0.083	69	0.011	92	0.058	67
<i>One_ins-107</i>	0.052	97	0.018	75	0.031	96
<i>One_KCT1-453</i>	0.082	71	0.029	60	0.035	90
<i>One_KPNA-422</i>	0.054	96	0.020	68	0.048	80
<i>One_LEI-87</i>	0.146	21	0.088	17	0.079	42
<i>One_leptin-92<sup>b</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_lpp1-44</i>	0.127	30	0.021	67	0.087	34
<i>One_MARCKS-241</i>	0.036	107	0.237	2	0.008	108
<i>One_metA-253</i>	0.398	1	0.024	64	0.041	85
<i>One_MHC2_190<sup>d</sup></i>	0.255	7	0.073	26	0.146	14
<i>One_MHC2_251<sup>d</sup></i>	0.255	7	0.073	26	0.146	14
<i>One_Mkpro-129</i>	0.150	19	0.030	58	0.063	58
<i>One_ODC1-196</i>	0.100	54	0.023	65	0.093	28
<i>One_Ots208-234</i>	0.205	15	0.013	85	0.142	15
<i>One_Ots213-181</i>	0.104	49	0.031	54	0.077	44
<i>One_p53-534</i>	0.121	35	0.012	89	0.200	5
<i>One_parp3-170<sup>b</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_pax7-248</i>	0.078	79	0.152	9	0.004	110
<i>One_PIP</i>	0.058	93	0.083	21	0.057	68
<i>One_ppie-74</i>	0.292	4	0.050	41	0.037	87
<i>One_PPM1K-118<sup>e</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_Prl2</i>	0.097	57	0.020	70	0.074	47
<i>One_psme2-354</i>	0.110	45	-0.009	109	0.122	20
<i>One_rab1a-76</i>	0.081	73	0.002	104	0.052	75
<i>One_RAG1-103</i>	0.142	24	-0.023	112	0.177	8
<i>One_RAG3-93</i>	0.065	87	0.016	80	0.079	41
<i>One_redd1-414</i>	0.160	17	0.013	86	0.067	52
<i>One_RFC2-102</i>	0.081	74	0.022	66	0.061	64
<i>One_RFC2-285</i>	0.056	95	0.089	16	0.095	27
<i>One_RH2op-395</i>	0.012	110	0.063	31	0.008	107
<i>One_rpo2j-261</i>	0.089	65	0.055	38	0.107	23
<i>One_sast-211</i>	0.059	91	0.065	30	0.075	46
<i>One_serpin-75</i>	0.040	104	0.020	72	0.015	105

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Table 8. Page 3 of 4.

Assay	$\theta_s$		$\theta_p$		$\theta_{\text{Pairs}}$	
	Statistic	Rank	Statistic	Rank	Statistic	Rank
<i>One_spf30-207</i>	0.085	68	0.028	61	0.090	31
<i>One_srp09-127</i>	0.120	37	0.027	62	0.086	35
<i>One_ssr-135</i>	0.059	92	0.154	8	0.016	103
<i>One_STC-410</i>	0.233	9	0.046	43	0.196	7
<i>One_STR07</i>	0.135	27	0.082	22	0.135	17
<i>One_SUMO1-6</i>	0.097	56	0.060	32	0.034	93
<i>One_sys1-230</i>	0.094	59	0.032	53	0.056	70
<i>One_taf12-248</i>	0.109	47	0.088	18	0.121	21
<i>One_Tf_ex11-750</i>	0.233	10	0.039	47	0.251	3
<i>One_Tf_in3-182</i>	0.212	13	0.084	20	0.295	2
<i>One_tshB-92</i>	0.045	101	0.200	3	0.046	81
<i>One_txnip-401</i>	0.149	20	0.010	93	0.147	13
<i>One_U1002-101</i>	0.045	103	0.014	84	0.049	79
<i>One_U1003-75</i>	0.221	12	-0.009	110	0.061	62
<i>One_U1004-183</i>	0.323	2	0.068	29	0.490	1
<i>One_U1009-91</i>	0.090	63	0.055	39	0.096	26
<i>One_U1010-81</i>	0.063	88	0.036	48	0.052	77
<i>One_U1012-68</i>	0.107	48	0.032	52	0.097	25
<i>One_U1013-108</i>	0.076	82	0.068	28	0.032	94
<i>One_U1014-74</i>	0.061	90	0.019	73	0.034	92
<i>One_U1016-115</i>	0.142	23	0.059	35	0.135	16
<i>One_U1017-62</i>	0.062	89	0.020	71	0.009	106
<i>One_U1021-57<sup>d</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_U1024-197</i>	0.083	70	0.058	36	0.088	33
<i>One_U1101</i>	0.075	83	0.168	5	0.053	73
<i>One_U1102-220</i>	0.116	38	0.002	103	0.037	88
<i>One_U1103</i>	0.135	28	0.261	1	0.062	61
<i>One_U1104-138</i>	0.045	102	0.060	33	0.060	65
<i>One_U1105</i>	0.135	26	0.005	99	0.090	32
<i>One_U1201-492</i>	0.092	62	-0.001	107	0.046	82
<i>One_U1202-1052</i>	0.115	40	0.130	11	0.078	43
<i>One_U1203-175</i>	0.072	85	0.031	56	0.053	74
<i>One_U1204-53</i>	0.078	81	0.059	34	0.065	54
<i>One_U1205-57</i>	0.039	105	0.106	13	0.034	91
<i>One_U1206-108</i>	0.088	66	0.077	25	0.071	50

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Table 8. Page 4 of 4.

Assay	$\theta_S$		$\theta_P$		$\theta_{\text{Pairs}}$	
	Statistic	Rank	Statistic	Rank	Statistic	Rank
<i>One_U1207-231<sup>e</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_U1208-67</i>	0.122	33	0.000	106	0.044	83
<i>One_U1209-111</i>	0.112	42	0.029	59	0.070	51
<i>One_U1210-173</i>	0.074	84	0.002	102	0.052	76
<i>One_U1211-97</i>	0.081	75	0.031	55	0.057	69
<i>One_U1212-106</i>	0.121	34	0.013	87	0.066	53
<i>One_U1214-107</i>	0.271	6	0.015	83	0.043	84
<i>One_U1215-82</i>	0.047	100	0.057	37	0.017	102
<i>One_U1216-230</i>	0.087	67	0.005	97	0.027	97
<i>One_U301-92</i>	0.079	78	0.019	74	0.063	59
<i>One_U401-224</i>	0.113	41	0.011	91	0.025	100
<i>One_U404-229</i>	0.144	22	0.009	94	0.080	39
<i>One_U502-167</i>	0.102	53	0.043	45	0.016	104
<i>One_U503-170</i>	0.092	61	0.070	27	0.054	72
<i>One_U504-141</i>	0.097	55	0.045	44	0.058	66
<i>One_U508-533</i>	0.115	39	0.015	82	0.102	24
<i>One_UCA-24<sup>e</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_vamp5-255</i>	0.104	50	0.003	101	0.076	45
<i>One_vatf-214</i>	0.103	52	0.033	51	0.165	10
<i>One_VIM-569</i>	0.078	80	0.165	6	0.080	40
<i>One_zn706-68</i>	0.006	111	0.004	100	0.006	109
<i>One_ZNF-61</i>	0.129	29	0.040	46	0.162	11
<i>One_Zp3b-49</i>	0.205	14	0.049	42	0.091	29

<sup>a</sup> These assays were dropped due to significant linkage and were not included in this analysis.

<sup>b</sup> These assays were dropped due to fixation in the 36 test populations and were not included in this analysis.

<sup>c</sup> These assays are linked and were included as a haplotype marker in this analysis.

<sup>d</sup> These assays are linked and were included as a haplotype marker in this analysis.

<sup>e</sup> These assays were dropped due to laboratory failure and were not included in this analysis.

<sup>f</sup> This assay was dropped due to failure to conform to Hardy-Weinberg expectations and was not included in this analysis.

<sup>g</sup> This assay was fixed for both populations in each pair and was given a rating equal to the worst.

Table 9.—Test statistics and rankings based upon log-likelihood ratio (G) tests that describe the power of 124 SNPs (Tables 2 and 3) to discriminate between 7 pairs of sockeye salmon populations of interest (Table 1).

Assay	Ualik-Pick		Becharof-Deer		Deer-Cinder		Broadway-Hatchery		Yentna-Susitna		Larson-Mama		McDonald-Hugh	
	G	Rank	G	Rank	G	Rank	G	Rank	G	Rank	G	Rank	G	Rank
<i>One_ACBP-79</i>	3.39	30	0.4	71	0.23	87	0.39	71	6.82	22	0.07	71	7.86	13
<i>One_agt-132</i>	0.52	66	0.32	74.5	9.31	18	1.38	43	0.33	83	1.12	40	6.18	19
<i>One_aldB-152</i>	5.4	17	0.46	66	0.99	70	4.14	17	3.46	35	0.27	58	5.53	21
<i>One_ALDOB-135<sup>a</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_apoe-83</i>	4.95	19	0.05	97.5	10.7	16	0	93	8.16	18	5.83	4	5.38	22
<i>One_bckB-137<sup>b</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_c3-98</i>	0	94.5	1.68	37	0.2	92.5	1.66	38	2.25	50	1.29	35	0.03	93
<i>One_ccd16-131</i>	N/A <sup>g</sup>	94.5	N/A <sup>g</sup>	104.5	N/A <sup>g</sup>	92.5	N/A <sup>g</sup>	93	N/A <sup>g</sup>	93	N/A <sup>g</sup>	90	N/A <sup>g</sup>	93
<i>One_CD9-269</i>	0	94.5	7.96	7	17.98	12	0.27	77	14.76	4	0.01	85	0.68	67
<i>One_cetn1-167</i>	0.22	80	0.24	83	1.06	69	0.85	59.5	2.96	39.5	2.99	11	0.36	78
<i>One_CFP1</i>	2.26	42	0.82	53	0.05	92.5	2.26	30	0.6	78	0.04	75.5	4.21	28
<i>One_cin-177</i>	4.9	20	1.2	47	11.1	15	0.01	91	0.22	87	2.47	21	3.17	34
<i>One_COI<sup>c</sup></i>	1.26	55	0.05	97.5	2.8	37	2.8	22	1.89	58	0.03	78.5	32.65	2
<i>One_CTGF-301</i>	N/A <sup>g</sup>	94.5	0.41	69.5	1.29	61.5	2.02	33	9.23	14.5	2.6	17.5	4.15	29
<i>One_Cytb_17<sup>c</sup></i>	1.26	55	0.05	97.5	2.8	37	2.8	22	1.89	58	0.03	78.5	32.65	2
<i>One_Cytb_26<sup>c</sup></i>	1.26	55	0.05	97.5	2.8	37	2.8	22	1.89	58	0.03	78.5	32.65	2
<i>One_dds-529</i>	3.42	28	1.52	40	0.22	89.5	4.33	14	10.08	12	0.74	44	0.05	93
<i>One_DDX5-86</i>	3.4	29	0.57	61	0.55	77	0	93	2.01	56	0.26	60	3.95	30
<i>One_E2-65</i>	3.38	31	1.63	39	2.66	40	0.85	59.5	0.15	91.5	0.21	66.5	0.01	93
<i>One_gadd45-269</i>	N/A <sup>g</sup>	94.5	N/A <sup>g</sup>	104.5	N/A <sup>g</sup>	92.5	N/A <sup>g</sup>	93	1.48	63.5	N/A <sup>g</sup>	90	1.37	55
<i>One_gdh-212</i>	7.42	12	0.6	59.5	0.53	78	0.7	62.5	2.57	46	0	90	0.56	70
<i>One_GHII-2165</i>	0.65	62	0.11	90.5	0.81	73.5	1.08	52	0.22	87	10.05	2	0.06	93
<i>One_ghsR-66</i>	0.98	59	6.48	10	2.69	39	0.96	55	0.22	87	2.53	20	2.35	38
<i>One_GPDH-201</i>	8.11	9	0.8	54.5	0.09	92.5	6.58	7	0.66	74	1.91	26	0.93	62

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Table 9. Page 2 of 6.

Assay	Ualik-Pick		Becharof-Deer		Deer-Cinder		Broadway-Hatchery		Yentna-Susitna		Larson-Mama		McDonald-Hugh	
	G	Rank	G	Rank	G	Rank	G	Rank	G	Rank	G	Rank	G	Rank
<i>One_GPDH2-187</i>	0.48	68	1.92	36	6.11	22	2.76	23	5.81	27.5	0.06	72.5	6.46	18
<i>One_GPH-414<sup>a</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_GTHa</i>	8.26	8	8.29	6	18.52	11	0.11	81	4.47	32	0	90	13.85	5
<i>One_HGFA-49</i>	0.46	69.5	0.38	72	3.5	32	5.73	8	6.52	24	0.54	49	8.89	11
<i>One_HpaI-71</i>	1.44	50	0.41	69.5	1.54	52	1.77	36	5	30	0.02	82	1.11	58.5
<i>One_HpaI-99</i>	0.2	82	1.19	48	2.54	42	1.01	54	0.37	81.5	1.34	32	3.81	32
<i>One_hsc71-220</i>	4.73	21	0.07	95	1.14	65	1.72	37	4.52	31	2.76	16	0.15	89.5
<i>One_Hsp47</i>	0.06	88	0.02	100	3.74	30	2.41	25	31.91	1	1.19	38	1.56	50
<i>One_Ig-90</i>	N/A <sup>g</sup>	94.5	N/A <sup>g</sup>	104.5	N/A <sup>g</sup>	92.5	N/A <sup>g</sup>	93	N/A <sup>g</sup>	93	N/A <sup>g</sup>	90	1.07	61
<i>One_IL8r-362</i>	5.12	18	30.05	2	34.25	4	3.85	18	1.83	59	0	90	2.1	43
<i>One_ins-107</i>	2.93	32	1.13	49.5	0	92.5	4.41	13	2.1	55	0.02	82	1.09	60
<i>One_KCT1-453</i>	0.03	89.5	2.89	21.5	3.34	34	1.14	50.5	3.78	34	4.55	7	0.13	91.5
<i>One_KPNA-422</i>	3.54	27	0.14	86	3.6	31	0.54	65	3.16	38	0.1	70	8.68	12
<i>One_LEI-87</i>	2.53	38	3.18	15	0.91	71	0.01	91	1.77	60.5	0.51	50	6.65	17
<i>One_leptin-92<sup>b</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_lpp1-44</i>	1.14	56	16.31	3	10.27	17	0.05	83	1	70	2.91	14	0.32	80
<i>One_MARCKS-241</i>	N/A <sup>g</sup>	94.5	2.68	26.5	0	92.5	N/A <sup>g</sup>	93	3.99	33	N/A <sup>g</sup>	90	3.09	35
<i>One_metA-253</i>	N/A <sup>g</sup>	94.5	1.34	45	1.37	58	N/A <sup>g</sup>	93	2.23	51.5	5.96	3	4.67	26
<i>One_MHC2_190<sup>d</sup></i>	20.19	3	158.28	1	37.4	3	53.39	2	9.29	13	14.01	1	37.13	1
<i>One_MHC2_251<sup>d</sup></i>	20.19	3	158.28	1	37.4	3	53.39	2	9.29	13	14.01	1	37.13	1
<i>One_Mkpro-129</i>	7.57	11	0.52	63	1.47	53	3.81	19	0.03	93	1.71	28	0.3	81.5
<i>One_ODC1-196</i>	25.2	2	0.01	101.5	0.17	92.5	8.78	6	1.48	63.5	0.62	47	2.12	42
<i>One_Ots208-234</i>	0.99	58	1.21	46	0.52	79	4.22	15	2.8	43	0.33	55	0.02	93
<i>One_Ots213-181</i>	2.12	44	2.22	30.5	1.24	63	1.28	47	6.78	23	1.96	23	0.04	93
<i>One_p53-534</i>	N/A <sup>g</sup>	94.5	2.89	21.5	1.41	54.5	N/A <sup>g</sup>	93	N/A <sup>g</sup>	93	N/A <sup>g</sup>	90	0.39	76

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Table 9. Page 3 of 6.

Assay	Ualik-Pick		Becharof-Deer		Deer-Cinder		Broadway-Hatchery		Yentna-Susitna		Larson-Mama		McDonald-Hugh	
	G	Rank	G	Rank	G	Rank	G	Rank	G	Rank	G	Rank	G	Rank
<i>One_parp3-170<sup>b</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_pax7-248</i>	0.15	84	2.84	24	0.81	73.5	1.88	35	9.05	16	1.03	42	0.07	93
<i>One_PIP</i>	0.56	64	0.64	56	23.14	9	0.02	88	2.79	44	0.35	54	3.89	31
<i>One_ppie-74</i>	N/A <sup>g</sup>	94.5	2.78	25	1.32	60	1.39	41.5	0.01	93	N/A <sup>g</sup>	90	0.23	85
<i>One_PPM1K-118<sup>e</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_Prl2</i>	0.66	61	0.11	90.5	6.86	20	0.04	85	0	93	0.72	45	0.41	73
<i>One_psme2-354</i>	0.01	92	7.71	8	15.93	13	4.91	12	0.16	90	0	90	10.57	6
<i>One_rab1a-76</i>	0.22	80	0	104.5	0.2	92.5	0.19	78	0.01	93	0.03	78.5	0	93
<i>One_RAG1-103</i>	2.87	33	1.35	43.5	1.08	67.5	4.15	16	14.11	6	1.51	29	1.27	57
<i>One_RAG3-93</i>	4.65	22	6.65	9	1.08	67.5	1.36	44	0.02	93	2.97	12.5	5.86	20
<i>One_redd1-414</i>	8.52	7	0.12	88.5	0.38	82	0.9	57.5	7.48	21	1.25	37	0.52	72
<i>One_RFC2-102</i>	1.82	46	2.92	19.5	1.2	64	0.01	91	17.87	2	0.03	78.5	9.33	10
<i>One_RFC2-285</i>	2.7	36	4.35	13.5	30.81	5	1.14	50.5	2.96	39.5	N/A <sup>g</sup>	90	1.92	46
<i>One_RH2op-395</i>	0	94.5	0.36	73	1.4	56	N/A <sup>g</sup>	93	1.31	66	N/A <sup>g</sup>	90	7.22	16
<i>One_rpo2j-261</i>	4.38	23	1.48	41	1.36	59	1.15	49	10.98	10	0.6	48	0.12	93
<i>One_sast-211</i>	0.29	78	2.15	33	5.29	24	0	93	9.23	14.5	0.26	60	2.85	36
<i>One_serpin-75</i>	1.1	57	0.18	84.5	0.66	76	13.02	4	0	93	N/A <sup>g</sup>	90	0.74	65
<i>One_spf30-207</i>	0.5	67	0	104.5	2.31	45.5	0	93	14.65	5	0.04	75.5	0.29	83
<i>One_srp09-127</i>	1.36	53	4.43	12	1.29	61.5	0.34	74	1.11	69	1.3	34	0.4	74.5
<i>One_ssrd-135</i>	1.28	54	3.1	17	0.9	72	1.34	45	5.28	29	1.86	27	4.71	25
<i>One_STC-410</i>	5.87	15	0.49	65	24.82	7	0.92	56	0.04	93	3.52	8	1.59	49
<i>One_STR07</i>	5.95	14	0.54	62	44.39	2	0.35	72.5	10.27	11	0.01	85	0.35	79
<i>One_SUMO1-6</i>	0.33	77	2.34	29	0.02	92.5	0.02	88	2.54	48	0.25	62	0.68	67
<i>One_sys1-230</i>	0.02	91	1.13	49.5	5.27	25	3.37	20	0	93	0.05	74	0.68	67
<i>One_taf12-248</i>	N/A <sup>g</sup>	94.5	N/A <sup>g</sup>	104.5	1.41	54.5	0.35	72.5	N/A <sup>g</sup>	93	0.31	56	0.57	69

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	Ualik-Pick		Becharof-Deer		Deer-Cinder		Broadway-Hatchery		Yentna-Susitna		Larson-Mama		McDonald-Hugh	
Assay	G	Rank	G	Rank	G	Rank	G	Rank	G	Rank	G	Rank	G	Rank
<i>One_Tf_ex11-750</i>	0.14	85	0.29	78.5	0	92.5	12.81	5	0.14	93	1.4	31	1.55	51
<i>One_Tf_in3-182</i>	0.11	86	0.94	52	3.18	35	39.95	3	0.29	85	N/A <sup>g</sup>	90	0	93
<i>One_tshB-92</i>	4.32	24	1.4	42	0.05	92.5	2.14	32	8.42	17	1.16	39	1.28	56
<i>One_txnip-401</i>	N/A <sup>g</sup>	94.5	0.29	78.5	2.74	38	N/A <sup>g</sup>	93	N/A <sup>g</sup>	93	N/A <sup>g</sup>	90	0.15	89.5
<i>One_U1002-101</i>	N/A <sup>g</sup>	94.5	N/A <sup>g</sup>	104.5	N/A <sup>g</sup>	92.5	N/A <sup>g</sup>	93	N/A <sup>g</sup>	93	N/A <sup>g</sup>	90	4.64	27
<i>One_U1003-75</i>	1.52	48	0.18	84.5	2.55	41	5.21	11	0.9	72	0.39	52	9.67	9
<i>One_U1004-183</i>	73.74	1	0.3	77	46.52	1	137.5	1	13.52	7	0.03	78.5	7.85	14
<i>One_U1009-91</i>	2.8	34	0.05	97.5	0.11	92.5	2.31	28	1.59	62	0.01	85	0.03	93
<i>One_U1010-81</i>	N/A <sup>g</sup>	94.5	0.6	59.5	2.95	36	0.3	76	11.54	9	0.37	53	2.25	41
<i>One_U1012-68</i>	10.9	5	14.36	4	0.43	81	0.9	57.5	0	93	0.26	60	1.44	52
<i>One_U1013-108</i>	0.85	60	0.08	94	0.51	80	0.15	79	0.12	93	5.08	5	2.34	39
<i>One_U1014-74</i>	0.54	65	0.61	57.5	12.05	14	2.39	26	2.91	42	0	90	1.93	45
<i>One_U1016-115</i>	11.29	4	2.22	30.5	1.62	50	1.22	48	0.44	79	0.22	65	4.75	24
<i>One_U1017-62</i>	N/A <sup>g</sup>	94.5	1.35	43.5	1.38	57	0	93	N/A <sup>g</sup>	93	N/A <sup>g</sup>	90	0	93
<i>One_U1021-57<sup>f</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_U1024-197</i>	2.49	39	0.31	76	4.03	28	0	93	2.13	54	0.18	68.5	9.79	8
<i>One_U1101</i>	0.45	71	2.18	32	0.17	92.5	0.58	64	15.23	3	3.47	10	5.17	23
<i>One_U1102-220</i>	0	94.5	2.68	26.5	0	92.5	0	93	0.82	73	0.29	57	2.41	37
<i>One_U1103</i>	1.43	51.5	0	104.5	2.02	47	0.05	83	2.59	45	2.6	17.5	26.37	3
<i>One_U1104-138</i>	0.61	63	2.92	19.5	N/A <sup>g</sup>	92.5	N/A <sup>g</sup>	93	N/A <sup>g</sup>	93	N/A <sup>g</sup>	90	N/A <sup>g</sup>	93
<i>One_U1105</i>	3.95	25	1.65	38	0.22	89.5	2.23	31	6.07	25	1.26	36	0.19	86
<i>One_U1201-492</i>	0.03	89.5	8.48	5	0	92.5	1.56	40	2	57	3.49	9	1.81	47
<i>One_U1202-1052</i>	0.37	75	2.88	23	4.84	26	0.05	83	0.65	75.5	1.98	22	0.06	93
<i>One_U1203-175</i>	0.08	87	0.09	93	28.28	6	2.58	24	0.92	71	2.87	15	2	44
<i>One_U1204-53</i>	0.46	69.5	0.44	67	2.31	45.5	0.03	86	2.54	48	4.69	6	0.91	63

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Assay	Ualik-Pick		Becharof-Deer		Deer-Cinder		Broadway-Hatchery		Yentna-Susitna		Larson-Mama		McDonald-Hugh	
	G	Rank	G	Rank	G	Rank	G	Rank	G	Rank	G	Rank	G	Rank
<i>One_U1205-57</i>	2.09	45	4.35	13.5	5.67	23	1.06	53	1.3	67	N/A <sup>g</sup>	90	1.65	48
<i>One_U1206-108</i>	8.04	10	0.26	81.5	1.13	66	2.36	27	0.18	89	1.92	25	0.3	81.5
<i>One_U1207-231<sup>e</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_U1208-67</i>	0.22	80	1.06	51	6.53	21	0.53	66	5.81	27.5	0.06	72.5	0.12	93
<i>One_U1209-111</i>	0.37	75	4.96	11	8.56	19	0.82	61	3.35	36	0.45	51	0.13	91.5
<i>One_U1210-173</i>	2.46	40	2.54	28	1.93	48	5.38	10	2.16	53	1.08	41	1.42	53
<i>One_U1211-97</i>	3.66	26	0.26	81.5	3.36	33	0.31	75	12.88	8	0.85	43	0	93
<i>One_U1212-106</i>	0.38	72.5	0.42	68	1.57	51	0.44	69	7.69	20	0	90	3.76	33
<i>One_U1214-107</i>	1.77	47	0	104.5	24.17	8	1.39	41.5	0.37	81.5	1.47	30	7.29	15
<i>One_U1215-82</i>	0.38	72.5	0.05	97.5	0.29	86	1.32	46	0.38	80	1.93	24	1.11	58.5
<i>One_U1216-230</i>	10.27	6	3.07	18	19.24	10	0	93	0.15	91.5	2.57	19	0.4	74.5
<i>One_U301-92</i>	1.45	49	0.12	88.5	0.17	92.5	2.29	29	1.77	60.5	1.31	33	0.09	93
<i>One_U401-224</i>	0.18	83	3.12	16	0.35	84	1.96	34	1.18	68	0.02	82	0.75	64
<i>One_U404-229</i>	N/A <sup>g</sup>	94.5	2.12	34	2.32	44	0.5	68	1.34	65	2.97	12.5	10.16	7
<i>One_U502-167</i>	5.82	16	0.51	64	0.67	75	0.52	67	5.9	26	N/A <sup>g</sup>	90	14.47	4
<i>One_U503-170</i>	2.54	37	2.1	35	1.8	49	1.64	39	0.62	77	0.21	66.5	0.28	84
<i>One_U504-141</i>	2.16	43	0.01	101.5	0.22	89.5	0.42	70	0.32	84	0.18	68.5	0.54	71
<i>One_U508-533</i>	2.79	35	0.1	92	0.22	89.5	N/A <sup>g</sup>	93	7.87	19	0	90	0.17	87.5
<i>One_UCA-24<sup>e</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_vamp5-255</i>	0.37	75	0.61	57.5	0.37	83	0.02	88	0.07	93	0.68	46	2.31	40
<i>One_vatf-214</i>	N/A <sup>g</sup>	94.5	0.8	54.5	0.33	85	2.85	21	2.54	48	N/A <sup>g</sup>	90	0.37	77
<i>One_VIM-569</i>	2.37	41	0.13	87	3.82	29	0.13	80	3.23	37	0.23	63.5	0.17	87.5
<i>One_zn706-68</i>	N/A <sup>g</sup>	94.5	N/A <sup>g</sup>	104.5	N/A <sup>g</sup>	92.5	N/A <sup>g</sup>	93	2.94	41	N/A <sup>g</sup>	90	N/A <sup>g</sup>	93
<i>One_ZNF-61</i>	6.54	13	0.28	80	2.53	43	5.61	9	0.65	75.5	0.23	63.5	0.07	93

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	<u>Ualik-Pick</u>		<u>Becharof-Deer</u>		<u>Deer-Cinder</u>		<u>Broadway-Hatchery</u>		<u>Yentna-Susitna</u>		<u>Larson-Mama</u>		<u>McDonald-Hugh</u>	
Assay	<i>G</i>	Rank	<i>G</i>	Rank	<i>G</i>	Rank	<i>G</i>	Rank	<i>G</i>	Rank	<i>G</i>	Rank	<i>G</i>	Rank
<i>One_Zp3b-49</i>	1.43	51.5	0.32	74.5	4.22	27	0.7	62.5	2.23	51.5	N/A <sup>g</sup>	90	1.41	54

<sup>a</sup> These assays were dropped due to significant linkage and were not included in this analysis.

<sup>b</sup> These assays were dropped due to fixation in the 36 test populations and were not included in this analysis.

<sup>c</sup> These assays are linked and were included as a haplotype marker in this analysis.

<sup>d</sup> These assays are linked and were included as a haplotype marker in this analysis.

<sup>e</sup> These assays were dropped due to laboratory failure and were not included in this analysis.

<sup>f</sup> This assay was dropped due to failure to conform to Hardy-Weinberg expectations and was not included in this analysis.

<sup>g</sup> Some SNPs were homozygous for both populations in some *G* tests and lack a test statistic. These SNPs were given a ranking equal to the worst rank for the test in question.

Table 10.—Rankings based upon  $f_{ORCA}$  that describe the power of 124 SNPs (Tables 2 and 3) to discriminate between 7 pairs of sockeye salmon populations of interest (Table 1).

Assay	Ualik-Pick	Becharof-Deer	Deer-Cinder	Broadway-Hatchery	Yentna-Susitna	Larson-Mama	McDonald-Hugh
<i>One_ACBP-79</i>	20	61	72	59	20	64	13
<i>One_agt-132</i>	46	45	16	48	62	29	16
<i>One_aldB-152</i>	16	60	51	10	31	47	21
<i>One_ALDOB-135<sup>a</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_apoe-83</i>	21	103	14	109	15	5	19
<i>One_bckB-137<sup>b</sup></i>	108	112	104	100	103	57	109
<i>One_c3-98</i>	110	34	84	39	57	92	92
<i>One_ccd16-131</i>	114	106	113	92	110	108	101
<i>One_CD9-269</i>	109	7	7	67	5	87	59
<i>One_cetn1-167</i>	78	50	55	68	44	8	79
<i>One_CFP1</i>	43	44	105	27	68	105	26
<i>One_cin-177</i>	27	32	18	71	73	16	31
<i>One_CO1<sup>c</sup></i>	44	110	22	30	66	58	1
<i>One_CTGF-301</i>	92	95	76	66	30	53	51
<i>One_Cytb_17<sup>c</sup></i>	44	110	22	30	66	58	1
<i>One_Cytb_26<sup>c</sup></i>	44	110	22	30	66	58	1
<i>One_dds-529</i>	30	26	63	13	12	38	99
<i>One_DDX5-86</i>	33	54	79	110	46	61	28
<i>One_E2-65</i>	23	27	30	45	89	59	114
<i>One_gadd45-269</i>	97	108	109	103	95	85	87
<i>One_gdh-212</i>	7	48	65	56	27	102	54
<i>One_GHII-2165</i>	67	77	66	47	75	2	100
<i>One_ghsR-66</i>	45	8	31	44	82	13	32
<i>One_GPDH-201</i>	11	40	91	6	76	18	42

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Assay	Ualik-Pick	Becharof-Deer	Deer-Cinder	Broadway-Hatchery	Yentna-Susitna	Larson-Mama	McDonald-Hugh
<i>One_GPDH2-187</i>	60	72	38	51	34	73	15
<i>One_GPH-414<sup>a</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_GTHa</i>	8	6	10	69	29	109	5
<i>One_HGFA-49</i>	86	66	28	11	16	49	8
<i>One_HpaI-71</i>	37	59	48	38	22	77	55
<i>One_HpaI-99</i>	100	51	44	37	105	33	25
<i>One_hsc71-220</i>	17	114	52	23	25	17	95
<i>One_Hsp47</i>	84	99	23	35	1	27	46
<i>One_Ig-90</i>	105	113	114	114	112	89	67
<i>One_IL8r-362</i>	22	2	6	24	51	76	36
<i>One_ins-107</i>	35	35	106	12	52	72	56
<i>One_KCT1-453</i>	82	13	34	65	43	32	89
<i>One_KPNA-422</i>	28	82	26	64	39	60	10
<i>One_LEI-87</i>	36	21	54	113	40	41	11
<i>One_leptin-92<sup>b</sup></i>	95	87	107	112	109	84	105
<i>One_lpp1-44</i>	39	3	15	73	54	21	66
<i>One_MARCKS-241</i>	93	85	111	95	69	110	77
<i>One_metA-253</i>	103	79	103	87	74	42	37
<i>One_MHC2_190<sup>d</sup></i>	3	1	3	2	19	1	2
<i>One_MHC2_251<sup>d</sup></i>	3	1	3	2	19	1	2
<i>One_Mkpro-129</i>	10	47	56	14	92	19	82
<i>One_ODC1-196</i>	2	92	80	5	59	40	41
<i>One_Ots208-234</i>	80	49	94	21	41	36	108
<i>One_Ots213-181</i>	40	36	40	32	21	14	102
<i>One_p53-534</i>	96	81	88	93	91	88	72

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Assay	Ualik-Pick	Becharof-Deer	Deer-Cinder	Broadway-Hatchery	Yentna-Susitna	Larson-Mama	McDonald-Hugh
<i>One_parp3-170<sup>b</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_pax7-248</i>	72	20	49	33	10	34	96
<i>One_PIP</i>	64	38	8	76	42	37	30
<i>One_ppie-74</i>	102	55	71	72	101	112	88
<i>One_PPM1K-118<sup>e</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_Prl2</i>	49	56	20	79	96	31	83
<i>One_psme2-354</i>	94	5	9	7	70	93	12
<i>One_rab1a-76</i>	68	88	97	85	106	114	113
<i>One_RAG1-103</i>	62	98	81	62	2	24	76
<i>One_RAG3-93</i>	19	22	78	31	104	50	17
<i>One_redd1-414</i>	13	78	67	46	18	26	86
<i>One_RFC2-102</i>	42	15	53	80	6	97	14
<i>One_RFC2-285</i>	75	65	17	54	86	99	43
<i>One_RH2op-395</i>	101	68	90	101	88	113	57
<i>One_rpo2j-261</i>	18	25	47	43	8	52	93
<i>One_sast-211</i>	71	84	59	105	55	74	33
<i>One_serpin-75</i>	83	105	96	29	113	80	74
<i>One_spf30-207</i>	69	75	32	99	3	70	94
<i>One_srp09-127</i>	79	28	77	88	72	111	84
<i>One_ssrd-135</i>	47	14	62	40	26	15	22
<i>One_STC-410</i>	14	43	5	58	100	9	47
<i>One_STR07</i>	9	53	2	70	9	94	85
<i>One_SUMO1-6</i>	59	18	86	97	35	45	62
<i>One_sys1-230</i>	90	30	25	18	99	71	60
<i>One_taf12-248</i>	104	111	101	94	111	75	68

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Table 10. Page 4 of 6.

Assay	Ualik-Pick	Becharof-Deer	Deer-Cinder	Broadway-Hatchery	Yentna-Susitna	Larson-Mama	McDonald-Hugh
<i>One_Tf_ex11-750</i>	88	58	102	4	71	30	44
<i>One_Tf_in3-182</i>	85	37	35	3	98	107	104
<i>One_tshB-92</i>	52	46	95	52	38	25	53
<i>One_txnlp-401</i>	106	96	85	98	114	101	80
<i>One_U1002-101</i>	89	90	112	77	108	95	34
<i>One_U1003-75</i>	74	80	43	8	63	35	6
<i>One_U1004-183</i>	1	83	1	1	11	98	9
<i>One_U1009-91</i>	25	89	87	19	45	79	98
<i>One_U1010-81</i>	111	73	60	82	24	51	39
<i>One_U1012-68</i>	15	4	69	53	97	90	35
<i>One_U1013-108</i>	50	62	64	74	90	23	52
<i>One_U1014-74</i>	57	42	12	25	33	82	45
<i>One_U1016-115</i>	4	12	45	34	64	44	18
<i>One_U1017-62</i>	113	109	98	104	107	96	111
<i>One_U1021-57<sup>f</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_U1024-197</i>	48	67	36	90	61	55	4
<i>One_U1101</i>	56	23	73	61	4	7	20
<i>One_U1102-220</i>	99	94	108	91	67	43	49
<i>One_U1103</i>	81	107	75	96	81	63	3
<i>One_U1104-138</i>	73	63	93	111	102	81	112
<i>One_U1105</i>	29	33	82	26	17	28	70
<i>One_U1201-492</i>	98	9	110	36	47	6	40
<i>One_U1202-1052</i>	61	11	24	75	53	10	110
<i>One_U1203-175</i>	58	97	4	17	49	3	38
<i>One_U1204-53</i>	70	71	39	78	36	4	58

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Assay	Ualik-Pick	Becharof-Deer	Deer-Cinder	Broadway-Hatchery	Yentna-Susitna	Larson-Mama	McDonald-Hugh
<i>One_U1205-57</i>	54	70	57	86	77	104	69
<i>One_U1206-108</i>	6	57	41	28	87	11	71
<i>One_U1207-231<sup>e</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_U1208-67</i>	65	29	19	60	13	67	64
<i>One_U1209-111</i>	53	19	21	63	28	56	91
<i>One_U1210-173</i>	38	17	29	15	37	39	106
<i>One_U1211-97</i>	26	64	27	57	7	46	90
<i>One_U1212-106</i>	55	41	42	81	23	78	23
<i>One_U1214-107</i>	63	104	13	102	93	83	24
<i>One_U1215-82</i>	51	74	68	41	79	20	61
<i>One_U1216-230</i>	5	10	11	108	83	12	73
<i>One_U301-92</i>	34	91	74	16	48	22	103
<i>One_U401-224</i>	76	16	83	22	50	86	50
<i>One_U404-229</i>	91	31	58	55	80	68	7
<i>One_U502-167</i>	41	69	89	89	60	69	27
<i>One_U503-170</i>	31	24	46	42	58	65	97
<i>One_U504-141</i>	24	100	70	50	78	54	63
<i>One_U508-533</i>	66	102	92	107	14	100	81
<i>One_UCA-24<sup>e</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_vamp5-255</i>	77	39	61	84	94	66	29
<i>One_vatf-214</i>	112	76	100	20	84	91	65
<i>One_VIM-569</i>	32	86	50	83	32	62	75
<i>One_zn706-68</i>	107	101	99	106	85	103	78
<i>One_ZNF-61</i>	12	52	33	9	56	48	107

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Assay	Ualik-Pick	Becharof-Deer	Deer-Cinder	Broadway-Hatchery	Yentna-Susitna	Larson-Mama	McDonald-Hugh
<i>One_Zp3b-49</i>	87	93	37	49	65	106	48

<sup>a</sup> These assays were dropped due to significant linkage and were not included in this analysis.

<sup>b</sup> These assays were dropped due to fixation in the 36 test populations and were not included in this analysis.

<sup>c</sup> These assays are linked and were included as a haplotype marker in this analysis.

<sup>d</sup> These assays are linked and were included as a haplotype marker in this analysis.

<sup>e</sup> These assays were dropped due to laboratory failure and were not included in this analysis.

<sup>f</sup> This assay was dropped due to failure to conform to Hardy-Weinberg expectations and was not included in this analysis.



Table 11.–Rankings based upon  $f_{ORCA}$  that describe the power of each of 124 SNPs (Tables 2 and 3) to discriminate among 36 sockeye salmon populations of interest (Table 1).

Assay	$f_{ORCA}$ rank	Assay	$f_{ORCA}$ rank
<i>One_ACBP-79</i>	41	<i>One_KCT1-453</i>	83
<i>One_agt-132</i>	25	<i>One_KPNA-422</i>	58
<i>One_aldB-152</i>	47	<i>One_LEI-87</i>	17
<i>One_ALDOB-135<sup>a</sup></i>	N/A	<i>One_leptin-92<sup>b</sup></i>	114
<i>One_apoe-83</i>	12	<i>One_lpp1-44</i>	26
<i>One_bckB-137<sup>b</sup></i>	112	<i>One_MARCKS-241</i>	104
<i>One_c3-98</i>	78	<i>One_metA-253</i>	7
<i>One_ccd16-131</i>	113	<i>One_MHC2_190<sup>d</sup></i>	1
<i>One_CD9-269</i>	74	<i>One_MHC2_251<sup>d</sup></i>	1
<i>One_cetn1-167</i>	56	<i>One_Mkpro-129</i>	11
<i>One_CFP1</i>	16	<i>One_ODC1-196</i>	29
<i>One_cin-177</i>	18	<i>One_Ots208-234</i>	38
<i>One_CO1<sup>c</sup></i>	2	<i>One_Ots213-181</i>	48
<i>One_CTGF-301</i>	105	<i>One_p53-534</i>	97
<i>One_Cytb_17<sup>c</sup></i>	2	<i>One_parp3-170<sup>b</sup></i>	N/A
<i>One_Cytb_26<sup>c</sup></i>	2	<i>One_pax7-248</i>	75
<i>One_dds-529</i>	79	<i>One_PIP</i>	84
<i>One_DDX5-86</i>	71	<i>One_ppie-74</i>	30
<i>One_E2-65</i>	86	<i>One_PPM1K-118<sup>e</sup></i>	N/A
<i>One_gadd45-269</i>	110	<i>One_Prl2</i>	43
<i>One_gdh-212</i>	42	<i>One_psme2-354</i>	21
<i>One_GHII-2165</i>	4	<i>One_rab1a-76</i>	73
<i>One_ghsR-66</i>	27	<i>One_RAG1-103</i>	90
<i>One_GPDH-201</i>	33	<i>One_RAG3-93</i>	93
<i>One_GPDH2-187</i>	63	<i>One_redd1-414</i>	6
<i>One_GPH-414<sup>a</sup></i>	N/A	<i>One_RFC2-102</i>	81
<i>One_GTHa</i>	28	<i>One_RFC2-285</i>	103
<i>One_HGFA-49</i>	69	<i>One_RH2op-395</i>	106
<i>One_HpaI-71</i>	14	<i>One_rpo2j-261</i>	64
<i>One_HpaI-99</i>	24	<i>One_sast-211</i>	100
<i>One_hsc71-220</i>	65	<i>One_serp1-75</i>	107
<i>One_Hsp47</i>	70	<i>One_spf30-207</i>	76
<i>One_Ig-90</i>	92	<i>One_srp09-127</i>	77
<i>One_IL8r-362</i>	62	<i>One_ssrd-135</i>	57
<i>One_ins-107</i>	61	<i>One_STC-410</i>	9

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Table 11. Page 2 of 2.

Assay	$f_{ORCA}$ rank	Assay	$f_{ORCA}$ rank
<i>One_STR07</i>	10	<i>One_U1203-175</i>	60
<i>One_SUMO1-6</i>	32	<i>One_U1204-53</i>	68
<i>One_sys1-230</i>	22	<i>One_U1205-57</i>	101
<i>One_taf12-248</i>	85	<i>One_U1206-108</i>	37
<i>One_Tf_ex11-750</i>	20	<i>One_U1207-231<sup>e</sup></i>	N/A
<i>One_Tf_in3-182</i>	45	<i>One_U1208-67</i>	13
<i>One_tshB-92</i>	98	<i>One_U1209-111</i>	54
<i>One_txnlp-401</i>	99	<i>One_U1210-173</i>	66
<i>One_U1002-101</i>	108	<i>One_U1211-97</i>	94
<i>One_U1003-75</i>	19	<i>One_U1212-106</i>	59
<i>One_U1004-183</i>	3	<i>One_U1214-107</i>	8
<i>One_U1009-91</i>	44	<i>One_U1215-82</i>	34
<i>One_U1010-81</i>	91	<i>One_U1216-230</i>	67
<i>One_U1012-68</i>	51	<i>One_U301-92</i>	55
<i>One_U1013-108</i>	50	<i>One_U401-224</i>	23
<i>One_U1014-74</i>	96	<i>One_U404-229</i>	72
<i>One_U1016-115</i>	5	<i>One_U502-167</i>	88
<i>One_U1017-62</i>	102	<i>One_U503-170</i>	46
<i>One_U1021-57<sup>f</sup></i>	N/A	<i>One_U504-141</i>	39
<i>One_U1024-197</i>	80	<i>One_U508-533</i>	89
<i>One_U1101</i>	35	<i>One_UCA-24<sup>e</sup></i>	N/A
<i>One_U1102-220</i>	52	<i>One_vamp5-255</i>	49
<i>One_U1103</i>	95	<i>One_vatf-214</i>	87
<i>One_U1104-138</i>	109	<i>One_VIM-569</i>	82
<i>One_U1105</i>	15	<i>One_zn706-68</i>	111
<i>One_U1201-492</i>	36	<i>One_ZNF-61</i>	31
<i>One_U1202-1052</i>	53	<i>One_Zp3b-49</i>	40

<sup>a</sup> These assays were dropped due to significant linkage and were not included in this analysis.

<sup>b</sup> These assays were dropped due to fixation in the 36 test populations and were not included in this analysis.

<sup>c</sup> These assays are linked and were included as a haplotype marker in this analysis.

<sup>d</sup> These assays are linked and were included as a haplotype marker in this analysis.

<sup>e</sup> These assays were dropped due to laboratory failure and were not included in this analysis.

<sup>f</sup> This assay was dropped due to failure to conform to Hardy-Weinberg expectations and was not included in this analysis.

Table 12.–Sum of rankings and final ADF&G rank for 124 SNPs (Tables 2 and 3) screened for 36 test populations (Table 1).

Assay	Sum of ranks	Final rank	Assay	Sum of ranks	Final rank
<i>One_ACBP-79</i>	1470.5	24	<i>One_KCT1-453</i>	1722	65
<i>One_agt-132</i>	1615	51	<i>One_KPNA-422</i>	1725	66
<i>One_aldB-152</i>	1565.5	43	<i>One_LEI-87</i>	1468.5	23
<i>One_ALDOB-135<sup>a</sup></i>	N/A	N/A	<i>One_leptin-92<sup>b</sup></i>	N/A	N/A
<i>One_apoe-83</i>	1068	6	<i>One_lpp1-44</i>	1278.5	10
<i>One_bckB-137<sup>b</sup></i>	N/A	N/A	<i>One_MARCKS-241</i>	2547	114
<i>One_c3-98</i>	2083	96	<i>One_metA-253</i>	1425	18
<i>One_ccd16-131</i>	2424	112	<i>One_MHC2_190<sup>e</sup></i>	991.5	5
<i>One_CD9-269</i>	1713	64	<i>One_MHC2_251<sup>e</sup></i>	704.5	1
<i>One_cetn1-167</i>	1986	91	<i>One_Mkpro-129</i>	1637	55
<i>One_CFP1</i>	1561	42	<i>One_ODC1-196</i>	1591.5	47
<i>One_cin-177</i>	1709.5	62	<i>One_Ots208-234</i>	1644.25	56
<i>One_COI<sup>c</sup></i>	894.5	4	<i>One_Ots213-181</i>	1509	32
<i>One_CTGF-301</i>	2039.5	94	<i>One_p53-534</i>	1995.5	93
<i>One_Cytb_17<sup>c</sup></i>	843.5	2	<i>One_parp3-170<sup>b</sup></i>	N/A	N/A
<i>One_Cytb_26<sup>c</sup></i>	874.5	3	<i>One_pax7-248</i>	1487	26
<i>One_dds-529</i>	1955.5	97	<i>One_PIP</i>	1776.5	74
<i>One_DDX5-86</i>	2323	108	<i>One_ppie-74</i>	2132	101
<i>One_E2-65</i>	1844.5	79	<i>One_PPM1K-118<sup>f</sup></i>	N/A	N/A
<i>One_gadd45-269</i>	2298.5	107	<i>One_Prl2</i>	1911.5	86
<i>One_gdh-212</i>	1665.5	58	<i>One_psme2-354</i>	1956	98
<i>One_GHII-2165</i>	1394.5	15	<i>One_rab1a-76</i>	2412	111
<i>One_ghsR-66</i>	1388.5	14	<i>One_RAG1-103</i>	1489.5	27
<i>One_GPDH-201</i>	1430	20	<i>One_RAG3-93</i>	1847	81
<i>One_GPDH2-187</i>	1608	50	<i>One_redd1-414</i>	1634	53
<i>One_GPH-414<sup>a,d</sup></i>	N/A	N/A	<i>One_RFC2-102</i>	1506	31
<i>One_GTHa<sup>d</sup></i>	1159.5	8	<i>One_RFC2-285</i>	1505	30
<i>One_HGFA-49</i>	1741	67	<i>One_RH2op-395</i>	2343.5	109
<i>One_HpaI-71</i>	1541.5	38	<i>One_rpo2j-261</i>	1429.5	19
<i>One_HpaI-99</i>	1746.5	69	<i>One_sast-211</i>	1778	76
<i>One_hsc71-220</i>	1711.5	63	<i>One_serpin-75</i>	2697	115
<i>One_Hsp47</i>	1520.5	34	<i>One_spf30-207</i>	1679	60
<i>One_Ig-90</i>	2164	103	<i>One_srp09-127</i>	1635.5	54
<i>One_IL8r-362</i>	1261.5	9	<i>One_ssrd-135</i>	1444	21
<i>One_ins-107</i>	2090.5	99	<i>One_STC-410</i>	1695	61

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Table 12. Page 2 of 2.

Assay	Sum of ranks	Final rank	Assay	Sum of ranks	Final rank
<i>One_STR07</i>	1528	36	<i>One_U1203-175</i>	1520	33
<i>One_SUMO1-6</i>	1858	82	<i>One_U1204-53</i>	1626	52
<i>One_sys1-230</i>	1844.5	80	<i>One_U1205-57</i>	2064	95
<i>One_taf12-248</i>	1915.5	87	<i>One_U1206-108</i>	1824	77
<i>One_Tf_ex11-750</i>	1538.5	37	<i>One_U1207-231<sup>f</sup></i>	N/A	N/A
<i>One_Tf_in3-182</i>	1410	16	<i>One_U1208-67</i>	1543.5	39
<i>One_tshB-92</i>	1666	59	<i>One_U1209-111</i>	1291.5	11
<i>One_txnlp-401</i>	1959	90	<i>One_U1210-173</i>	1646	57
<i>One_U1002-101</i>	2111.5	100	<i>One_U1211-97</i>	1992	92
<i>One_U1003-75</i>	1417.5	17	<i>One_U1212-106</i>	1758.5	72
<i>One_U1004-183</i>	1123	7	<i>One_U1214-107</i>	1550	41
<i>One_U1009-91</i>	1504.5	29	<i>One_U1215-82</i>	2393.5	110
<i>One_U1010-81</i>	1500.5	28	<i>One_U1216-230</i>	1831	78
<i>One_U1012-68</i>	1317.5	12	<i>One_U301-92</i>	1888	85
<i>One_U1013-108</i>	1876	83	<i>One_U401-224</i>	1753.5	70
<i>One_U1014-74</i>	1606.5	48	<i>One_U404-229</i>	1579	46
<i>One_U1016-115</i>	1334	13	<i>One_U502-167</i>	1471	25
<i>One_U1017-62</i>	2221.5	104	<i>One_U503-170</i>	1567.5	44
<i>One_U1021-57<sup>g</sup></i>	N/A	N/A	<i>One_U504-141</i>	1925.5	88
<i>One_U1024-197</i>	1607	49	<i>One_U508-533</i>	2286.5	106
<i>One_U1101</i>	1547	40	<i>One_UCA-24<sup>d</sup></i>	N/A	N/A
<i>One_U1102-220</i>	2241	105	<i>One_vamp5-255</i>	1933	89
<i>One_U1103</i>	1579	45	<i>One_vatf-214</i>	1757.5	71
<i>One_U1104-138</i>	2151.5	102	<i>One_VIM-569</i>	1768.5	73
<i>One_U1105</i>	1521	35	<i>One_zn706-68</i>	2472	113
<i>One_U1201-492</i>	1884.5	84	<i>One_ZNF-61</i>	1457.5	22
<i>One_U1202-1052</i>	1741.5	68	<i>One_Zp3b-49</i>	1777	75

<sup>a</sup> These assays were dropped due to significant linkage and were not included in this analysis.

<sup>b</sup> These assays were dropped due to fixation in the 36 test populations and were not included in this analysis.

<sup>c</sup> These assays are linked and were included as a haplotype marker in some analyses.

<sup>d</sup> These two linked assays have nearly identical allele frequencies for the 36 test populations and are expected to provide similar test statistics and rankings for all measures except laboratory performance. These were exchanged to provide additional overlap among markers run by stakeholder laboratories (i.e., CDFO). See text for details.

<sup>e</sup> These assays are linked and were included as a haplotype marker in some analyses.

<sup>f</sup> These assays were dropped due to laboratory failure and were not included in this analysis.

<sup>g</sup> This assay was dropped due to failure to conform to Hardy-Weinberg expectations and was not included in this analysis.

Table 13.–Cumulative performance measure (mean individual assignment accuracy), contribution to performance measure and rank based upon performance measure for 124 SNPs (Table 2) screened for 36 test populations (Table 1) from a Backwards Elimination Locus Selection (BELS) algorithm. See text for details.

Assay	Performance Measure	Contribution	Rank
<i>One_ACBP-79</i>	0.73	0.01	33
<i>One_agt-132</i>	0.80	0.00	49
<i>One_aldB-152</i>	0.68	0.01	26
<i>One_ALDOB-135<sup>a</sup></i>	N/A	N/A	N/A
<i>One_apoe-83</i>	0.34	0.03	6
<i>One_bckB-137</i>	0.89	0.00	110
<i>One_c3-98</i>	0.87	0.00	77
<i>One_ccd16-131</i>	0.89	0.00	112
<i>One_CD9-269</i>	0.78	0.00	42
<i>One_cetn1-167</i>	0.66	0.01	24
<i>One_CFP1</i>	0.88	0.00	95
<i>One_cin-177</i>	0.86	0.00	73
<i>One_CO1<sup>b</sup></i>	0.14	0.07	2
<i>One_CTGF-301</i>	0.88	0.00	84
<i>One_Cytb_17<sup>b</sup></i>	0.14	0.07	2
<i>One_Cytb_26<sup>b</sup></i>	0.14	0.07	2
<i>One_dds-529</i>	0.87	0.00	81
<i>One_DDX5-86</i>	0.76	0.00	37
<i>One_E2-65</i>	0.76	0.00	39
<i>One_gadd45-269</i>	0.89	0.00	107
<i>One_gdh-212</i>	0.87	0.00	83
<i>One_GHII-2165</i>	0.46	0.02	10
<i>One_ghsR-66</i>	0.52	0.02	13
<i>One_GPDH-201</i>	0.86	0.00	75
<i>One_GPDH2-187</i>	0.82	0.00	55
<i>One_GPH-414<sup>a</sup></i>	N/A	N/A	N/A
<i>One_GTHa</i>	0.63	0.01	21
<i>One_hsc71-220</i>	0.78	0.00	43
<i>One_HGFA-49</i>	0.88	0.00	87
<i>One_HpaI-71</i>	0.54	0.02	14
<i>One_HpaI-99</i>	0.56	0.02	15
<i>One_Hsp47</i>	0.60	0.01	18
<i>One_Ig-90</i>	0.89	0.00	111
<i>One_IL8r-362</i>	0.80	0.00	48

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Table 13. Page 2 of 4.

Assay	Performance Measure	Contribution	Rank
<i>One_KCT1-453</i>	0.77	0.00	41
<i>One_KPNA-422</i>	0.83	0.00	60
<i>One_LEI-87</i>	0.84	0.00	62
<i>One_leptin-92</i>	0.89	0.00	108
<i>One_lpp1-44</i>	0.48	0.02	11
<i>One_MARCKS-241</i>	0.89	0.00	104
<i>One_metA-253</i>	0.88	0.00	89
<i>One_MHC2_190<sup>c</sup></i>	0.00	0.14	1
<i>One_MHC2_251<sup>c</sup></i>	0.00	0.14	1
<i>One_Mkpro-129</i>	0.59	0.01	17
<i>One_ODC1-196</i>	0.66	0.01	23
<i>One_Ots208-234</i>	0.81	0.00	51
<i>One_Ots213-181</i>	0.79	0.00	45
<i>One_p53-534</i>	0.89	0.00	99
<i>One_parp3-170<sup>d</sup></i>	N/A	N/A	N/A
<i>One_pax7-248</i>	0.80	0.00	47
<i>One_pIns</i>	0.89	0.00	96
<i>One_PIP</i>	0.84	0.00	61
<i>One_ppie-74</i>	0.89	0.00	109
<i>One_PPM1K-118<sup>e</sup></i>	N/A	N/A	N/A
<i>One_Prl2</i>	0.70	0.01	29
<i>One_psme2-354</i>	0.75	0.01	36
<i>One_rab1a-76</i>	0.85	0.00	65
<i>One_RAG1-103</i>	0.82	0.00	53
<i>One_RAG3-93</i>	0.85	0.00	66
<i>One_redd1-414</i>	0.41	0.03	8
<i>One_RFC2-102</i>	0.67	0.01	25
<i>One_RFC2-285</i>	0.87	0.00	78
<i>One_RH2op395</i>	0.89	0.00	102
<i>One_rpo2j-261</i>	0.83	0.00	56
<i>One_sast-211</i>	0.88	0.00	86
<i>One_serpin</i>	0.88	0.00	85
<i>One_spf30-207</i>	0.81	0.00	52
<i>One_srp09-127</i>	0.89	0.00	97
<i>One_ssrd-135</i>	0.87	0.00	82
<i>One_STC-410</i>	0.38	0.03	7
<i>One_STR07</i>	0.50	0.02	12

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Table 13. Page 3 of 4.

Assay	Performance Measure	Contribution	Rank
<i>One_taf12-248</i>	0.89	0.00	98
<i>One_SUMO1-6</i>	0.86	0.00	70
<i>One_sys1-230</i>	0.01	0.01	20
<i>One_Tf_ex11-750</i>	0.04	0.04	5
<i>One_Tf_in3-182</i>	0.02	0.02	9
<i>One_tshB-92</i>	0.00	0.00	94
<i>One_txnip-401</i>	0.00	0.00	103
<i>One_U1002-101</i>	0.00	0.00	106
<i>One_U1003-75</i>	0.05	0.05	3
<i>One_U1004-183</i>	0.04	0.04	4
<i>One_U1009-91</i>	0.00	0.00	59
<i>One_U1010-81</i>	0.00	0.00	92
<i>One_U1012-68</i>	0.01	0.01	16
<i>One_U1013-108</i>	0.00	0.00	50
<i>One_U1014-74</i>	0.00	0.00	57
<i>One_U1016-115</i>	0.01	0.01	19
<i>One_U1017-62</i>	0.00	0.00	113
<i>One_U1021-57<sup>f</sup></i>	N/A	N/A	N/A
<i>One_U1024-197</i>	0.00	0.00	88
<i>One_U1101</i>	0.00	0.00	54
<i>One_U1102-220</i>	0.01	0.01	32
<i>One_U1103</i>	0.00	0.00	76
<i>One_U1104-138</i>	0.00	0.00	105
<i>One_U1105</i>	0.01	0.01	30
<i>One_U1201-492</i>	0.01	0.01	35
<i>One_U1202-1052</i>	0.00	0.00	79
<i>One_U1203-175</i>	0.00	0.00	44
<i>One_U1204-53</i>	0.00	0.00	69
<i>One_U1205-57</i>	0.00	0.00	93
<i>One_U1206-108</i>	0.00	0.00	74
<i>One_U1207-231<sup>e</sup></i>	N/A	N/A	N/A
<i>One_U1208-67</i>	0.00	0.00	72
<i>One_U1209-111</i>	0.01	0.01	22
<i>One_U1210-173</i>	0.00	0.00	90
<i>One_U1211-97</i>	0.00	0.00	58
<i>One_U1212-106</i>	0.01	0.01	34
<i>One_U1214-107</i>	0.00	0.00	38

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Table 13. Page 4 of 4.

Assay	Performance Measure	Contribution	Rank
<i>One_U1215-82</i>	0.89	0.00	101
<i>One_U1216-230</i>	0.70	0.01	28
<i>One_U301-92</i>	0.86	0.00	71
<i>One_U401-224</i>	0.87	0.00	80
<i>One_U404-229</i>	0.77	0.00	40
<i>One_U502-167</i>	0.88	0.00	91
<i>One_U503-170</i>	0.72	0.01	31
<i>One_U504-141</i>	0.84	0.00	63
<i>One_U508-533</i>	0.85	0.00	68
<i>One_UCA-24<sup>e</sup></i>	N/A	N/A	N/A
<i>One_vamp5-255</i>	0.79	0.00	46
<i>One_vatf-214</i>	0.89	0.00	100
<i>One_VIM-569</i>	0.85	0.00	67
<i>One_zn706-68</i>	0.89	0.00	114
<i>One_ZNF-61</i>	0.69	0.01	27
<i>One_Zp3b-49</i>	0.84	0.00	64

<sup>a</sup> These assays were dropped due to significant linkage and were not included in this analysis.

<sup>b</sup> These assays are linked and were included as a haplotype marker in this analysis.

<sup>c</sup> These assays are linked and were included as a haplotype marker in this analysis.

<sup>d</sup> These assays were dropped due to fixation in the 36 test populations and were not included in this analysis.

<sup>e</sup> These assays were dropped due to laboratory failure and were not included in this analysis.

<sup>f</sup> This assay was dropped due to failure to conform to Hardy-Weinberg expectations and was not included in this analysis.



Table 14.–Log-likelihood ratio test statistics ( $G$ ) divided by degrees of freedom for 124 SNPs (Tables 2 and 3) screened for 36 test populations (Table 1) from a hierarchical analysis of allele frequency homogeneity among populations within 9 fine-scale regions (Table 1). Fine-scale regions are abbreviated as follows: Russia (R), Norton Sound and western Bristol Bay (NS/WBB), Eastern Bristol Bay (EBB), North Peninsula (NP), Chignik and Kodiak (C/K), Cook Inlet (CI), Copper River and northern Southeast Alaska (CR/NSE), Southern Southeast Alaska (SSE), and British Columbia and Washington (BC/WA).

Assay	R	NS/WBB	EBB	NP	C/K	CI	CR/NSE	SSE	BC/WA
<i>One_ACBP-79</i>	12.89	1.81	2.66	2.87	24.51	9.53	1.98	4.84	15.44
<i>One_agt-132</i>	8.73	2.97	2.71	10.58	1.01	15.91	44.57	3.61	67.97
<i>One_aldB-152</i>	24.41	2.17	6.73	18.84	5.14	24.65	17.75	2.85	24.98
<i>One_ALDOB-135<sup>a</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_apoe-83</i>	22.32	26.00	0.88	29.82	0.40	11.57	2.05	19.80	64.40
<i>One_bckB-137<sup>b</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_c3-98</i>	7.39	4.74	3.43	12.81	7.46	15.23	41.38	0.05	2.35
<i>One_ccd16-131</i>	0.00	0.00	0.00	0.00	0.00	0.00	4.39	3.43	0.00
<i>One_CD9-269</i>	20.09	6.69	16.27	8.78	0.56	9.67	7.12	0.46	7.90
<i>One_cetn1-167</i>	17.26	39.25	0.93	23.19	0.54	15.13	19.55	0.82	4.75
<i>One_CFP1</i>	1.13	0.81	0.48	16.82	1.13	12.13	29.99	2.12	66.87
<i>One_cin-177</i>	1.20	12.66	2.99	30.81	4.52	6.30	3.54	5.35	30.90
<i>One_COI<sup>c</sup></i>	4.33	0.35	0.32	4.26	2.55	8.20	3.89	10.46	20.27
<i>One_CTGF-301</i>	0.00	9.50	2.26	0.70	3.26	5.53	2.13	2.82	9.84
<i>One_Cytb_17<sup>c</sup></i>	4.33	0.35	0.32	4.26	2.55	8.20	3.89	10.46	20.27
<i>One_Cytb_26<sup>c</sup></i>	4.33	0.35	0.32	4.26	2.55	8.20	3.89	10.46	20.27
<i>One_dds-529</i>	5.73	6.54	2.15	4.53	3.93	8.33	35.99	0.75	10.46
<i>One_DDX5-86</i>	4.10	4.48	1.04	0.79	0.90	13.60	26.14	16.87	10.64
<i>One_E2-65</i>	20.04	12.74	3.40	6.50	0.66	25.88	6.93	1.23	24.73
<i>One_gadd45-269</i>	0.97	0.00	0.00	0.00	0.00	0.68	0.00	1.07	0.00
<i>One_gdh-212</i>	5.07	14.31	7.51	0.28	2.12	1.19	11.27	2.43	52.50
<i>One_GHII-2165</i>	11.32	3.45	1.65	6.82	0.82	15.18	80.25	0.85	20.93
<i>One_ghsR-66</i>	11.63	55.95	28.79	13.97	24.58	7.48	31.11	1.19	20.87
<i>One_GPDH-201</i>	9.63	10.99	0.33	4.57	15.52	4.37	17.69	4.62	23.75
<i>One_GPDH2-187</i>	16.64	1.98	1.91	8.44	3.76	7.57	4.75	9.47	11.23
<i>One_GPH-414<sup>a</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_GTHa</i>	60.03	3.29	3.00	8.93	56.05	7.82	14.06	17.57	33.57
<i>One_HGFA-49</i>	10.25	8.50	2.67	5.00	4.50	1.85	2.98	4.57	12.98
<i>One_HpaI-71</i>	16.76	5.59	7.27	41.53	6.39	19.75	13.48	2.45	91.20
<i>One_HpaI-99</i>	5.37	3.38	1.40	5.93	44.04	7.24	15.01	2.56	26.45
<i>One_hsc71-220</i>	23.08	8.36	3.21	28.03	17.07	22.33	2.80	19.95	25.38

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Table 14. Page 2 of 4.

Assay	R	NS/WBB	EBB	NP	C/K	CI	CR/NSE	SSE	BC/WA
<i>One_Hsp47</i>	3.77	17.09	4.99	16.76	12.45	10.64	6.52	14.80	34.47
<i>One_Ig-90</i>	0.00	0.00	0.00	2.20	0.00	0.00	5.41	1.54	18.28
<i>One_IL8r-362</i>	7.06	33.75	17.55	7.81	2.01	4.78	21.33	3.31	0.35
<i>One_ins-107</i>	32.68	3.22	0.79	5.03	2.61	8.92	11.34	0.92	12.95
<i>One_KCT1-453</i>	10.95	33.70	2.98	10.93	1.80	11.10	0.25	4.45	15.90
<i>One_KPNA-422</i>	3.23	17.29	1.78	0.76	9.51	7.35	3.71	9.22	14.25
<i>One_LEI-87</i>	28.90	19.12	2.18	6.58	0.69	14.25	10.29	3.57	18.66
<i>One_leptin-92<sup>b</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_lpp1-44</i>	9.89	60.50	10.34	11.26	33.29	5.80	0.95	0.68	14.00
<i>One_MARCKS-241</i>	5.11	0.00	1.15	11.66	2.50	5.44	8.21	1.98	9.38
<i>One_metA-253</i>	0.00	0.00	0.93	0.00	2.50	1.96	12.91	11.29	66.93
<i>One_MHC2_190<sup>d</sup></i>	17.93	14.68	13.71	11.78	4.23	10.02	9.30	7.12	12.94
<i>One_MHC2_251<sup>d</sup></i>	17.93	14.68	13.71	11.78	4.23	10.02	9.30	7.12	12.94
<i>One_Mkpro-129</i>	10.86	9.28	4.03	2.37	7.07	20.88	28.62	8.19	36.77
<i>One_ODC1-196</i>	3.88	20.63	0.90	1.30	5.46	10.26	8.94	2.03	43.37
<i>One_Ots208-234</i>	1.08	5.38	1.45	13.56	2.28	5.85	8.90	0.73	34.37
<i>One_Ots213-181</i>	6.69	11.36	4.53	3.69	4.72	4.86	5.88	2.27	26.65
<i>One_p53-534</i>	0.00	0.00	1.52	2.13	17.81	4.84	12.72	3.47	16.32
<i>One_parp3-170<sup>b</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_pax7-248</i>	46.23	4.50	8.62	26.26	6.96	8.97	25.16	4.19	16.08
<i>One_PIP</i>	22.78	1.78	0.49	10.90	0.07	9.82	15.40	2.23	13.57
<i>One_ppie-74</i>	10.37	0.00	2.52	2.14	6.84	5.92	3.02	9.67	102.87
<i>One_PPM1K-118<sup>e</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_Prl2</i>	15.15	33.50	3.60	5.56	0.64	10.44	11.37	7.44	44.37
<i>One_psme2-354</i>	2.40	2.99	7.54	7.70	7.29	12.60	34.13	9.10	21.43
<i>One_rab1a-76</i>	1.31	9.08	4.85	7.88	25.91	20.87	14.00	13.61	22.75
<i>One_RAG1-103</i>	0.00	7.23	0.92	2.97	4.05	36.33	5.60	4.59	17.07
<i>One_RAG3-93</i>	10.88	3.31	2.64	19.39	8.71	14.78	13.53	3.40	17.55
<i>One_redd1-414</i>	29.63	10.60	6.49	24.63	19.87	19.73	0.79	6.55	26.62
<i>One_RFC2-102</i>	7.02	8.79	2.36	16.56	0.86	44.87	3.28	19.30	38.30
<i>One_RFC2-285</i>	6.82	2.35	2.24	12.76	0.76	6.84	9.67	9.32	2.88
<i>One_RH2op-395</i>	0.12	4.43	1.49	3.01	0.00	0.65	4.93	4.33	1.63
<i>One_rpo2j-261</i>	7.28	6.01	1.58	14.50	15.67	27.18	15.53	16.81	35.43
<i>One_sast-211</i>	1.16	2.10	3.93	5.39	1.39	7.26	21.80	3.76	2.32
<i>One_serpin-75</i>	6.18	2.66	4.98	3.36	8.30	6.20	7.72	0.47	13.09
<i>One_spf30-207</i>	24.09	10.10	4.19	12.44	1.30	15.68	6.30	0.19	56.20

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Table 14. Page 3 of 4.

Assay	R	NS/WBB	EBB	NP	C/K	CI	CR/NSE	SSE	BC/WA
<i>One_srp09-127</i>	4.51	0.81	4.80	6.99	1.19	4.52	9.87	0.77	10.54
<i>One_ssrd-135</i>	7.13	9.21	2.37	19.05	2.98	5.07	20.38	9.98	19.79
<i>One_STC-410</i>	27.25	8.26	19.24	26.89	3.60	11.39	0.43	39.58	1.43
<i>One_STR07</i>	3.39	5.54	10.34	3.50	1.66	15.12	54.10	24.02	27.15
<i>One_SUMO1-6</i>	22.02	18.09	1.64	3.01	0.01	4.39	4.74	0.65	24.43
<i>One_sys1-230</i>	36.93	2.86	4.12	13.53	3.74	5.66	29.15	0.36	8.88
<i>One_taf12-248</i>	0.92	1.45	2.87	0.81	3.86	1.82	6.66	1.30	11.92
<i>One_Tf_ex11-750</i>	45.93	15.14	8.82	19.24	6.86	21.73	7.87	2.94	9.74
<i>One_Tf_in3-182</i>	2.75	2.88	17.15	11.36	22.07	2.72	2.80	0.03	23.19
<i>One_tshB-92</i>	7.64	31.63	0.50	1.58	1.67	9.93	5.93	0.77	8.73
<i>One_txnip-401</i>	0.00	0.91	1.06	0.00	0.00	0.65	13.36	13.63	7.38
<i>One_U1002-101</i>	0.00	0.00	0.00	0.00	0.00	0.00	7.96	8.60	0.00
<i>One_U1003-75</i>	58.23	6.22	3.39	11.79	14.06	18.25	14.33	6.11	20.80
<i>One_U1004-183</i>	27.21	24.10	1.91	14.74	74.95	68.95	20.16	5.87	22.06
<i>One_U1009-91</i>	18.37	11.24	7.64	15.54	2.78	3.02	30.25	15.31	33.67
<i>One_U1010-81</i>	7.34	0.00	6.05	0.26	2.53	7.36	4.30	6.96	19.85
<i>One_U1012-68</i>	14.50	13.21	11.78	0.09	5.07	35.02	39.50	5.42	48.40
<i>One_U1013-108</i>	23.55	0.56	2.77	11.14	1.35	13.98	2.26	25.82	13.63
<i>One_U1014-74</i>	8.18	9.69	2.53	7.98	4.21	7.77	6.85	1.01	17.62
<i>One_U1016-115</i>	38.40	7.13	2.72	5.11	11.40	18.82	26.83	16.03	36.37
<i>One_U1017-62</i>	5.54	0.00	0.91	2.20	0.68	12.27	5.68	1.64	11.22
<i>One_U1021-57<sup>f</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_U1024-197</i>	10.94	4.51	1.50	5.51	0.49	2.21	12.96	9.64	32.59
<i>One_U1101</i>	13.18	7.55	7.30	2.21	2.73	7.11	4.35	10.77	74.97
<i>One_U1102-220</i>	8.86	4.16	14.34	19.23	6.28	8.19	0.37	9.31	10.15
<i>One_U1103</i>	6.32	0.82	0.92	4.41	3.48	2.37	36.34	19.26	10.94
<i>One_U1104-138</i>	0.00	7.32	1.89	1.10	0.00	0.00	0.00	0.00	0.00
<i>One_U1105</i>	32.03	17.16	11.44	7.54	1.33	26.95	3.65	8.15	8.00
<i>One_U1201-492</i>	22.33	9.76	12.18	7.78	10.80	13.86	3.10	2.54	75.70
<i>One_U1202-1052</i>	33.23	5.67	3.97	4.05	0.92	11.92	2.67	0.28	18.21
<i>One_U1203-175</i>	12.56	1.97	0.39	0.24	3.79	14.07	0.85	16.19	17.19
<i>One_U1204-53</i>	7.83	11.95	1.28	1.74	0.98	17.05	0.22	15.51	14.31
<i>One_U1205-57</i>	2.63	3.13	6.25	1.00	8.75	2.51	26.65	3.86	7.50
<i>One_U1206-108</i>	41.67	9.01	0.59	1.23	1.35	6.19	21.16	25.49	26.41
<i>One_U1207-231<sup>e</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_U1208-67</i>	3.36	7.40	3.09	13.18	16.55	15.60	31.43	0.07	24.71
<i>One_U1209-111</i>	1.38	30.75	6.80	13.85	14.12	4.00	13.92	0.51	19.76

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Assay	R	NS/WBB	EBB	NP	C/K	CI	CR/NSE	SSE	BC/WA
<i>One_U1210-173</i>	6.72	9.23	3.65	30.52	6.46	9.93	6.80	4.17	6.00
<i>One_U1211-97</i>	2.77	3.05	2.65	11.73	7.45	17.07	0.96	0.28	0.00
<i>One_U1212-106</i>	24.66	7.25	7.06	11.17	5.89	12.46	58.15	6.62	51.47
<i>One_U1214-107</i>	27.38	24.27	1.50	8.48	2.64	4.80	14.05	8.63	93.53
<i>One_U1215-82</i>	6.45	8.43	0.10	55.15	1.16	2.37	19.00	4.26	9.11
<i>One_U1216-230</i>	12.17	12.08	9.78	0.80	21.67	7.40	0.33	6.65	37.13
<i>One_U301-92</i>	1.11	15.98	4.54	1.39	4.39	4.94	24.47	23.16	12.99
<i>One_U401-224</i>	8.38	4.01	1.82	9.83	1.11	1.17	17.77	6.08	16.32
<i>One_U404-229</i>	0.91	0.00	6.55	5.73	0.34	4.46	38.87	8.00	6.97
<i>One_U502-167</i>	0.00	5.89	2.93	2.20	1.09	4.37	11.03	7.27	42.73
<i>One_U503-170</i>	29.31	14.79	5.32	36.69	2.21	24.25	13.37	6.10	19.16
<i>One_U504-141</i>	11.30	5.82	12.64	0.39	3.70	6.14	29.59	17.21	64.50
<i>One_U508-533</i>	2.17	0.86	4.13	4.15	0.00	16.62	29.26	4.21	26.10
<i>One_UCA-24<sup>e</sup></i>	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A
<i>One_vamp5-255</i>	1.38	34.58	0.28	16.87	3.49	26.03	27.93	1.61	18.58
<i>One_vatf-214</i>	5.80	0.80	0.73	1.23	1.44	14.24	9.05	8.26	11.94
<i>One_VIM-569</i>	40.20	16.58	0.36	18.06	2.93	17.90	5.19	0.16	8.59
<i>One_zn706-68</i>	0.00	0.00	0.00	1.10	2.68	1.34	0.00	0.00	0.00
<i>One_ZNF-61</i>	28.86	41.23	1.10	17.48	7.50	8.61	2.11	0.11	20.99
<i>One_Zp3b-49</i>	0.90	2.92	0.22	9.16	13.07	19.40	1.04	6.15	22.49
Average	12.58	9.70	4.31	9.55	6.81	11.15	13.78	6.85	23.72

<sup>a</sup> These assays were dropped due to significant linkage and were not included in this analysis.

<sup>b</sup> These assays were dropped due to fixation in the 36 test populations and were not included in this analysis.

<sup>c</sup> These assays are linked and were included as a haplotype marker in this analysis.

<sup>d</sup> These assays are linked and were included as a haplotype marker in this analysis.

<sup>e</sup> These assays were dropped due to laboratory failure and were not included in this analysis.

<sup>f</sup> This assay was dropped due to failure to conform to Hardy-Weinberg expectations and was not included in this analysis.

Table 15.–The percentage of variation attributable to allele frequency differences within populations ( $G_{WP}$ ), among populations within fine-scale regions ( $G_{PF}$ ), among fine-scale regions within broad-scale regions ( $G_{FB}$ ), and among broad-scale regions within the total ( $G_{BT}$ ) for 124 SNPs (Tables 2 and 3) screened for 36 test populations (Table 1) of sockeye salmon.

Assay	$G_{WP}$	$G_{PF}$	$G_{FB}$	$G_{BT}$
<i>One_ACBP-79</i>	88%	3%	0%	8%
<i>One_agt-132</i>	89%	7%	0%	4%
<i>One_aldB-152</i>	91%	6%	1%	1%
<i>One_ALDOB-135<sup>a</sup></i>	N/A	N/A	N/A	N/A
<i>One_apoe-83</i>	78%	8%	3%	11%
<i>One_bckB-137<sup>b</sup></i>	N/A	N/A	N/A	N/A
<i>One_c3-98</i>	93%	4%	1%	1%
<i>One_ccd16-131</i>	100%	0%	0%	0%
<i>One_CD9-269</i>	95%	4%	0%	1%
<i>One_cetn1-167</i>	90%	6%	1%	3%
<i>One_CFP1</i>	85%	7%	2%	6%
<i>One_cin-177</i>	92%	4%	3%	1%
<i>One_CO1<sup>c</sup></i>	76%	14%	3%	7%
<i>One_CTGF-301</i>	96%	2%	1%	1%
<i>One_Cytb_17<sup>c</sup></i>	76%	14%	3%	7%
<i>One_Cytb_26<sup>c</sup></i>	76%	14%	3%	7%
<i>One_dds-529</i>	95%	3%	1%	1%
<i>One_DDX5-86</i>	94%	3%	1%	2%
<i>One_E2-65</i>	92%	5%	2%	1%
<i>One_gadd45-269</i>	100%	0%	0%	0%
<i>One_gdh-212</i>	92%	4%	1%	3%
<i>One_GHII-2165</i>	71%	6%	3%	21%
<i>One_ghsR-66</i>	88%	9%	2%	2%
<i>One_GPDH-201</i>	91%	3%	1%	5%
<i>One_GPDH2-187</i>	87%	3%	1%	10%
<i>One_GPH-414<sup>a</sup></i>	N/A	N/A	N/A	N/A
<i>One_GTHa</i>	89%	7%	2%	2%
<i>One_HGFA-49</i>	96%	2%	1%	1%
<i>One_HpaI-71</i>	84%	8%	2%	5%
<i>One_HpaI-99</i>	72%	7%	11%	10%
<i>One_hsc71-220</i>	90%	7%	1%	2%
<i>One_Hsp47</i>	89%	5%	1%	5%
<i>One_Ig-90</i>	88%	4%	1%	7%
<i>One_IL8r-362</i>	92%	5%	1%	2%

-continued-

Table 15. Page 2 of 4.

Assay	$G_{WP}$	$G_{PR}$	$G_{RB}$	$G_{BT}$
<i>One_ins-107</i>	94%	4%	1%	1%
<i>One_KCT1-453</i>	92%	5%	2%	1%
<i>One_KPNA-422</i>	94%	3%	2%	1%
<i>One_LEI-87</i>	86%	5%	1%	8%
<i>One_leptin-92<sup>b</sup></i>	N/A	N/A	N/A	N/A
<i>One_lpp1-44</i>	87%	7%	1%	4%
<i>One_MARCKS-241</i>	96%	3%	1%	1%
<i>One_metA-253</i>	63%	11%	6%	20%
<i>One_MHC2_190<sup>d</sup></i>	73%	18%	4%	5%
<i>One_MHC2_251<sup>d</sup></i>	73%	18%	4%	5%
<i>One_Mkpro-129</i>	86%	5%	2%	7%
<i>One_ODC1-196</i>	90%	4%	1%	5%
<i>One_Ots208-234</i>	81%	4%	1%	14%
<i>One_Ots213-181</i>	90%	3%	0%	7%
<i>One_p53-534</i>	88%	3%	2%	6%
<i>One_parp3-170<sup>b</sup></i>	N/A	N/A	N/A	N/A
<i>One_pax7-248</i>	92%	6%	0%	2%
<i>One_PIP</i>	94%	3%	2%	1%
<i>One_ppie-74</i>	69%	25%	3%	3%
<i>One_PPM1K-118<sup>e</sup></i>	N/A	N/A	N/A	N/A
<i>One_Prl2</i>	90%	6%	3%	1%
<i>One_psme2-354</i>	89%	4%	4%	2%
<i>One_rab1a-76</i>	92%	6%	1%	1%
<i>One_RAG1-103</i>	86%	8%	0%	6%
<i>One_RAG3-93</i>	93%	4%	1%	2%
<i>One_redd1-414</i>	85%	6%	1%	8%
<i>One_RFC2-102</i>	92%	7%	0%	1%
<i>One_RFC2-285</i>	95%	2%	1%	2%
<i>One_RH2op-395</i>	99%	1%	0%	0%
<i>One_rpo2j-261</i>	91%	6%	2%	1%
<i>One_sast-211</i>	94%	3%	2%	2%
<i>One_serpin-75</i>	96%	2%	1%	0%
<i>One_spf30-207</i>	91%	6%	0%	2%
<i>One_srp09-127</i>	89%	2%	2%	7%
<i>One_ssrd-135</i>	94%	4%	1%	1%
<i>One_STC-410</i>	78%	5%	5%	12%

-continued-

Table 15. Page 3 of 4.

Assay	$G_{WP}$	$G_{PR}$	$G_{RB}$	$G_{BT}$
<i>One_STR07</i>	87%	5%	5%	3%
<i>One_SUMO1-6</i>	91%	3%	1%	5%
<i>One_sys1-230</i>	91%	4%	1%	4%
<i>One_taf12-248</i>	90%	2%	3%	5%
<i>One_Tf_ex11-750</i>	78%	5%	0%	16%
<i>One_Tf_in3-182</i>	80%	5%	2%	13%
<i>One_tshB-92</i>	95%	4%	0%	0%
<i>One_txnip-401</i>	86%	3%	3%	7%
<i>One_U1002-101</i>	95%	2%	1%	1%
<i>One_U1003-75</i>	79%	7%	2%	12%
<i>One_U1004-183</i>	69%	11%	3%	17%
<i>One_U1009-91</i>	91%	5%	2%	2%
<i>One_U1010-81</i>	94%	4%	2%	1%
<i>One_U1012-68</i>	89%	8%	0%	2%
<i>One_U1013-108</i>	92%	4%	1%	2%
<i>One_U1014-74</i>	94%	3%	1%	2%
<i>One_U1016-115</i>	86%	7%	3%	4%
<i>One_U1017-62</i>	93%	4%	1%	2%
<i>One_U1021-57<sup>f</sup></i>	N/A	N/A	N/A	N/A
<i>One_U1024-197</i>	91%	5%	0%	4%
<i>One_U1101</i>	92%	6%	1%	1%
<i>One_U1102-220</i>	89%	3%	3%	5%
<i>One_U1103</i>	87%	6%	4%	3%
<i>One_U1104-138</i>	95%	2%	1%	1%
<i>One_U1105</i>	87%	6%	1%	5%
<i>One_U1201-492</i>	91%	7%	1%	1%
<i>One_U1202-1052</i>	89%	4%	0%	7%
<i>One_U1203-175</i>	93%	3%	1%	3%
<i>One_U1204-53</i>	92%	4%	1%	2%
<i>One_U1205-57</i>	96%	3%	0%	1%
<i>One_U1206-108</i>	91%	6%	0%	3%
<i>One_U1207-231<sup>e</sup></i>	N/A	N/A	N/A	N/A
<i>One_U1208-67</i>	88%	5%	2%	5%
<i>One_U1209-111</i>	89%	5%	2%	4%
<i>One_U1210-173</i>	92%	4%	0%	3%
<i>One_U1211-97</i>	92%	3%	0%	4%
<i>One_U1212-106</i>	88%	7%	2%	2%

-continued-

Table 15. Page 4 of 4.

Assay	$G_{WP}$	$G_{PR}$	$G_{RB}$	$G_{BT}$
<i>One_U1214-107</i>	74%	15%	4%	7%
<i>One_U1215-82</i>	95%	4%	0%	1%
<i>One_U1216-230</i>	91%	5%	2%	2%
<i>One_U301-92</i>	92%	4%	3%	1%
<i>One_U401-224</i>	89%	2%	2%	6%
<i>One_U404-229</i>	86%	4%	1%	9%
<i>One_U502-167</i>	89%	9%	1%	2%
<i>One_U503-170</i>	91%	7%	1%	2%
<i>One_U504-141</i>	90%	7%	2%	1%
<i>One_U508-533</i>	89%	6%	1%	4%
<i>One_UCA-24<sup>e</sup></i>	N/A	N/A	N/A	N/A
<i>One_vamp5-255</i>	90%	7%	1%	2%
<i>One_vatf-214</i>	90%	3%	4%	4%
<i>One_VIM-569</i>	92%	5%	1%	2%
<i>One_zn706-68</i>	94%	0%	0%	0%
<i>One_ZNF-61</i>	87%	6%	0%	6%
<i>One_Zp3b-49</i>	81%	4%	1%	14%

<sup>a</sup> These assays were dropped due to significant linkage and were not included in this analysis.

<sup>b</sup> These assays were dropped due to fixation in the 36 test populations and were not included in this analysis.

<sup>c</sup> These assays are linked and were included as a haplotype marker in this analysis.

<sup>d</sup> These assays are linked and were included as a haplotype marker in this analysis.

<sup>e</sup> These assays were dropped due to laboratory failure and were not included in this analysis.

<sup>f</sup> This assay was dropped due to failure to conform to Hardy-Weinberg expectations and was not included in this analysis.



Table 16.–Proposed test collections for marker selection in chum salmon.

Region	Population	Collection	Sample size	Lat.	Long.
Japan	Tokachi River	CMTOKA02	80	42.6950	-143.6653
	Gakko River early	CMGAKK03E	80	39.0525	-139.8864
Russia	Amur River summer	CMAMU01	95	53.1100	-140.7400
	Palana River	CMPALA98	95	59.0667	-159.8333
Kotzebue Sound	Kiana River	CMKIAN04	95	66.9728	-160.4269
	Kelly Lake	CMKEL91	95	67.9187	-162.3501
Norton Sound	American River	CMAMER04	95	65.4245	-165.7849
	Unalakleet River	CMUNAL04	95	63.8703	-160.7859
Yukon summer	Anvik River	CMOTT93	95	63.2425	-160.6972
	Nulato River	CMNUL03	95	64.7356	-158.1870
Yukon fall	Pelly River	CMPEL93	84	62.5500	-136.7500
	Kluane River	CMKLUA01	95	61.6222	-139.3912
Kuskokwim summer	George River	CMGEO96	95	61.8975	-157.7135
	Goodnews River weir	CMGOO91	95	59.1028	-161.5610
Kuskokwim fall	Windy Fork	CMWINDF08	95	62.6944	-154.5926
	Big River	CMBIGR08	95	62.6063	-155.0135
Western Bristol Bay	Togiak River	CMTOG93	95	59.0783	-160.3372
	Mulchatna River	CMMUL94	95	59.6449	-157.1168
Eastern Bristol Bay	Naknek River (Big Cr)	CMBRIB93	80	58.2926	-157.5340
	Meshik River	CMMES92	78	56.7910	-158.6617
Alaska Peninsula	Frosty Creek	CMFRO92	95	55.1933	-162.8604
	Canoe Bay Creek	CMCAN92	95	55.7250	-161.2188
Southcentral Alaska	Lake Creek	CMLAK96	95	61.9060	-150.9089
	Olsen Creek	CMPWS95A	95	60.7596	-146.1747
Southeast Alaska	Chilkat River - 24Mile	CM24MI06	95	59.4204	-135.9495
	North Arm Creek	CMNARM06S	95	56.6855	-132.3081
British Columbia	Kitimat River	CMKITIM06	95	54.0000	-128.6667
	Kitwanga River	CMKITW06	95	55.1000	-128.0834
Washington	Nisqually River Hatchery	CMNISQ04	95	47.0959	-122.6960
	Elwha River	CMELWH04	95	48.1452	-123.5640



## FIGURES

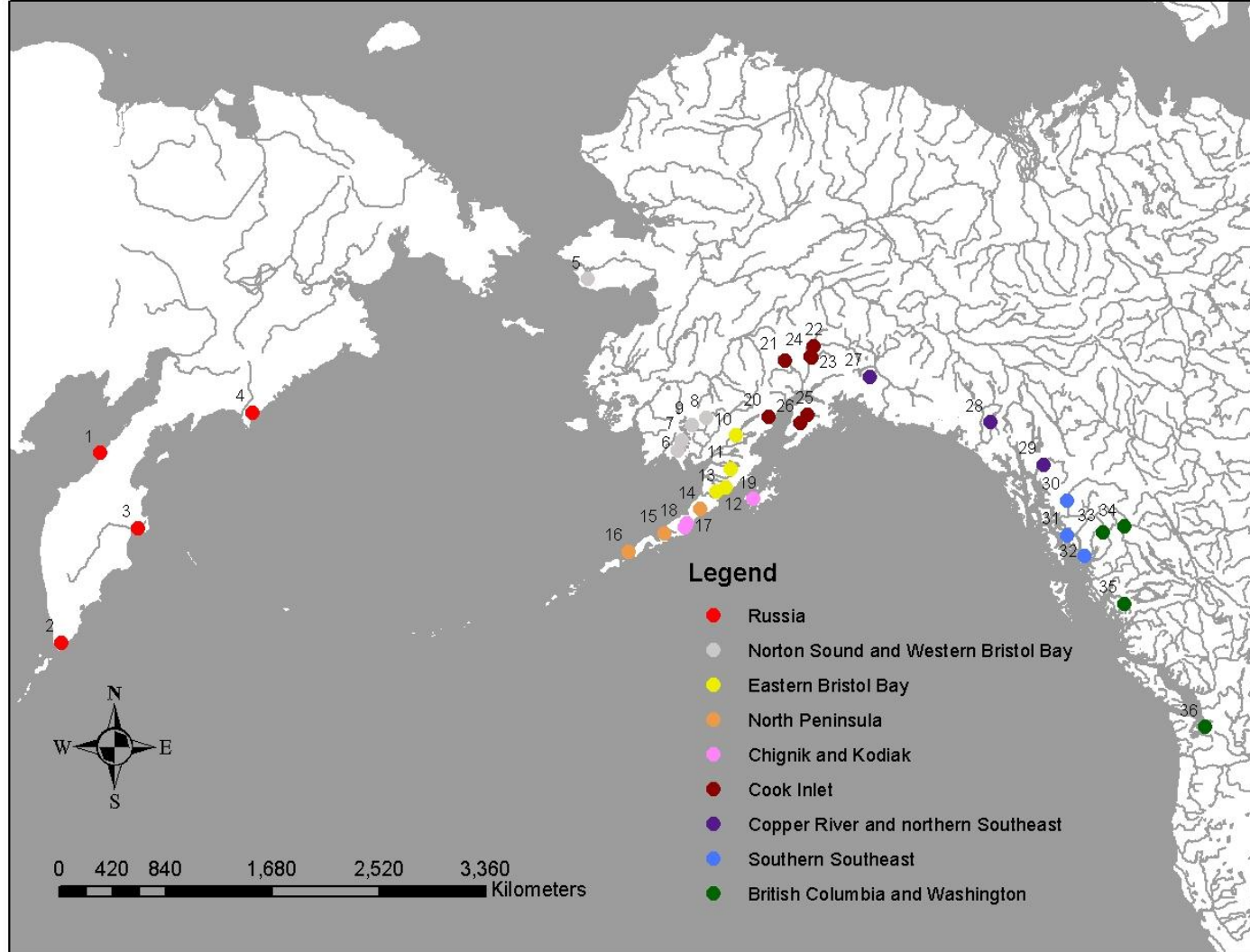


Figure 1.—Backbone collections from 9 fine-scale geographic regions (Table 1) genotyped for 124 SNPs (Table 2). See text and Table 1 for details.

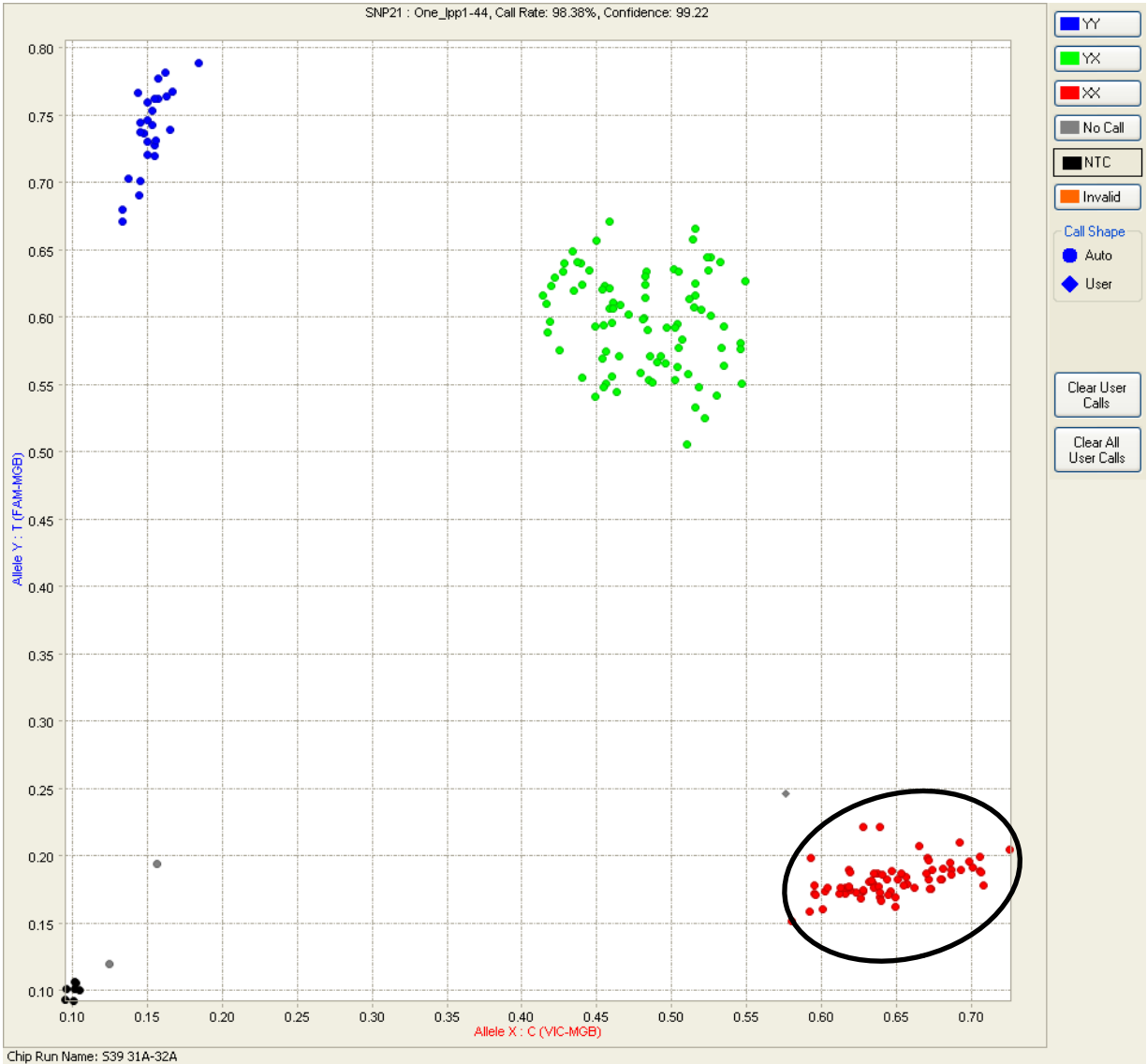


Figure 2.—An image from our genotyping software exhibiting an assay with tight genotype clusters (circle), i.e., one in which the spatial variation among individuals of a genotype is low. Individuals homozygous for genotype AA are colored red, heterozygous for AB are green, and homozygous for BB are blue. Individuals that failed to amplify or are of an uncertain genotype are grey and no template controls (no DNA) are black.

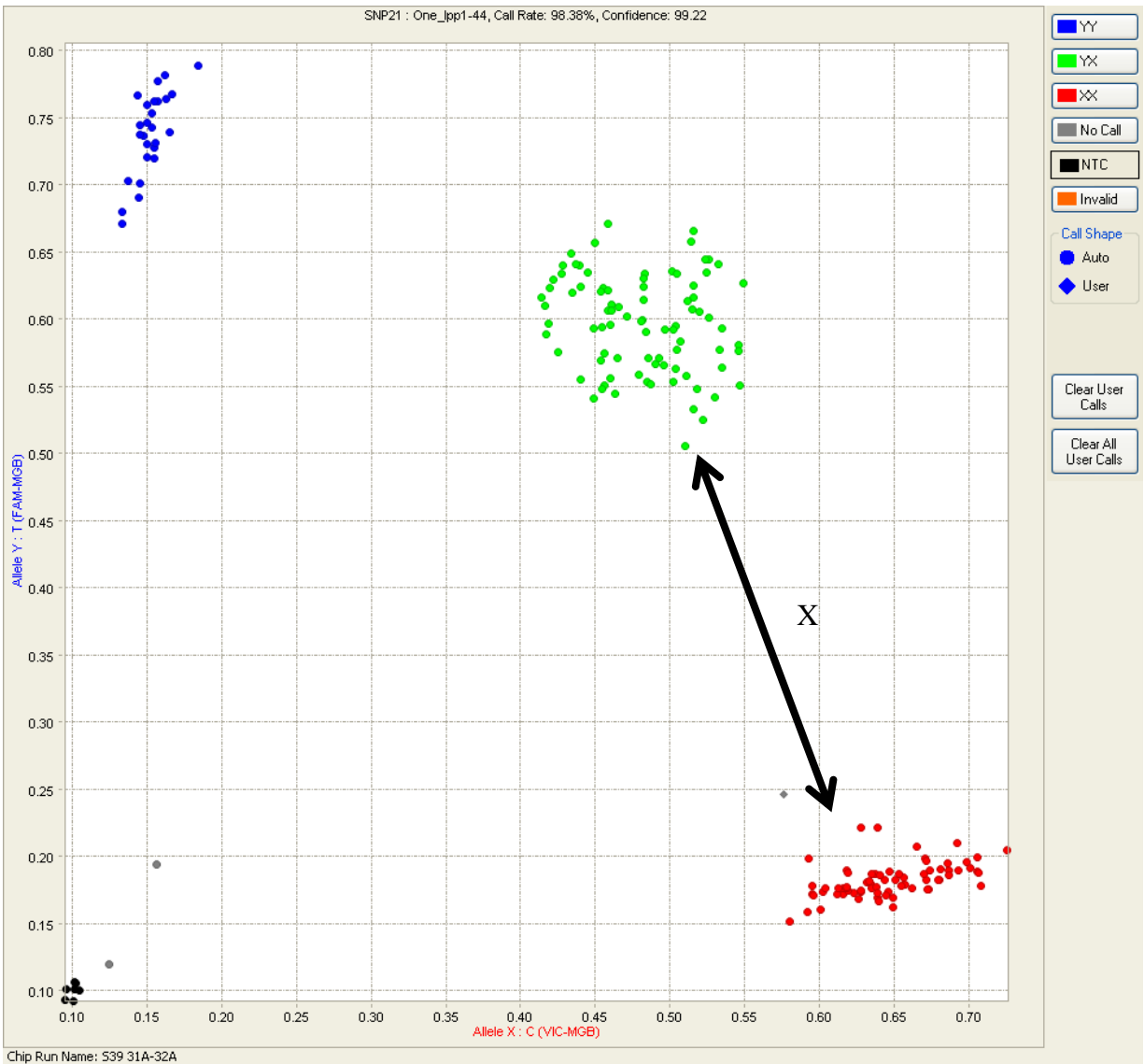


Figure 3.—An image from our genotyping software exhibiting an assay with widely separated genotype clusters, i.e., one in which the separation between cluster edges (X) is high.

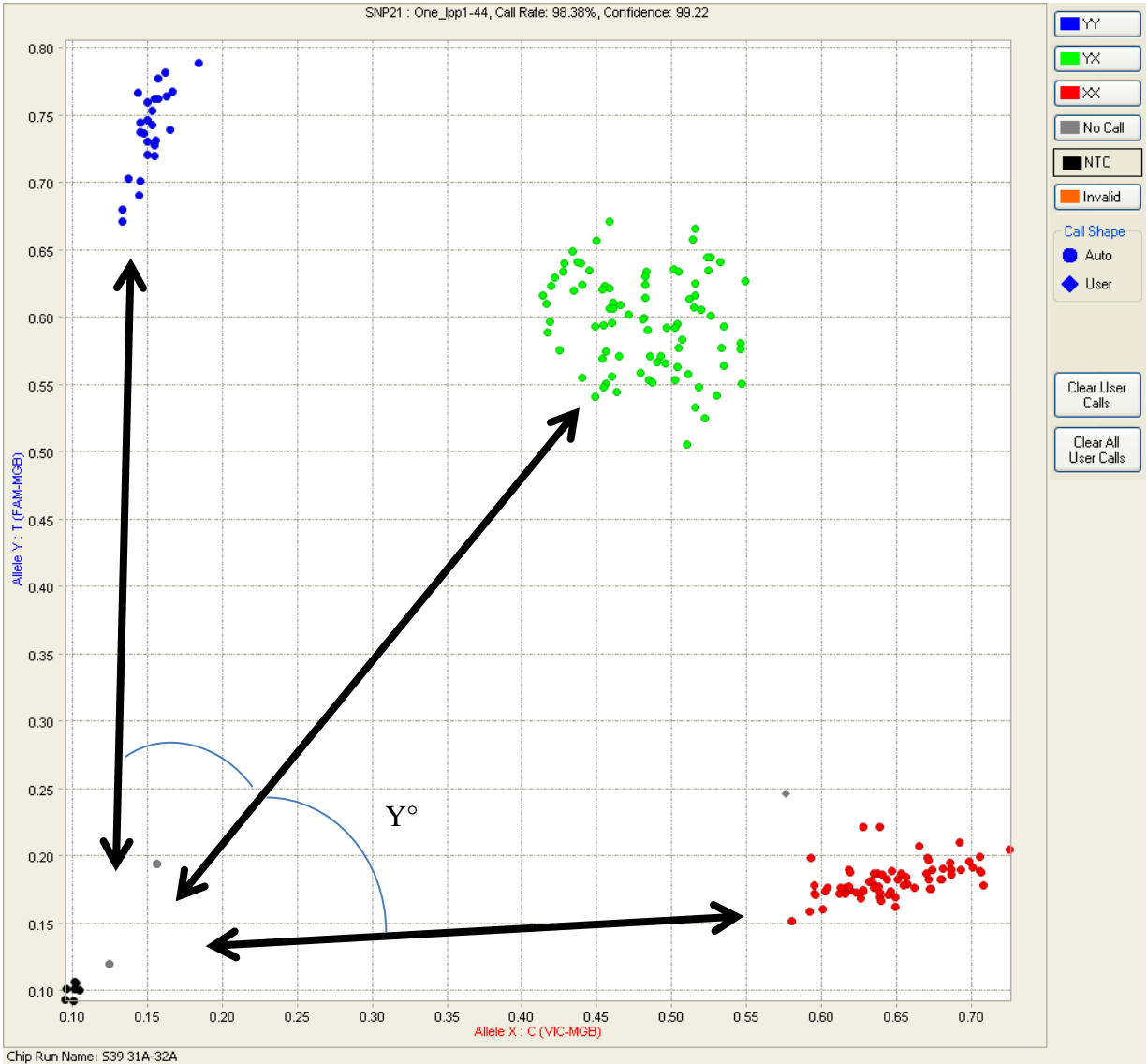


Figure 4.—An image from our genotyping software exhibiting an assay with widely separated genotype cluster alignment, i.e., one in which the degree of separation between cluster axes from the origin ( $Y$ ) is high.

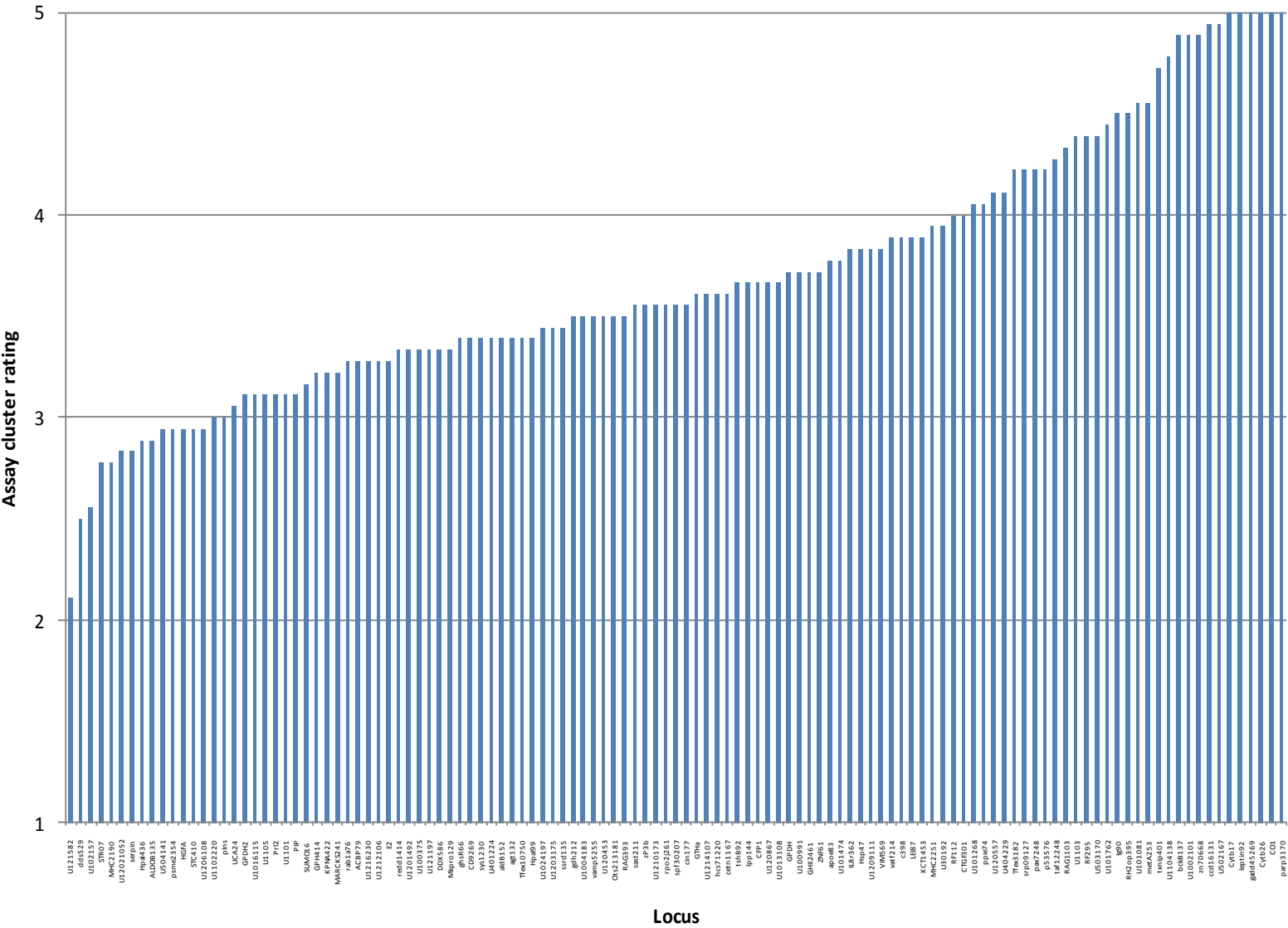
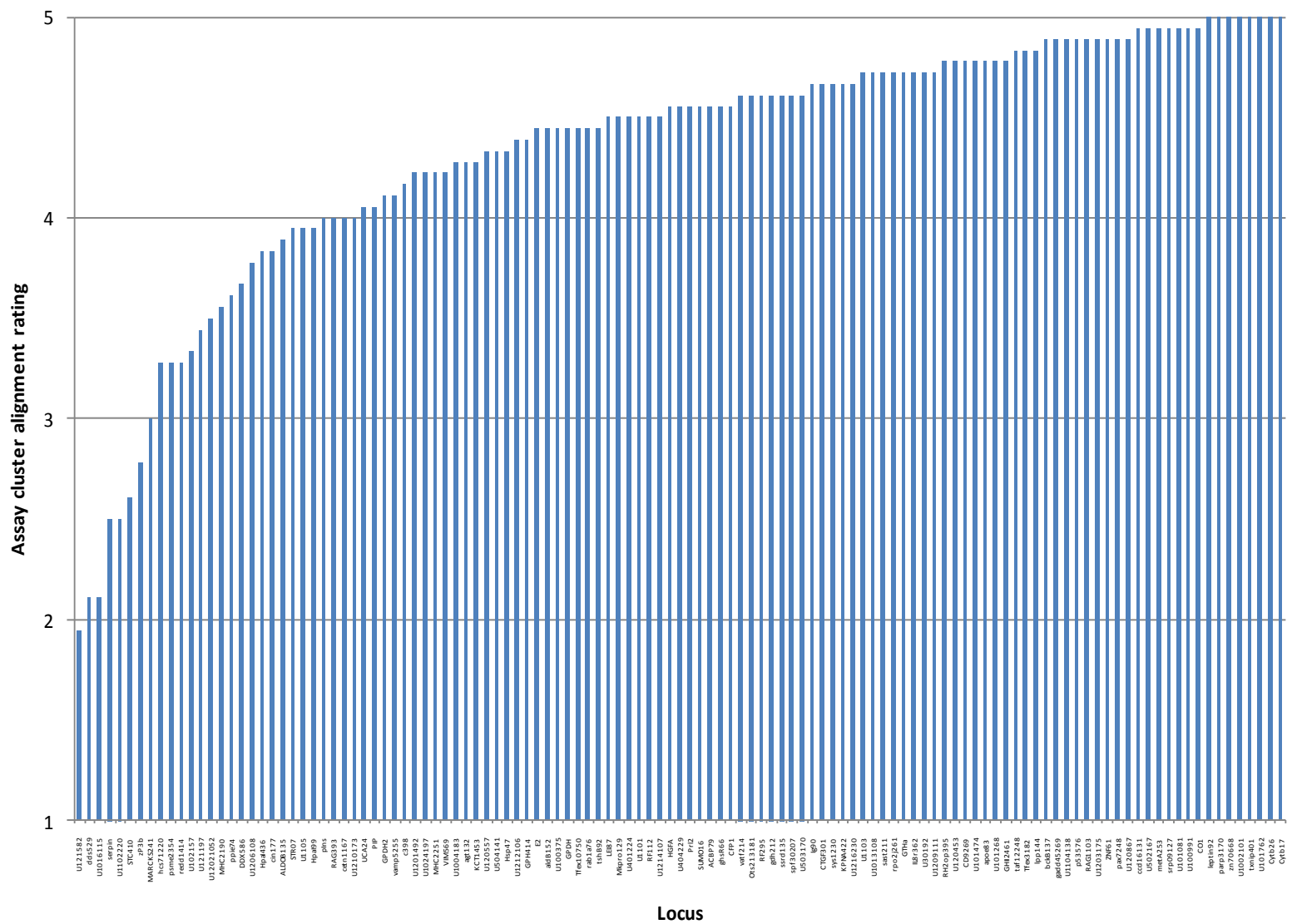




Figure 6.—The average space between cluster rating of 124 SNPs (Tables 2 and 3) screened for 36 test populations (Table 1) ordered from lowest rating (Left) to highest rating (Right).



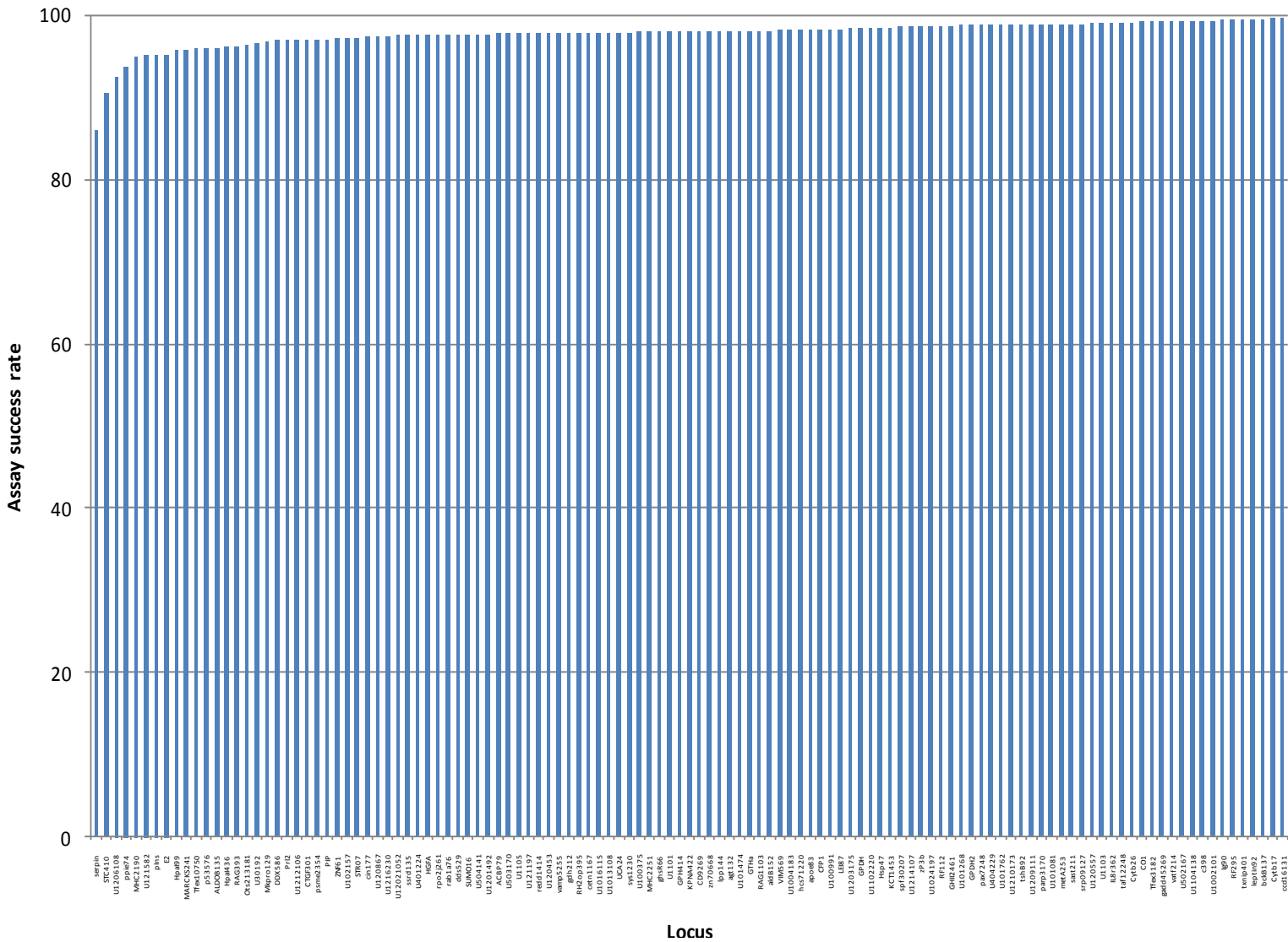


Figure 8.—The average success rate (%) of 124 SNPs (Tables 2 and 3) screened for 36 test populations (Table 1) ordered from lowest rating (Left) to highest rating (Right).

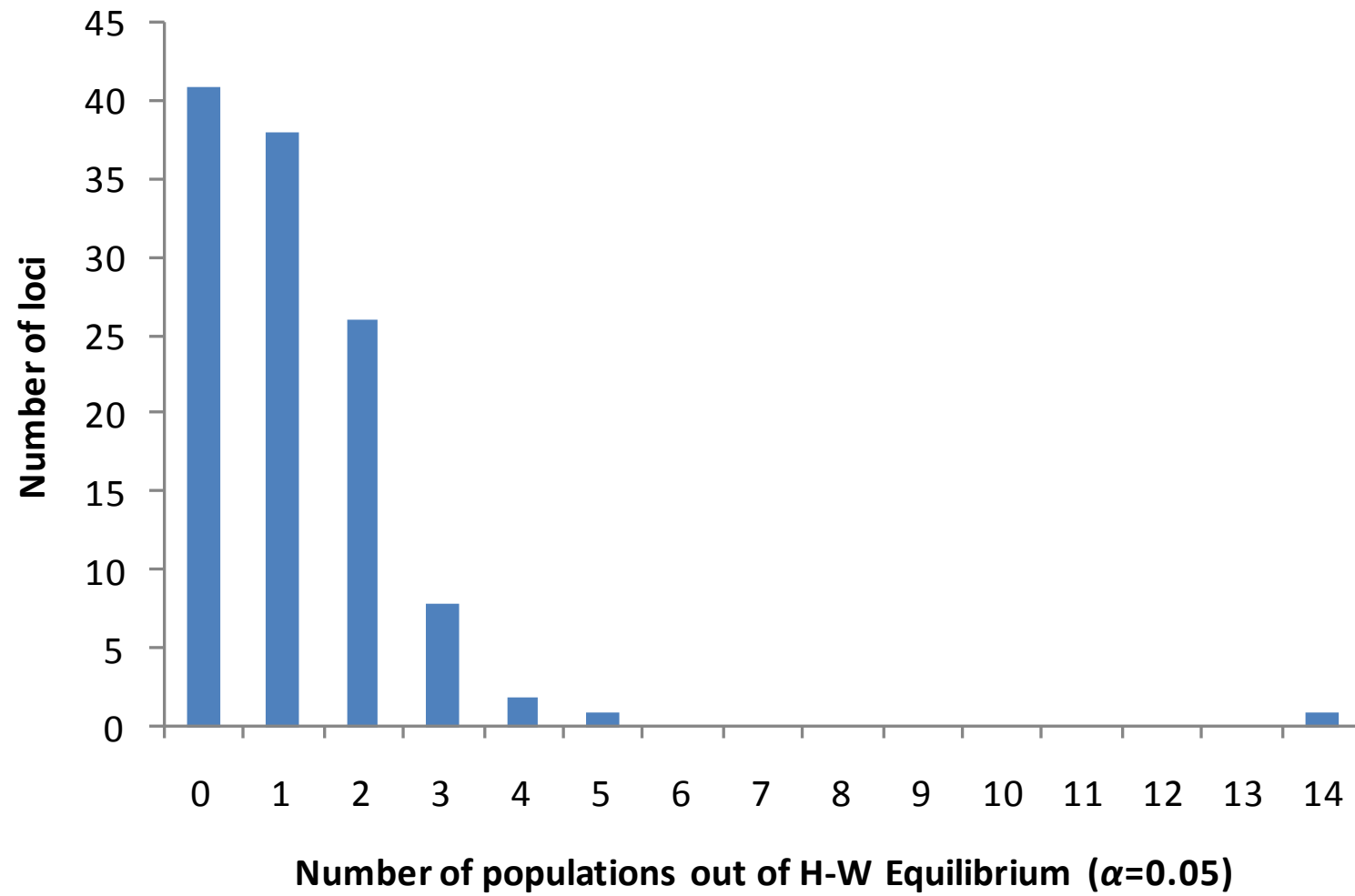
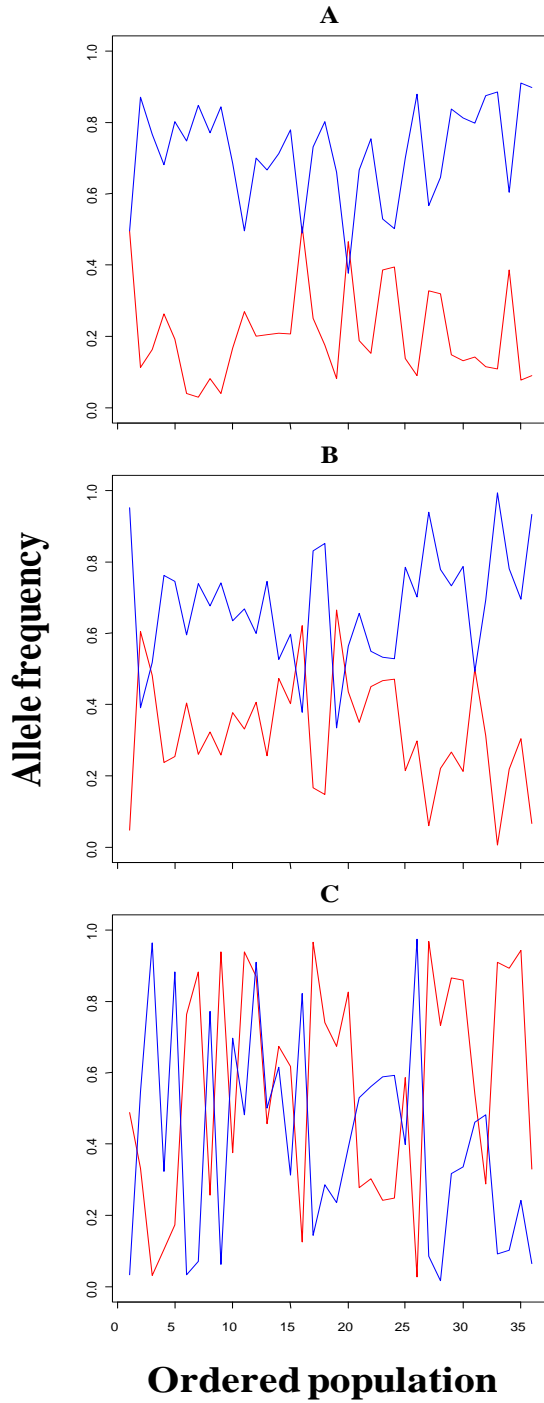
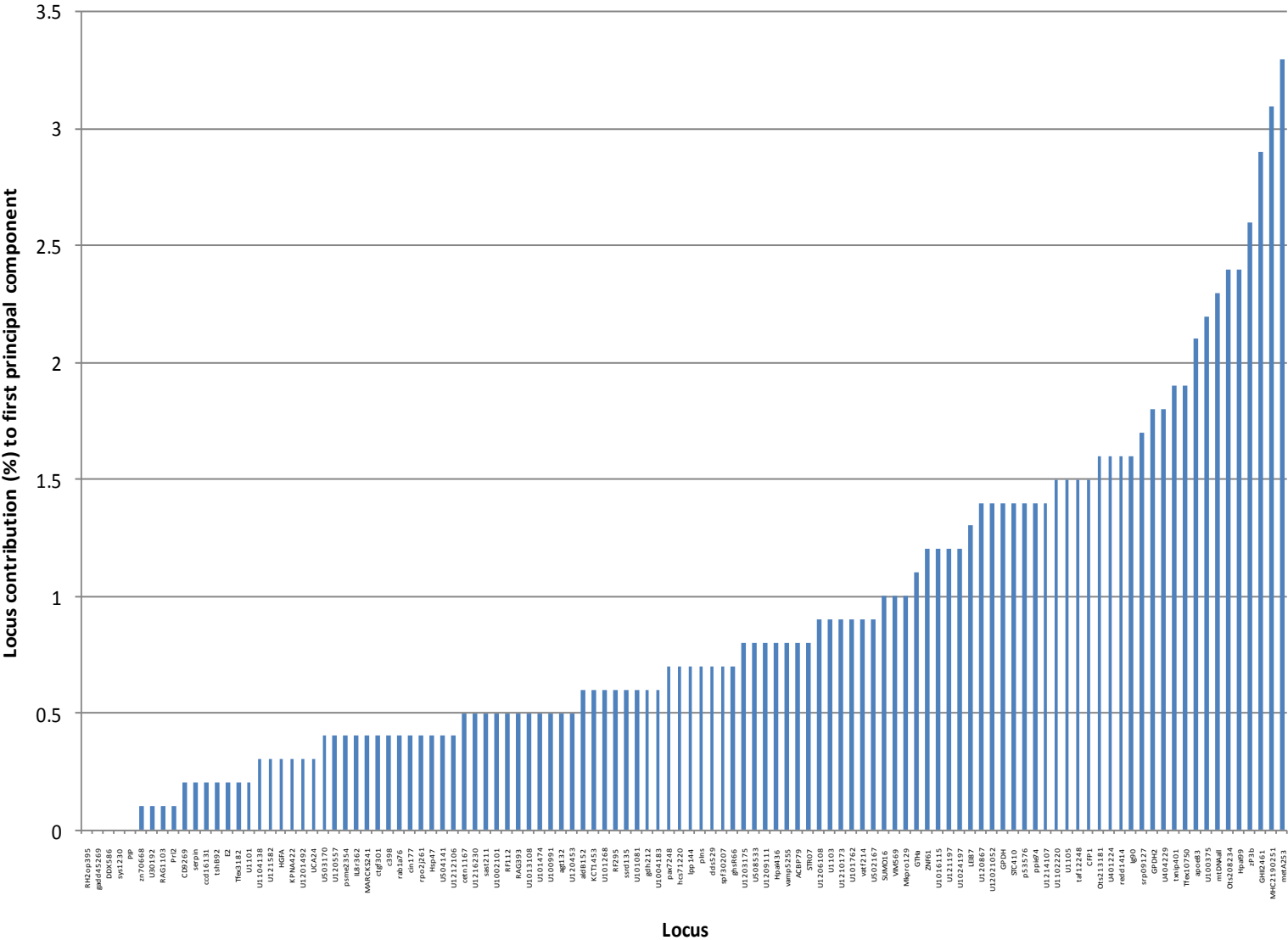


Figure 9.—Number of markers that were out of H-W equilibrium ( $\alpha = 0.05$ ) for 0 to 14 populations. By chance, the one would expect 1.8 populations to be out of H-W expectation at this criterion (i.e.,  $36 \text{ populations} \times 0.05$ ).



**Minor allele frequency for SNP1**  
**Major allele frequency for SNP2**

Figure 10.—Minor (SNP1 in red) and major (SNP2 in blue) allele frequencies for three pairs of SNPs exhibiting linkage disequilibrium: *One\_aldB-152/One\_ALDOB-135* (A), *One\_GPH-414/ One\_GTHa* (B), and *One\_MHC2\_190/ One\_MHC2\_251* (C). Note that the frequencies of these two alleles for the first two pairs of loci are mirror images of each other, indicating close-to-perfect phasing, whereas the last locus set does not. The lack of tight phasing between linked loci is indicative of locus pairs that are useful in MSA as a combined set.



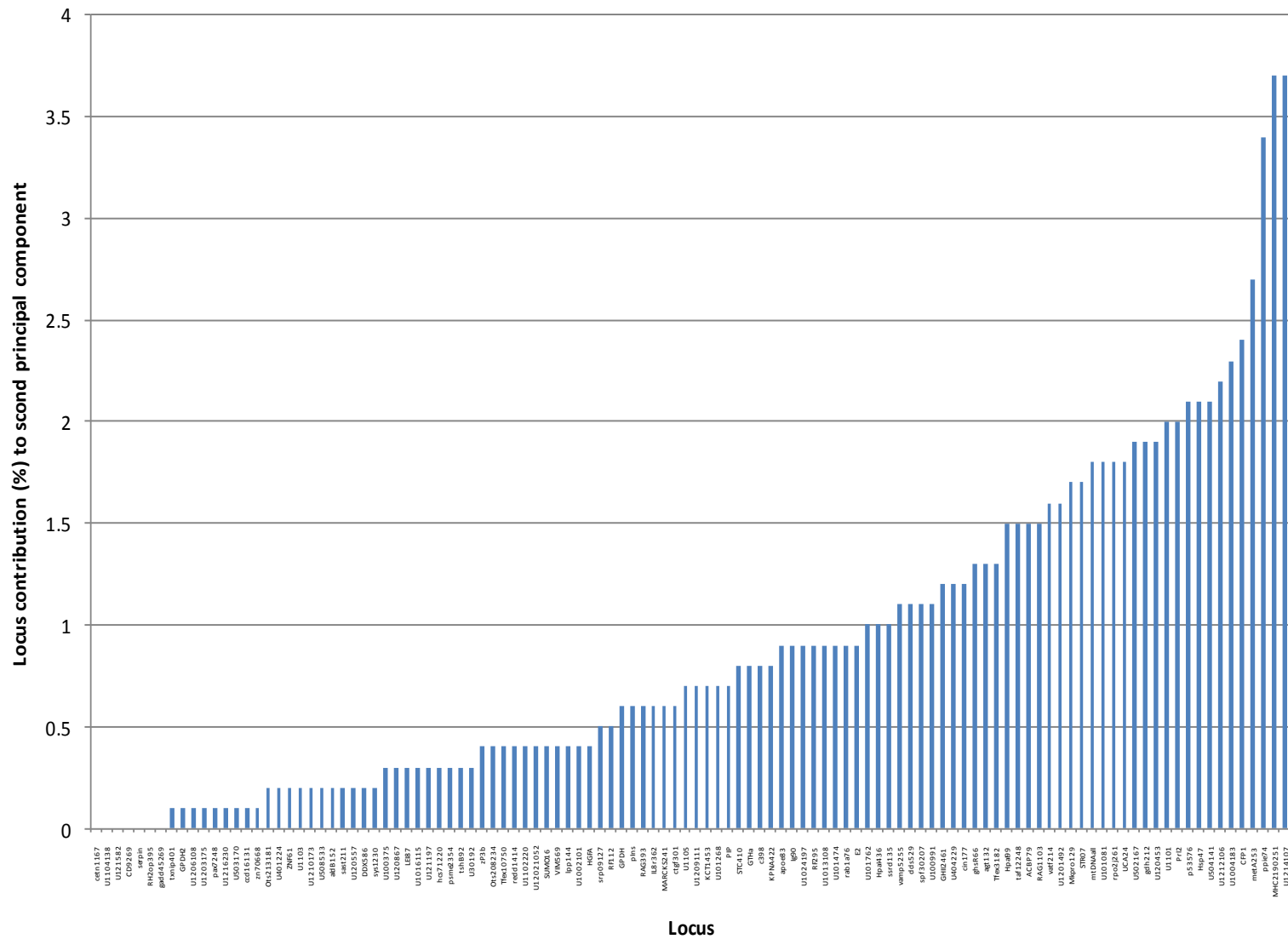


Figure 12.—The contribution to principal component 2 (%) for each of 114 SNPs included in a principle component analysis of 36 test populations (Table 1) ordered from lowest contribution (Left) to highest contribution (Right). See text and Table 7 for details.

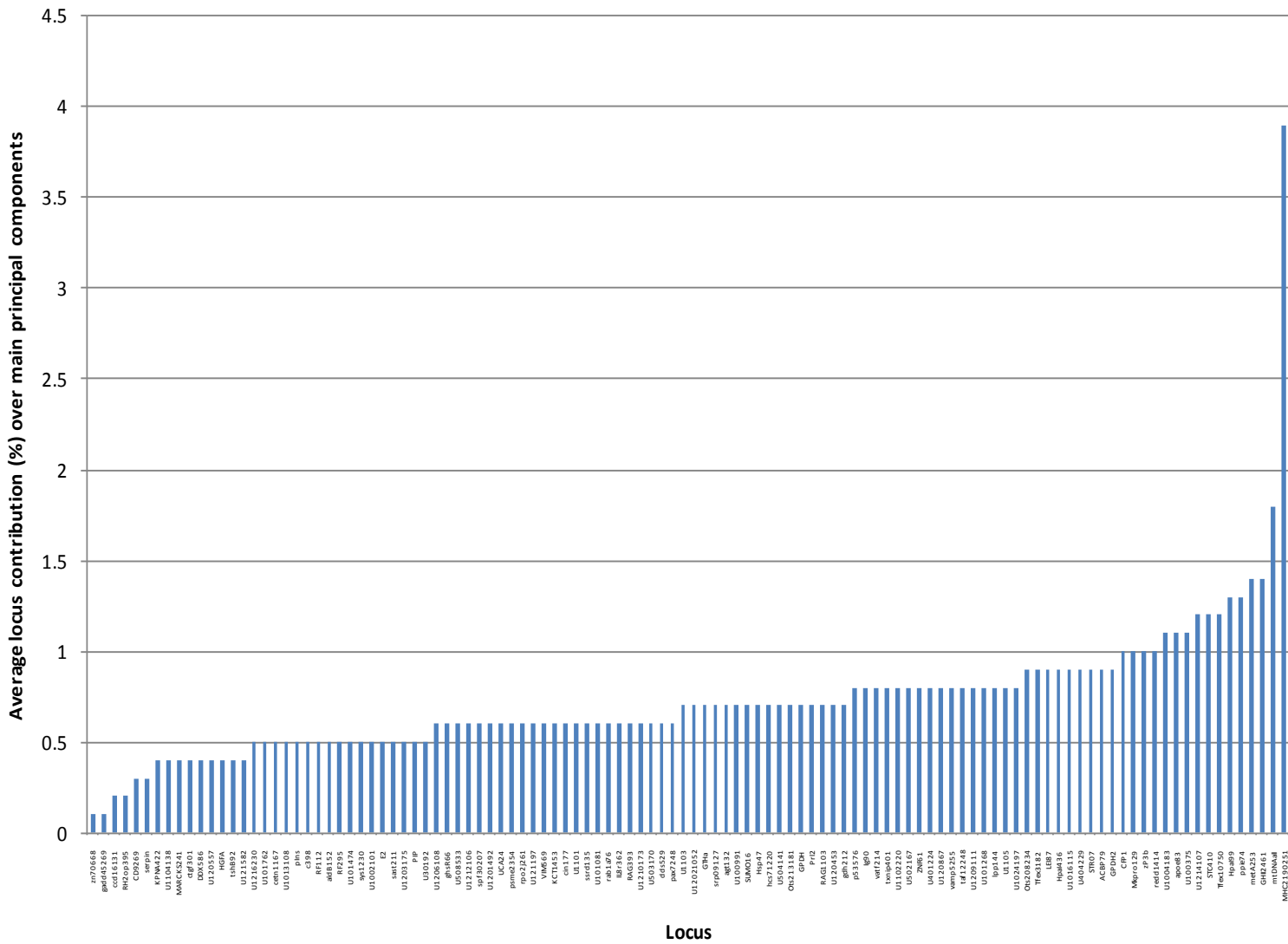


Figure 13.—The average contribution (%) to the first 12 principal components that explained 80% of the total variation for each of 114 SNPs included in a principle component analysis of 36 test populations (Table 1) ordered from lowest contribution (Left) to highest contribution (Right). See text and Table 7 for details.



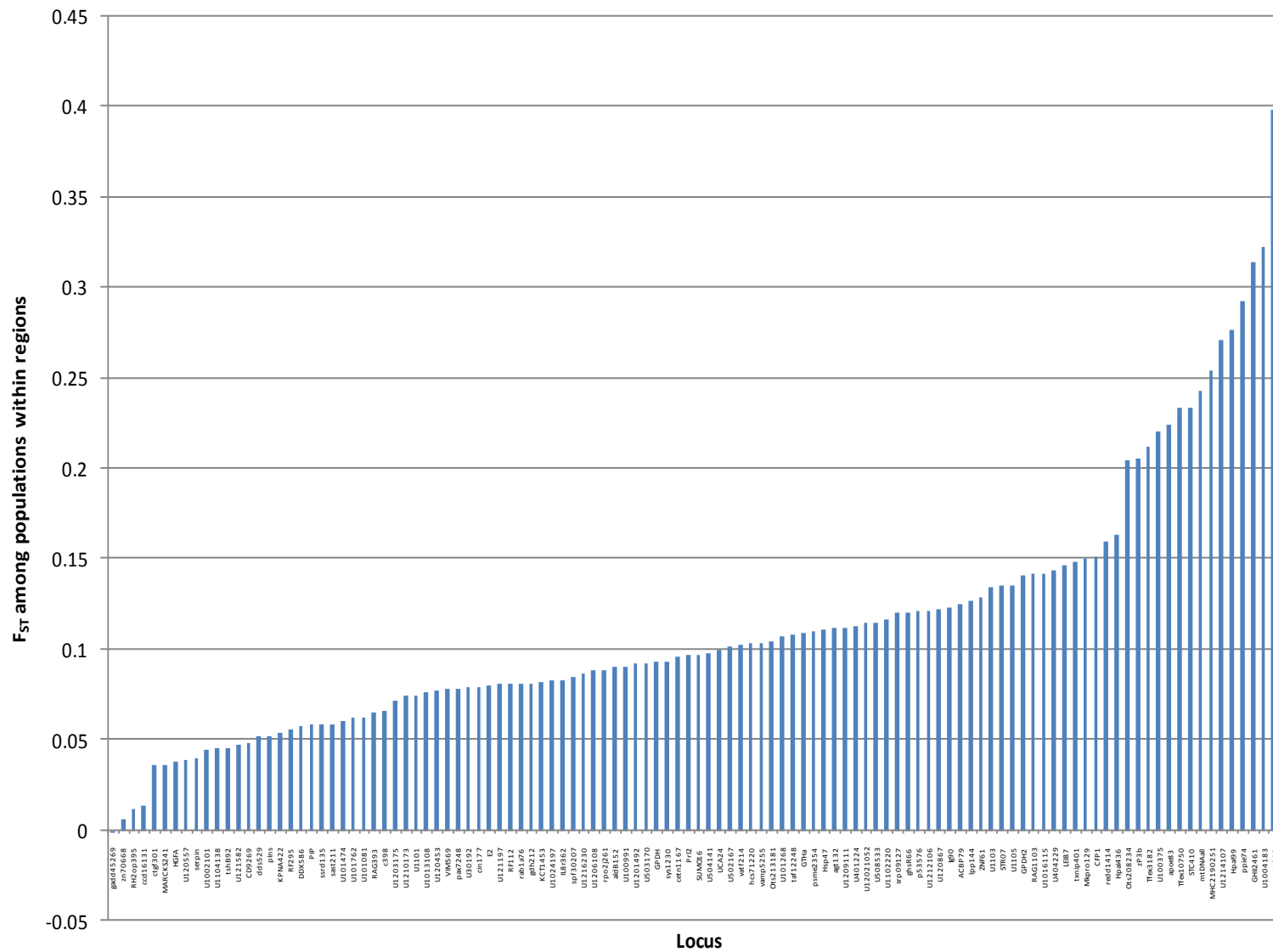
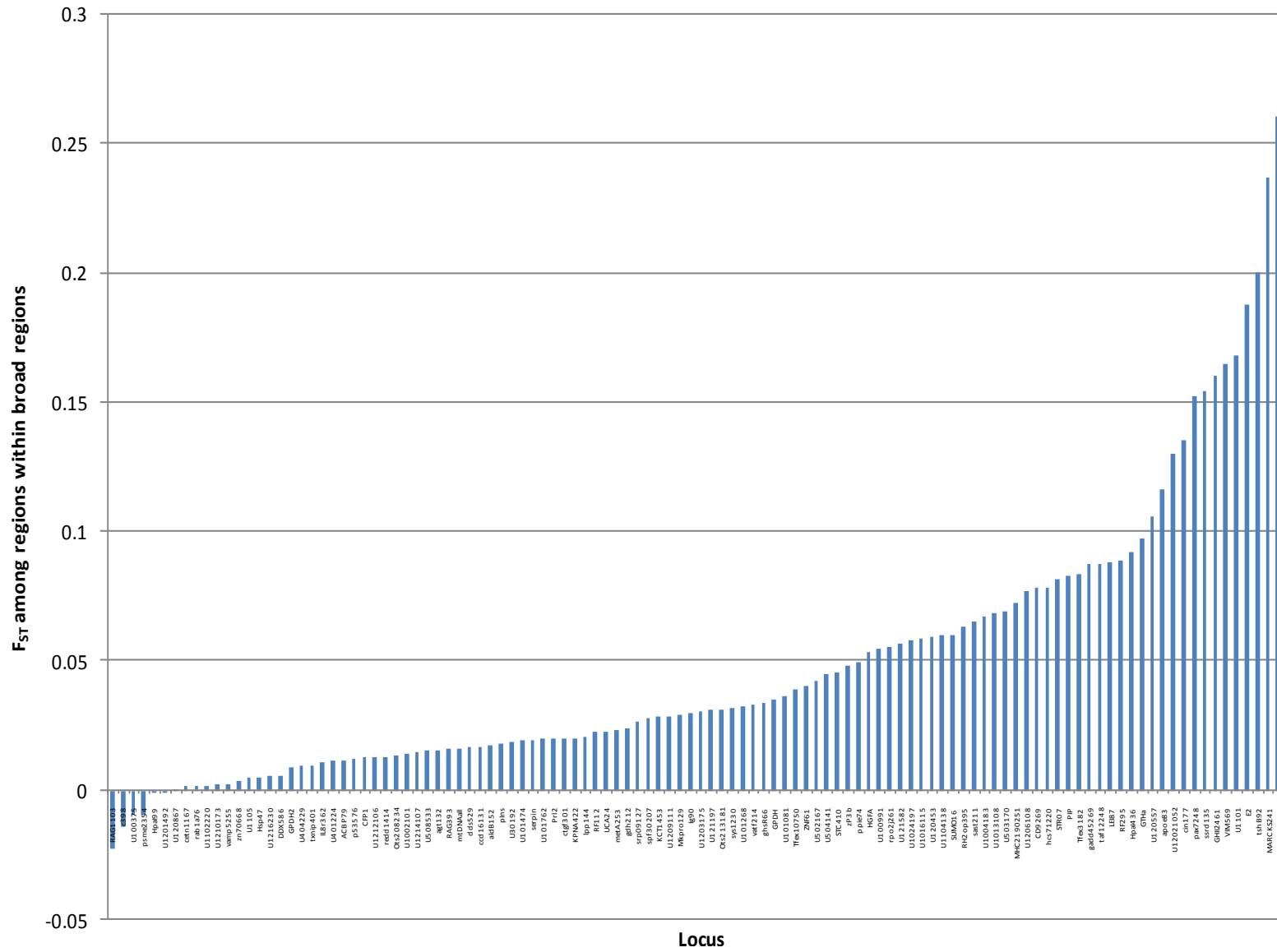


Figure 14.—Weir and Cockerham’s  $F_{ST}$  (1984) among populations within fine-scale regions ( $\theta_s$ ) of 36 test populations (Table 1) ordered from lowest (Left) to highest (Right). See text and Table 8 for details.



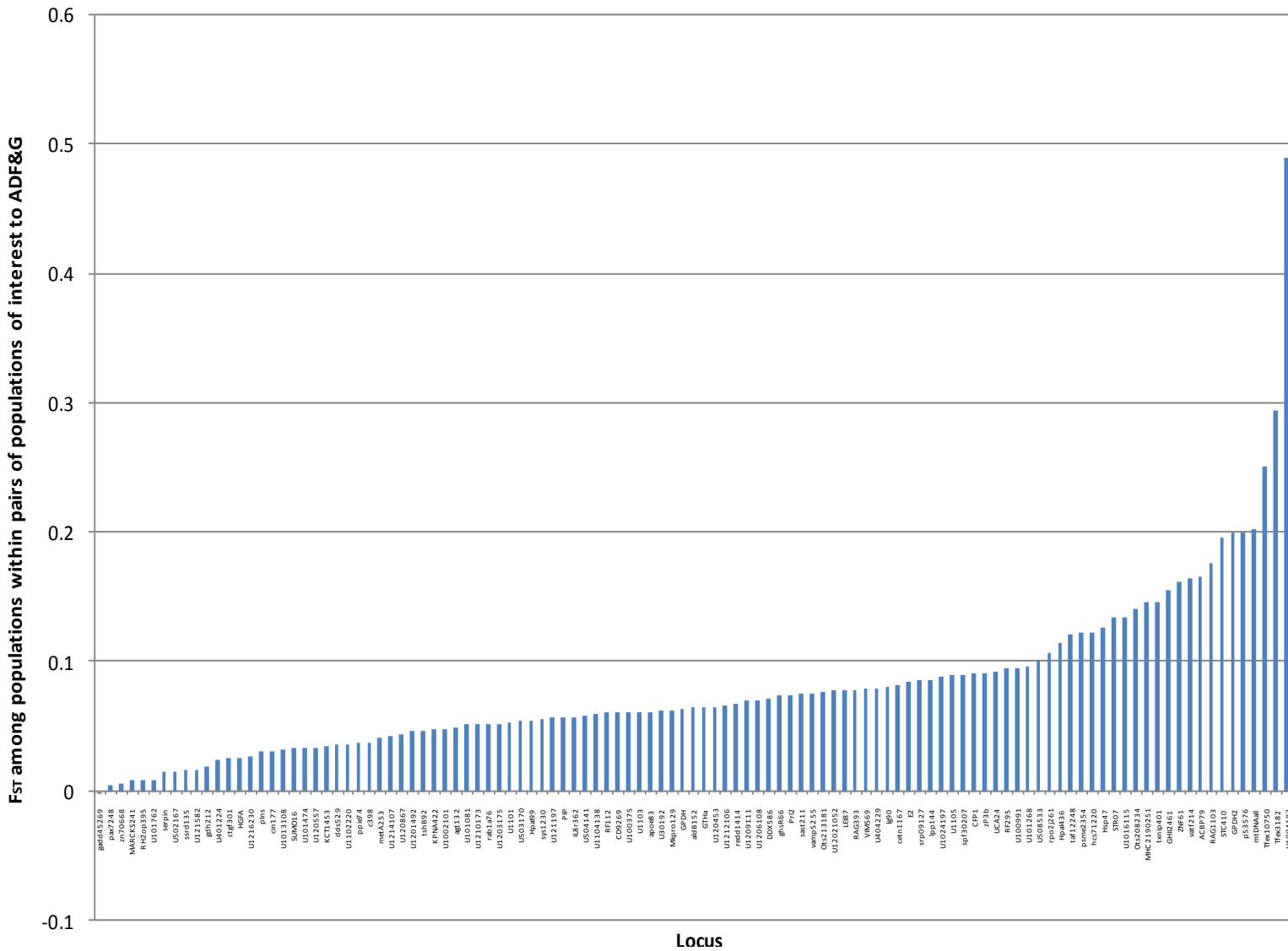


Figure 16.—Weir and Cockerham’s  $F_{ST}$  (1984) between populations within pairs of populations ( $\theta_{\text{pairs}}$ ) of 36 test populations (Table 1) ordered from lowest (Left) to highest (Right). See text and Table 8 for details.

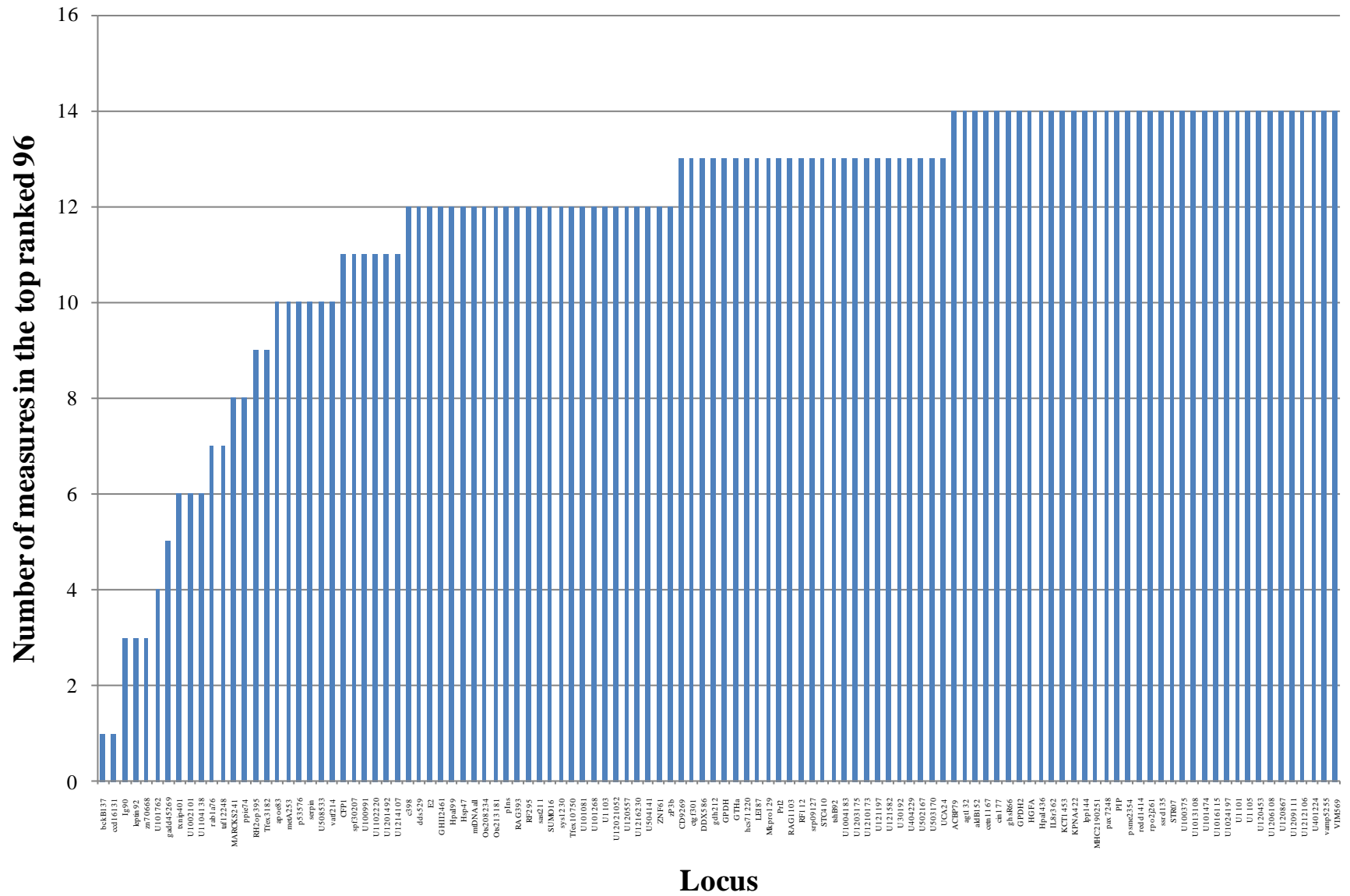


Figure 17.—The number of measures of differentiation (of 14 total) between 7 pairs of populations (Table 1) that each of 124 SNPs (Table 2) ranked in the top 96 for, ordered from lowest (Left) to highest (Right). See text and Tables 9 and 10 for details.

Highest sum of ranks – Lowest sum of ranks

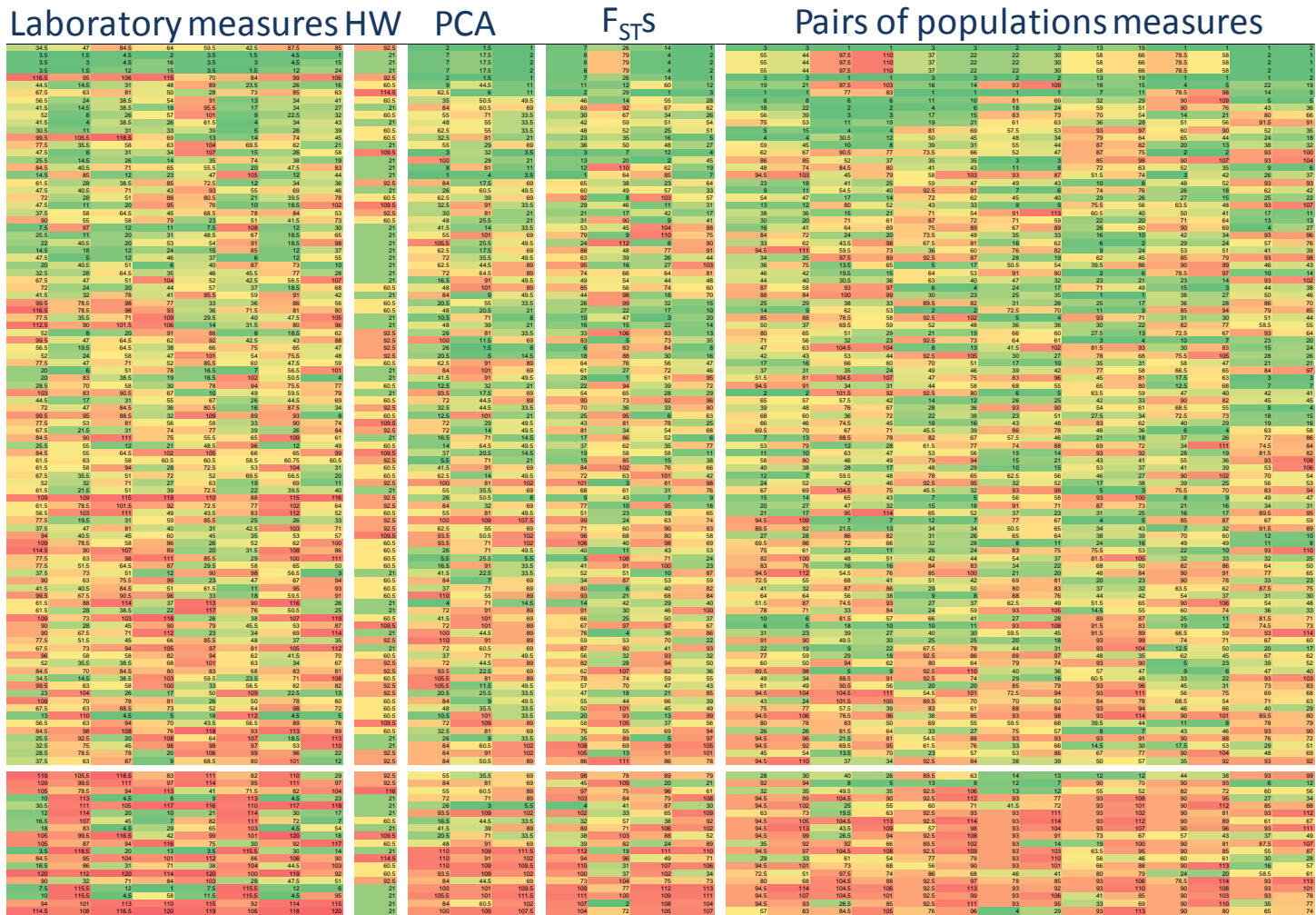


Figure 18.—A color-coded representation of rankings ordered by sum of ranks vertically (Top – Bottom: lowest sum of ranks to highest sum of ranks) and category of judges horizontally (Left – Right: Laboratory measures, Hardy-Weinberg, principle component analyses,  $F_{ST}$ 's and measures of differentiation between pairs of populations of interest) for the 115 SNPs that were not removed from consideration. Note the horizontal break separating the 96 SNPs with the lowest sum of ranks above and the 19 SNPs selected for removal below the break.

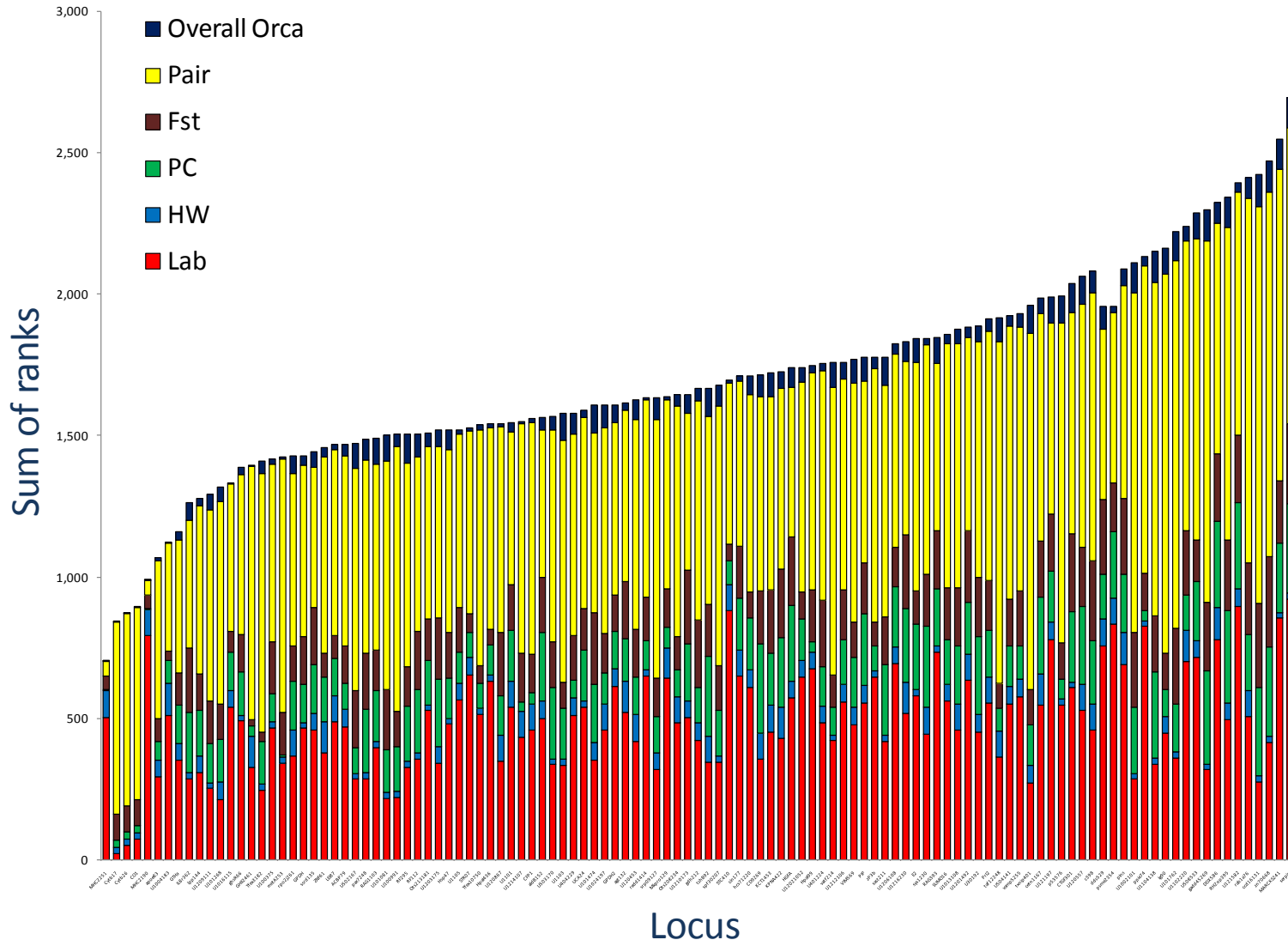


Figure 19.—The sum of rankings for the 115 SNPs that were not removed from consideration and color-coded by category of judge: Overall Orca = Overall  $f_{\text{ORCA}}$  measure, Pair = 14 measures of pairwise differentiation,  $F_{\text{ST}}$  = three measures of  $\theta_{\text{ST}}$ , PC = 3 measures from principal component analysis, HW = Hardy-Weinberg Equilibrium measure, and Lab = 8 measures of laboratory performance.

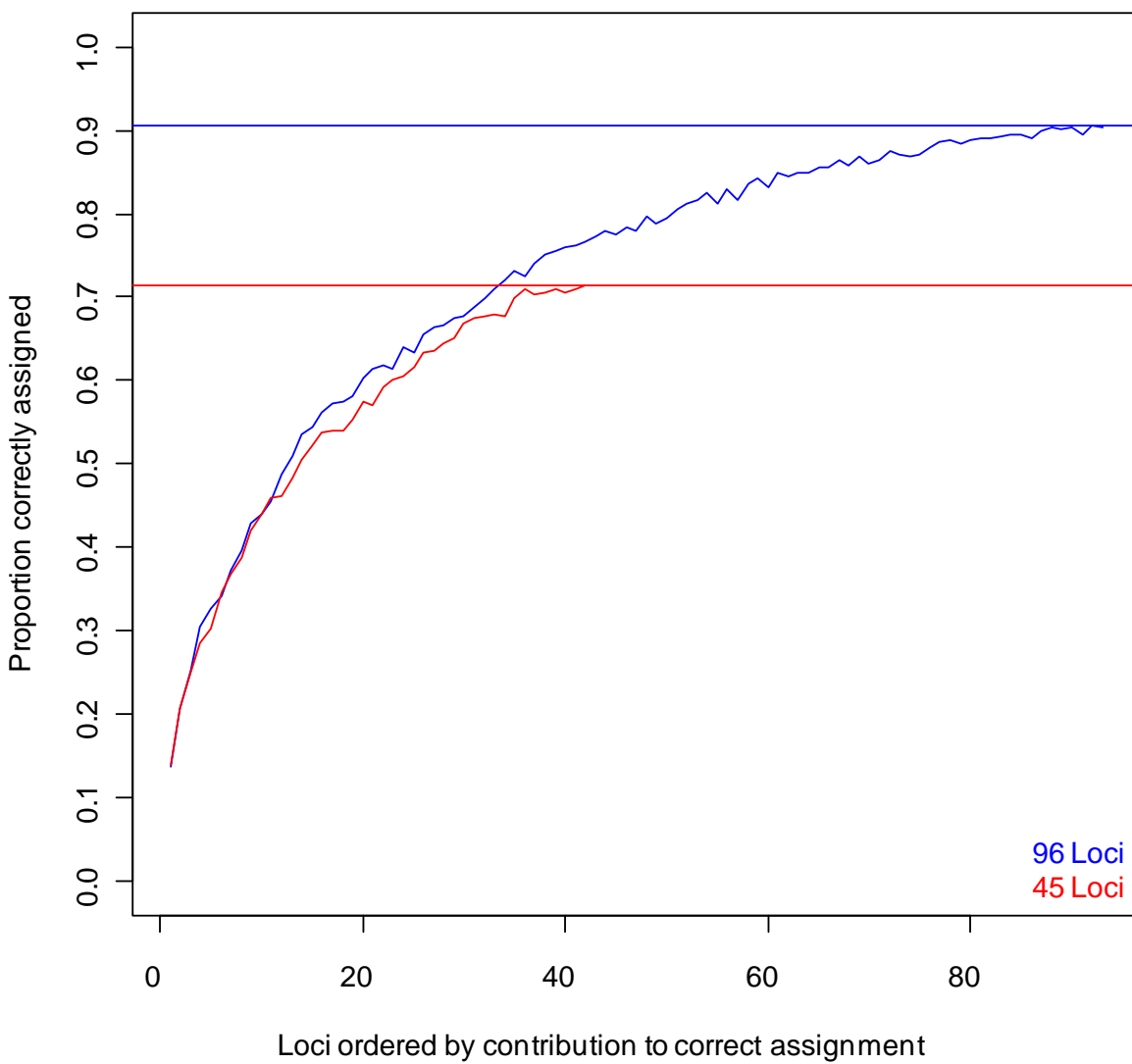


Figure 20.—Assignment curves based upon  $f_{\text{ORCA}}$  for the 45 SNPs that comprised our previous baseline for sockeye salmon and the 96 SNPs chosen to represent our future baseline. See text for details.

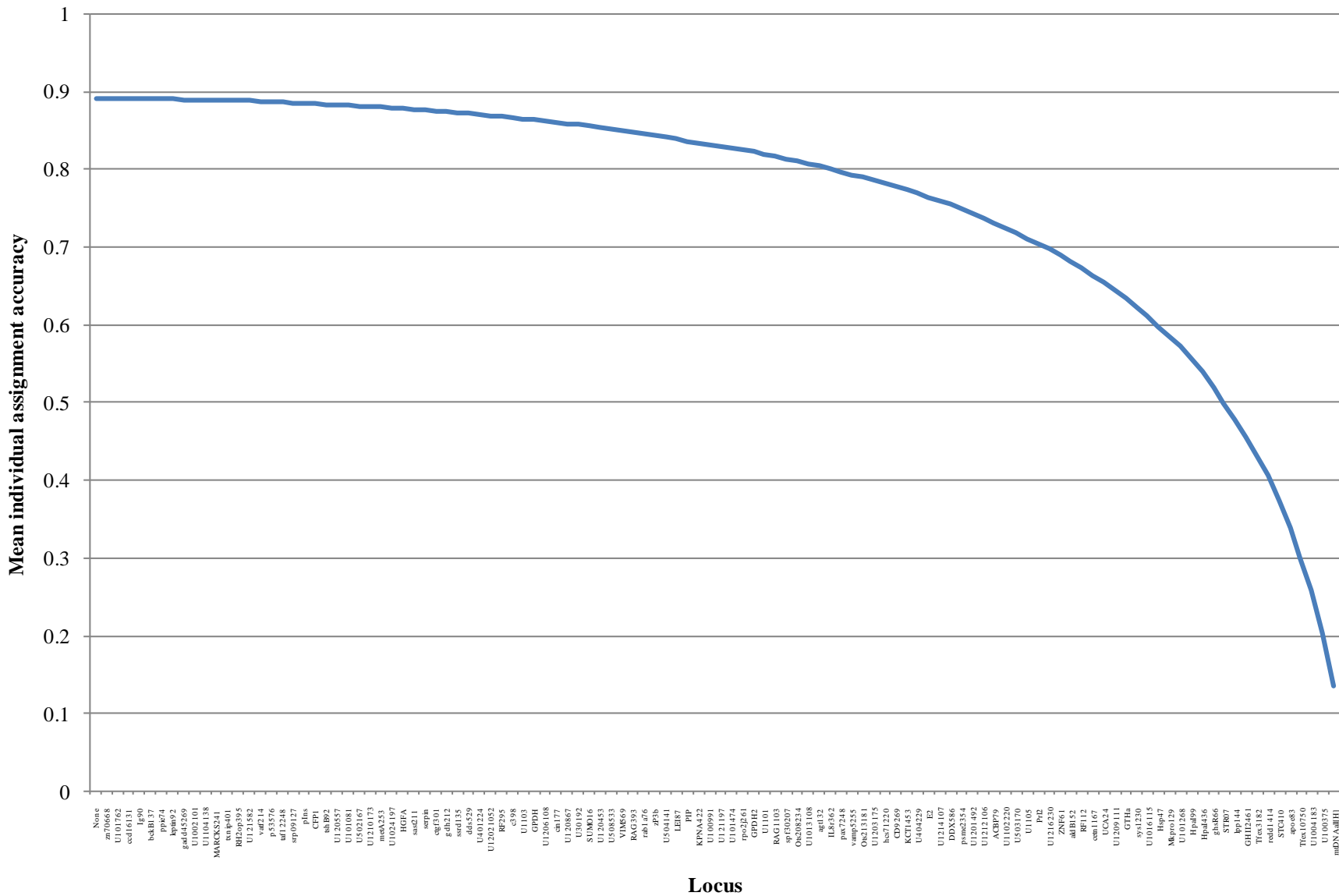


Figure 21.—Cumulative performance measure (mean individual assignment accuracy) for 124 SNPs (Table 2) screened for 36 test populations (Table 1) from a Backwards Elimination Locus Selection (BELS) algorithm. See text for details.



## **APPENDIX**

Appendix A.—Allele frequencies of 115 SNP markers screened for 36 populations in 9 fine-scale regional groupings.

# Allele frequencies of 115 SNP markers screened for 36 populations in 9 fine-scale regional groupings

The following plots of frequencies for the first alphabetical allele (A, C, G, T) for the 36 ‘Backbone’ populations denote fine-scale reporting group membership (Table 1) by symbol and color. Individual loci are ordered alphabetically, except for the combined mtDNA and MHC markers, which are at the end. Population numbers correspond to those in Table 1 and Figure 1 and are ordered geographically (generally clockwise West to East). A smoothed lowess curve was fitted to the estimates of allele frequency for each population. Note that the y-axis scale varies.

Haplotype/phenotype frequencies are plotted for the combined mtDNA and MHC markers rather than for each SNP independently. For these figures, the size of the dot represents frequency (i.e., larger dots represent higher frequency).

Note that the following 9 markers are not included:

SNP	Cause for removal
<i>One_ALDOB-135</i>	Linkage
<i>One_bckB-137</i>	Fixation
<i>One_GPH-414</i>	Linkage
<i>One_leptin-92</i>	Fixation
<i>One_parp3-170</i>	Fixation
<i>One_PPM1K-118</i>	Laboratory failure
<i>One_U1021-57</i>	Failed to conform to HWE
<i>One_U1207-231</i>	Laboratory failure
<i>One_UCA-24</i>	Laboratory failure

<sup>1</sup>This document serves as a record of communication between the Alaska Department of Fish and Game Commercial Fisheries Division and the Western Alaska Salmon Stock Identification Program Technical Committee. As such, these documents serve diverse ad hoc information purposes and may contain basic, uninterpreted data. The contents of this document have not been subjected to review and should not be cited or distributed without the permission of the authors or the Commercial Fisheries Division.

