

Genetic structure of chum and pink salmon in Prince William Sound and Southeast Alaska



Gene Conservation Laboratory
Alaska Department of Fish and Game
Alaska Board of Fisheries
March 8, 2019

Outline

- Background
- Chum results
- Pink results



Alaska Hatchery Research Program

- 1) What is the genetic structure of pink and chum in PWS and SEAK?
- 2) What is the extent and annual variability of straying?
- 3) What is the impact on fitness (productivity) of natural pink and chum stocks due to straying hatchery pink and chum salmon?

Understanding Genetic Structure

- Differences between populations:
 - Influenced by: selection, mutation, genetic drift, migration

Understanding Genetic Structure

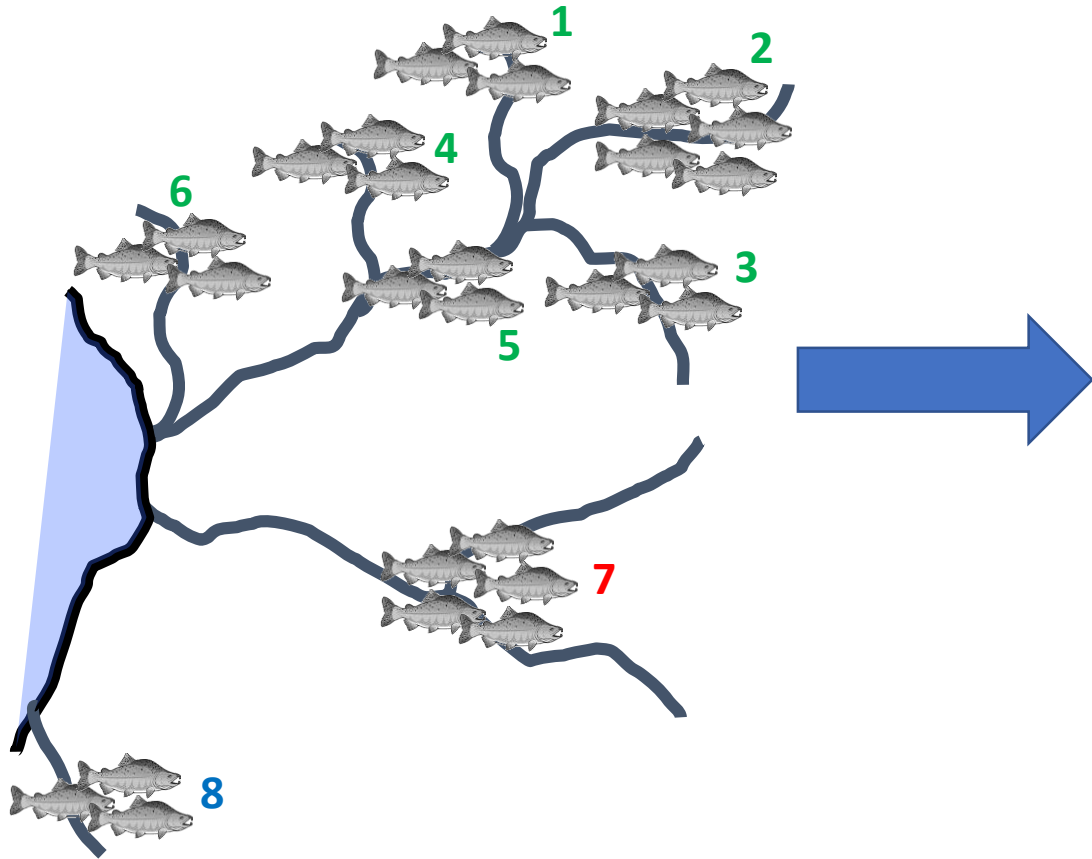
- Differences between populations:
 - Influenced by: selection, mutation, *genetic drift*, *migration*

genetic drift ~ **homing**

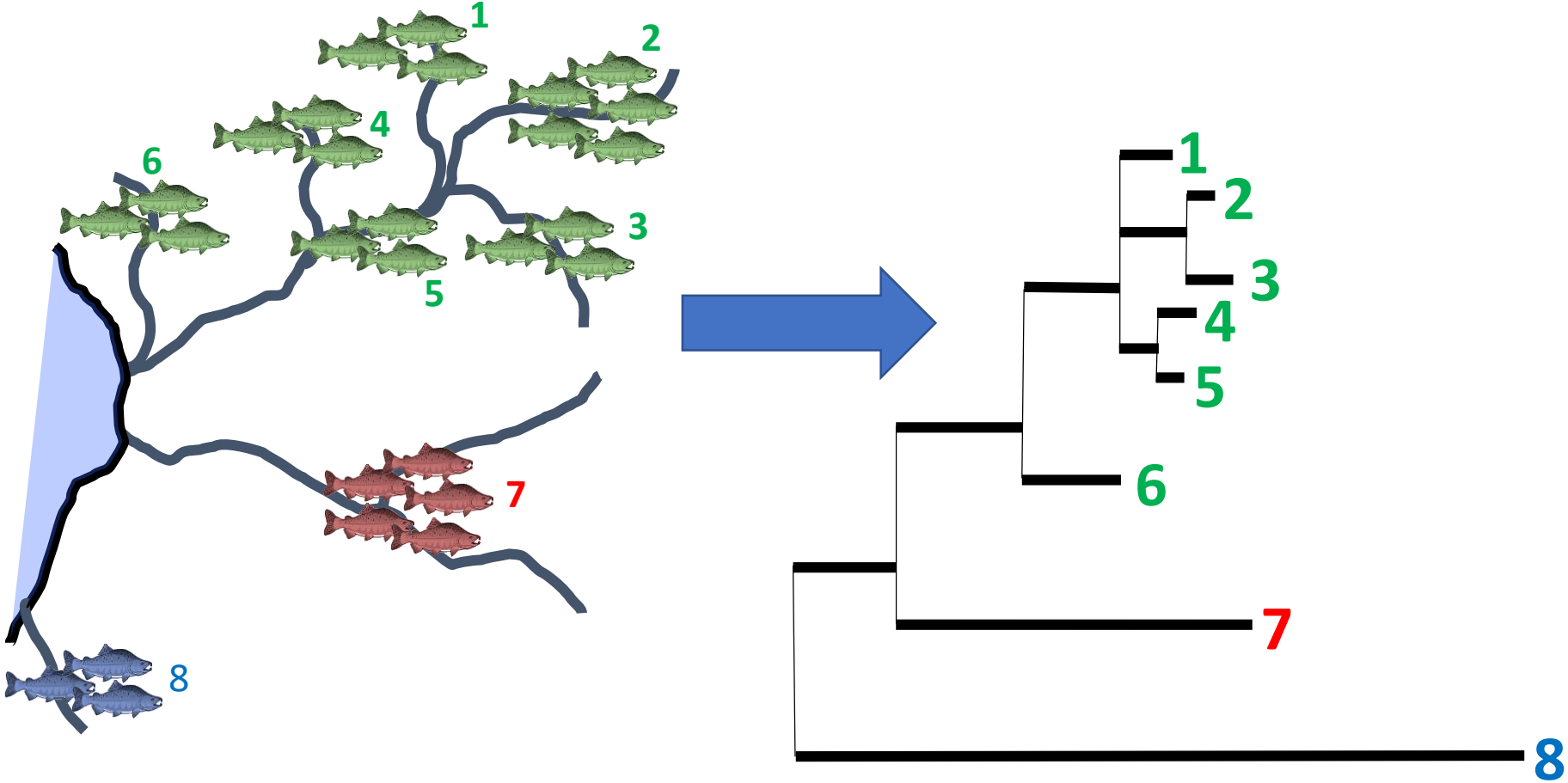
migration ~ **straying**

- *Measuring the balance between these within a species across an area*
- Measured by quantifying pairwise genetic differences
- Visualize using genetic trees

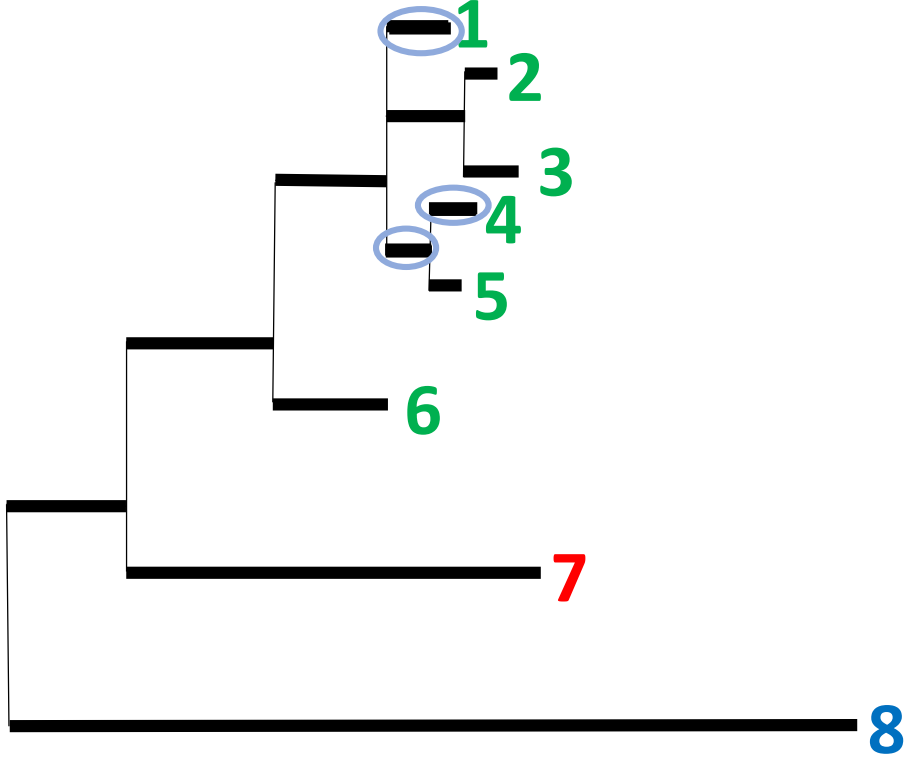
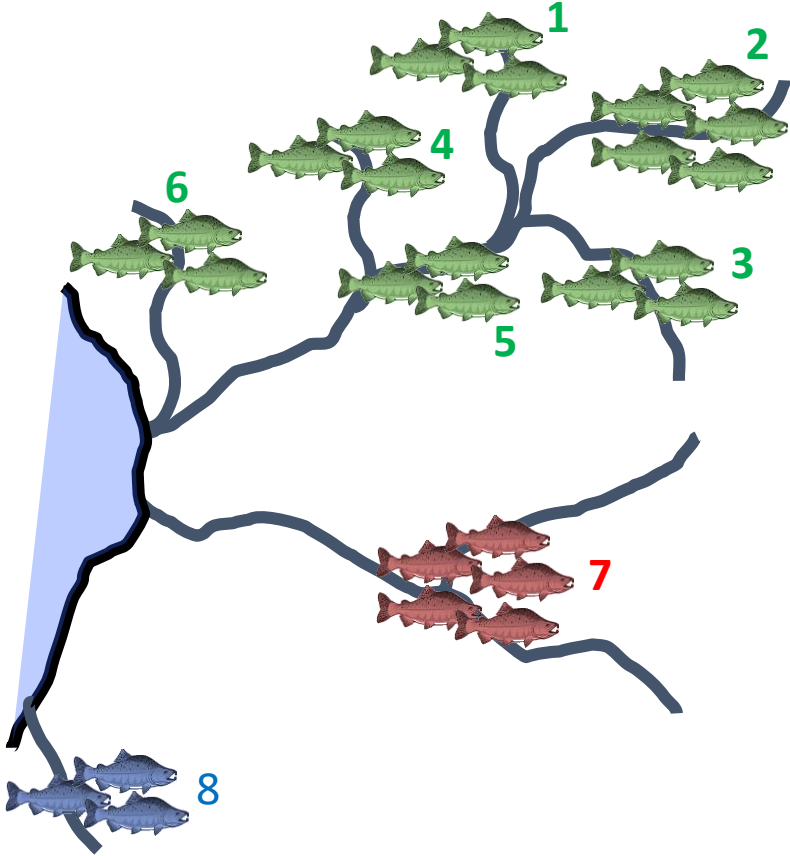
Population Structure: An example



Population Structure: An example

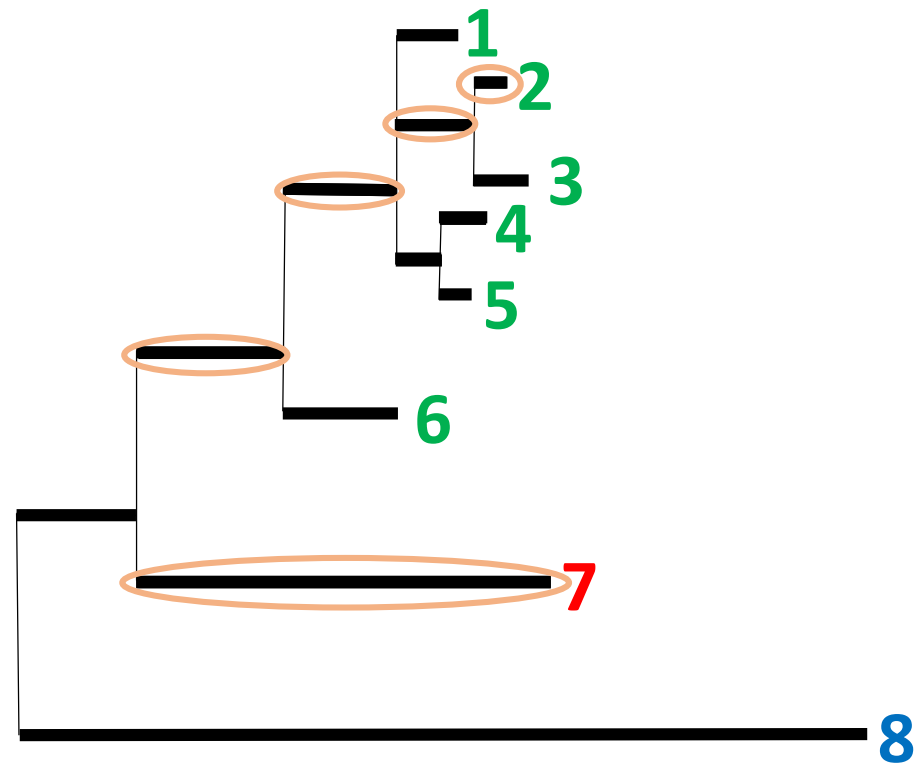
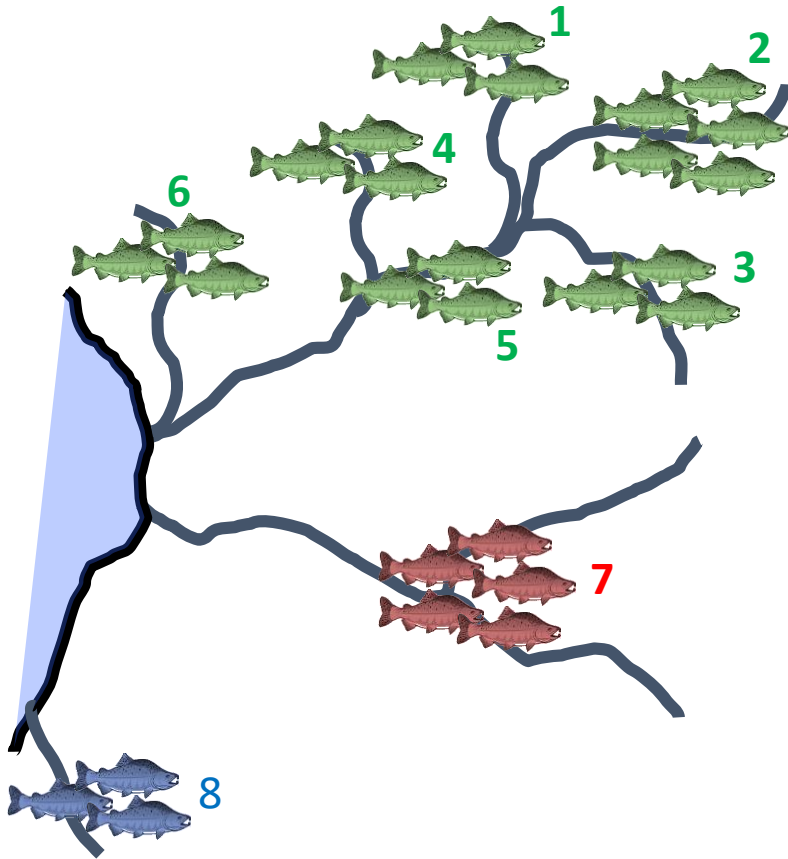


Population Structure: An example



Difference between 1 and 4: + + =

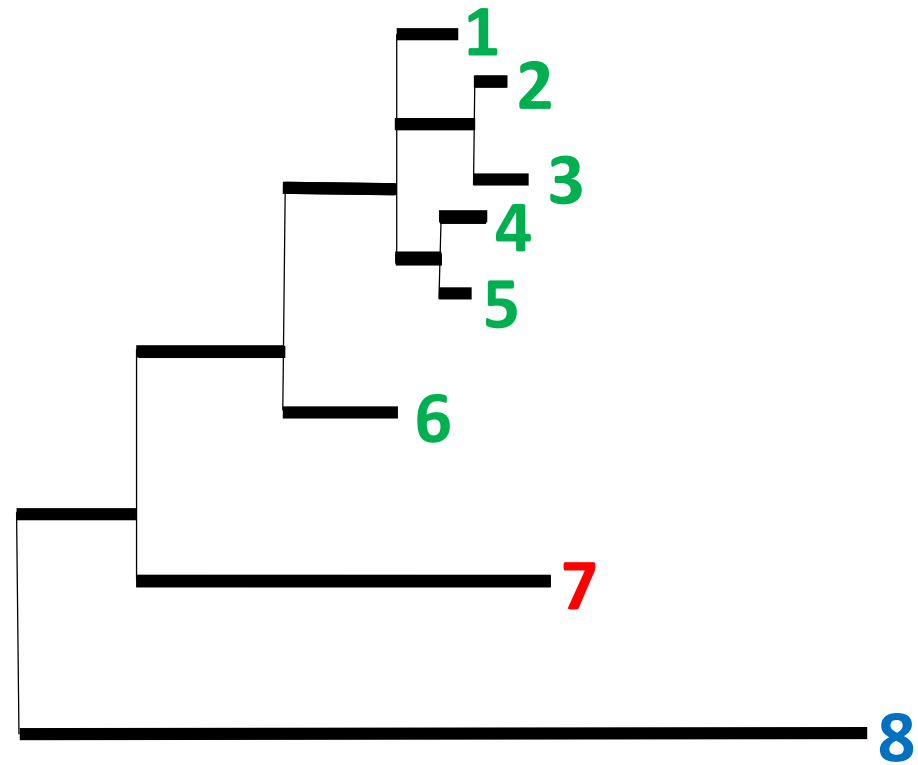
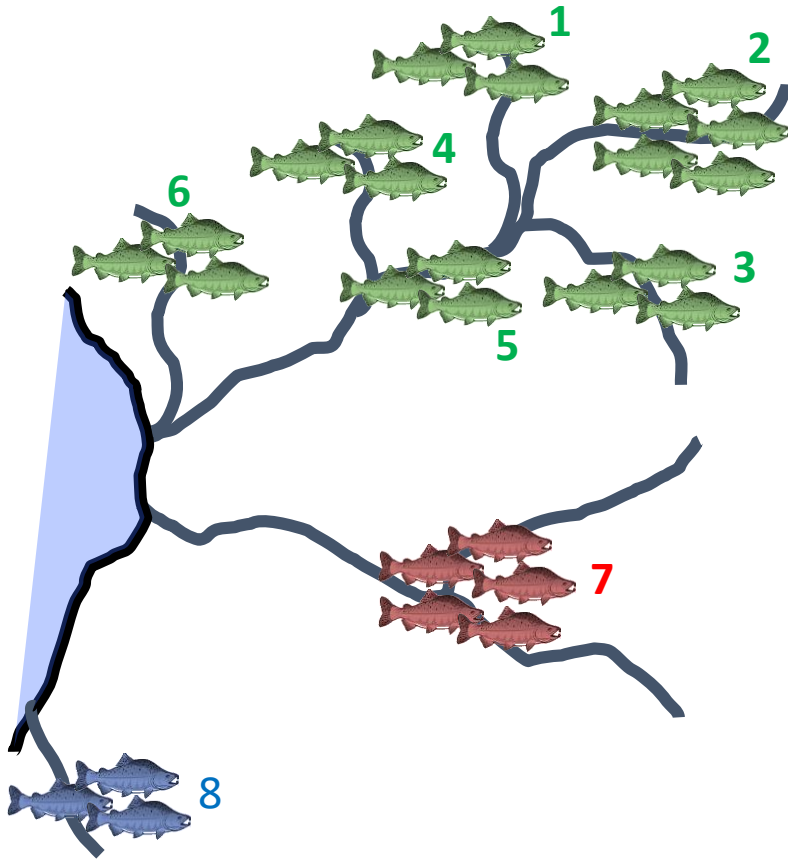
Population Structure: An example



Difference between 1 and 4:

Difference between 2 and 7: + + + + =

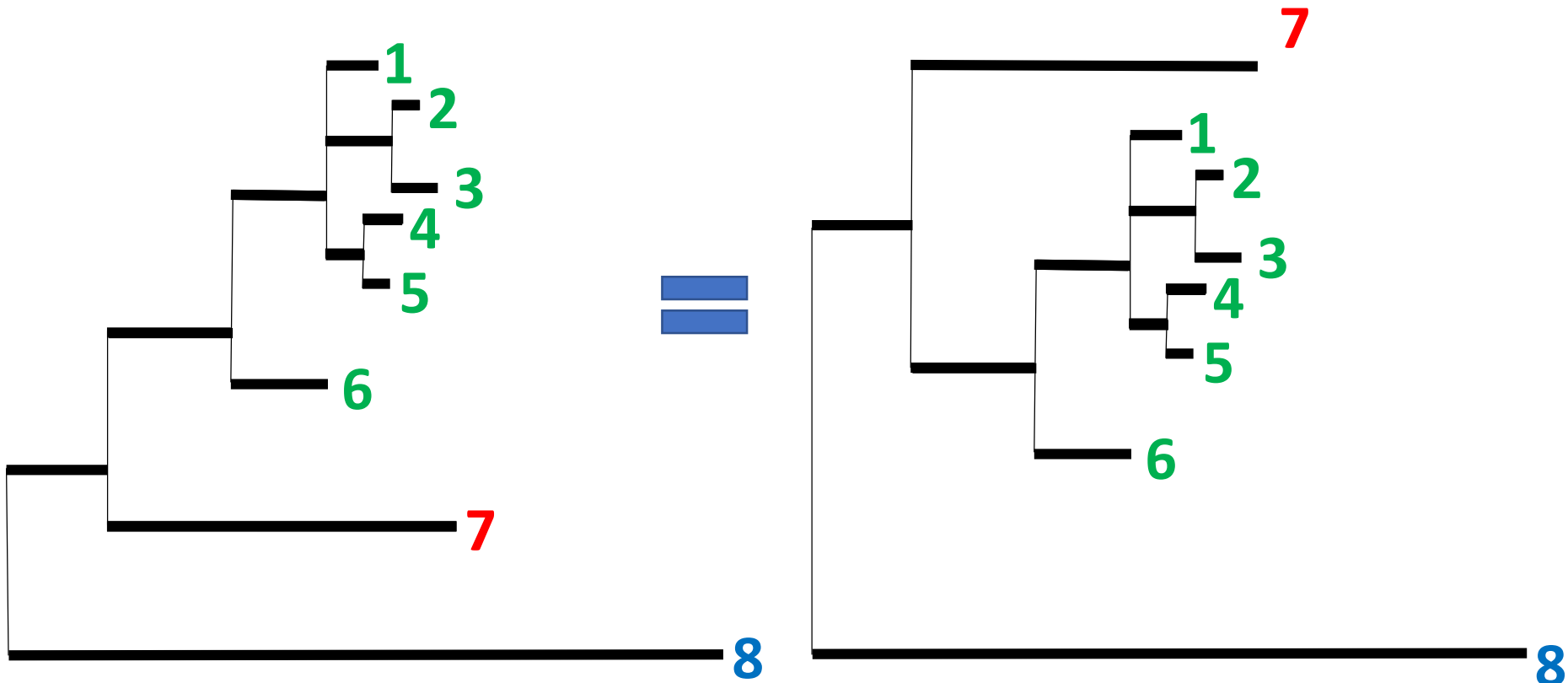
Population Structure: An example



Difference between 1 and 4:

Difference between 2 and 7:

Population Structure: An example



Chum salmon in Prince William Sound and Southeast Alaska



Sara Gilk-Baumer and William D. Templin

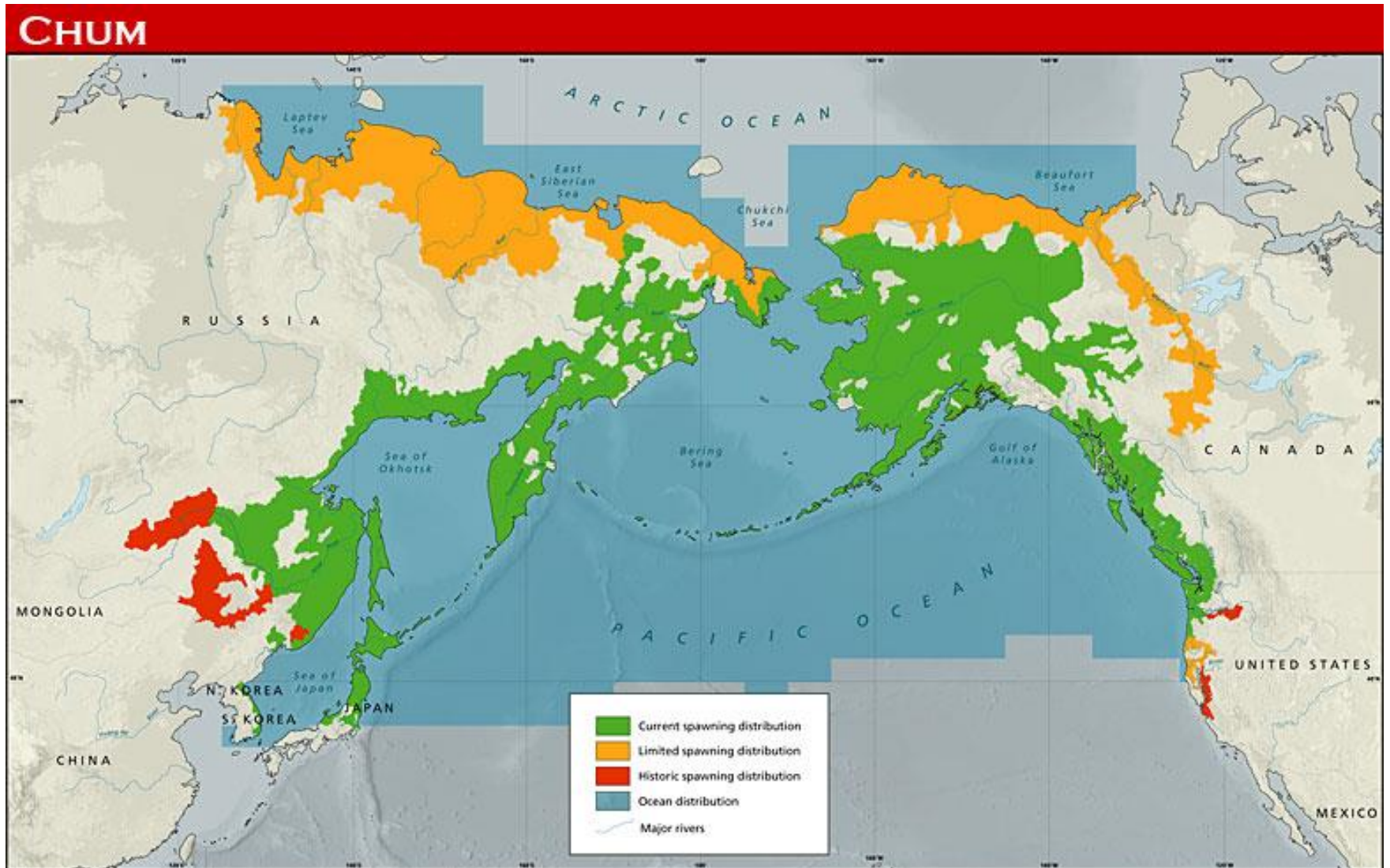
Alaska Department of Fish and Game, Gene Conservation Lab

Life History of Chum Salmon

- Migrate as juveniles to ocean
- Typically 2-4 years spent at sea
- Two run timings: summer & fall



Distribution of Chum Salmon



http://www.salmonnation.org/fish/meet_species.html

Previous work (a sampling)

Determining Continent of Origin of Chum Salmon (*Oncorhynchus keta*) Using Genetic Stock Identification Techniques: Status of Allozyme Baseline in Asia

Gary A. Winans and Paul B. Aebbersold

Northwest Fisheries Science Center, National Marine Fisheries Service, Seattle, WA 98112-2097, USA

Shigehiko Urawa

Hokkaido Salmon Hatchery, Fisheries Agency of Japan, Sapporo 062, Japan

and Nataly V. Varnavskaya

Kamchatka-TINRO, Petropavlovsk, Russia

Genetic Relationships Among Chum Salmon Populations in Southeast Alaska and Northern British Columbia

C.M. Kondzela, C.M. Guthrie, S.L. Hawkins, C.D. Russell, and J.H. Helle

Auke Bay Laboratory, Alaska Fisheries Science Center, National Marine Fisheries Service, National Oceanographic and Atmospheric Administration, 11305 Glacier Highway, Juneau, AK 99801-8626, U.S.A.

and A.J. Garret

School of Fisheries and Ocean Sciences, University of Alaska Fairbanks, 11120 Glacier Highway, Juneau, AK 99801, U.S.A.

Population structure and stock identification of chum salmon (*Oncorhynchus keta*) from British Columbia determined with microsatellite DNA variation

Terry D. Beacham, Brian Spilsted, Khal D. Le, and Michael Wetklo

Microsatellite Stock Identification of Chum Salmon on a Pacific Rim Basis

TERRY D. BEACHAM,* JOHN R. CANDY, AND C. WALLACE

Fisheries and Oceans Canada, Pacific Biological Station,
3190 Hammond Bay Road, Nanaimo, British Columbia V9T 6N7, Canada

SHIGEHICO URAWA¹ AND SHUNPEI SATO

National Salmon Resources Center, Fisheries Research Agency, Toyohira-ku, Sapporo 062-0922, Japan

NATALIA V. VARNAVSKAYA

Kamchatka Fishery and Oceanography Research Institute,
18 Naberezhnaya Street; Petropavlovsk-Kamchatsky 683000, Russia

KHAL D. LE AND MICHAEL WETKLO

Fisheries and Oceans Canada, Pacific Biological Station,
3190 Hammond Bay Road, Nanaimo, British Columbia V9T 6N7, Canada

Environmental Biology of Fishes 69: 37–50, 2004.
© 2004 Kluwer Academic Publishers. Printed in the Netherlands.

Genetic population structure of chum salmon in the Pacific Rim inferred from mitochondrial DNA sequence variation

Shunpei Sato^a, Hiroyuki Kojima^b, Junko Ando^a, Hironori Ando^a, Richard L. Wilmoth^c, Lisa W. Seeb^d, Vladimir Efremov^e, Larry LeClair^f, Wally Buchholz^g, Deuk-Hee Jin^h, Shigehiko Urawaⁱ, Masahide Kaeriyama^j, Akihisa Urano^{k,l} & Syuiti Abe^{k,l}

^aDivision of Biological Science, Graduate School of Science, Hokkaido University, Sapporo 060-0810, Japan

^bGraduate School of Science and Engineering, Hokkaido Tokai University, Sapporo 005-8601, Japan

^cAuke Bay Laboratory, Alaska Fisheries Science Center, NOAA, Juneau, U.S.A.

^dAlaska Department of Fish and Game, Anchorage, U.S.A.

^eRussian Academy of Science, Vladivostok, Russia

^fWashington Department of Fish and Wildlife, Olympia, Washington, U.S.A.

^gU.S. Fish and Wildlife Service, Anchorage, AK, U.S.A.

^hKangnung National University, Kangnung, Korea

ⁱSalmon Resources Center, Sapporo 062-0922, Japan

^jField Science Center, Hokkaido University, Sapporo 060-0811, Japan

^kLaboratory of Animal Cytogenetics, Center for Advanced Science and Technology, Hokkaido University, Sapporo 060-0810, Japan (e-mail: sabe@ees.hokudai.ac.jp)

^lLaboratory of Breeding Science, Graduate School of Fisheries Sciences, Hokkaido University, Hakodate 041-8611, Japan

Received 17 April 2003

Accepted 27 April 2003

Chum Salmon Genetic Diversity in the Northeastern Pacific Ocean Assessed with Single Nucleotide Polymorphisms (SNPs): Applications to Fishery Management

Maureen P. Small^a

Washington Department of Fish and Wildlife, Molecular Genetics Lab,
1111 Washington Street Southeast, Olympia, Washington 98501, USA

Serena D. Rogers Olive

Alaska Department of Fish and Game, Division of Commercial Fisheries,
Gene Conservation Laboratory, 333 Raspberry Road, Anchorage, Alaska 99518, USA

Lisa W. Seeb, James E. Seeb, and Carita E. Pascal

School of Aquatic and Fishery Sciences, University of Washington, 1122 Northeast Boat Street,
Box 355020, Seattle, Washington 98195, USA

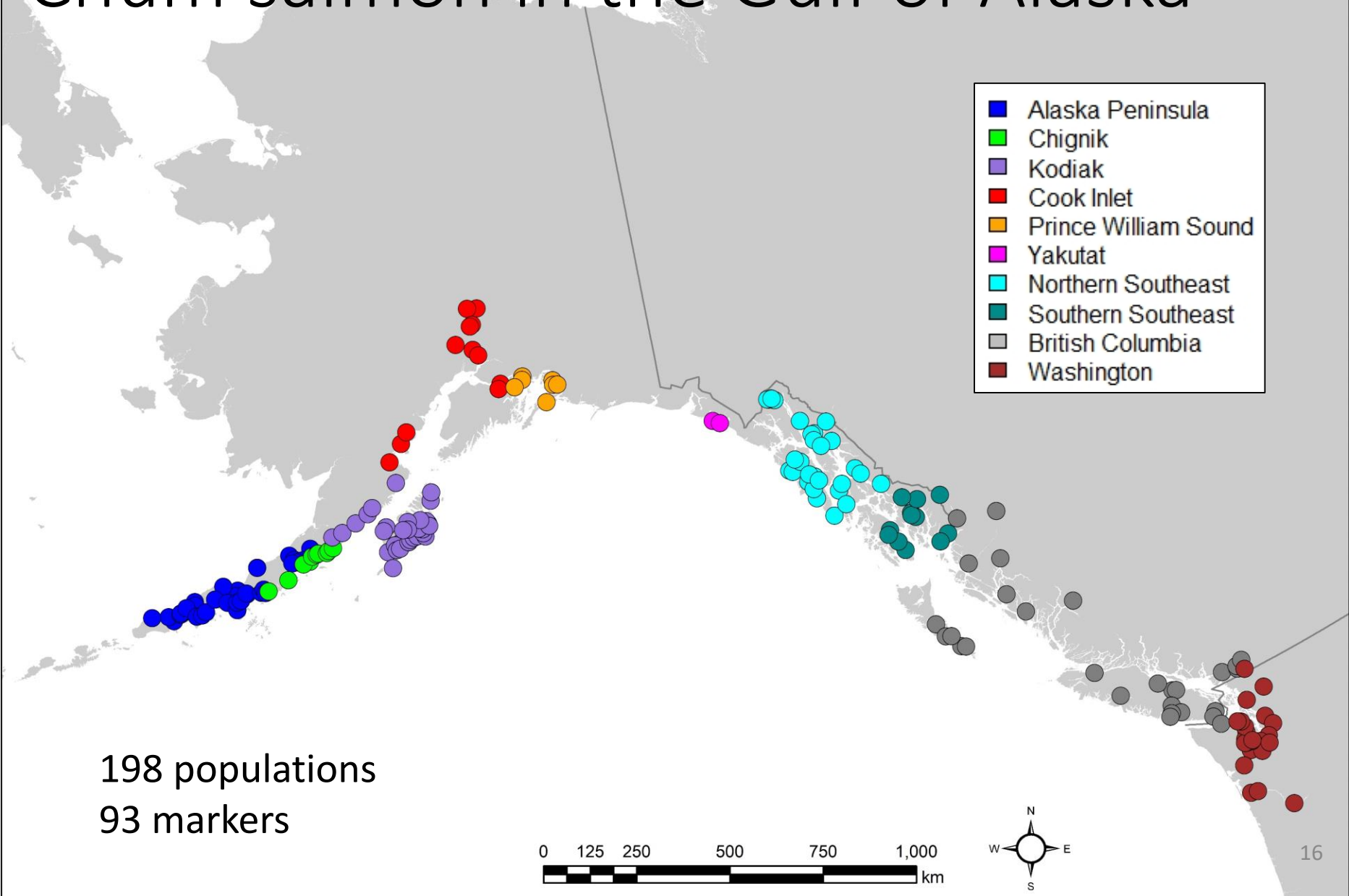
Kenneth I. Warheit

Washington Department of Fish and Wildlife, Molecular Genetics Lab,
1111 Washington Street Southeast, Olympia, Washington 98501, USA; and School of Aquatic and Fishery Sciences,
University of Washington, 1122 Northeast Boat Street, Box 355020, Seattle, Washington 98195, USA

William Templin

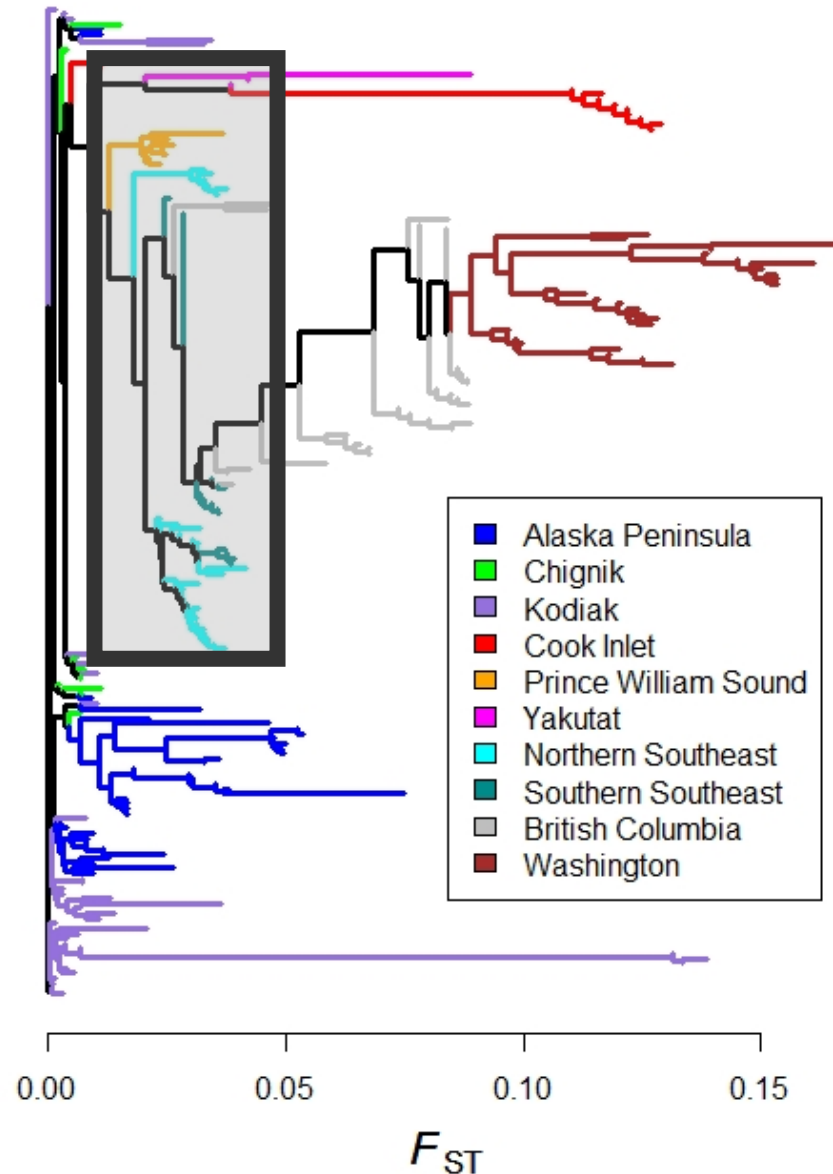
Alaska Department of Fish and Game, Division of Commercial Fisheries,
Gene Conservation Laboratory, 333 Raspberry Road, Anchorage, Alaska 99518, USA

Chum salmon in the Gulf of Alaska



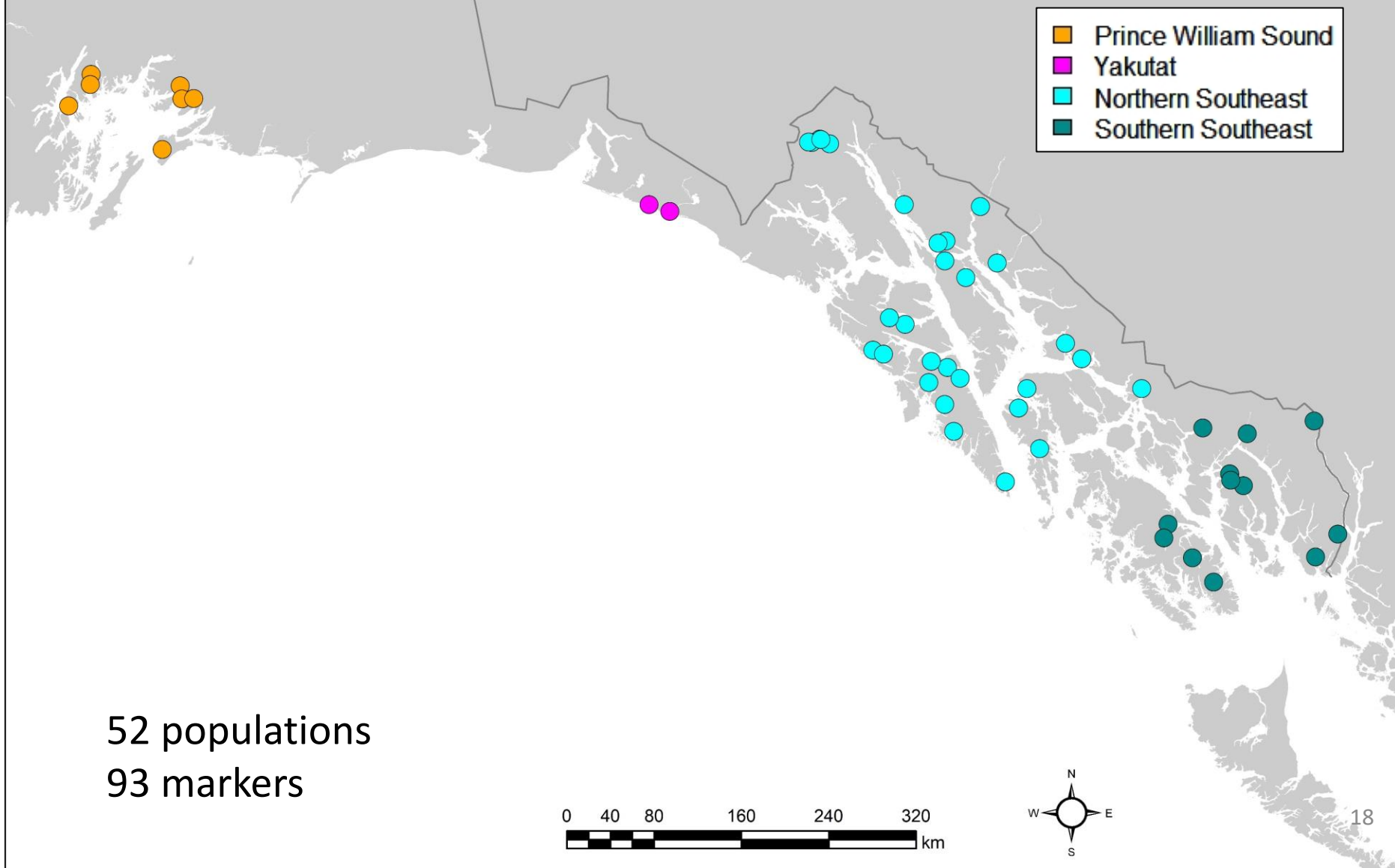
Chum salmon in the Gulf of Alaska

198 populations
93 markers



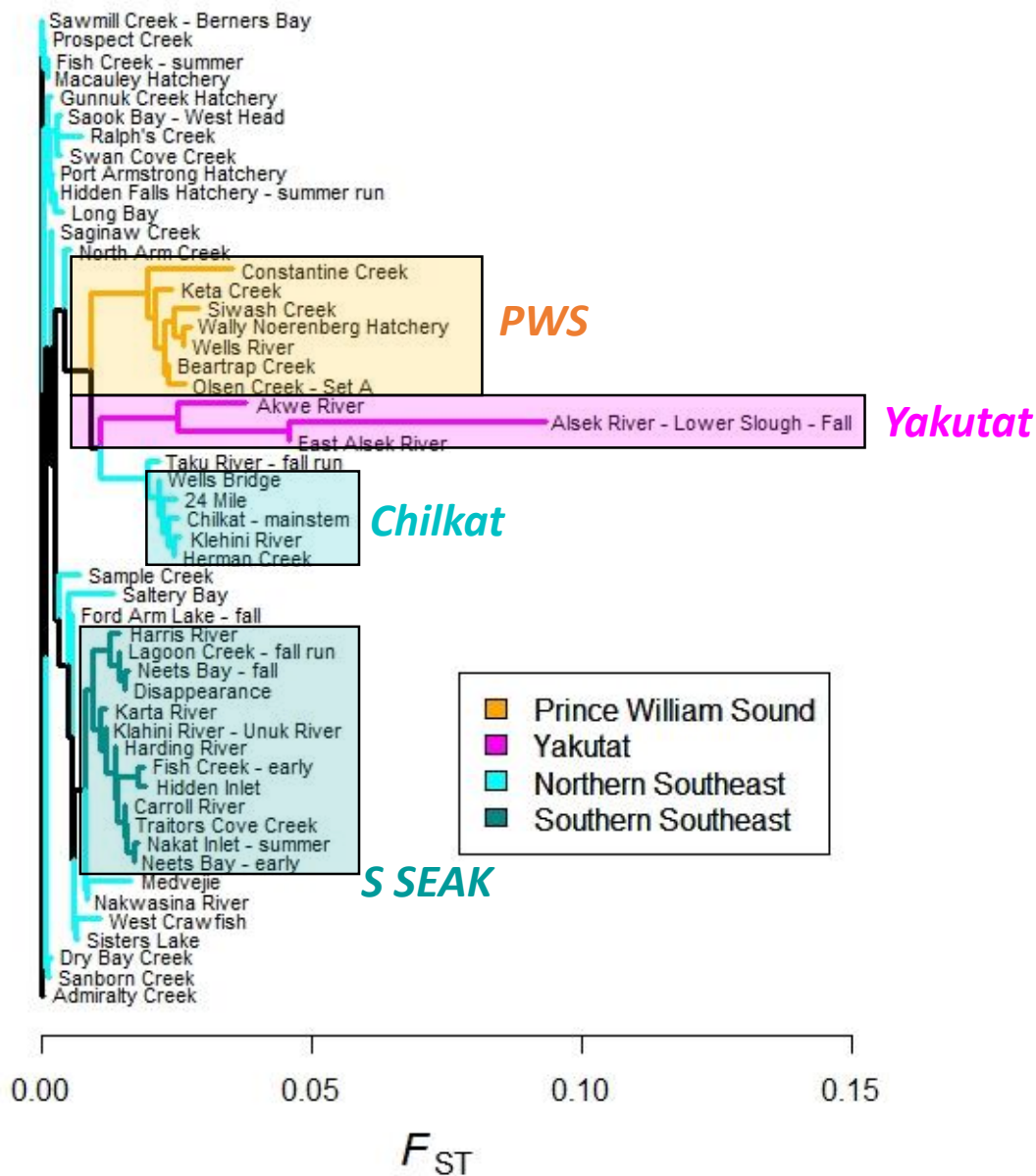
PWS to SEAK

Chum salmon in PWS and SEAK



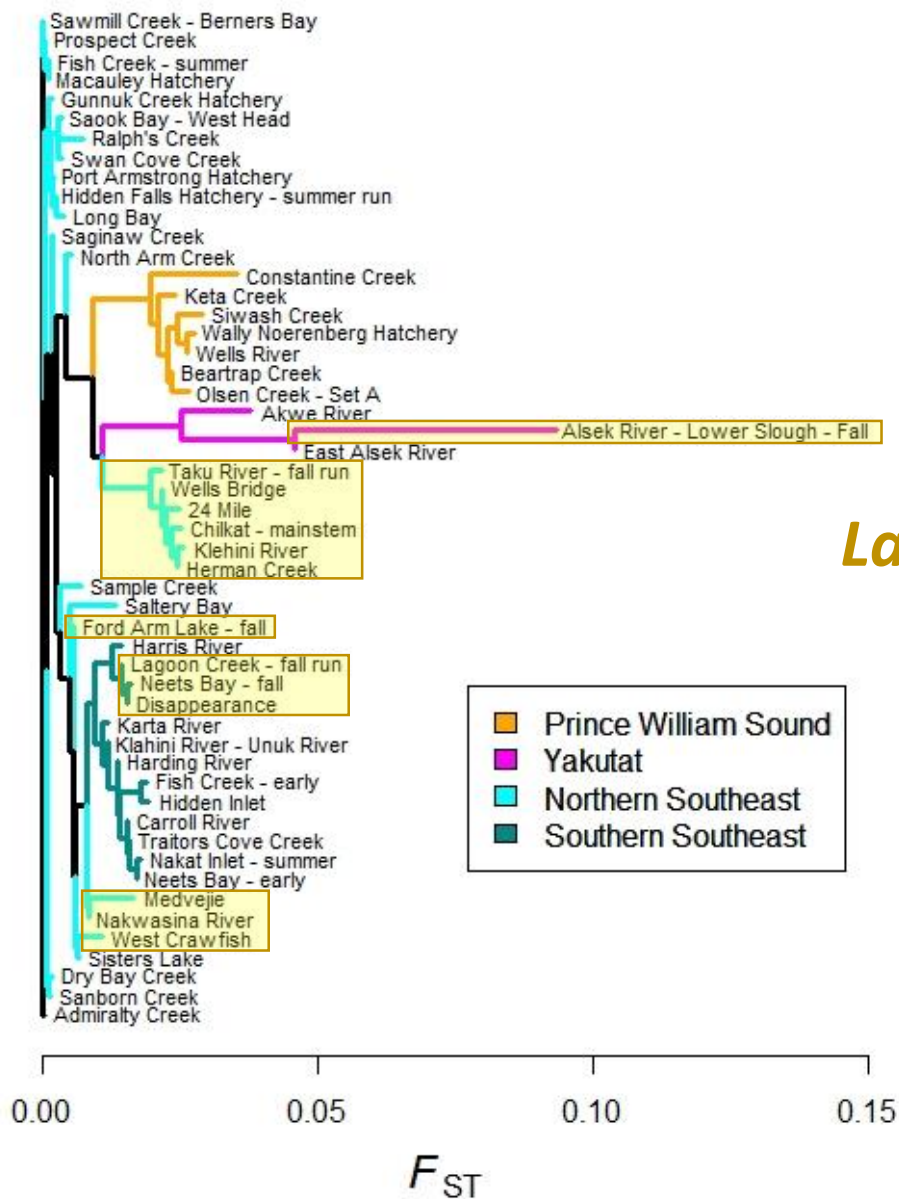
Chum salmon in PWS and SEAK

52 populations
93 markers



Chum salmon in PWS and SEAK

52 populations
93 markers



Late run timing

Conclusions: Chum salmon structure in PWS and SEAK

- Generally correlated with geography
- Some differentiation by run timing
- Similar to other studies



Population structure of pink salmon in Prince William Sound



Wei Cheng^{1,2}, Christopher Habicht¹, William D. Templin¹, Zachary D. Grauvogel¹, and Anthony J. Gharrett²

¹Alaska Department of Fish and Game, Gene Conservation Laboratory

²University of Alaska Fairbanks, College of Fisheries and Ocean Sciences

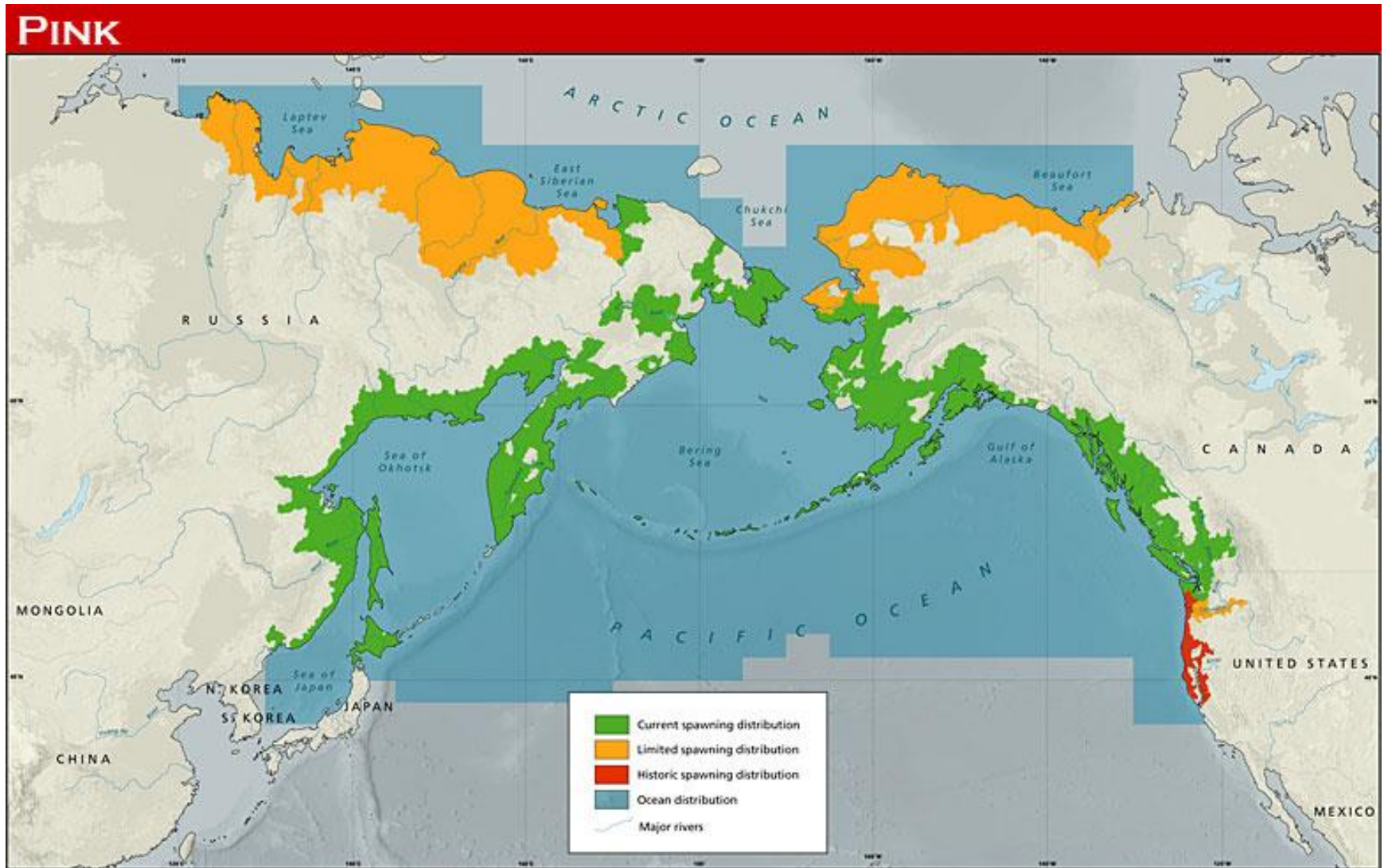
Life History of Pink Salmon

- **Two-year life cycle**
 - **Odd year**
 - **Even year**
- **Limited freshwater life history**



<https://www.n-sea.org/pink-salmon>

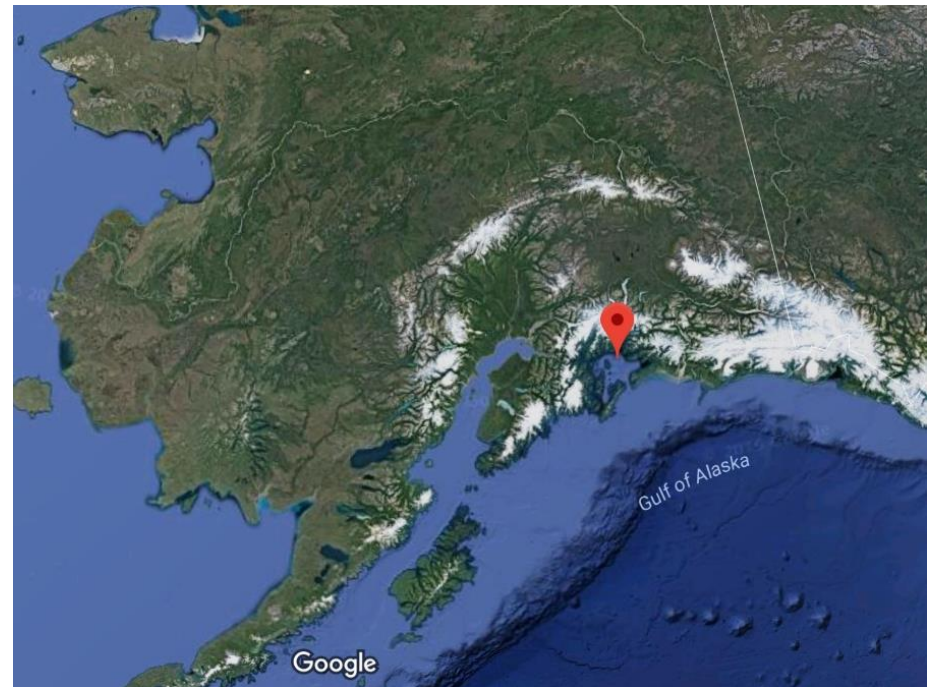
Distribution of Pink Salmon



http://www.salmonnation.org/fish/meet_species.html

PWS Pink Salmon

- **Number of streams in Prince William Sound (PWS)**
 - Over 800 streams
- **Variation in run timing across streams**



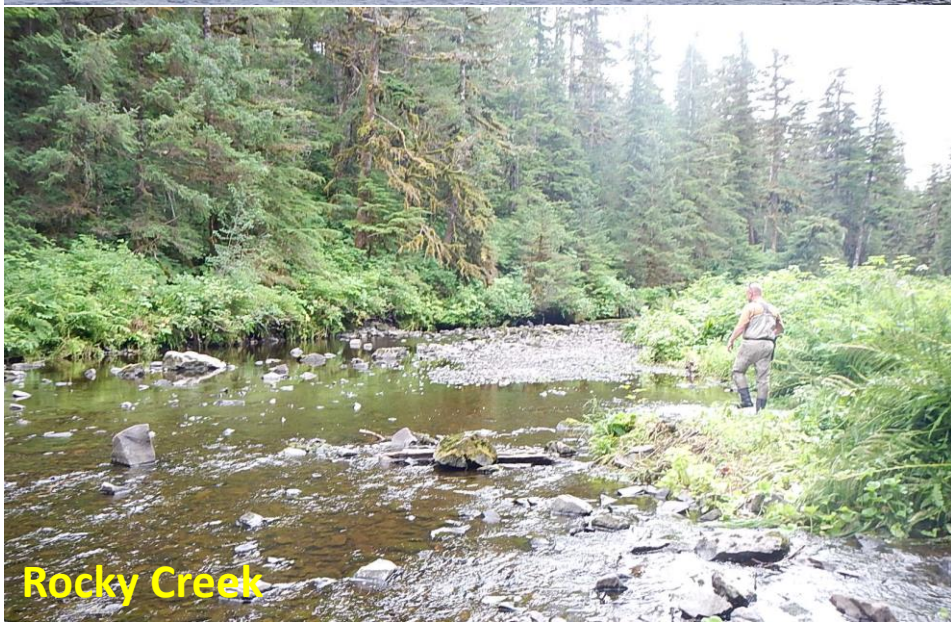
Variability in spawning habitat



Swanson Creek



McCleod Creek



Rocky Creek



Duck River

Previous Studies: Pink Salmon in PWS

Genetic Characterization of Prince William Sound Pink Salmon Populations

Report
to
Alaska Department of Fish and Game
Feb. 15, 1977
by
Jim Seeb
and
Lisa Wishard

INFORMATIONAL LEAFLET NO. 181

SEPARATION OF SOME PINK SALMON (*Oncorhynchus gorbuscha* Walbaum)
SUB-POPULATIONS IN PRINCE WILLIAM SOUND, ALASKA BY LENGTH-WEIGHT
RELATIONSHIPS AND HORIZONTAL STARCH GEL ELECTROPHORESIS

By
Richard B. Nickerson

Ecology of Freshwater Fish 1999: 8: 122-140
Printed in Denmark · All rights reserved

Copyright © Munksgaard 1999

ECOLOGY OF
FRESHWATER FISH
ISSN 0906-6691

Allozyme and mitochondrial DNA variation describe ecologically important genetic structure of even-year pink salmon inhabiting Prince William Sound, Alaska

Seeb JE, Habicht C, Templin WD, Seeb LW, Shaklee JB, Utter FM. Allozyme and mitochondrial DNA variation describe ecologically important genetic structure of even-year pink salmon inhabiting Prince William Sound, Alaska. *Ecology of Freshwater Fish 1999: 8: 122-140*. © Munksgaard, 1999

Abstract – Allozyme and mitochondrial DNA (mtDNA) data were obtained from pink salmon throughout Prince William Sound, Alaska, from two hatchery, five upstream, and 20 tidal locations distributed among five management regions collected during 1994. Screening for allozymes included 66 loci for 92 to 100 fish per sample. Thirty-four loci had variant allele frequencies >0.01 in one or more collections and were used for population analyses. Eight haplotypes were detected after screening 40 fish per collection for variation at the NDS/ND6 region of mtDNA using six restriction enzymes. Significant and apparently stable differences detected by both data sets permit rejecting a null hypothesis of panmixia and support managing native populations in Prince William Sound at the regional level. Distinctions between upstream and tidal collections were detected within Lagoon Creek (allozymes) and Koppen Creek (mtDNA). Significant regional heterogeneity was detected within upstream (allozymes and mtDNA) and tidal (allozymes) collections; however, upstream collections were more divergent from each other than were tidal collections. The absence of distinction of Armin F. Koernig Hatchery from almost all regions was consistent with multiple origins of this stock. Conversely, Solomon Gulch Hatchery in the East Region was distinct from all regions but East, consistent with a more restricted origin and influence.

**J. E. Seeb¹, C. Habicht¹,
W. D. Templin¹, L. W. Seeb¹,
J. B. Shaklee², F. M. Utter³**

¹Alaska Department of Fish & Game, Commercial Fisheries Division, Anchorage, Alaska, ²Washington Department of Fish & Wildlife, Olympia, ³School of Fisheries, University of Washington, Seattle, Washington, USA

Key words: allozyme; mtDNA; genetics; pink salmon

J. E. Seeb, Alaska Department of Fish & Game, Commercial Fisheries Division, Anchorage, AK 99518, USA

Accepted for publication April 9, 1999

Un resumen en español se incluye detrás del texto principal de este artículo.

Study Design

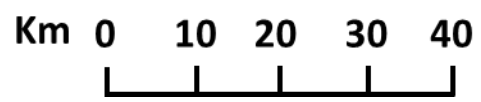
		Contemporary	Historical
Odd Year	Natural		
	Hatchery		
Even Year	Natural		
	Hatchery		

Study Design

		Contemporary	Historical
Odd Year	Natural	✓	<i>(pending)</i>
	Hatchery	✓	<i>(pending)</i>
Even Year	Natural		
	Hatchery		

Prince William Sound

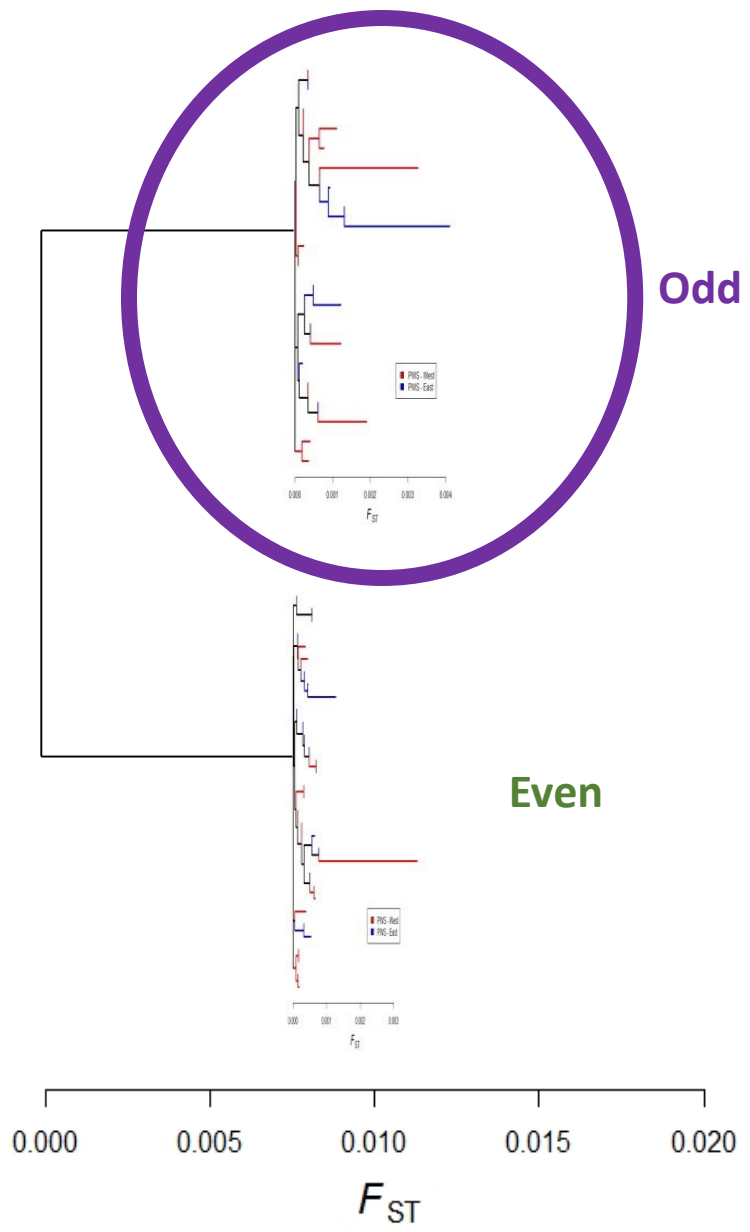
- ▲ Hatchery
- East
- West



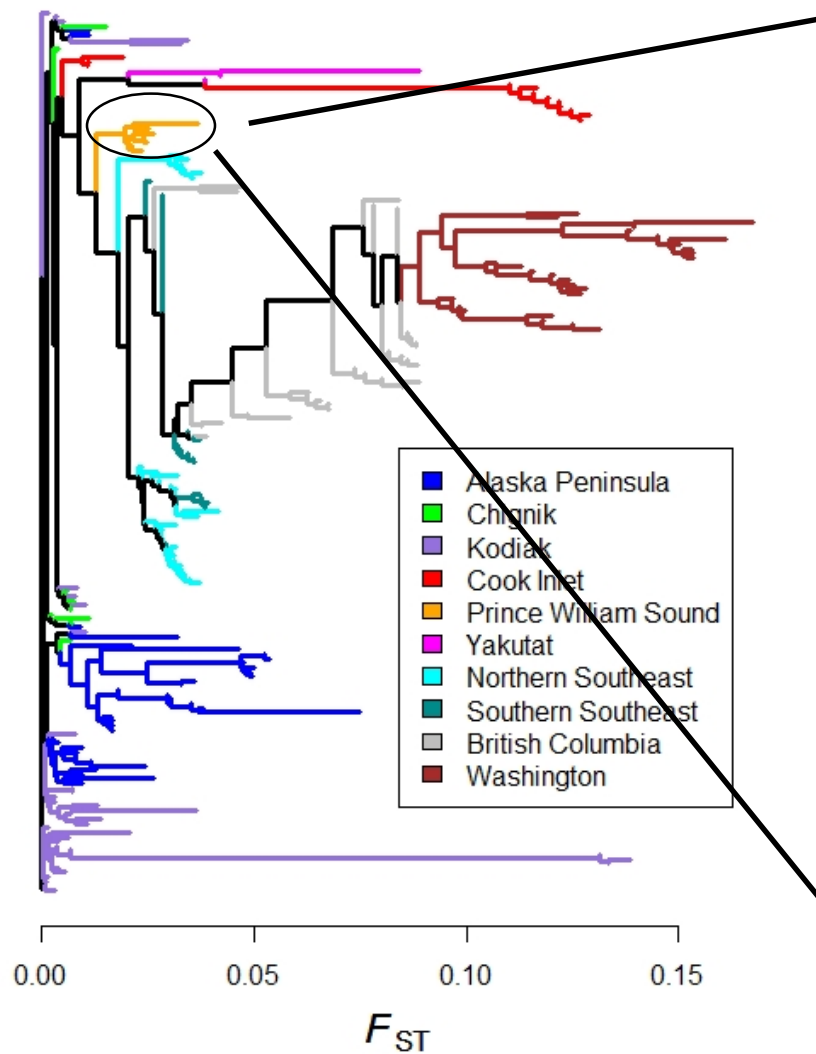
Population Structure Analyses

- **Calculate genetic differences among collections**
- **Test for significance of these differences**
- **Visualize the relationships among collections**

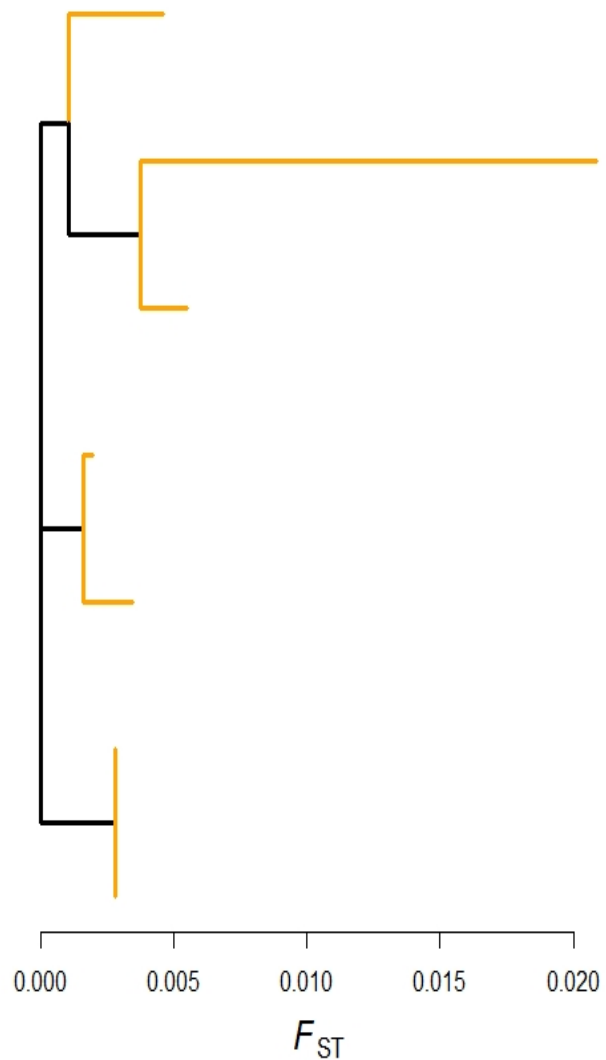
PWS Pink Salmon



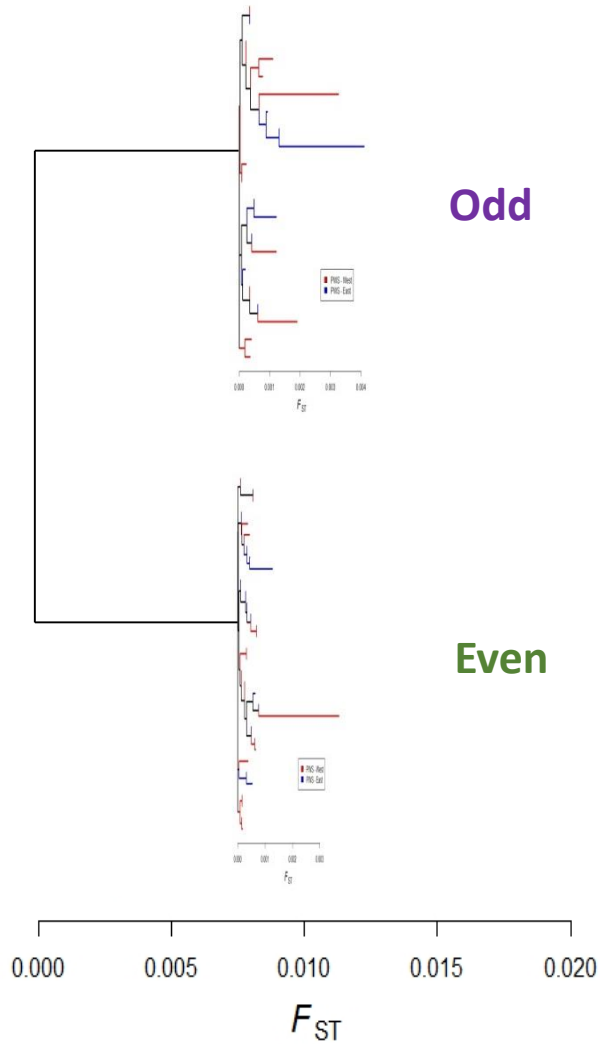
GOA Chum Salmon



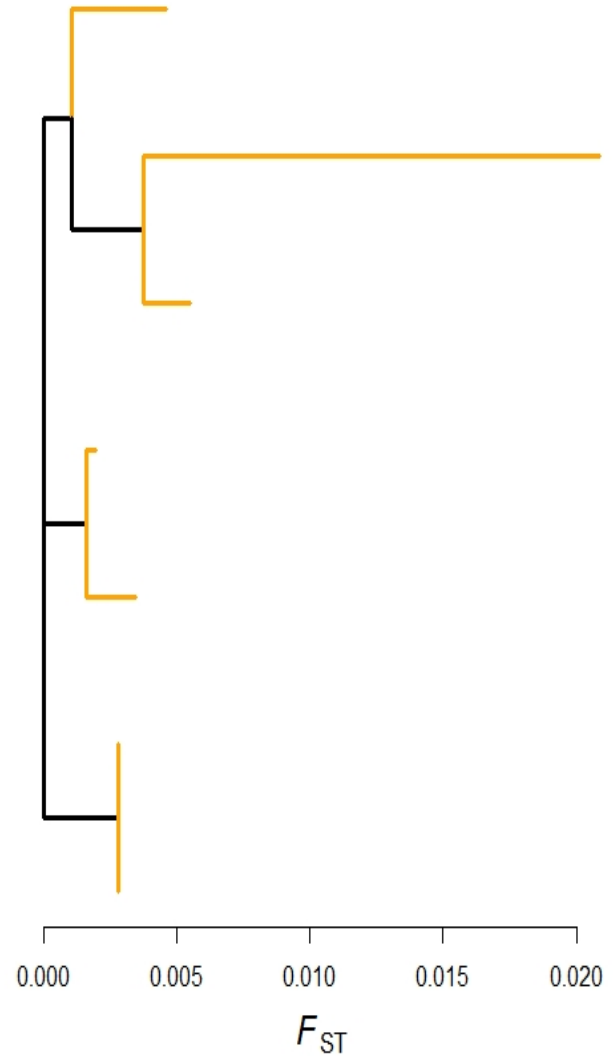
PWS Chum Salmon



PWS Pink Salmon



PWS Chum Salmon



Alaska Hatchery Research Group

Technical
Document:¹
#14

Title: Population Genetic Structure of Odd-Year Pink Salmon from Prince William Sound Based on a Single Year (2013) Version: 1.0

Authors: W. Cheng, C. Habicht, W. D. Templin, Z. D. Grauvogel, S. D. Moffitt, R. E. Brenner, R. P. Josephson, and A. J. Gharrett

Date: May 13, 2016

Abstract

Pink salmon (*Oncorhynchus gorbuscha*) are commercially and ecologically important. In Prince William Sound (PWS), Alaska, pink salmon are the most abundant salmon harvested and generating the highest total value. An understanding of their population genetic structure is useful for conservation and management, especially given the magnitude of the hatchery program in the sound. We analyzed the population genetic structure of pink salmon from four hatcheries and 19 natural spawning areas in PWS and one hatchery in Kodiak Management Area (KMA) by genotyping 16 microsatellite loci for nearly 3000 pink salmon sampled in 2013. Across all populations in PWS, the number of alleles observed per locus ranged from 11 (*Os7e*) to 87 (*Ok10I*), and the total for all loci was 726. The fixation index (F_{ST}), a measure of population differentiation, was 0.002 over all loci and the F_{ST} of individual loci ranged from 0.001 to 0.003. Significant difference was detected among those populations from PWS, which means that pink salmon in PWS are not from a single large homogeneous population. The KMA collection was the most divergent. Within PWS, Solomon Gulch Hatchery in the northeastern PWS was distinct from all other collections and suggested that it had not received many migrants from other PWS areas. Early-run fish from Snug Harbor Creek were distinct from other samples.

Key words: Pink salmon, odd-year, Prince William Sound, population genetic structure, microsatellite.

Population structure of odd-broodline Asian pink salmon and its contrast to the even-broodline structure

S. L. HAWKINS*, N. V. VARNAVKAYA†, E. A. MATZAK‡, V. V. EFREMOV¶, C. M. GUTHRIE III*, R. L. WILMOT*, H. MAYAMA§, F. YAMAZAKI|| AND A. J. GHARRETT##**

*National Marine Fisheries Service, Auke Bay Laboratory, 11305 Glacier Hwy, Juneau, AK 99801-8626, U.S.A., †Kamchatka Scientific Research Institute of Fisheries and Oceanography, KamchatNIRO, Petropavlovsk-Kamchatsky 683602, Kamchatka, Russia, ‡Pacific Research Fisheries Centre (TINRO-CENTRE), 4 Shevchenko Alley, Vladivostok 690600, Russia, §Russian Academy of Sciences, Far East Branch, Institute of Marine Biology, Vladivostok 690041, Russia, ¶National Salmon Resources Center, Nakanoshima, Sapporo 062-0922, Japan, ||Hokkaido University, Laboratory of Genetics and Embryology, Faculty of Fisheries, Hakodate 041, Japan and ##Division of Fisheries, University of Alaska Fairbanks, 11120 Glacier Hwy, Juneau, AK 99801, U.S.A.

(Received 6 June 2001, Accepted 28 November 2001)

Most of the variation (99%) of Asian odd-broodline pink salmon *Oncorhynchus gorbuscha*, based on data at 32 variable (46 total) allozyme loci from 35 populations, occurred within populations. The remaining interpopulation variation was attributable to: (1) differences between northern (the northern Sea of Okhotsk, eastern Kamchatka Peninsula and western Kamchatka Peninsula) and southern (Hokkaido Island, Kuril Islands and Sakhalin Island) populations; (2) differences between the southern areas; (3) low variation among populations within some areas. The pattern contrasted strongly with that observed for Asian even-broodline populations, which had a strong structure, possibly related to geographic and oceanographic influences. Isolation-by-distance analyses of each of the two broodlines showed a stronger relationship ($\times 4.8$) among even- than odd-broodline populations. Allele frequency differences between even- and odd-broodlines reflected the reproductive isolation of the broodlines. However, there were no fixed frequency differences which, considered with the differing population structures, suggests that migration-drift equilibrium has not yet obtained in one or both broodlines. The structural differences also suggest it is likely that the even- and odd-broodlines are of different ages and that one is derived from the other. Allozyme data do not provide a genealogical basis for identifying the ancestral lineage.

Key words: *Oncorhynchus gorbuscha*; pink salmon; population structure; allozyme; isolation by distance.

Genetic Interpretation of Broad-Scale Microsatellite Polymorphism in Odd-Year Pink Salmon

JEFFREY B. OLSEN*¹ AND LISA W. SEEB

Alaska Department of Fish and Game, Genetics Program
333 Raspberry Road, Anchorage, Alaska 99518, USA

PAUL BENTZEN

Marine Molecular Biotechnology Laboratory, University of Washington
3707 Brooklyn Avenue Northeast, Seattle, Washington 98105-6715, USA

JAMES E. SEEB

Alaska Department of Fish and Game, Genetics Program

Abstract.—We examined genetic variation at five microsatellite loci in 12 odd-year populations and one even-year population of pink salmon *Oncorhynchus gorbuscha* from six geographic regions of North America. The degree of polymorphism varied widely among loci. The total number of alleles in the odd-year samples varied from 4 (*Onej3*) to 53 (*Ssa85*). A probability test revealed significant heterogeneity in allele frequencies among all odd-year samples and among pooled odd-year samples from six regions. We compared estimates of a standard index of population structure (θ) based on variance in allele frequency with a new index suggested for microsatellites (θ_{ST}) based on variance in allele size. Our results suggest θ is a better estimator of intralineage (odd-year \times odd-year) population structure, whereas θ_{ST} is best suited for estimating interlineage (odd-year \times even-year) population structure. The difference in performance of θ and θ_{ST} for estimating intralineage and interlineage population structure suggests high migration rates and possibly recent low divergence times are dominant influences on genetic population structure in odd-year pink salmon. We showed statistical support for genetic isolation by distance and geographically correlated allele frequency clines, suggesting broad-scale gene flow is best described by a linear stepping-stone model. An analysis of molecular variation showed weak but significant regional structuring under two different population grouping schemes. Our results suggest broad-scale population aggregations of odd-year pink salmon are temporally stable but that differentiation is weak, presumably due to migration.

Electrophoretic Characterization of Odd-Year Pink Salmon (*Oncorhynchus gorbuscha*) Populations from the Pacific Coast of Russia, and Comparison with Selected North American Populations

James B. Shaklee

Washington Department of Fish and Wildlife, 600 Capital Way N, Olympia, WA 98501-1091, U.S.A.

and Natalya V. Varnavskaya

Kamchatka Branch of Pacific Research Institute of Fisheries and Oceanography (KoTRINRO), Petropavlovsk-Kamchatsky, 683602, Naberejnaja 18, Russia

Shaklee, J.B., and N.V. Varnavskaya. 1994. Electrophoretic characterization of odd-year pink salmon (*Oncorhynchus gorbuscha*) populations from the Pacific Coast of Russia, and comparison with selected North American populations. *Can. J. Fish. Aquat. Sci.* 51(Suppl. 1): 158–171.

We collected and electrophoretically analyzed a total of 558 fish from eight locations along the Pacific Coast of Russia. We successfully screened 44 enzyme-coding loci: 14 loci were polymorphic at the 0.95 level in at least one collection, an additional eight were polymorphic at the 0.99 level but not at the 0.95 level, and the remaining 22 were either monomorphic or exhibited only very rare variation in these collections. Contingency χ^2 tests using the 23 most variable loci revealed significant heterogeneity among all eight collections ($p = 0.028$) but little or no significant heterogeneity among collections within areas (northeastern Kamchatka peninsula, $p = 0.180$; southwestern Kamchatka, $p = 0.533$; and mainland adjacent to the northwestern Sea of Okhotsk, $p = 0.071$). Multidimensional scaling and minimum spanning tree analyses using genetic distances among collections indicated that geographic proximity of spawning sites was not associated with genetic similarity. The eight odd-year pink salmon (*Oncorhynchus gorbuscha*) collections from Russia were compared with 16 collections from North America (southeastern Alaska, British Columbia, and Washington) using data for 33 loci. The Russian populations differed from the North American populations in their patterns of allelic variation at many loci. The amount of genetic differentiation among populations from different rivers in Russia was comparable to that seen within similar-sized areas in North America.

Abstract.—Population structure of pink salmon (*Oncorhynchus gorbuscha*) from British Columbia and Washington was examined with a survey of microsatellite variation to describe the distribution of genetic variation. Variation at 16 microsatellite loci was surveyed for approximately 46,500 pink salmon sampled from 146 locations in the odd-year broodline and from 116 locations in the even-year broodline. An index of genetic differentiation, F_{ST} , over all populations and loci in the odd-year broodline was 0.005, with individual locus values ranging from 0.002 to 0.025. Population differentiation was less in the even-year broodline, with a F_{ST} value of 0.002 over all loci, and with individual locus values ranging from 0.001 to 0.005. Greater genetic diversity was observed in the odd-year broodline. Differentiation in pink salmon allele frequencies between broodlines was approximately 5.5 times greater than regional differentiation within

Population structure of pink salmon (*Oncorhynchus gorbuscha*) in British Columbia and Washington, determined with microsatellites

Terry D. Beacham (contact author)¹

Brenda McIntosh¹

Cathy MacConnachie¹

Brian Spilsted²

Bruce A. White³

E-mail address for contact author: Terry.Beacham@dfg-mpo.gc.ca

¹ Fisheries and Oceans Canada

Pacific Biological Station
3190 Hammond Bay Road
Nanaimo, B. C., Canada V9T 6N7

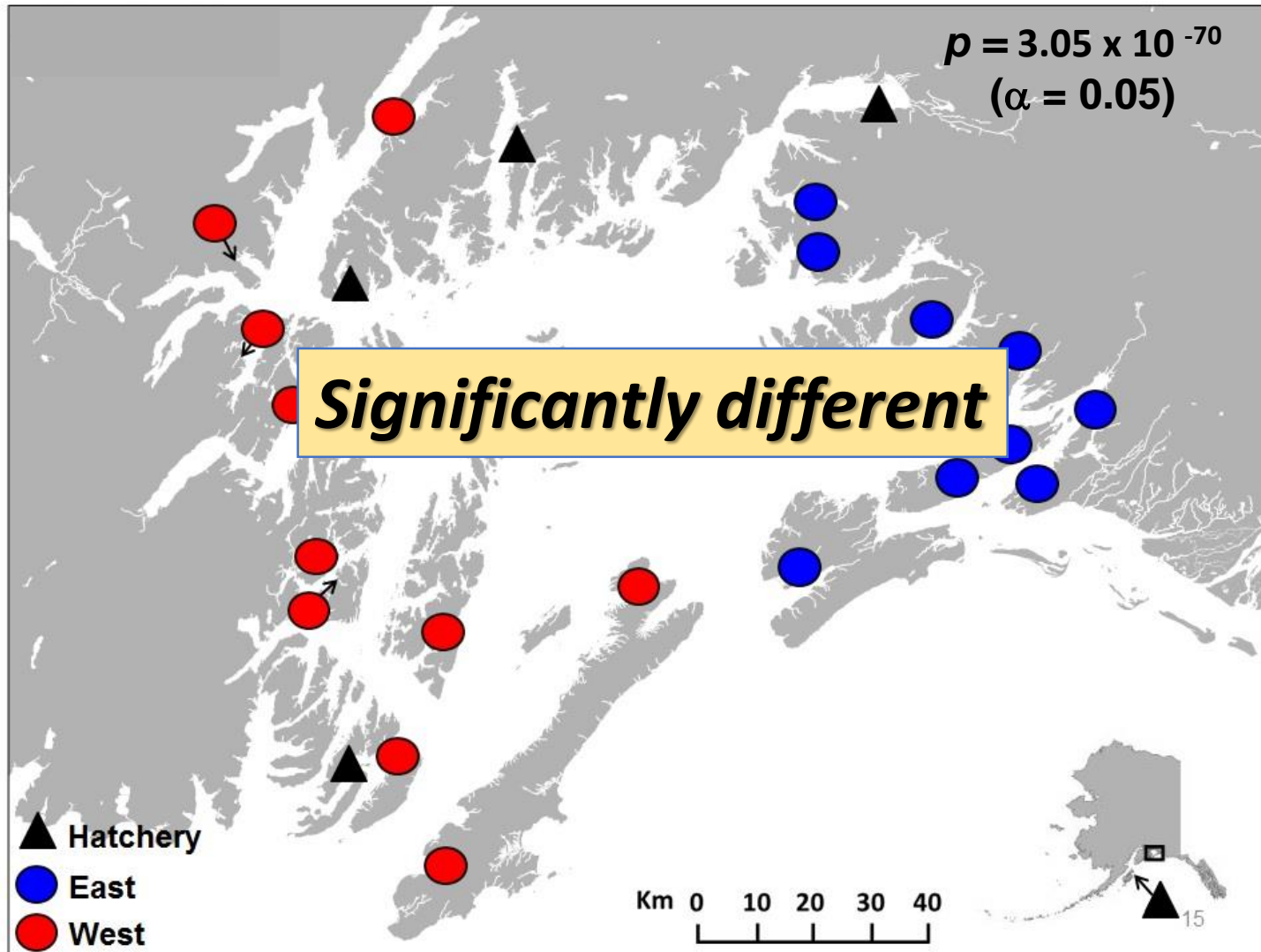
² Fisheries and Oceans Canada
417-2nd Avenue West
Prince Rupert, B. C. Canada V8J 1G8

³ Pacific Salmon Commission
600-1155 Robson Street
Vancouver, B. C., Canada V6E 1B5

Population Structure Analyses

- Calculate genetic differences among collections
- **Test for significance of these differences**

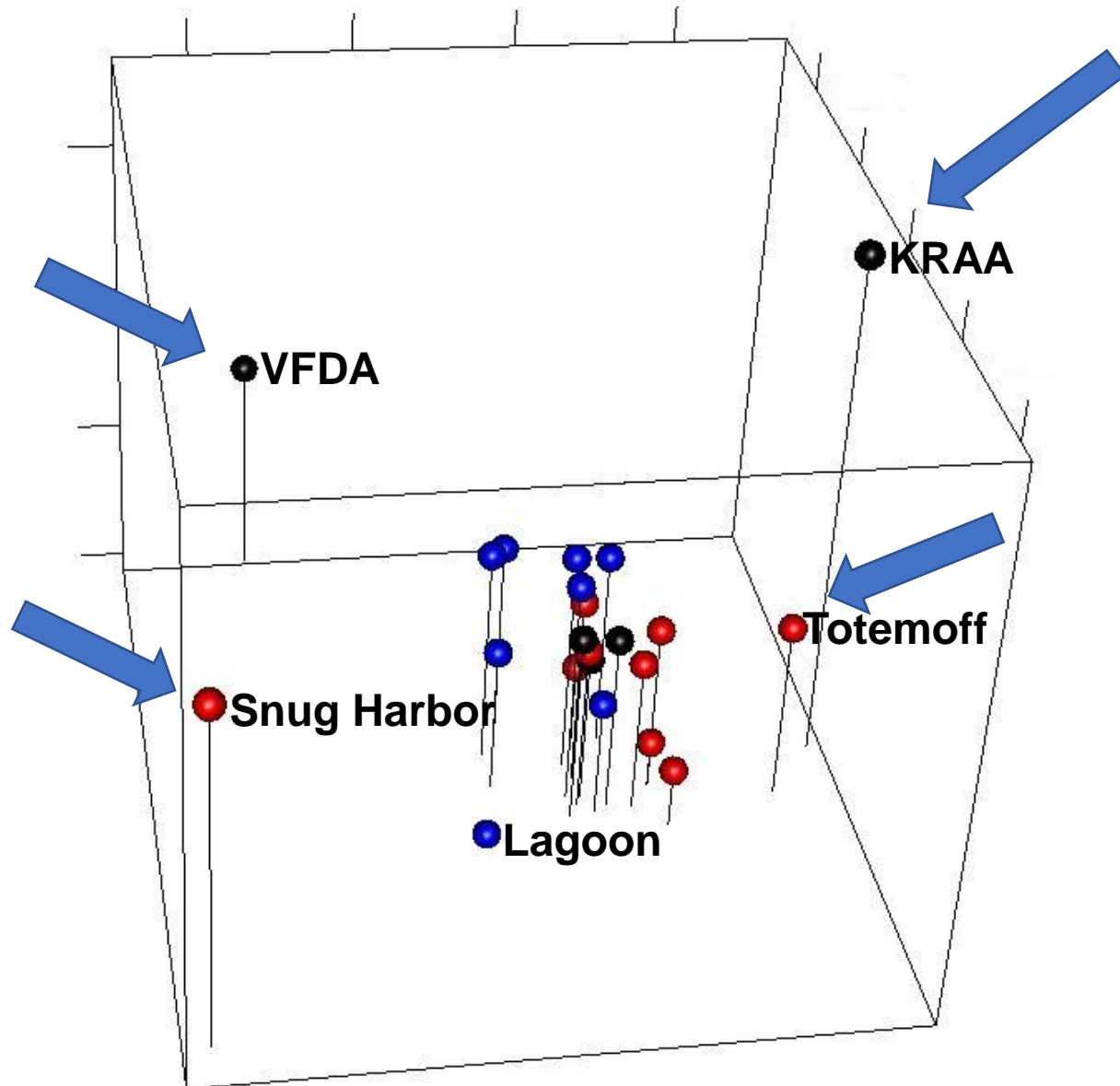
Testing for Differences: among Prince William Sound



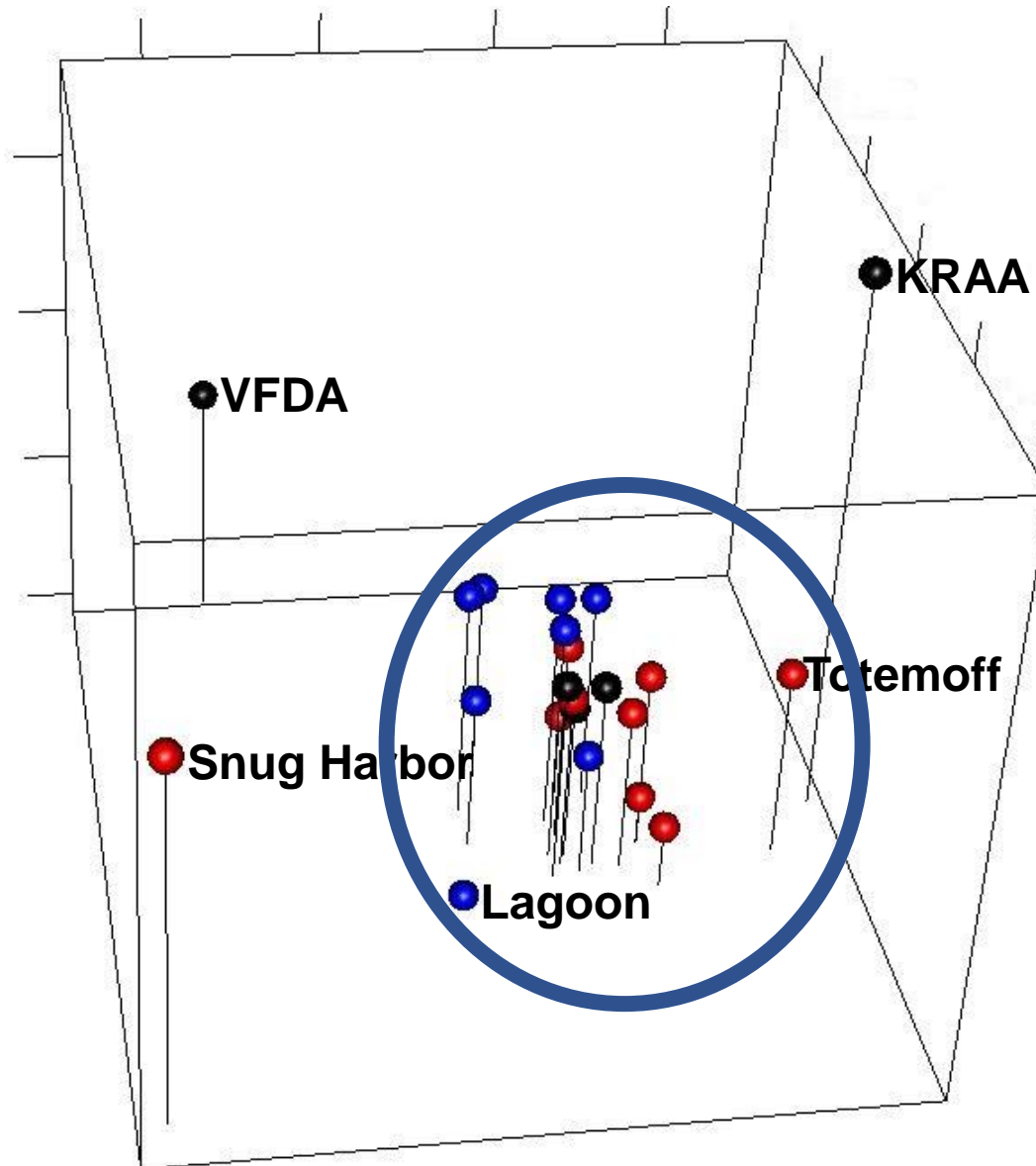
Population Structure Analyses

- Calculate genetic differences among collections
- Test for significance of these differences
- **Visualize the relationships among collections**

Visualizing the Relationships among Collections

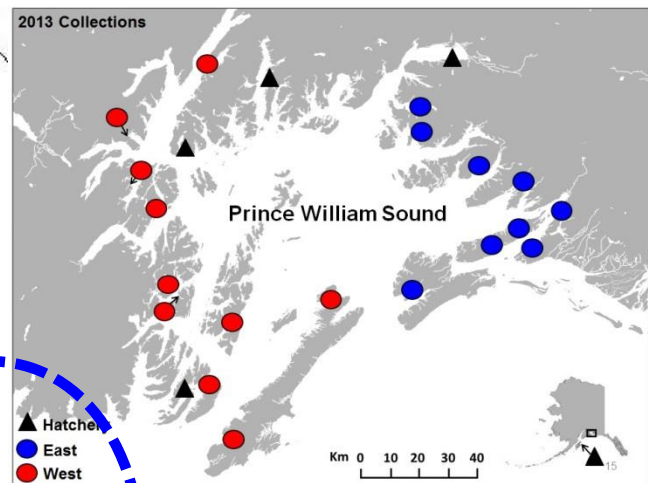
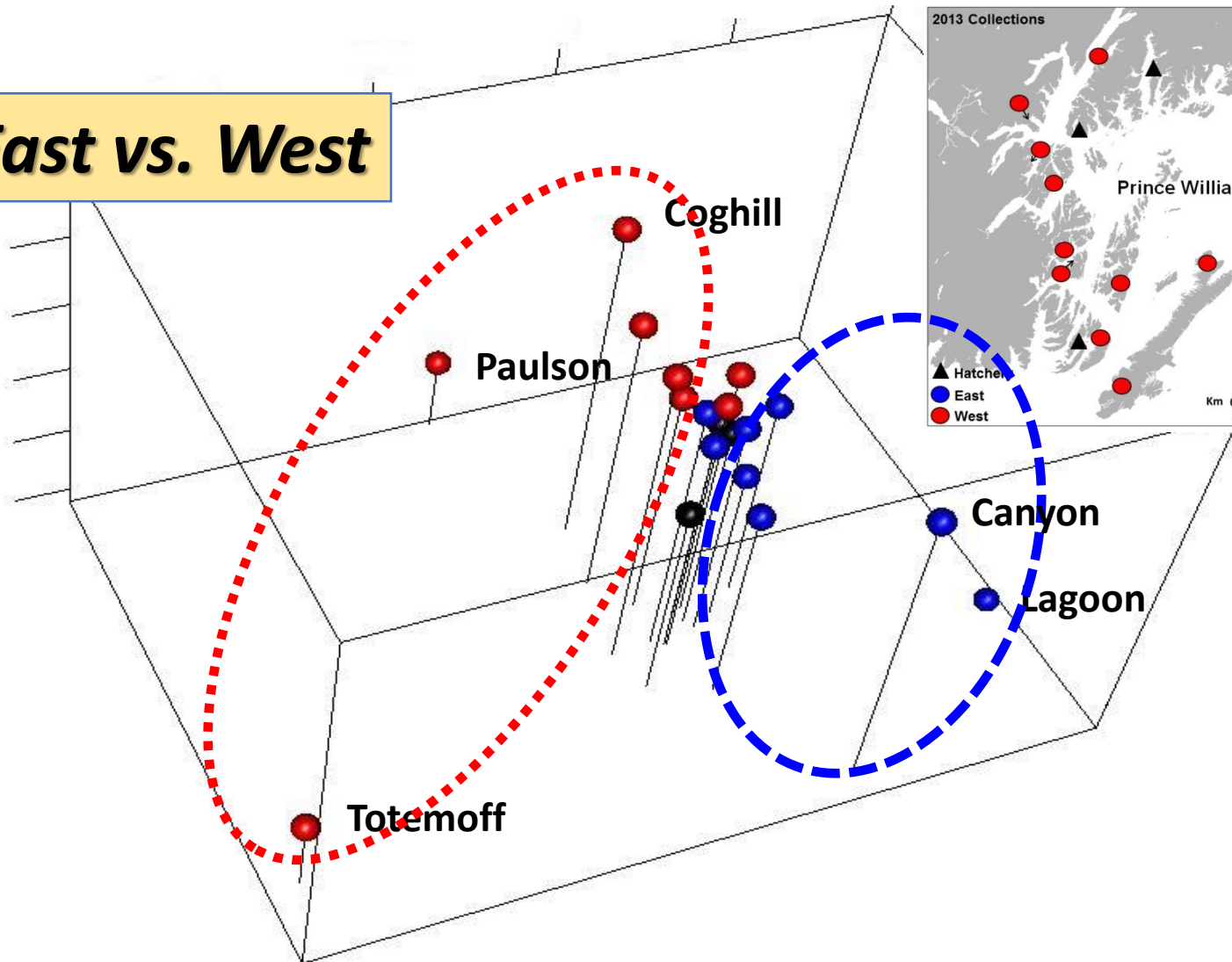


Visualizing the Relationships among Collections



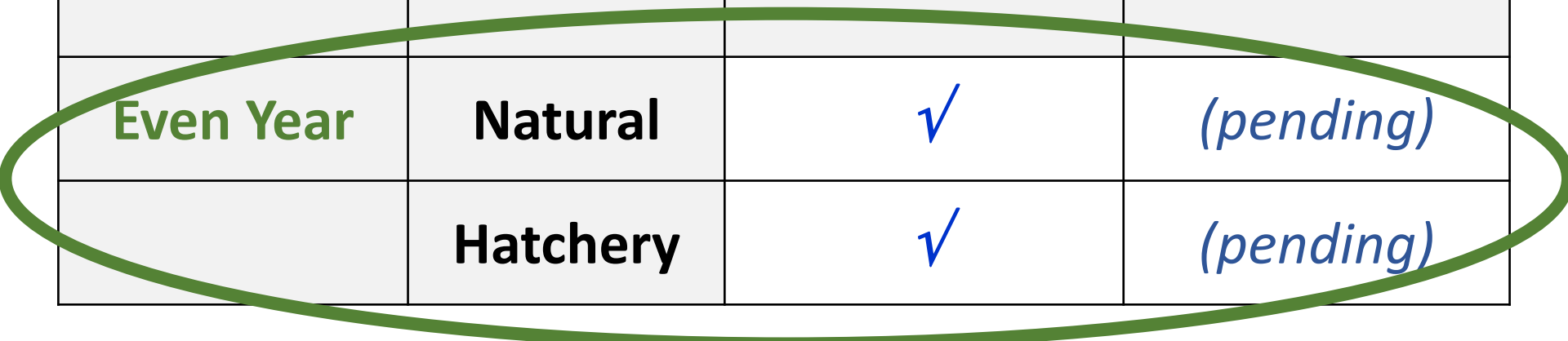
Visualizing Relationships among Collections – Zooming in

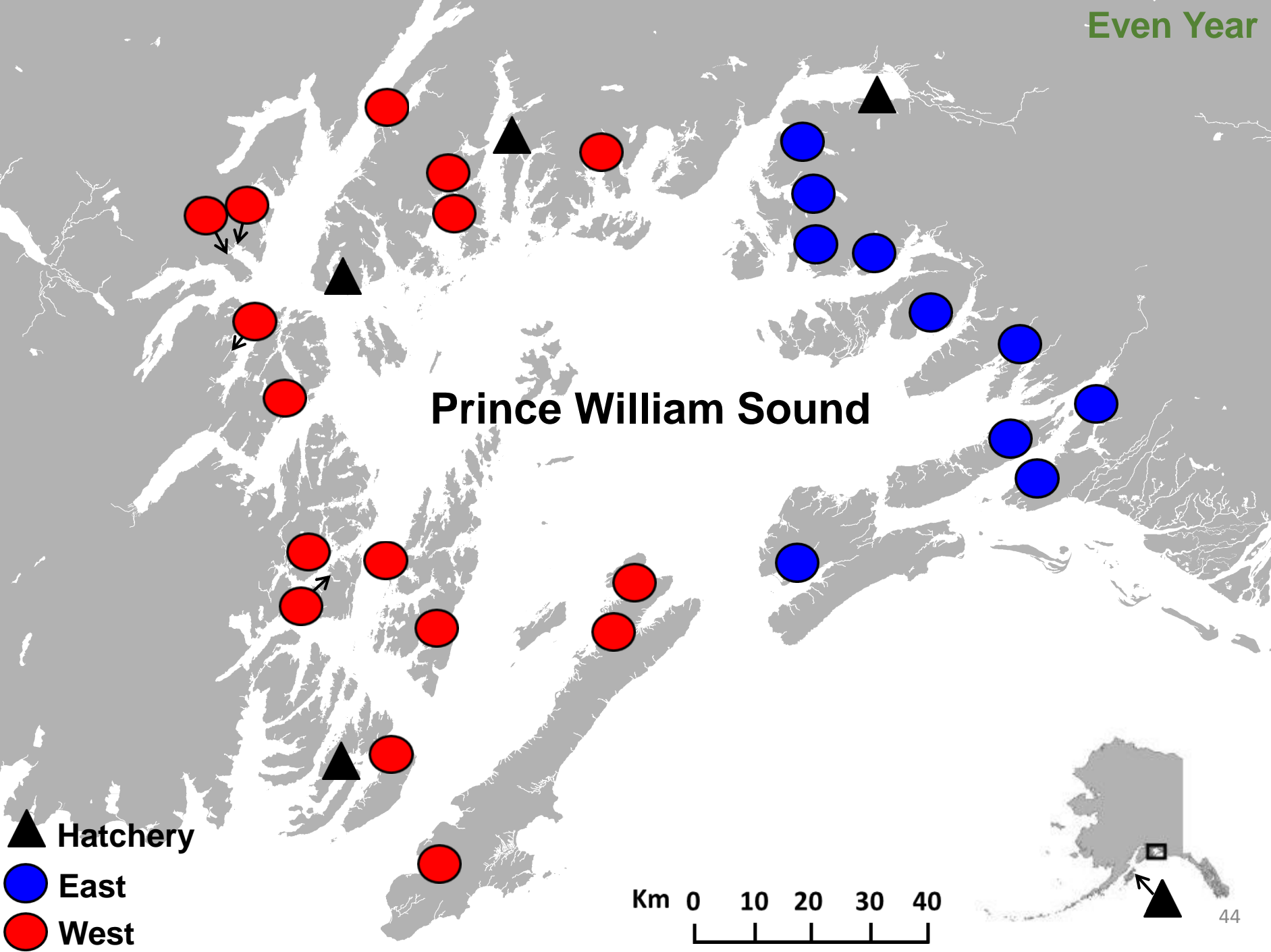
East vs. West



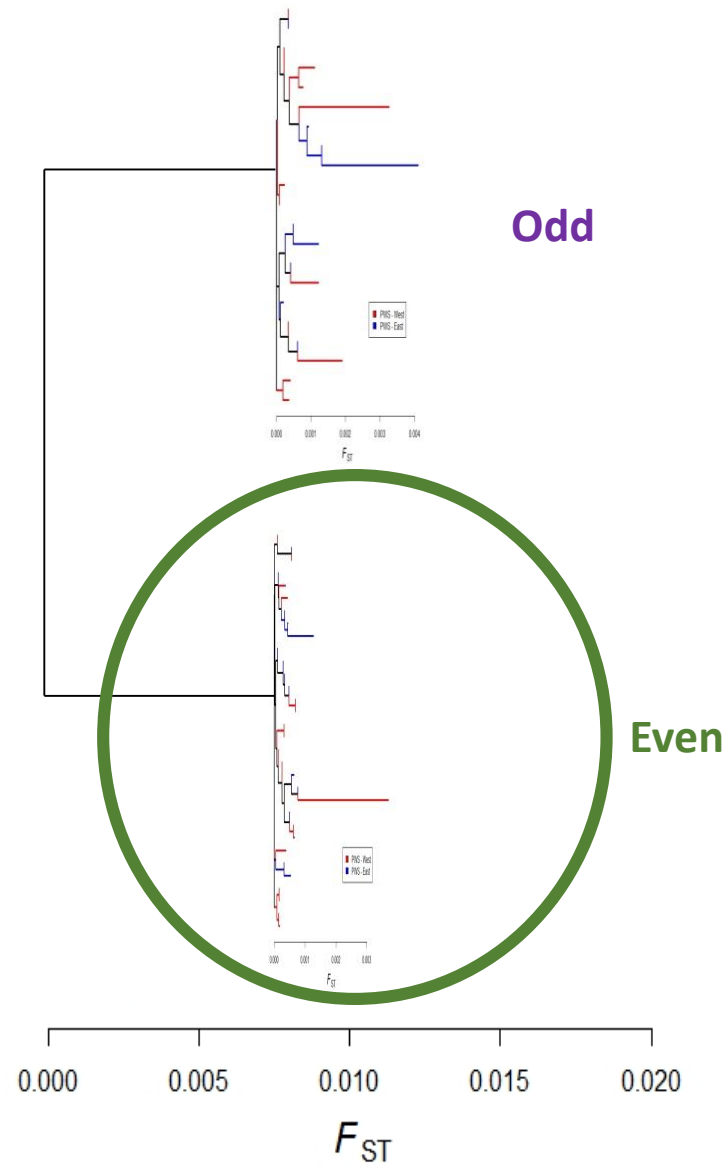
Study Design

		Contemporary	Historical
Odd Year	Natural	✓	<i>(pending)</i>
	Hatchery	✓	<i>(pending)</i>
Even Year	Natural	✓	<i>(pending)</i>
	Hatchery	✓	<i>(pending)</i>





PWS Pink Salmon



Abstract

Pink salmon (*Oncorhynchus gorbuscha*) are commercially and ecologically important. In Prince William Sound (PWS), Alaska, pink salmon are the most abundant Pacific salmon species and generate the highest total value for commercial fishery. Pink salmon have a fixed two-year life cycle, which has created reproductively separate broodlines in even- and odd-years. An understanding of their population genetic structure is useful for conservation and management, especially given the magnitude of the hatchery program in the sound. We analyzed the population genetic structure of pink salmon from four hatcheries and 26 natural spawning areas in PWS and one hatchery in Kodiak Management Area (KMA) by genotyping 16 microsatellite loci for nearly 6,554 pink salmon sampled in 2014. The fixation index (F_{ST}), a measure of population divergence, was 0.001 over all loci and the F_{ST} of individual loci ranged from 0.001 to 0.002. Significant differences were detected among those populations from PWS, which meant that pink salmon in PWS were not from a single large homogeneous population. The early fish collection from Snug Harbor Creek was the most divergent. The KMA collection was the second most divergent. Solomon Gulch Hatchery in the northeastern PWS was distinct from collections from other PWS districts, which suggested that it had not exchanged many migrants with other districts. The population structure of even-year pink salmon collected in 2014 was not as strong as odd-year pink salmon collected in 2013, where the F_{ST} over all loci was an order of magnitude higher.

Key words: Pink salmon, even-year, hatchery, Prince William Sound, population genetic

Allozyme and mitochondrial DNA variation describe ecologically important genetic structure of even-year pink salmon inhabiting Prince William Sound, Alaska

Seeb JE, Habicht C, Templin WD, Seeb LW, Shaklee JB, Utter FM. Allozyme and mitochondrial DNA variation describe ecologically important genetic structure of even-year pink salmon inhabiting Prince William Sound, Alaska. Ecology of Freshwater Fish 1999; 8: 122-140. © Munksgaard, 1999

Abstract – Allozyme and mitochondrial DNA (mtDNA) data were obtained from pink salmon throughout Prince William Sound, Alaska, from two hatchery, five upstream, and 20 tidal locations distributed among five management regions collected during 1994. Screening for allozymes included 66 loci for 92 to 100 fish per sample. Thirty-four loci had variant allele frequencies >0.01 in one or more collections and were used for population analyses. Eight haplotypes were detected after screening 40 fish per collection for variation at the ND5/ND6 region of mtDNA using six restriction enzymes. Significant and apparently stable differences detected by both data sets permit rejecting a null hypothesis of panmixia and support managing native populations in Prince William Sound at the regional level. Distinctions between upstream and tidal collections were detected within Lagoon Creek (allozymes) and Koppen Creek (mtDNA). Significant regional heterogeneity was detected within upstream (allozymes and mtDNA) and tidal (allozymes) collections; however, upstream collections were more divergent from each other than were tidal collections. The absence of distinction of Armin F. Koernig Hatchery from almost all regions was consistent with multiple origins of this stock. Conversely, Solomon Gulch Hatchery in the East Region was distinct from all regions but East, consistent with a more restricted origin and influence.

J. E. Seeb¹, C. Habicht¹,
W. D. Templin¹, L. W. Seeb¹,
J. B. Shaklee², F. M. Utter³

¹Alaska Department of Fish & Game, Commercial Fisheries Division, Anchorage, Alaska; ²Washington Department of Fish & Wildlife, Olympia; ³School of Fisheries, University of Washington, Seattle, Washington, USA

Key words: allozyme, mtDNA, genetics, pink salmon

J. E. Seeb, Alaska Department of Fish & Game, Commercial Fisheries Division, Anchorage, AK 99518, USA

Accepted for publication April 9, 1999

Un resumen en español se incluye detrás del texto principal de este artículo.

Abstract—Population structure of pink salmon (*Oncorhynchus gorbuscha*) from British Columbia and Washington was examined with a survey of microsatellite variation to describe the distribution of genetic variation. Variation at 16 microsatellite loci was surveyed for approximately 46,500 pink salmon sampled from 146 locations in the odd-year broodline and from 116 locations in the even-year broodline. An index of genetic differentiation, F_{ST} , over all populations and loci in the odd-year broodline was 0.005, with individual locus values ranging from 0.002 to 0.025. Population differentiation was less in the even-year broodline, with a F_{ST} value of 0.002 over all loci, and with individual locus values ranging from 0.001 to 0.005. Greater genetic diversity was observed in the odd-year broodline. Differentiation in pink salmon allele frequencies between broodlines was approximately 5.5 times greater than regional differentiation within

Population structure of pink salmon (*Oncorhynchus gorbuscha*) in British Columbia and Washington, determined with microsatellites

Terry D. Beacham (contact author)¹

Brenda McIntosh¹

Cathy MacConnachie¹

Brian Spilsted²

Bruce A. White³

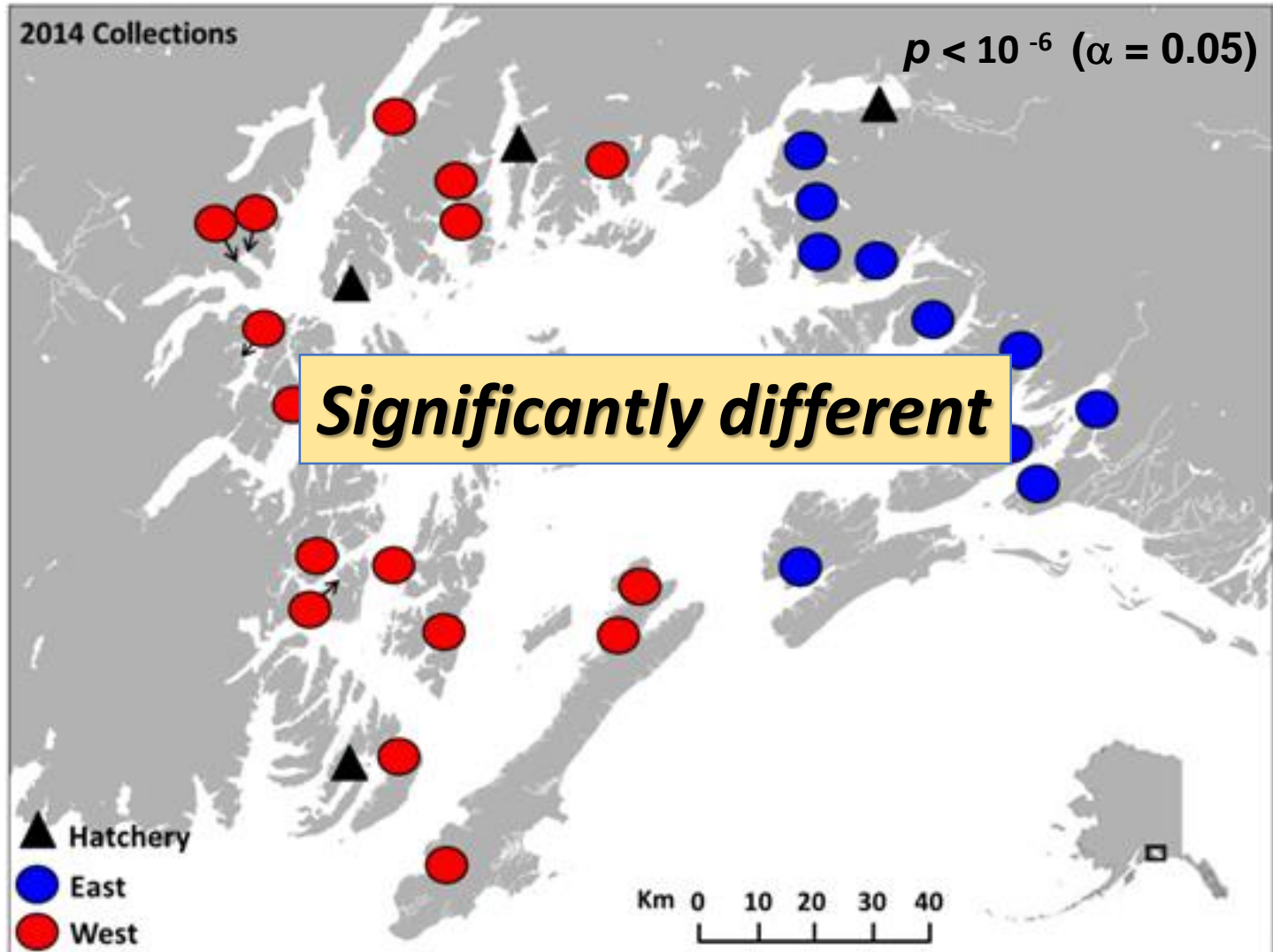
E-mail address for contact author: Terry.Beacham@dfg-mpo.gc.ca

¹ Fisheries and Oceans Canada
Pacific Biological Station
3190 Hammond Bay Road
Nanaimo, B. C., Canada V9T 6N7

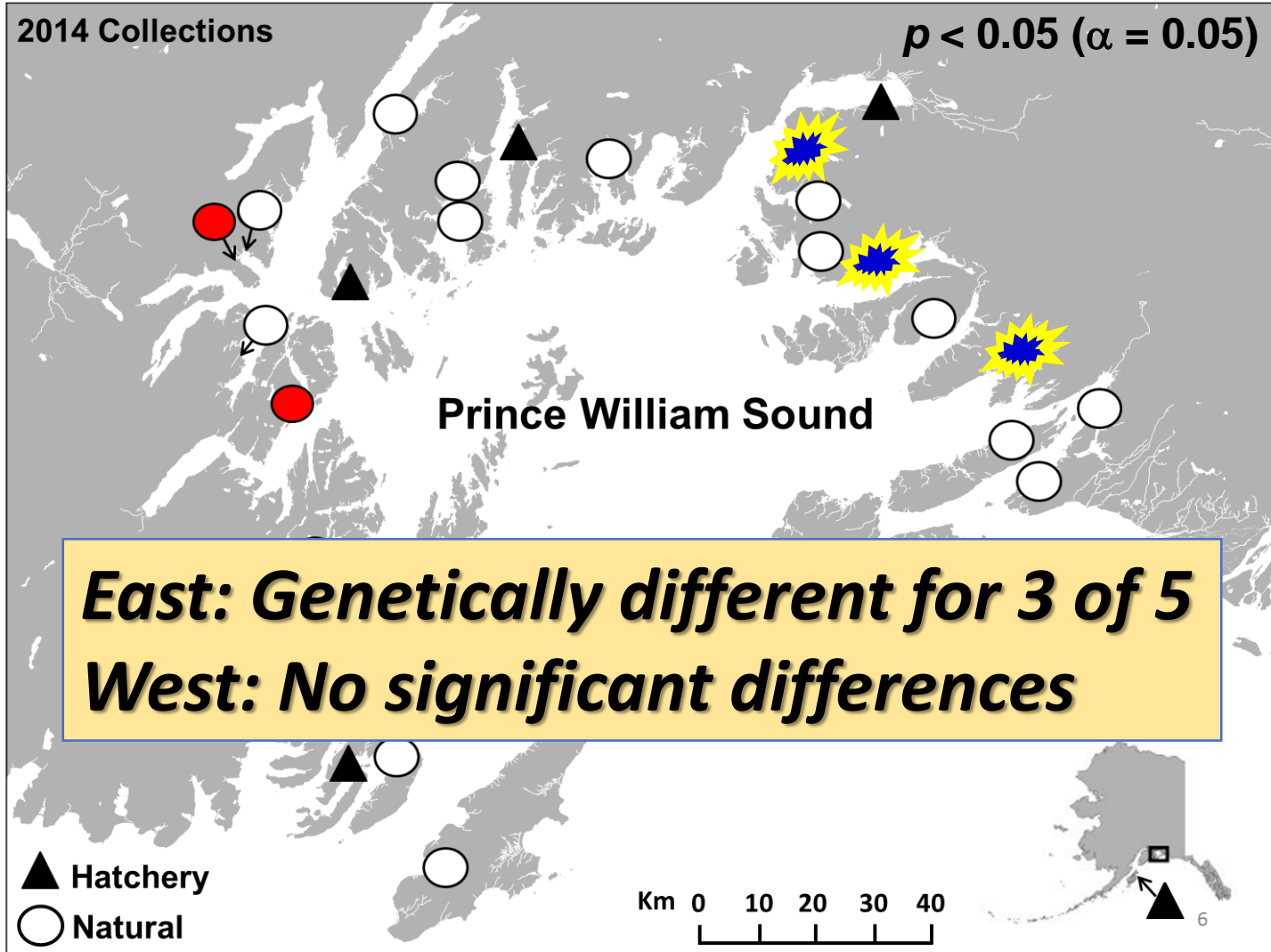
² Fisheries and Oceans Canada
417-2nd Avenue West
Prince Rupert, B. C. Canada V8J 1G8

³ Pacific Salmon Commission
600-1155 Robson Street
Vancouver, B. C., Canada V6E 1B5

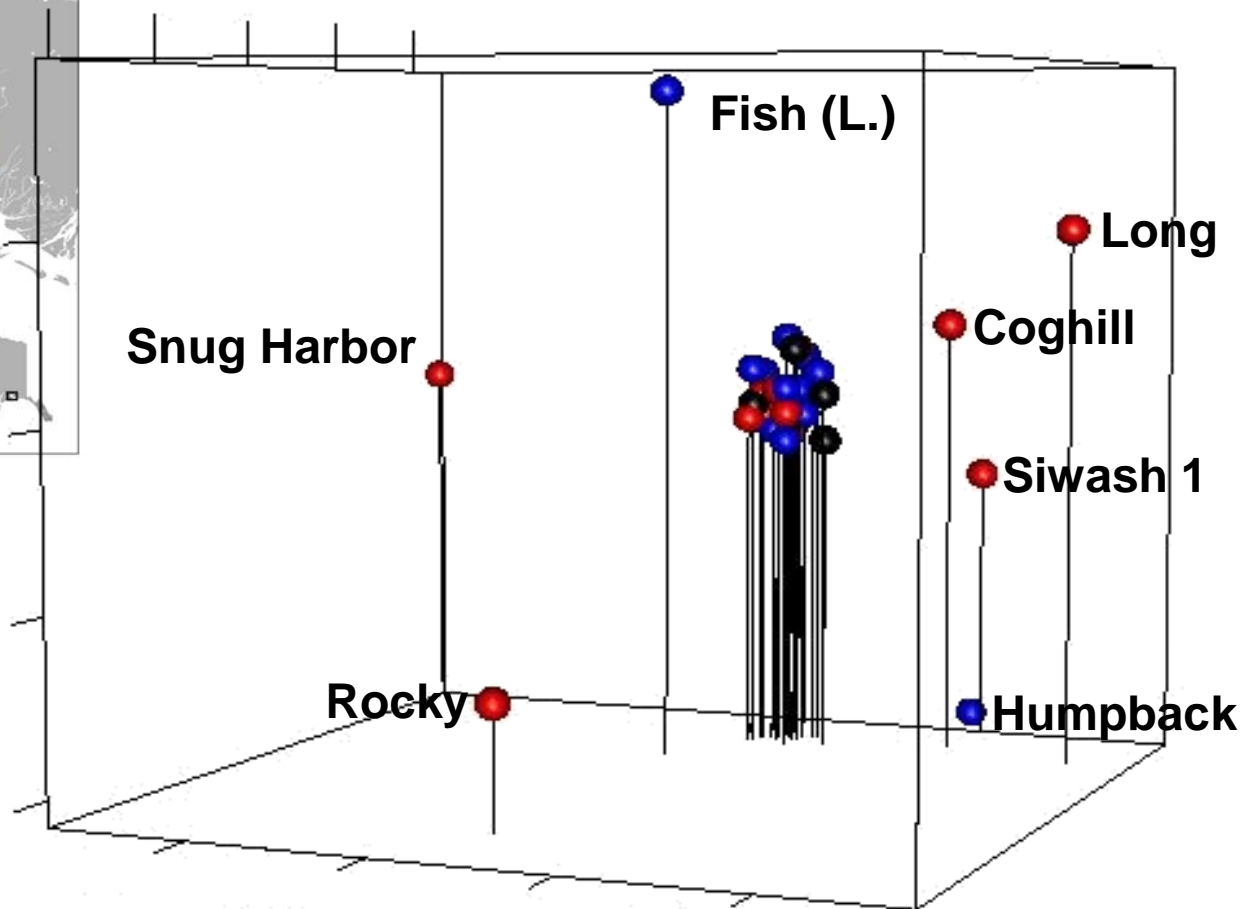
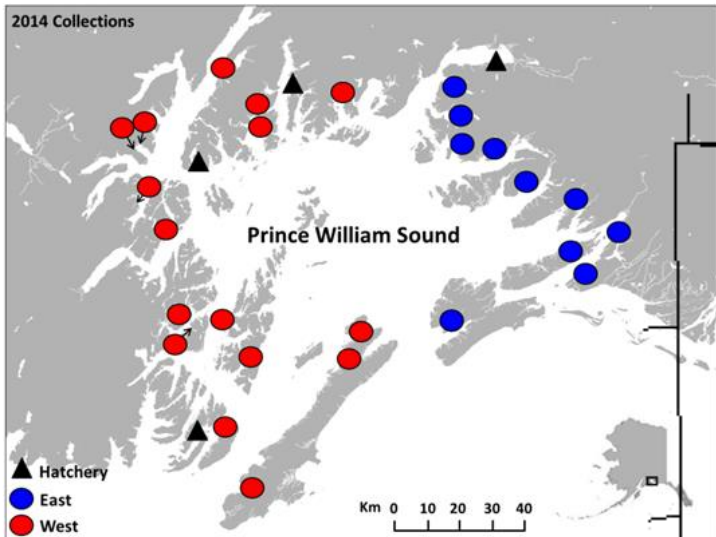
Testing for Differences: among Prince William Sound



Testing for Differences: Between Early and Late Collections



Visualizing the Relationships among Collections



Conclusions to date: Pink salmon structure in PWS

- Genetic variation among pink salmon populations in PWS is very small
 - ✓ Odd year – small
 - ✓ Even year – even smaller

- Kodiak vs. Prince William Sound (PWS) [data not shown]
 - ✓ Significantly different in both lineages

Conclusions to date: Pink salmon structure in PWS

- **Genetic difference within PWS**
 - ✓ Significantly different in both lineages

- **Within lineage patterns**
 - ✓ **Odd year:**
 - ✓ East vs. West
 - ✓ Early vs. Late?
 - ✓ **Even year:**
 - ✓ Early vs. Late (eastern side only)

Future Work

- **Historical samples**
 - ✓ **1991 – 1997**
 - ✓ **No otolith information**

- **Investigate the mechanisms driving the structure**

Acknowledgements

- **Hatcheries**
 - PWSAC, VFDA, KRAA
- **Prince William Sound Science Center**
- **Fisheries and Oceans Canada**
 - Pacific Biological Station
- **Alaska Department of Fish and Game**
- **Alaska Hatchery Research Program Science Panel**
- **University of Alaska Fairbanks**

What is the extent and annual variability of straying?



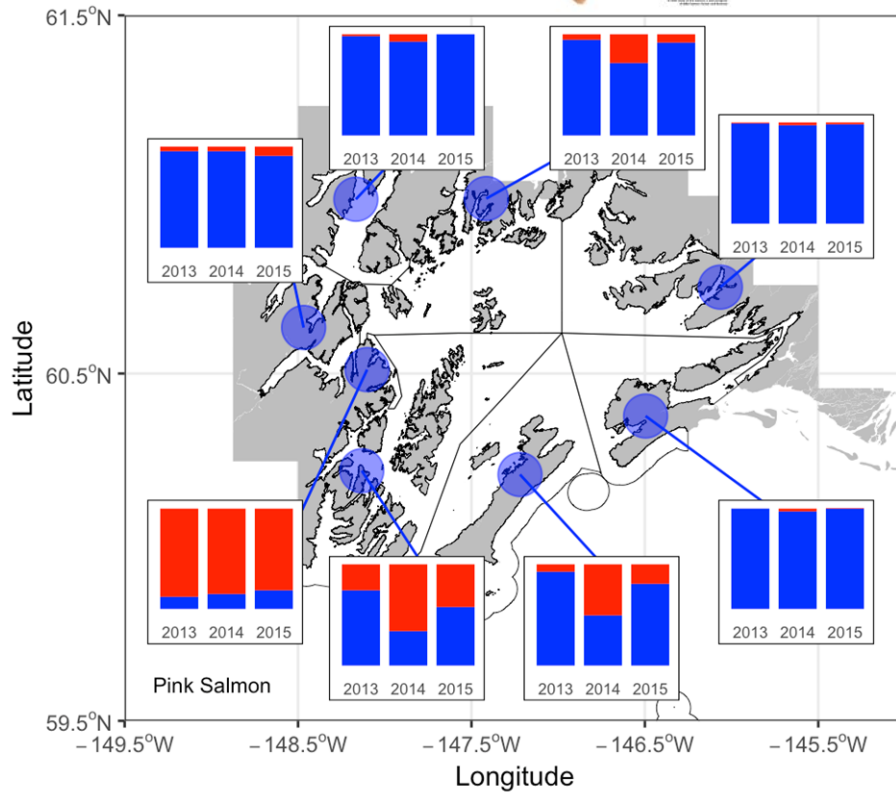
C. Habicht and W. D. Templin

Alaska Department of Fish and Game Gene Conservation Lab

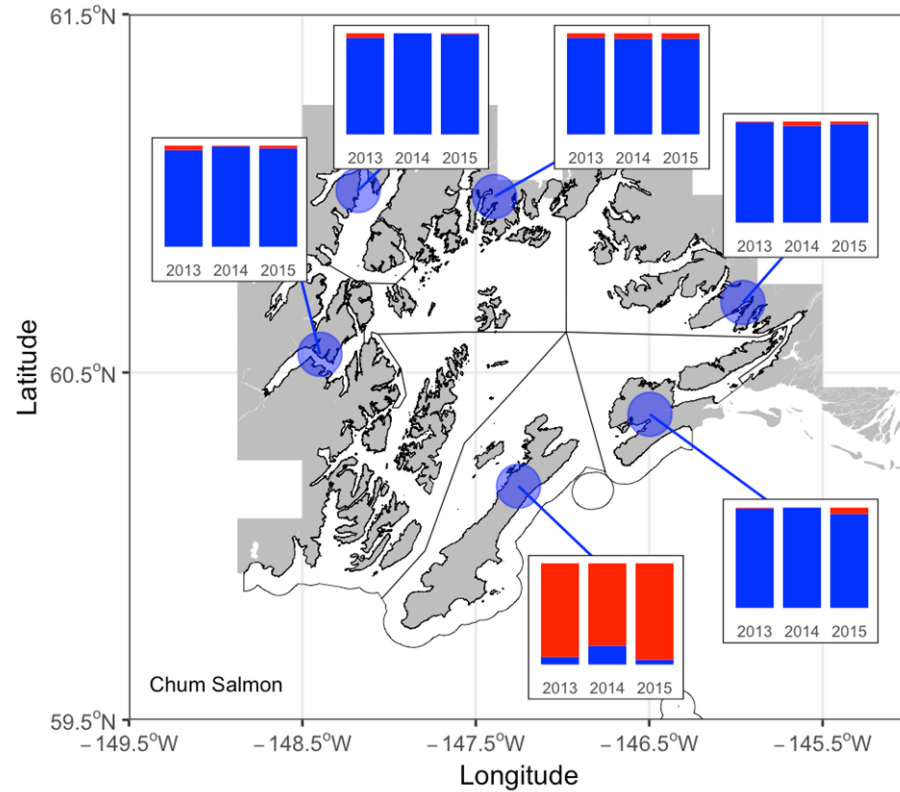
Alaska Board of Fisheries, Hatchery Committee Meeting

March 8, 2019

PWS: Stream results, district averages



0.1% - 89.9%

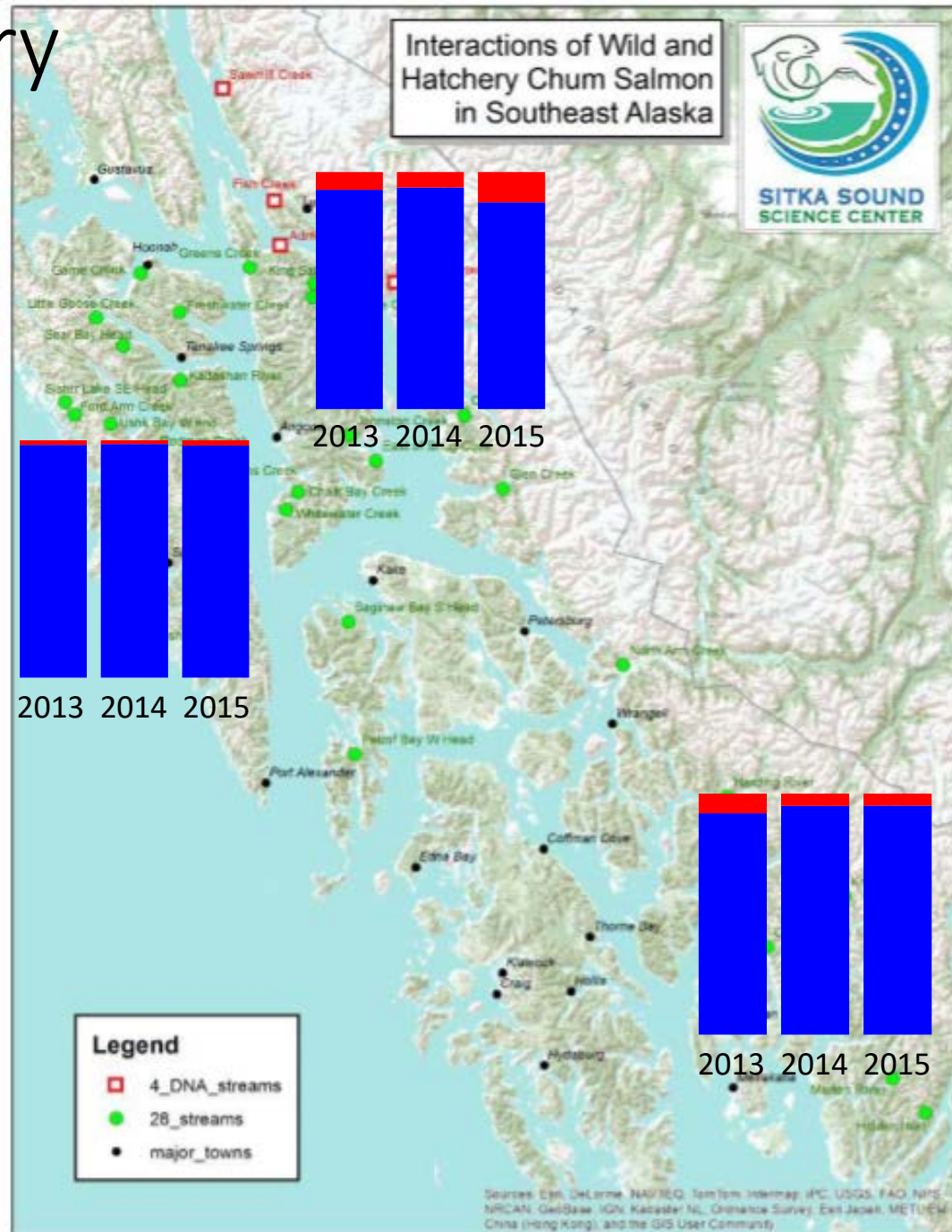


0.0% - 84.6%

Overall PWS hatchery fractions in spawning streams

Species	2013	2014	2015
Pink	4.4%	14.8%	9.5%
Chum	2.8%	3.2%	3.1%

SEAK: Hatchery fraction by stream:
1.5% - 12.7%



Overall SEAK hatchery fractions in spawning streams

Species	2013	2014	2015
Chum	7.3%	5.4%	9.2%

PWS: Run Size and Harvest Rates



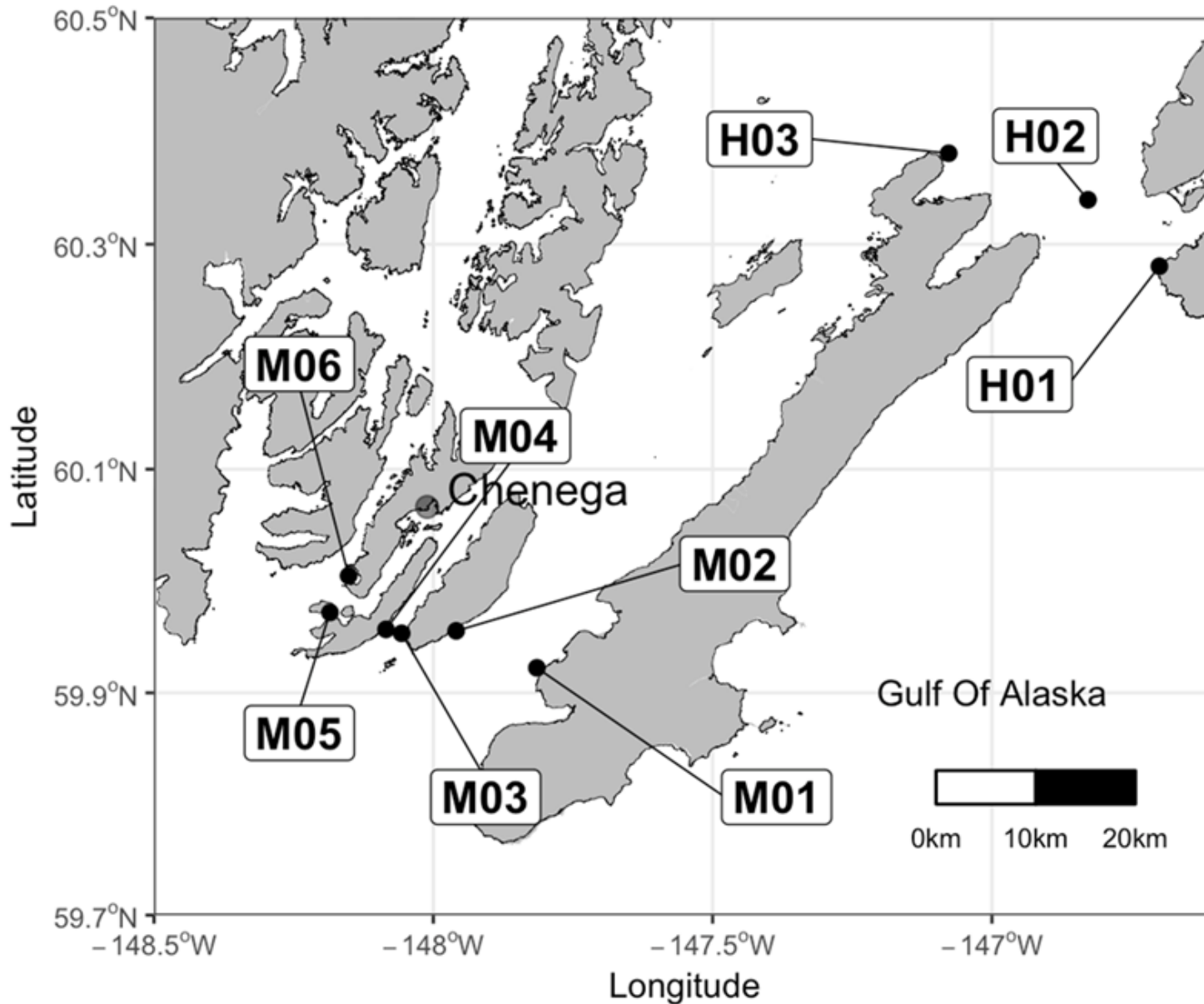
C. Habicht and W. D. Templin

Alaska Department of Fish and Game Gene Conservation Lab

Alaska Board of Fisheries, Hatchery Committee Meeting

March 8, 2019

Ocean Sampling: PWS



Ocean sampling 2013–2015 (PWS only)

- Proportions of hatchery fish in **run**
- Results (7,800 samples):
 - Pink salmon: 55 - 86%
 - Chum salmon: 51 - 73%

Species Common Name	Year	Hatchery Proportion	SE
Pink Salmon	2013	0.679	.016
	2014	0.864	.03
	2015	0.549	.004
Chum Salmon	2013	0.725	.019
	2014	0.511	.029
	2015	0.688	.015

Wild and Hatchery run size estimates

- Preliminary PWS run size estimates; 2013-2015
(Thousands)

Species Year	Natural spawners	Hatchery strays	Total spawners	Natural run	Hatchery run	Total run
Pink salmon						
2013	15,698	701	16,399	33,096	69,888	102,985
2014	5,130	741	5,872	6,960	42,757	49,718
2015	37,972	4,009	41,981	63,531	77,335	140,866
Chum salmon						
2013	894	50	944	1,141	3,007	4,148
2014	925	49	975	1,175	1,228	2,404
2015	890	28	919	1,128	2,484	3,612

Knudsen et al. (2016). Interactions of Wild and Hatchery Pink Salmon and Chum Salmon in Prince William Sound and Southeast Alaska.

Natural and Hatchery harvest rate estimates:
PWS pink salmon

	Estimated Harvest Rates	
Year	Hatchery	Natural
2013	0.99	0.53
2014	0.98	0.26
2015	0.95	0.40

AHRP Fitness Study: PWS Pink Salmon



Emily Lescak, K. Shedd, D. Prince, H. Hoyt, T. Dann, C. Habicht

Alaska Department of Fish and Game Gene Conservation Lab

Alaska Board of Fisheries Hatchery Committee

March 8, 2019

Alaska Hatchery Research Program

- 1) What is the genetic structure of pink and chum in PWS and SEAK?
- 2) What is the extent and annual variability of straying?
- 3) What is the impact on fitness (productivity) of natural pink and chum stocks due to straying hatchery pink and chum salmon?

Hatchery/Natural Fitness

Steelhead

433

Differential reproductive success of sympatric, naturally spawning hatchery and wild steelhead trout (*Oncorhynchus mykiss*) through the adult stage

Jennifer E. McLean, Paul Bentzen, and Thomas P. Quinn

MOLECULAR ECOLOGY

Molecular Ecology (2011) 20, 1860–1869 doi: 10.1111/j.1365-2942.2011.02684.x

Reduced reproductive success of hatchery coho salmon in the wild: insights into most likely mechanisms

VÉRONIQUE THÉRIAULT,* GREGORY R. MOYER,[†] LAURA S. JACKSON,[‡] MICHAEL S. BLOUIN[‡] and MICHAEL A. BANKS*

Genetic Effects of Captive Breeding Cause a Rapid, Cumulative Fitness Decline in the Wild

Hitoshi Araki,¹ Becky Cooper, Michael S. Blouin

Molecular Ecology (2007) 16, 953–966 doi: 10.1111/j.1365-2942.2006.01286.x

Effective population size of steelhead trout: influence of variance in reproductive success, hatchery programs, and genetic compensation between life-history forms

HITOSHI ARAKI,¹ ROBIN S. WAPLES,² WILLIAM R. ARDEN,³ BECKY COOPER¹ and MICHAEL S. BLOUIN⁴

biology letters

Conservation biology

Carry-over effect of captive breeding reduces reproductive fitness of wild-born descendants in the wild

Hitoshi Araki¹, Becky Cooper and Michael S. Blouin

Transactions of the American Fisheries Society

Publication details, including instructions for authors and subscription information: <http://www.tandfonline.com/loi/taf2>

Diminished Reproductive Success of Steelhead from a Hatchery Supplementation Program (Little Sheep Creek, Imnaha Basin, Oregon)

Erwan A. Bertrson¹, Richard W. Carmichael², Michael W. Fleisher³, Eric J. Ward⁴ & Paul Moran⁴

PNAS

Genetic adaptation to captivity can occur in a single generation

Mark R. Christie¹, Melanie L. Marino¹, Rod A. French², and Michael S. Blouin¹

Chinook

North American Journal of Fisheries Management 20: 1472–1485, 2000
© Copyright by the American Fisheries Society 2000
DOI: 10.1577/M00-105.1

[Article]

Use of Parentage Analysis to Determine Reproductive Success of Hatchery-Origin Spring Chinook Salmon Outplanted into Shilike Creek, Oregon

JASON BAUMSTEGER¹

DAVID M. HANCO² AND DOUGLAS E. OLSON³

ROBERT SPATEROLY² AND GHOFF FITZGERALD³

WILLIAM R. ARDEN⁴

MOLECULAR ECOLOGY

Molecular Ecology (2012) 21, 5236–5250 doi: 10.1111/j.1365-2942.2012.02684.x

Factors influencing the relative fitness of hatchery and wild spring Chinook salmon (*Oncorhynchus tshawytscha*) in the Wenatchee River, Washington, USA

Kevin S. Williamson, Andrew R. Murdoch, Todd N. Pearsons, Eric J. Ward, and Michael J. Ford

MOLECULAR ECOLOGY

Molecular Ecology (2012) 21, 5236–5250 doi: 10.1111/j.1365-2942.2012.02684.x

Supportive breeding boosts natural population abundance with minimal negative impacts on fitness of a wild population of Chinook salmon

MAUREEN A. HESS,¹ CRAIG D. RAB,¹ JASON L. VOGEL,¹ JEFF J. STEPHENSON,² DOUG D. NELSON¹ and SHAWN R. NARUM³

Evolutionary Applications

Evolutionary Applications (2011) 4, 1172–1181 doi: 10.1111/j.1520-4101.2011.01720.x

Reproductive success of captive bred and naturally spawned Chinook salmon colonizing newly accessible habitat

Joseph H. Anderson,^{1,2*} Paul L. Faulds,³ William L. Atlas⁴ and Thomas P. Quinn¹

Evolutionary Applications

ORIGINAL ARTICLE

Reproductive success of captive bred and naturally spawned Chinook salmon colonizing newly accessible habitat

Joseph H. Anderson,^{1,2*} Paul L. Faulds,³ William L. Atlas⁴ and Thomas P. Quinn¹

Evolutionary Applications

Reproductive success of captive bred and naturally spawned Chinook salmon colonizing newly accessible habitat

Joseph H. Anderson,^{1,2*} Paul L. Faulds,³ William L. Atlas⁴ and Thomas P. Quinn¹

Coho

2243

Changes in run timing and natural smolt production in a naturally spawning coho salmon (*Oncorhynchus kisutch*) population after 60 years of intensive hatchery supplementation

Michael J. Ford, Howard Fuss, Brant Boelts, Eric LaHood, Jeffrey Hard, and Jason Miller

MOLECULAR ECOLOGY

Molecular Ecology (2011) 20, 1860–1869 doi: 10.1111/j.1365-2942.2011.02684.x

Reduced reproductive success of hatchery coho salmon in the wild: insights into most likely mechanisms

VÉRONIQUE THÉRIAULT,* GREGORY R. MOYER,[†] LAURA S. JACKSON,[‡] MICHAEL S. BLOUIN[‡] and MICHAEL A. BANKS*

Supplementation of wild salmonids with captive-bred fish is a common practice for both commercial and conservation purposes. However, evidence for lower fitness of captive-reared fish relative to wild fish has accumulated in recent years, diminishing the apparent effectiveness of supplementation as a management tool. To date, the mechanisms responsible for these fitness declines remain unknown. In this study, we showed with molecular parentage analysis that hatchery coho salmon (*Oncorhynchus kisutch*) had lower reproductive success than wild fish once they reproduced in the wild. This effect was more pronounced in males than in same-aged females. Hatchery spawned fish that were released as unfed fry (age 0), as well as hatchery fish raised for one year in the hatchery (released as smolts, age 1), both experienced lower lifetime reproductive success (RS) than wild fish. However, the subset of hatchery males that returned as 2-year olds (age 2) did not exhibit the same fitness decrease as males that returned as 3-year olds. Thus, we report three lines of evidence pointing to the absence of sexual selection in the hatchery as a contributing mechanism for fitness declines of hatchery fish in the wild: 0) hatchery fish released as unfed fry that survived to adulthood still had low RS relative to wild fish, 0) age-1 male hatchery fish consistently showed a lower relative RS than female hatchery fish (suggesting a role for sexual selection), and 0) age-2 males, which use a sneaker mating strategy, did not show the same decline as 3-year olds, which compete differently for females (again, implicating sexual selection).

Keywords: captive breeding, parentage analysis, reproductive success, salmonids, sexual selection, supplementation

Received 20 January 2010; revision received 14 January 2011; accepted 18 January 2011

Chum

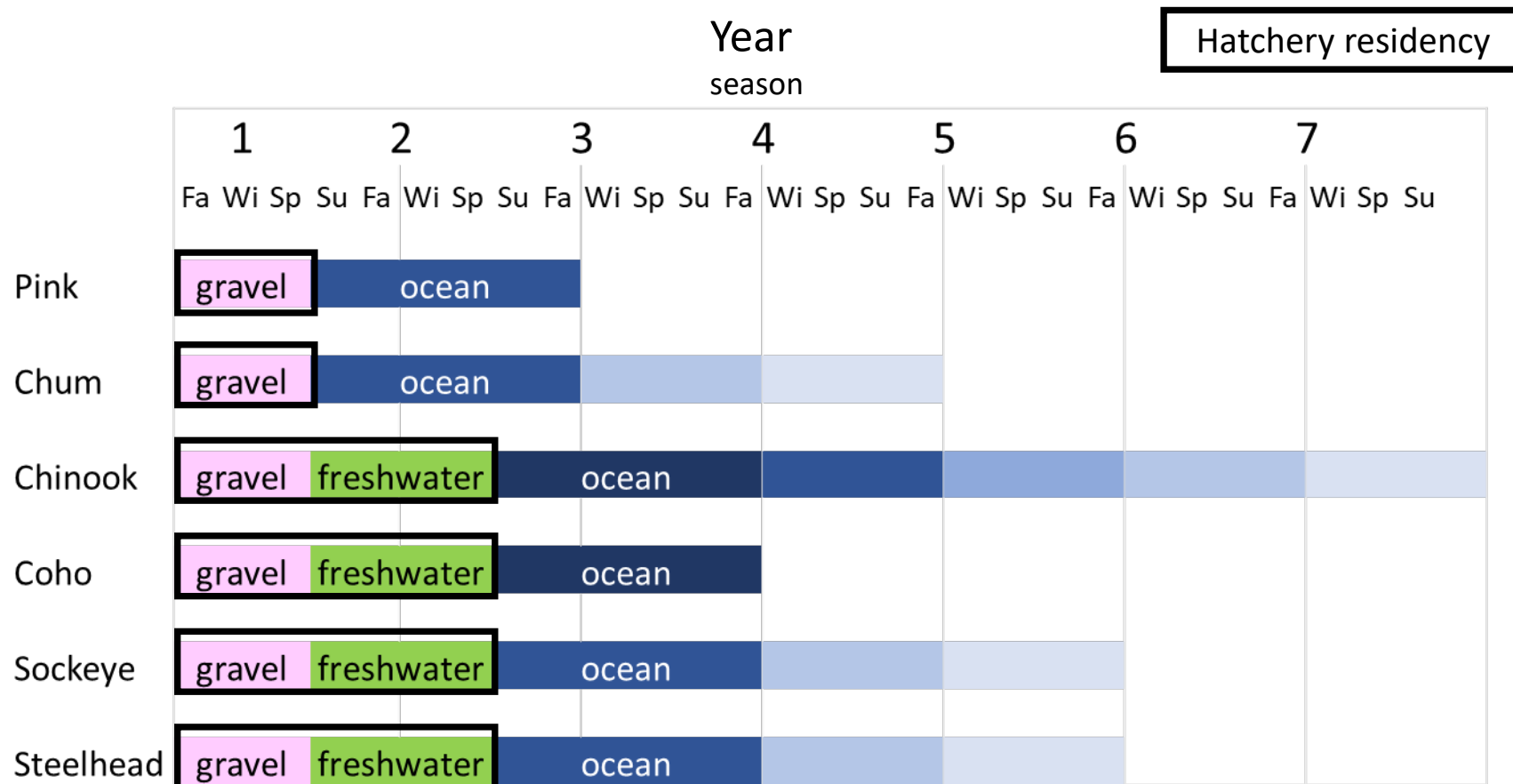
701

Reproductive behavior and relative reproductive success of natural- and hatchery-origin Hood Canal summer chum salmon (*Oncorhynchus keta*)

Barry A. Berejikian, Donald M. Van Doornik, Julie A. Scheurer, and Richard Bush

Abstract: Estimates of the relative fitness of hatchery- and natural-origin salmon can help determine the value of hatchery stocks in contributing to recovery efforts. This study compared the adult life reproductive success of natural-origin summer chum salmon (*Oncorhynchus keta*) with that of first- to third-generation hatchery-origin salmon in an experiment that included four replicate breeding groups. Hatchery- and natural-origin chum salmon exhibited similar reproductive success. Hatchery- and natural-origin males obtained similar access to mating females, and females of both types exhibited similar breeding behaviors and duration. Male body size was positively correlated with access to mating females and reproductive success. The estimate of relative reproductive success (hatchery:natural = 0.83) in this study were similar to those in other studies of other anadromous salmonids to which the hatchery population was founded from the local natural population and much higher than those in studies that evaluated the lifetime relative reproductive success of medical hatchery populations.

Hatchery/Natural Fitness



AHRP Streams in PWS

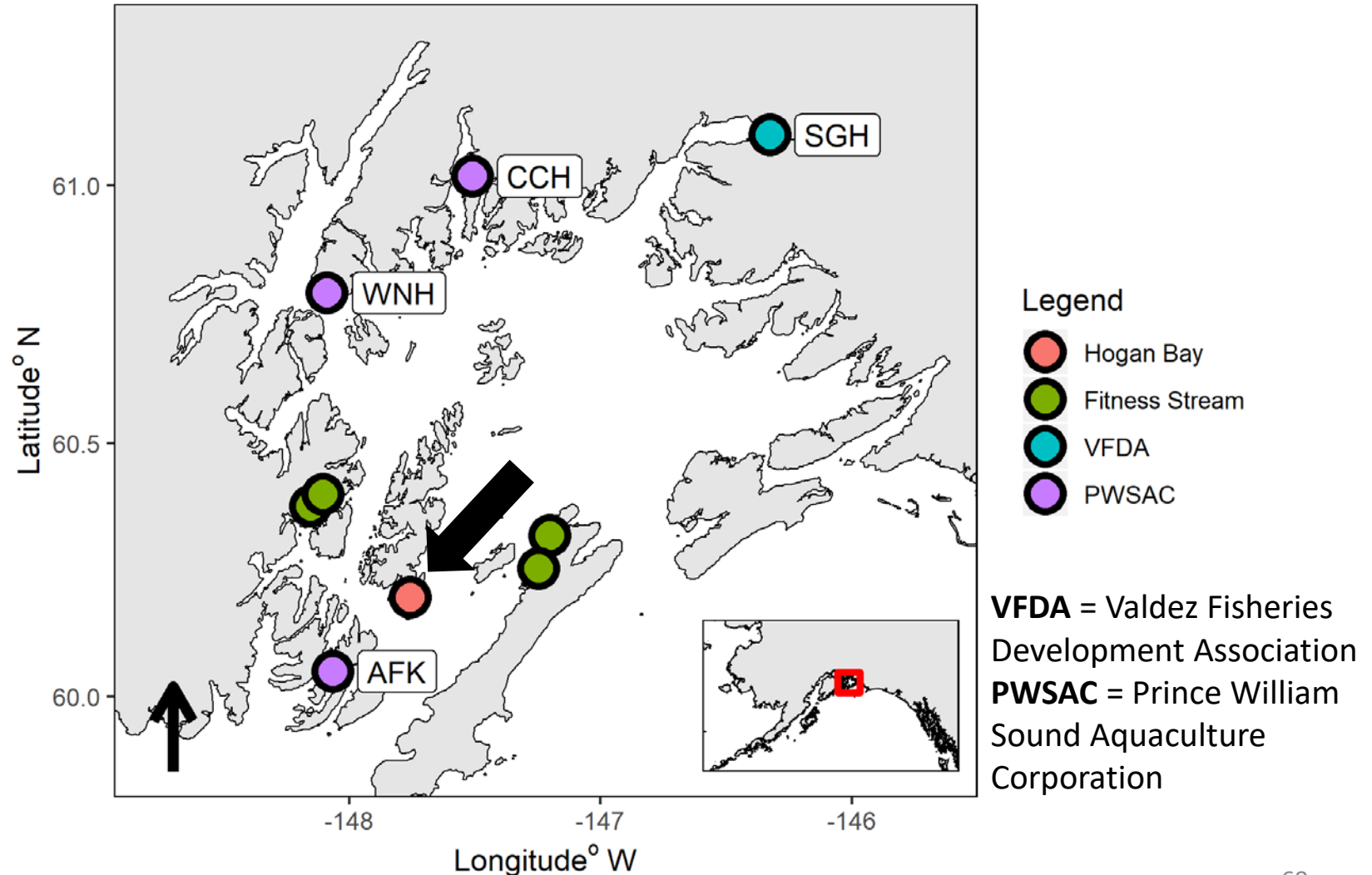


Figure 1 – Lescak et al. *in prep*

Fitness = Reproductive Success

Parent



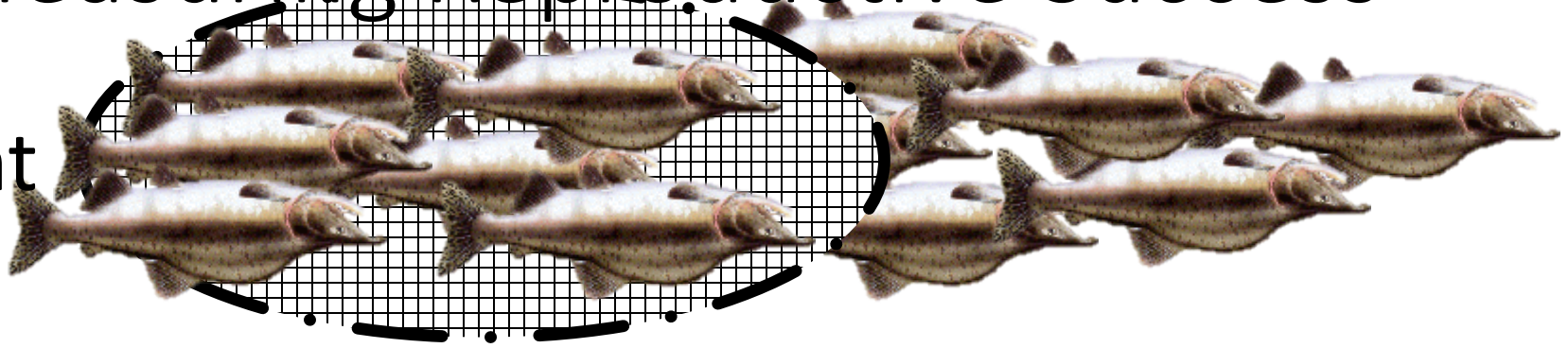
Measuring Reproductive Success

Parent



Measuring Reproductive Success

Parent



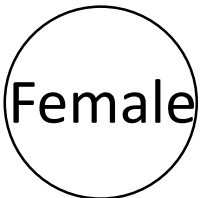
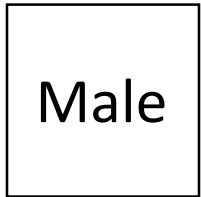
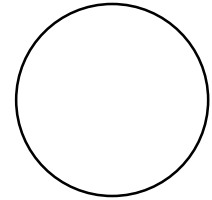
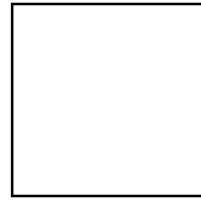
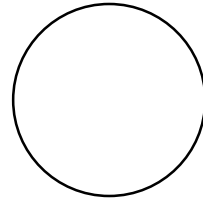
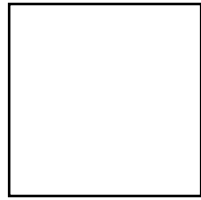
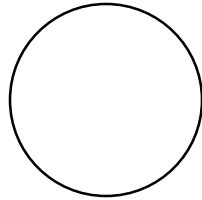
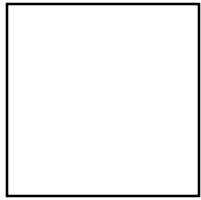
Measuring Reproductive Success

Parent



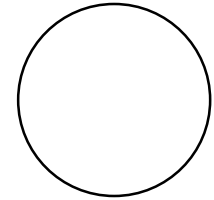
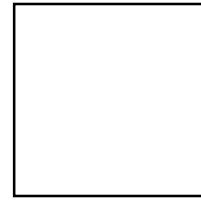
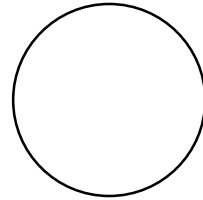
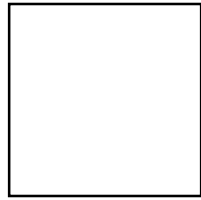
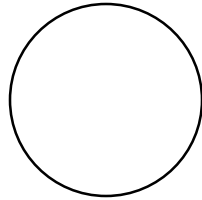
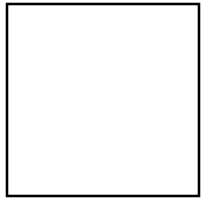
Measuring Reproductive Success

P



Measuring Reproductive Success

P

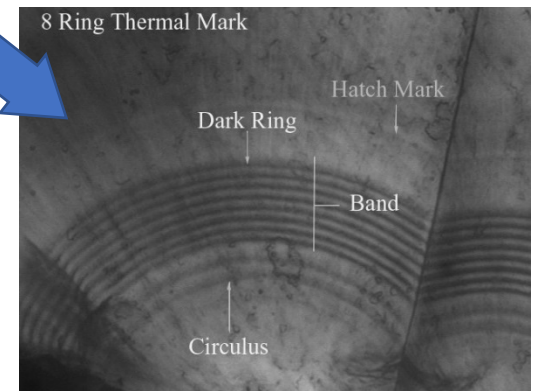
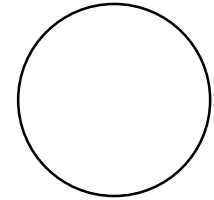
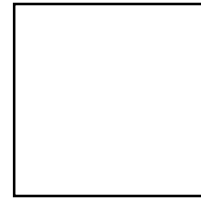
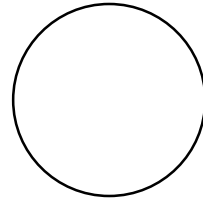
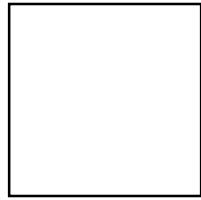
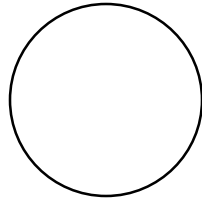
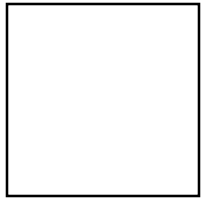


Male

Female

Measuring Reproductive Success

P



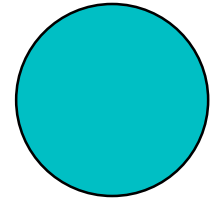
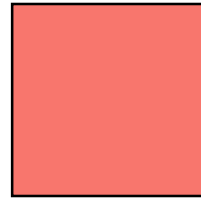
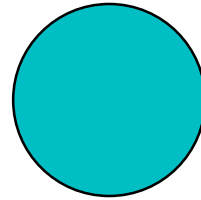
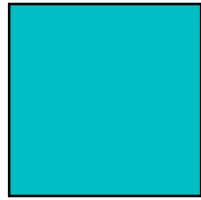
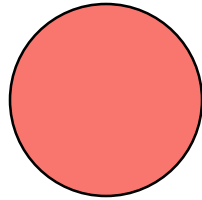
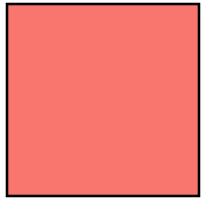
Hatchery-origin

Male

Female

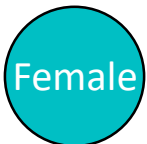
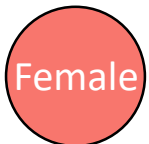
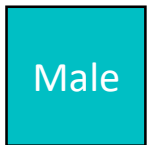
Measuring Reproductive Success

P



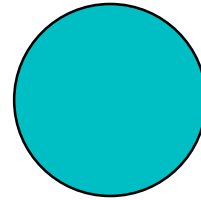
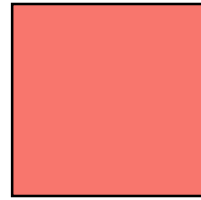
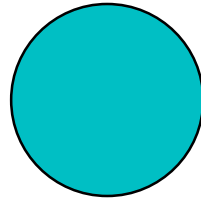
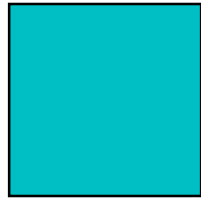
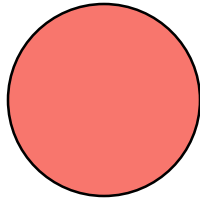
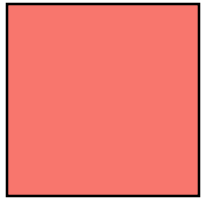
Natural

Hatchery



Measuring Reproductive Success

P

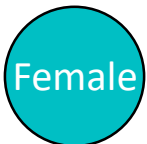
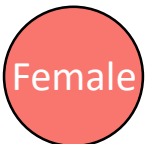
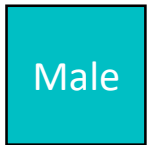


O



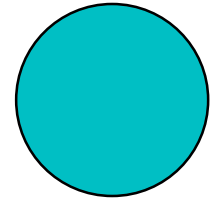
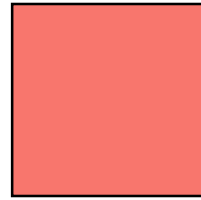
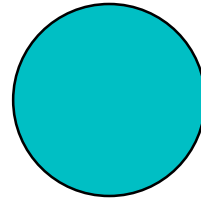
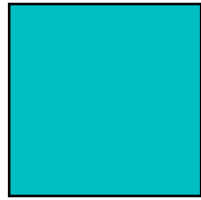
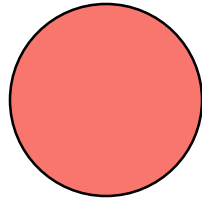
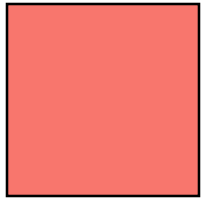
Natural

Hatchery



Measuring Reproductive Success

P

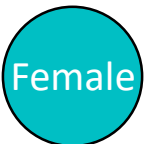
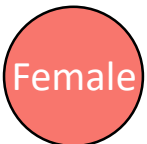


O



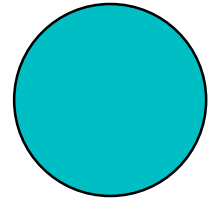
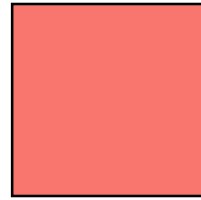
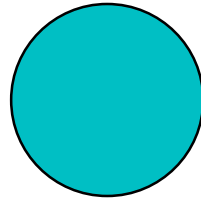
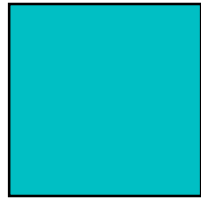
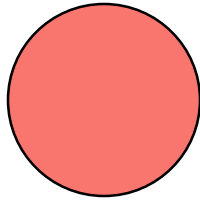
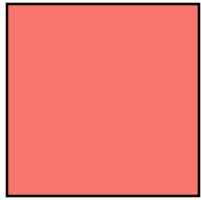
Natural

Hatchery

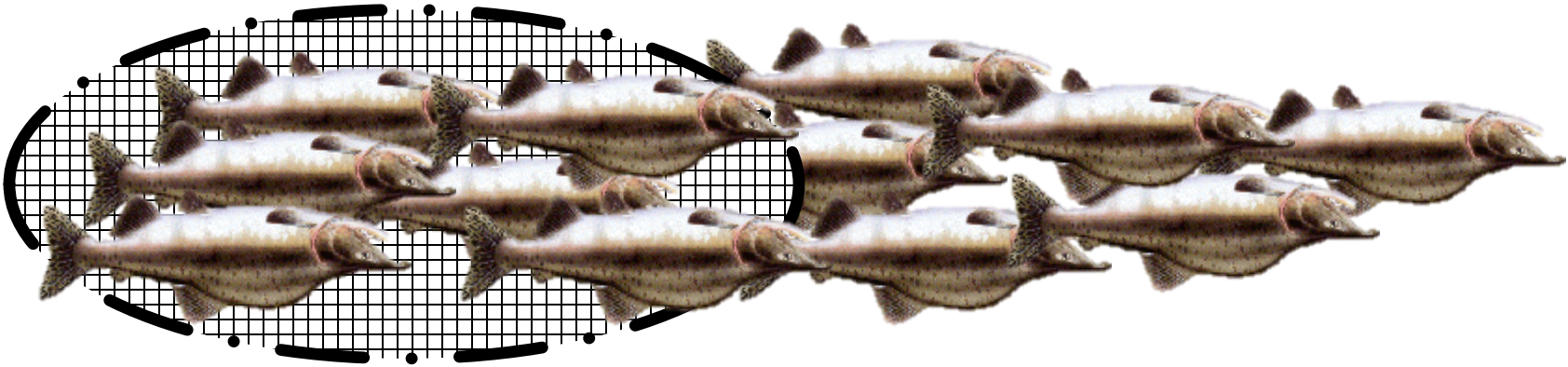


Measuring Reproductive Success

P

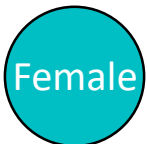
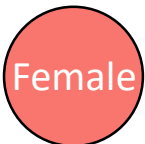
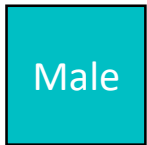


O



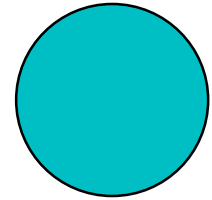
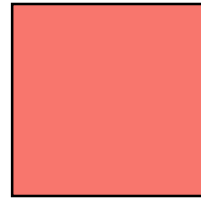
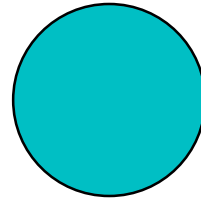
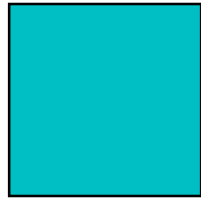
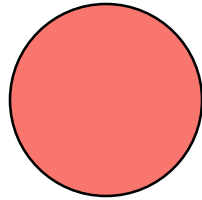
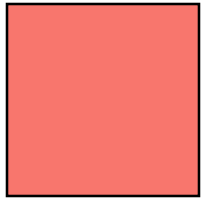
Natural

Hatchery

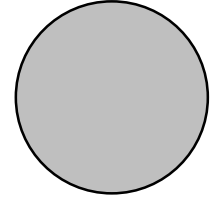
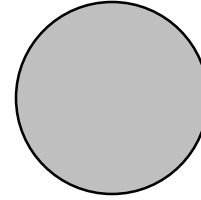
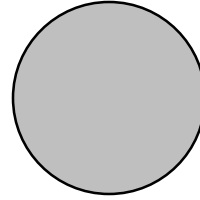
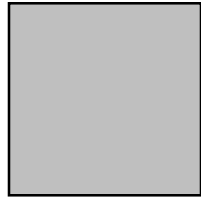
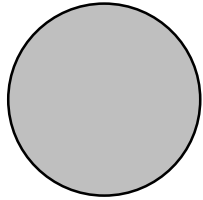


Measuring Reproductive Success

P

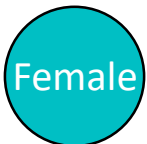
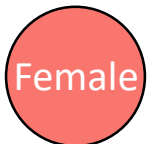
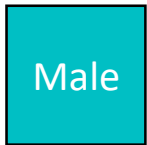


O



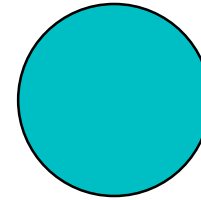
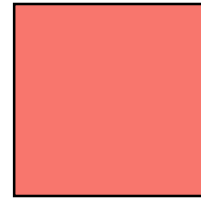
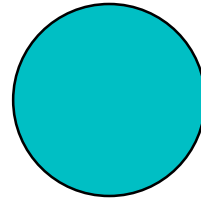
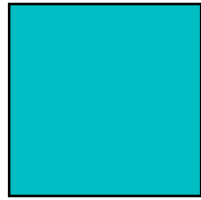
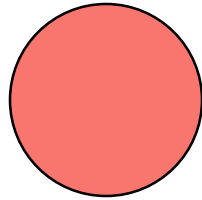
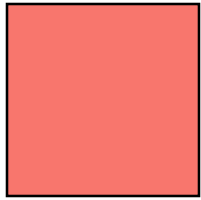
Natural

Hatchery

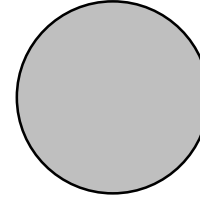
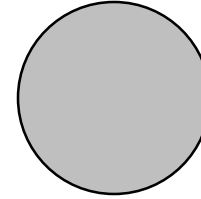
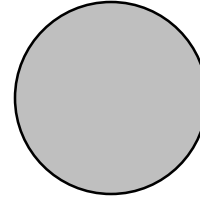
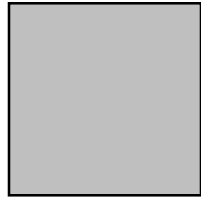
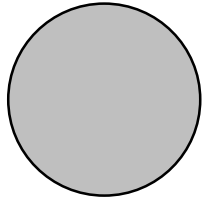


Measuring Reproductive Success

P

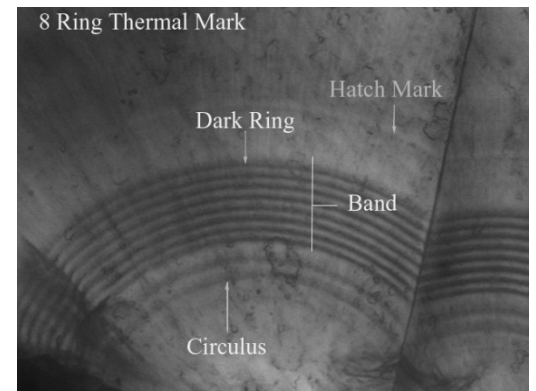
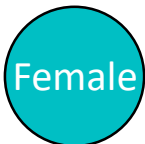
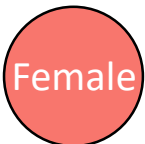
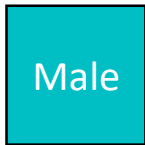


O



Natural

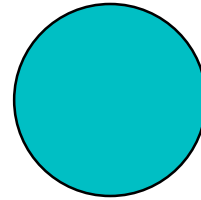
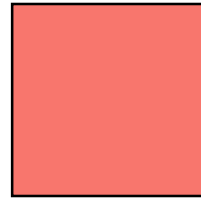
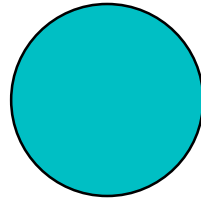
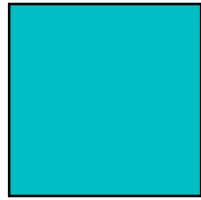
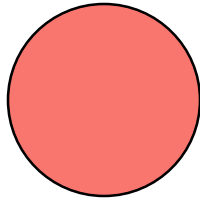
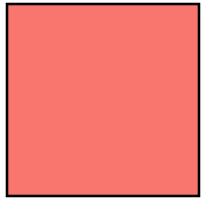
Hatchery



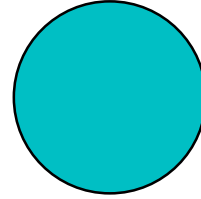
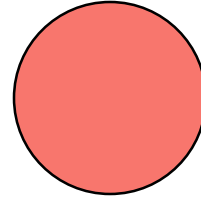
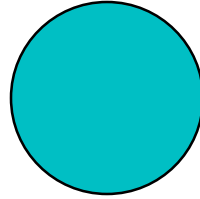
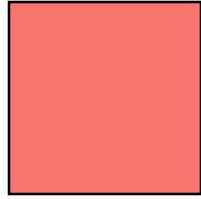
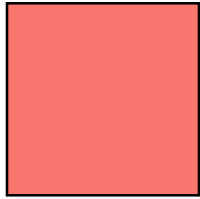
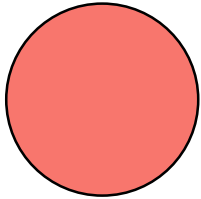
Hatchery-origin

Measuring Reproductive Success

P

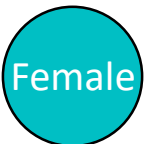
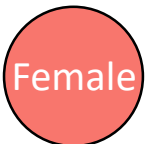
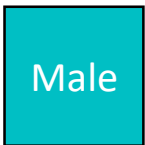


O

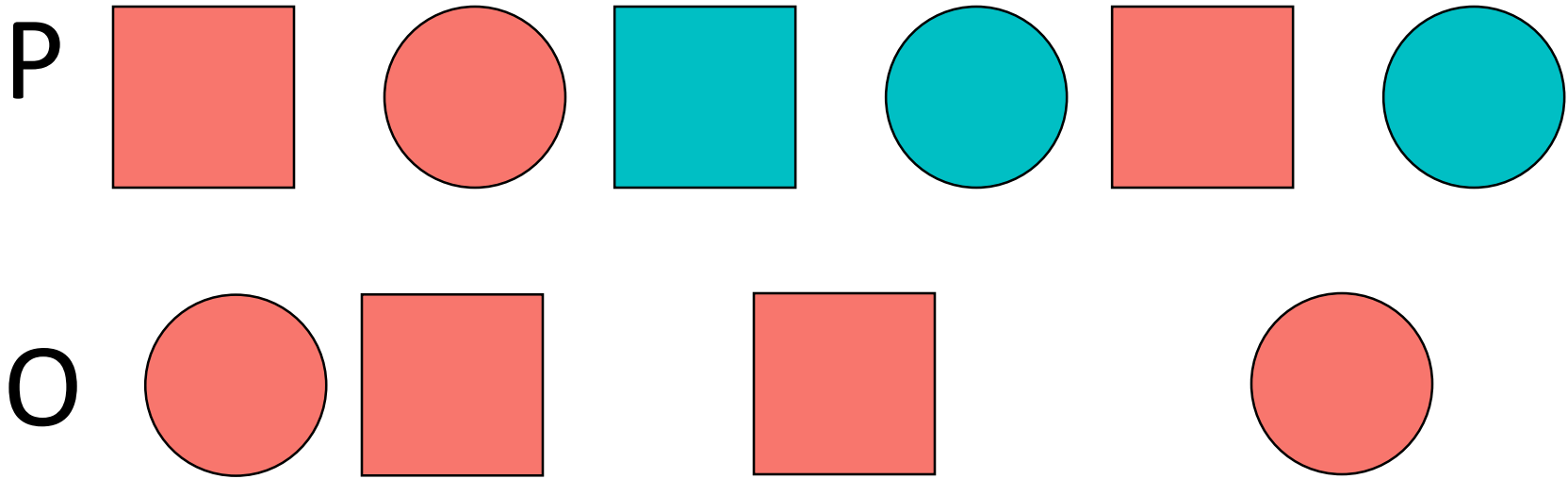


Natural

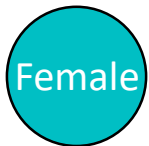
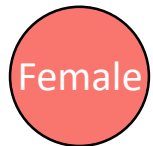
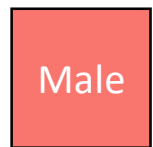
Hatchery



Measuring Reproductive Success



Natural Hatchery



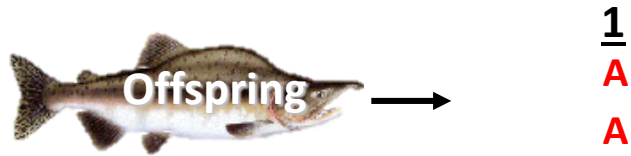
Hatchery-origin fish are not genotyped in the offspring generation because they have a known origin.

Genetic markers for parentage analysis

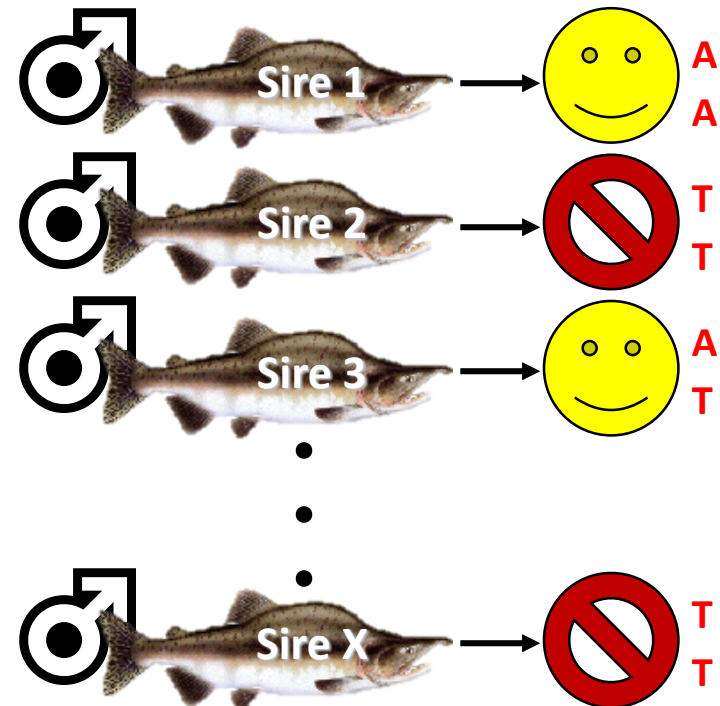
	→	CTATGTA T AAATGTTAATAATAACTAGCTAACC CTATGTA A AAATGTTAATAATAACTAGCTAACC	<i>T allele</i> <i>A allele</i>
	→	CTATGTA A AAATGTTAATAATAACTAGCTAACC CTATGTA A AAATGTTAATAATAACTAGCTAACC	<i>A allele</i> <i>A allele</i>
	→	CTATGTA A AAATGTTAATAATAACTAGCTAACC CTATGTA T AAATGTTAATAATAACTAGCTAACC	<i>A allele</i> <i>T allele</i>
	→	CTATGTA T AAATGTTAATAATAACTAGCTAACC CTATGTA T AAATGTTAATAATAACTAGCTAACC	<i>T allele</i> <i>T allele</i>

Genetic markers for parentage analysis

Markers

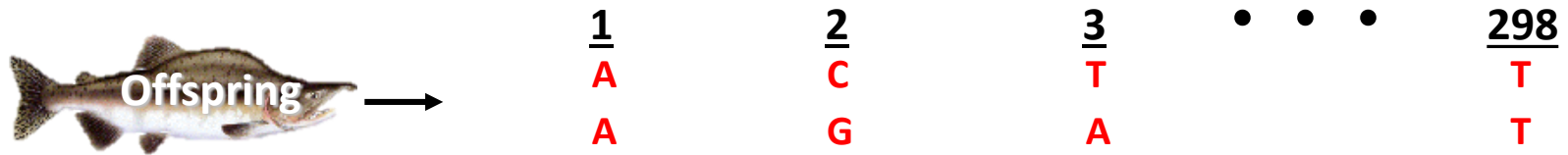


Potential sires (♂)

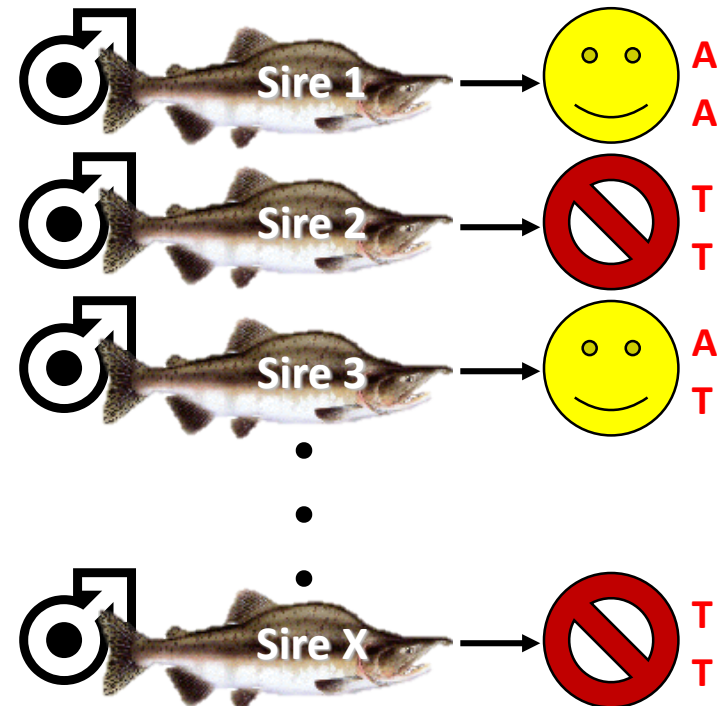


Genetic markers for parentage analysis

Markers

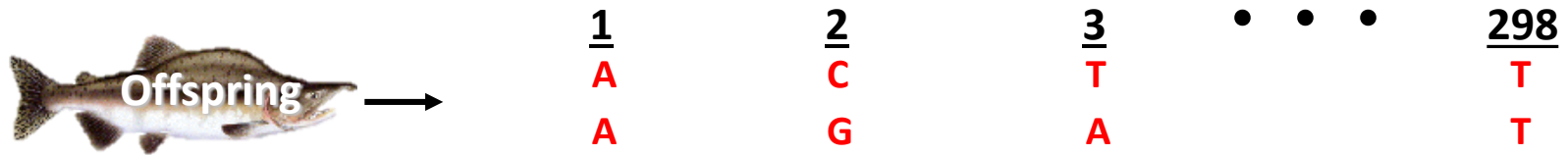


Potential sires (♂)

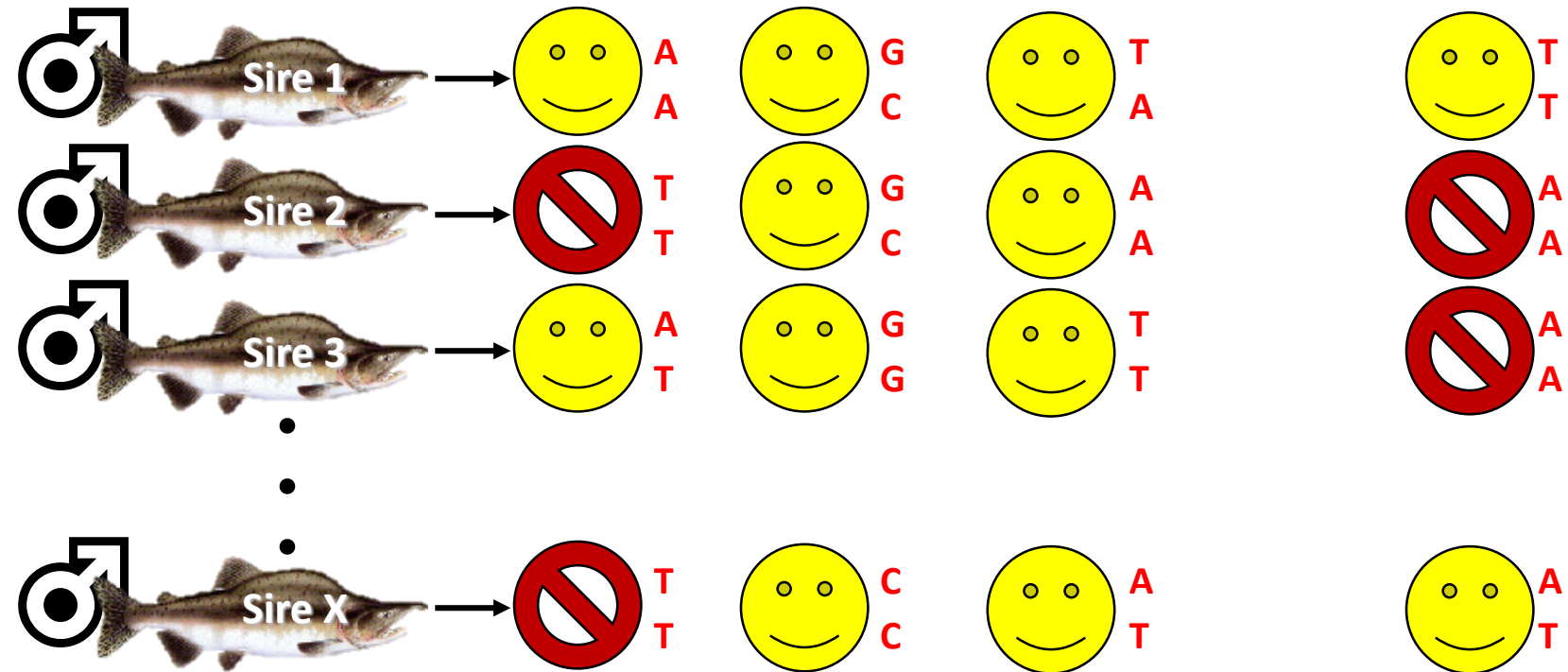


Genetic markers for parentage analysis

Markers

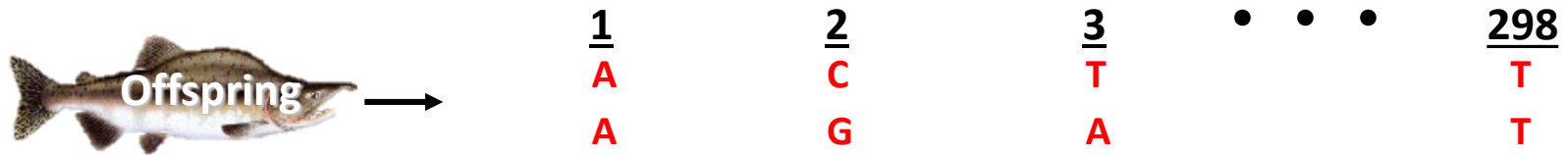


Potential sires (♂)

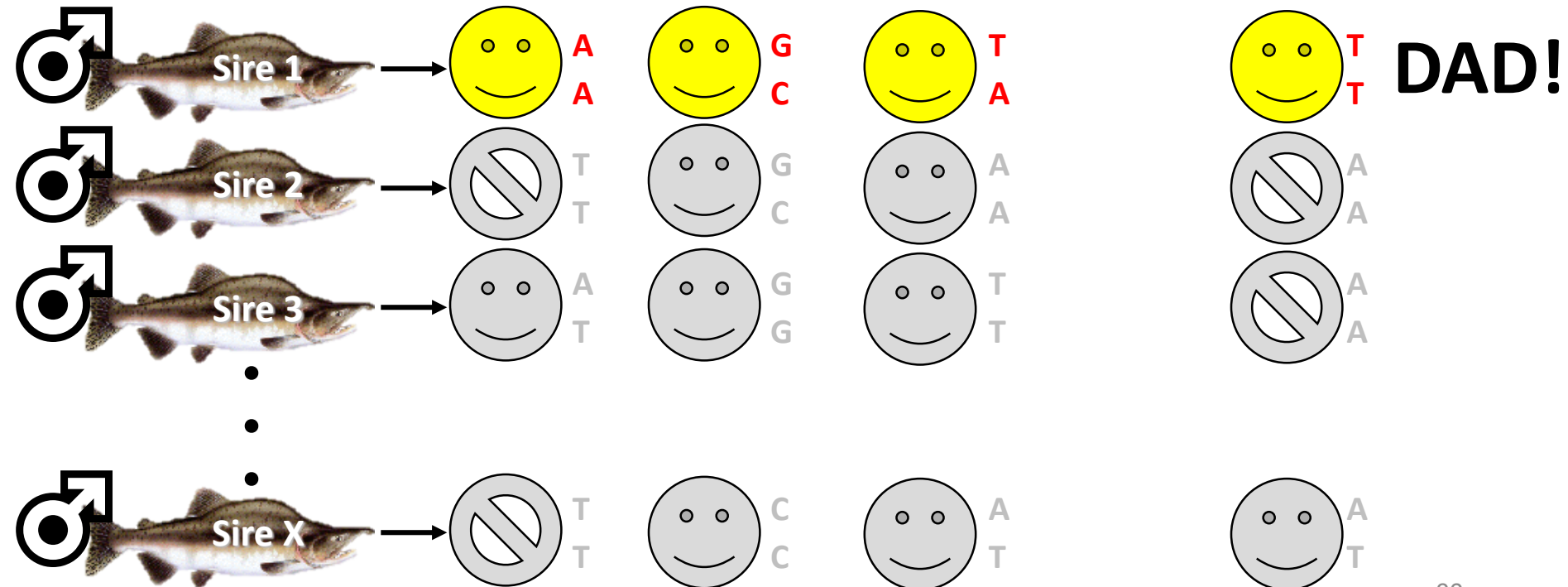


Genetic markers for parentage analysis

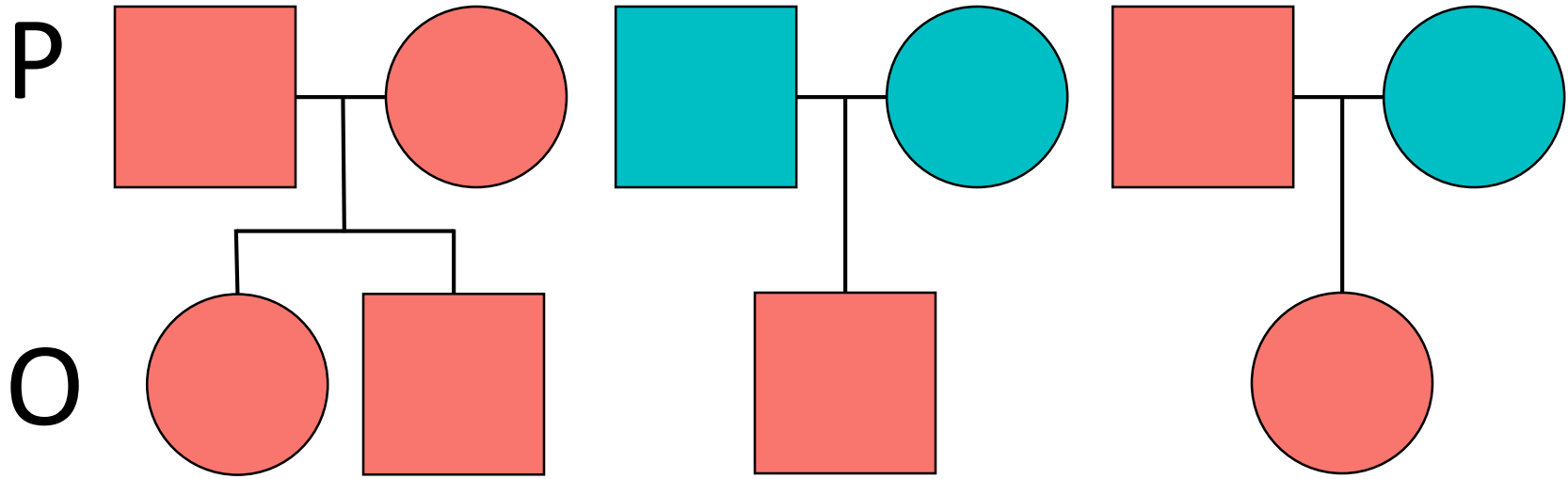
Markers



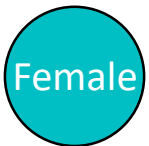
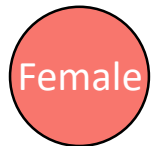
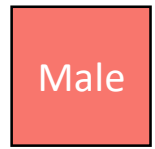
Potential sires (♂)



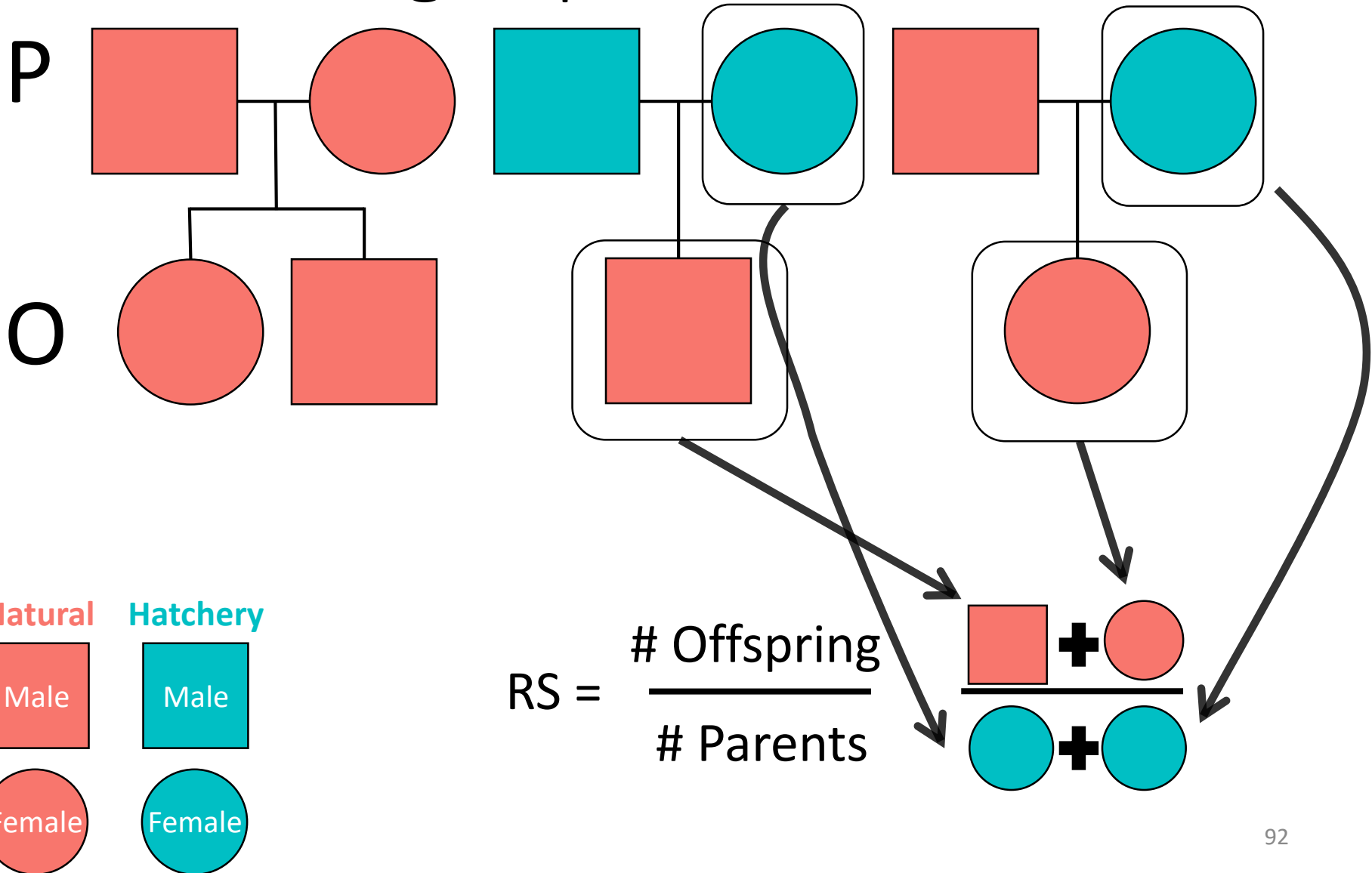
Measuring Reproductive Success



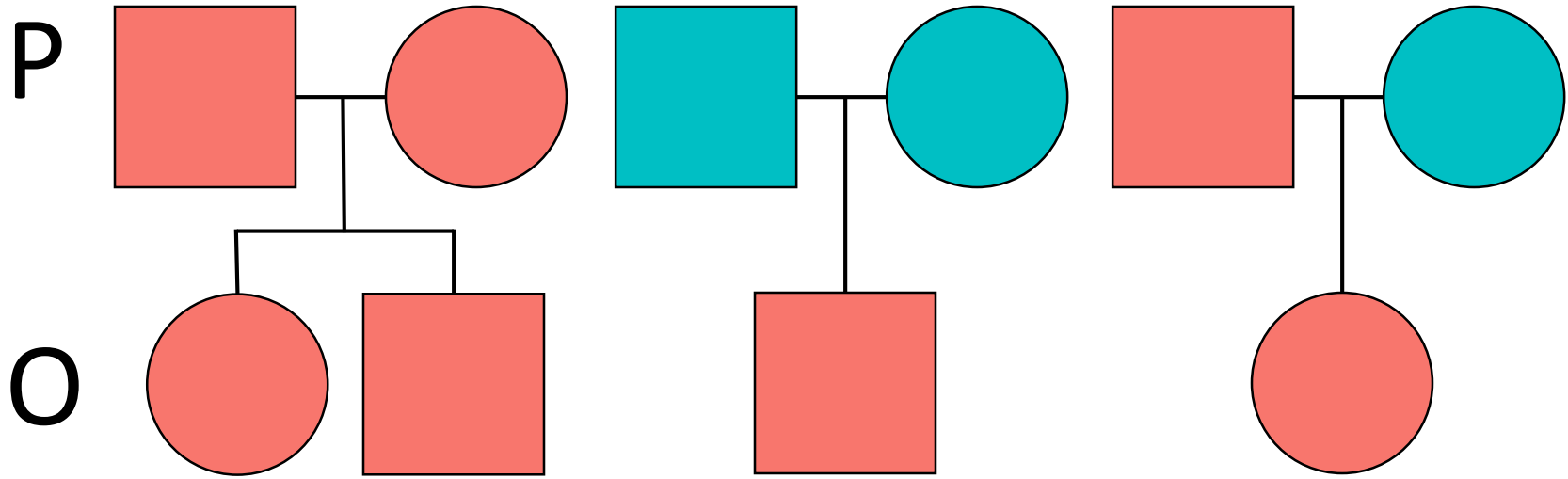
Natural Hatchery



Measuring Reproductive Success

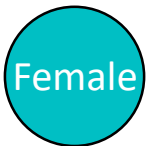
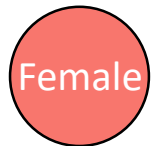


Measuring Reproductive Success

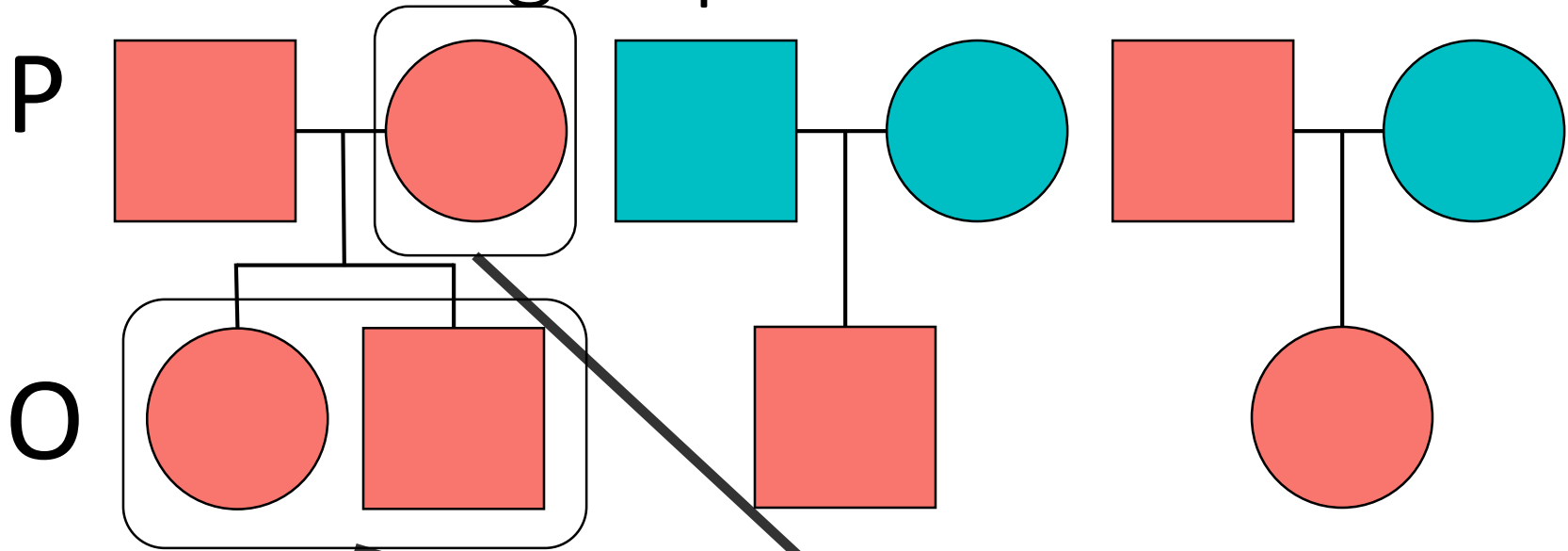


$$RS_{H \text{ Female}} = 1$$

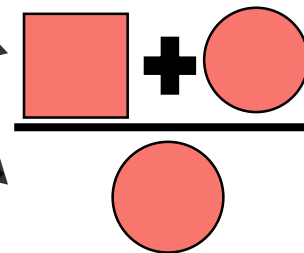
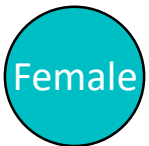
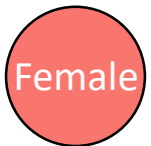
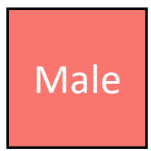
Natural Hatchery



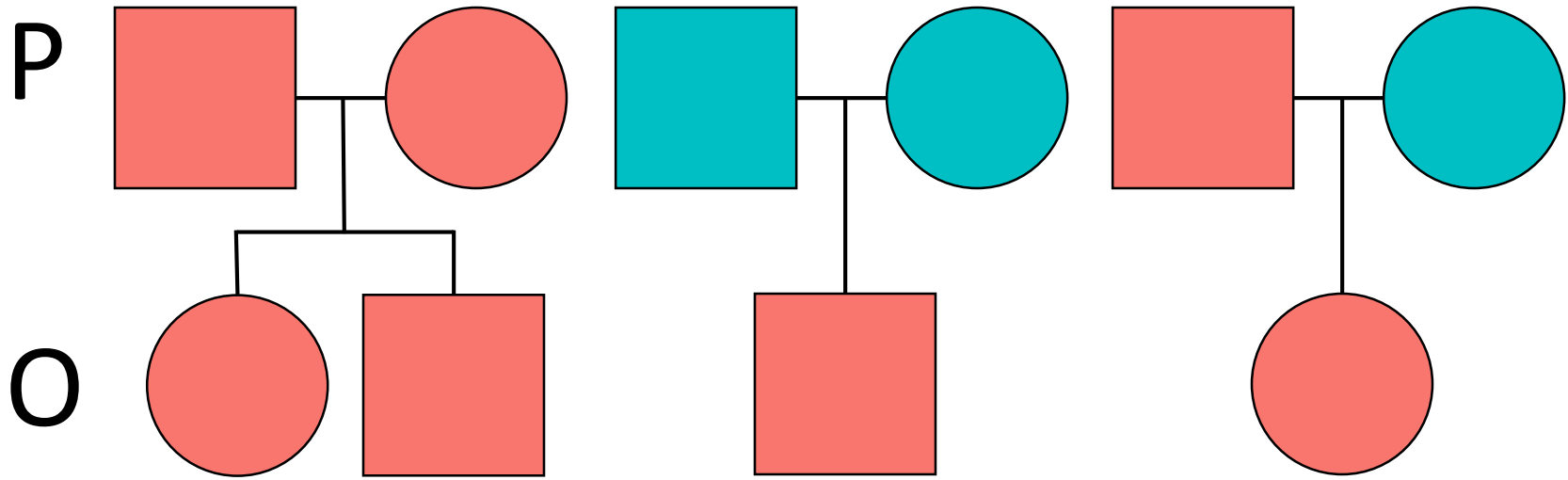
Measuring Reproductive Success



Natural Hatchery



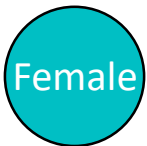
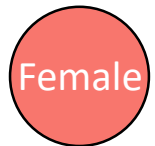
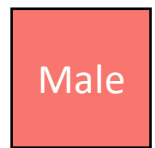
Measuring Reproductive Success



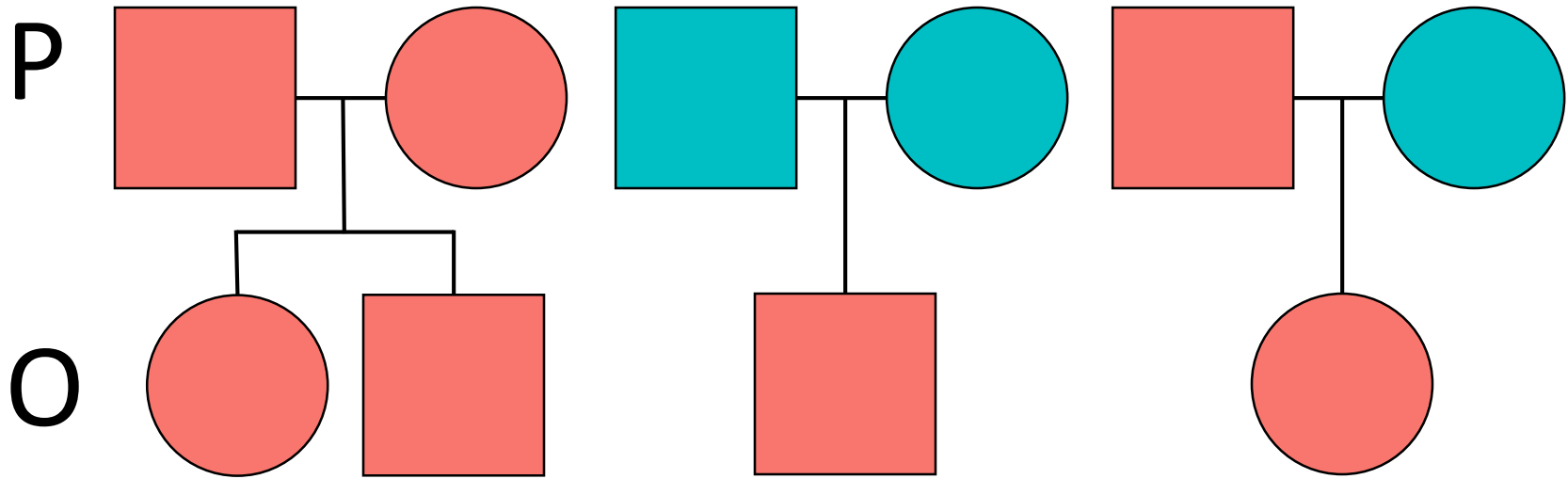
$$RS_{N \text{ Female}} = 2$$

$$RS_{H \text{ Female}} = 1$$

Natural Hatchery



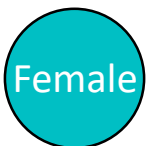
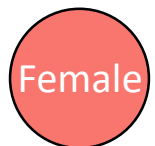
Measuring Reproductive Success



$$RS_{N \text{ Female}} = 2$$

$$RS_{H \text{ Female}} = 1$$

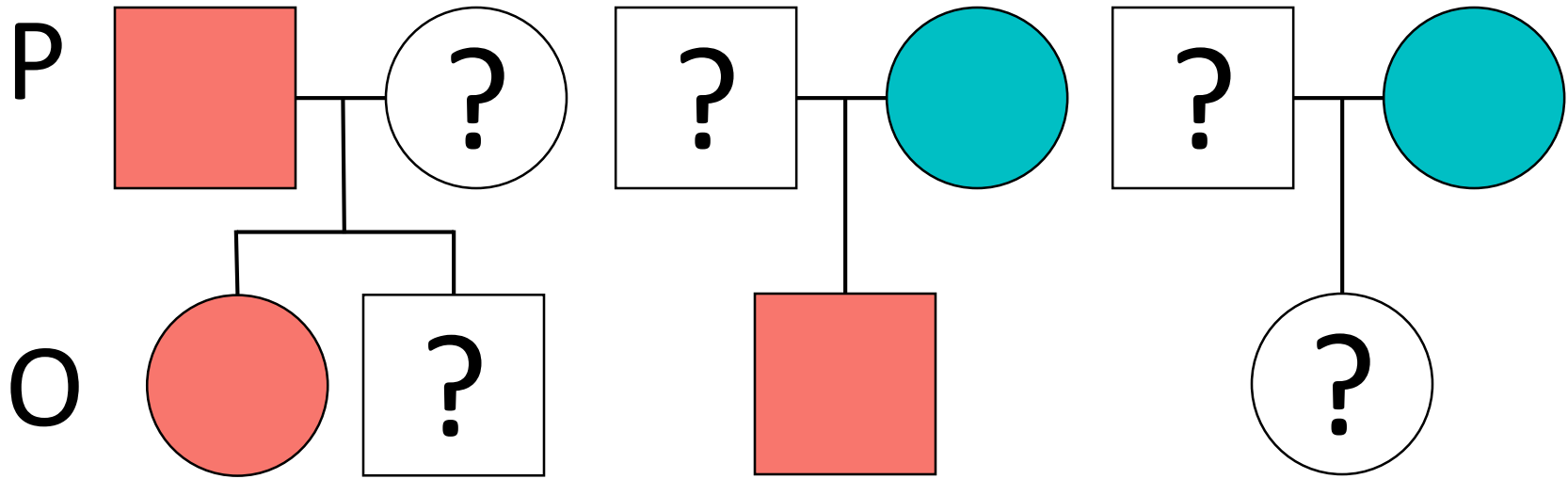
Natural Hatchery



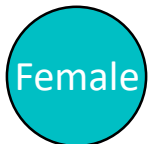
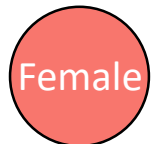
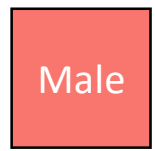
Relative Reproductive Success (RRS)

$$RRS = \frac{1}{2} = 0.5$$

Measuring Reproductive Success



Natural Hatchery



Relative Reproductive Success (RRS)

$$RRS = \frac{\overline{RS}_{\text{Hatchery}}}{\overline{RS}_{\text{Natural}}}$$

Analyzed Samples: Even-Lineage

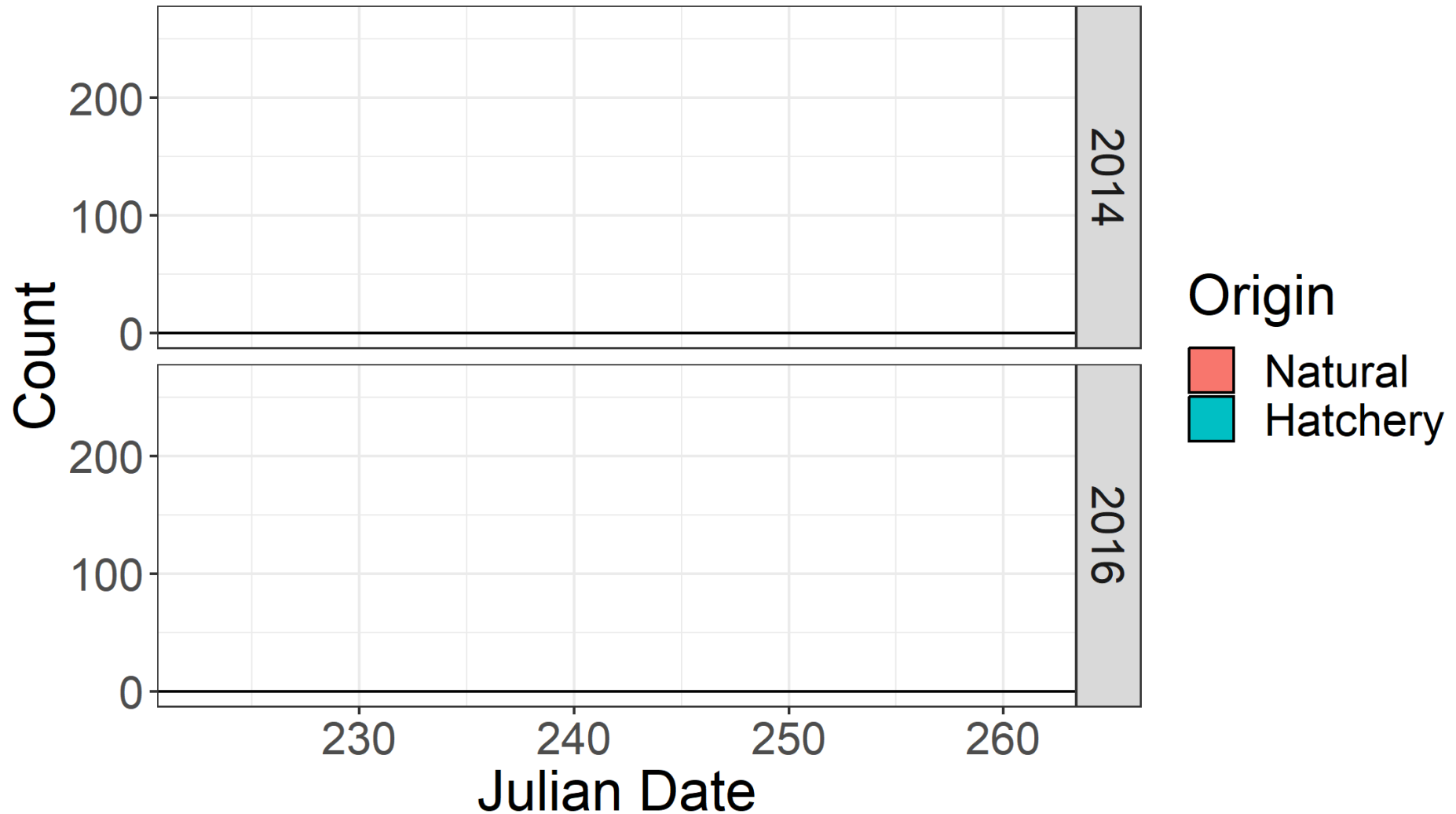


Figure 2b – Lescak et al. *in prep*

Analyzed Samples: Even-Lineage

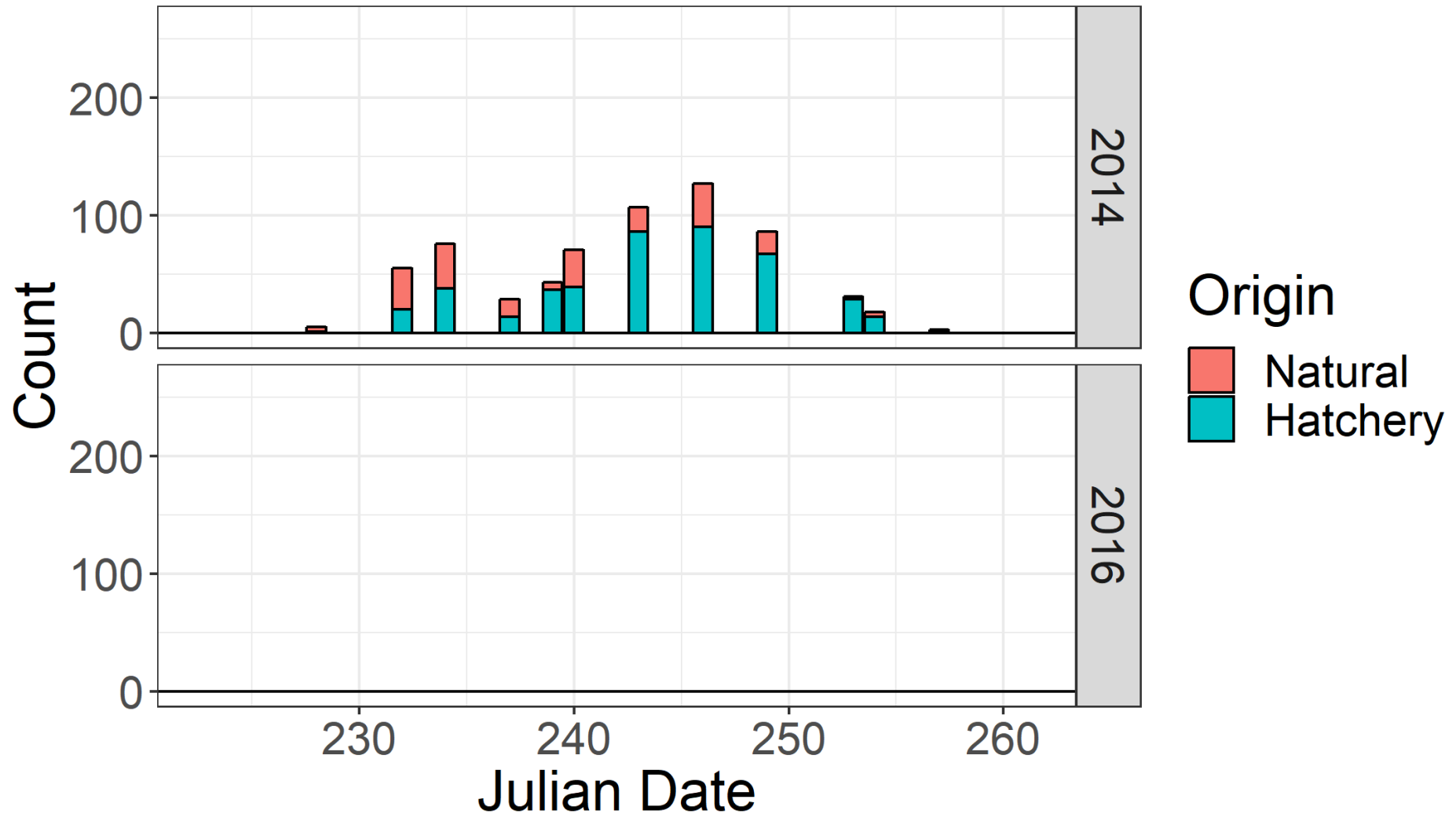


Figure 2b – Lescak et al. *in prep*

Analyzed Samples: Even-Lineage

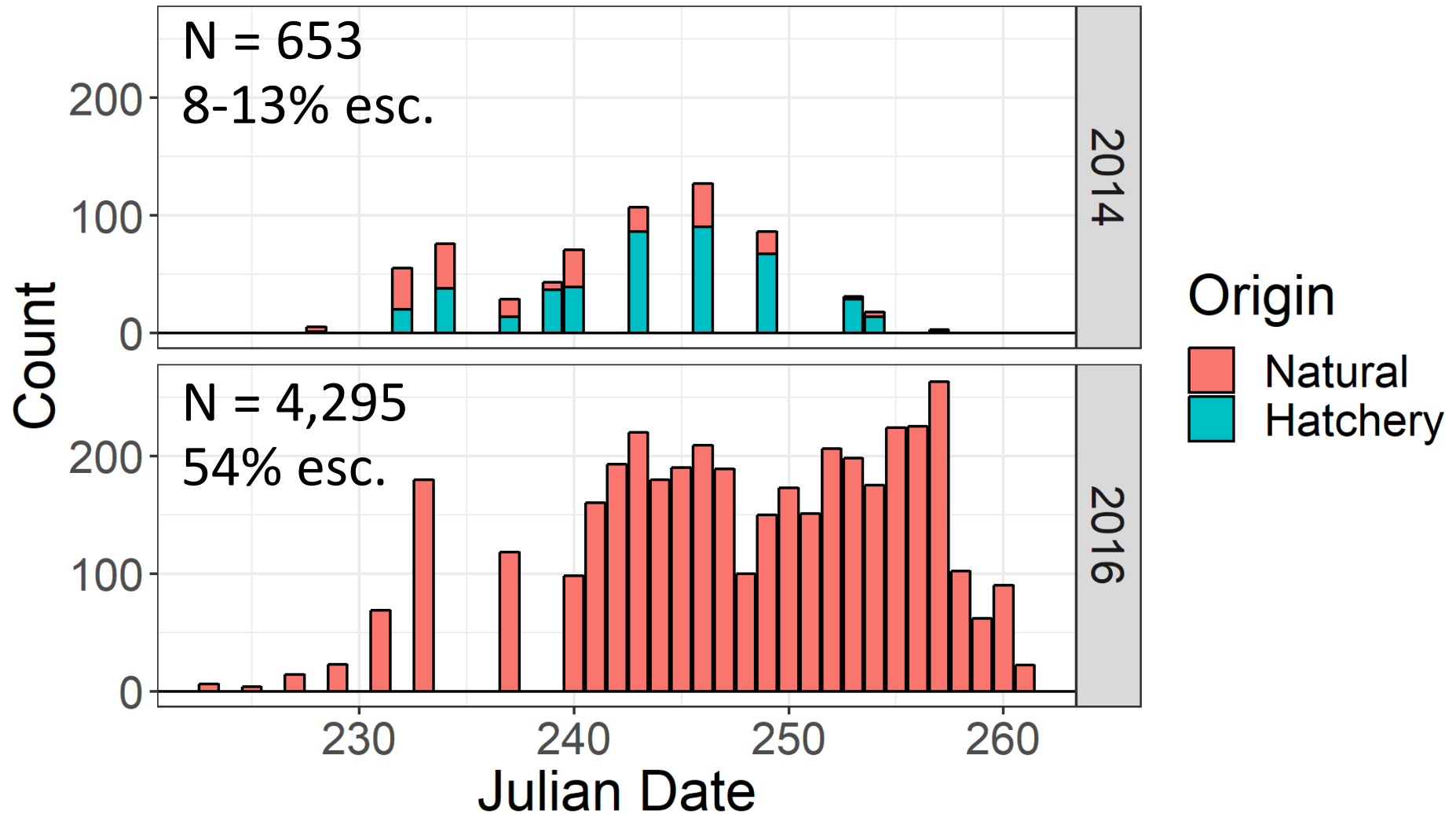


Figure 2b – Lescak et al. *in prep*

Pedigree Results: Even-Lineage

- 451 offspring (11%) assigned to 184 parents
 - 208 → natural-origin parents
 - 265 → hatchery-origin parents
 - 202 – AFI
 - 41 – WNI
 - 22 – CCH
 - 0 – SGH

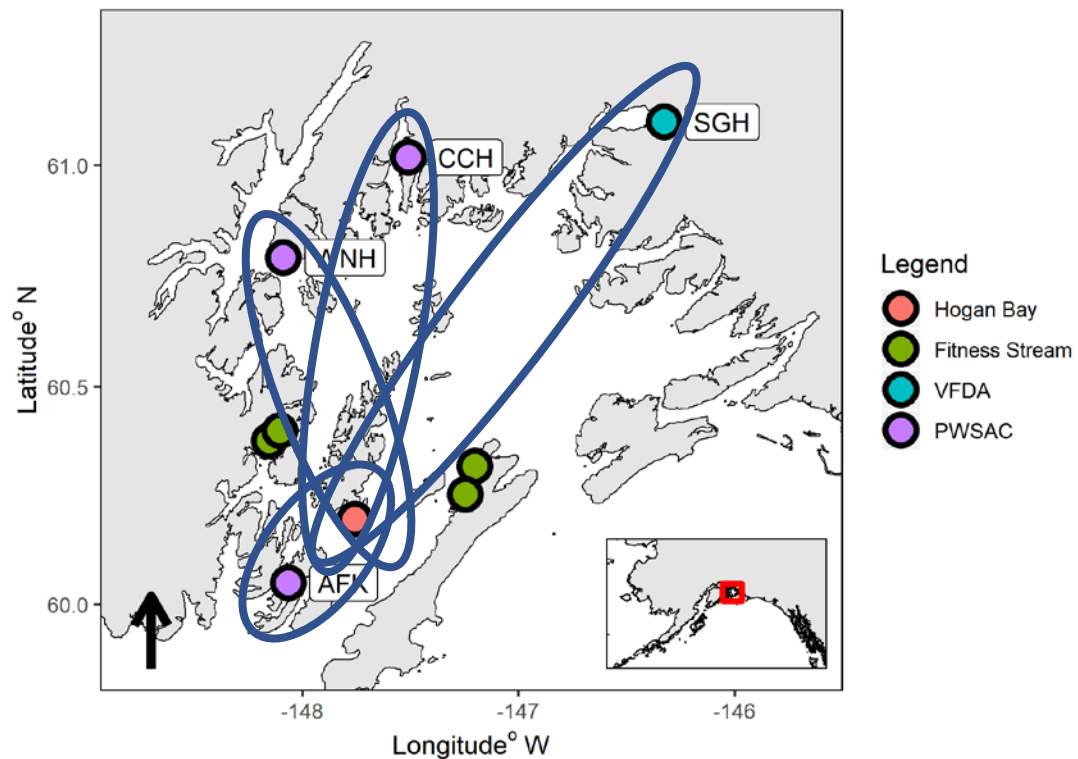


Figure 1 – Lescak et al. *in prep*

RS Distribution: Even-Lineage

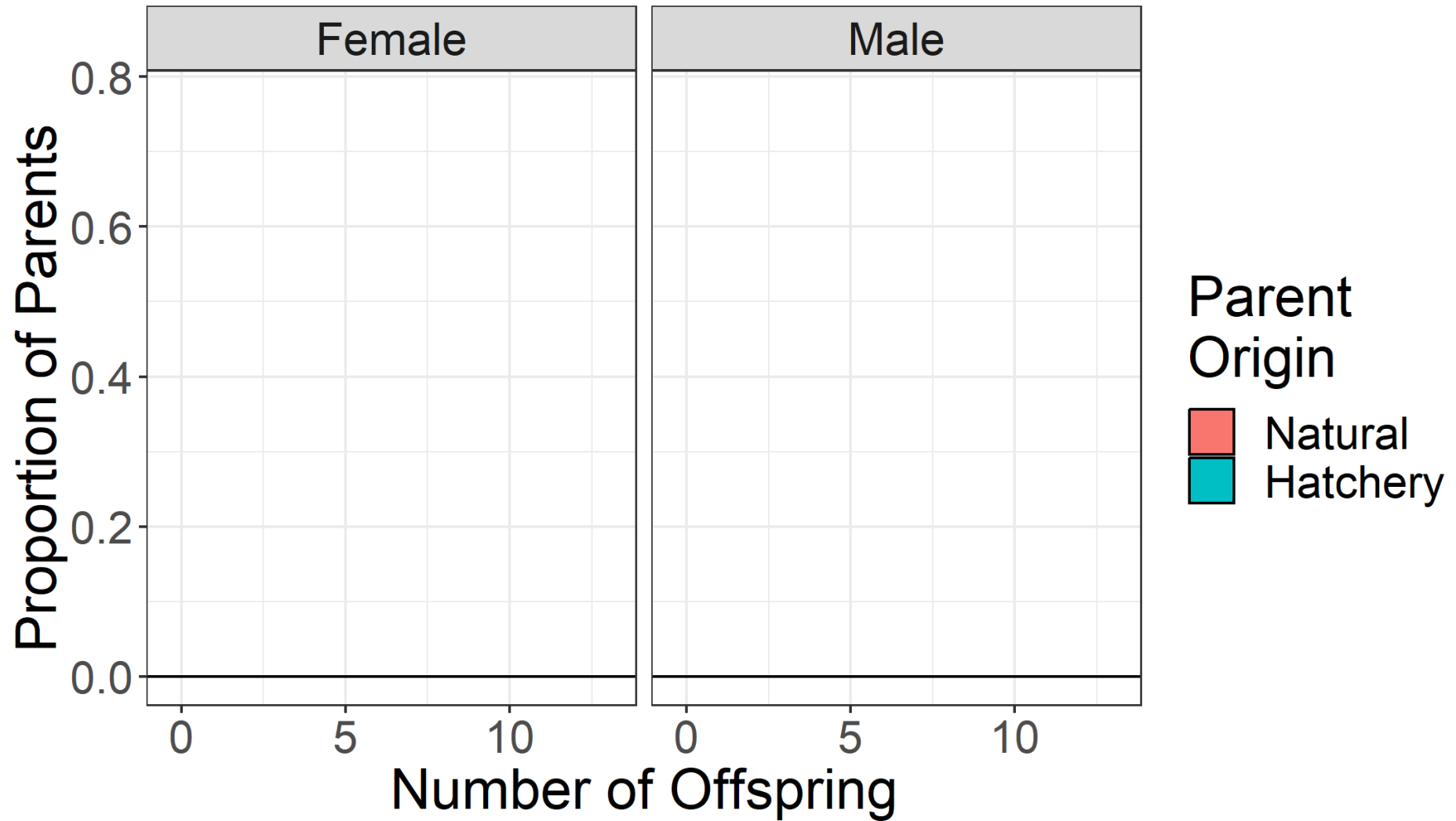


Figure 3b – Lescak et al. *in prep*

RS Distribution: Even-Lineage

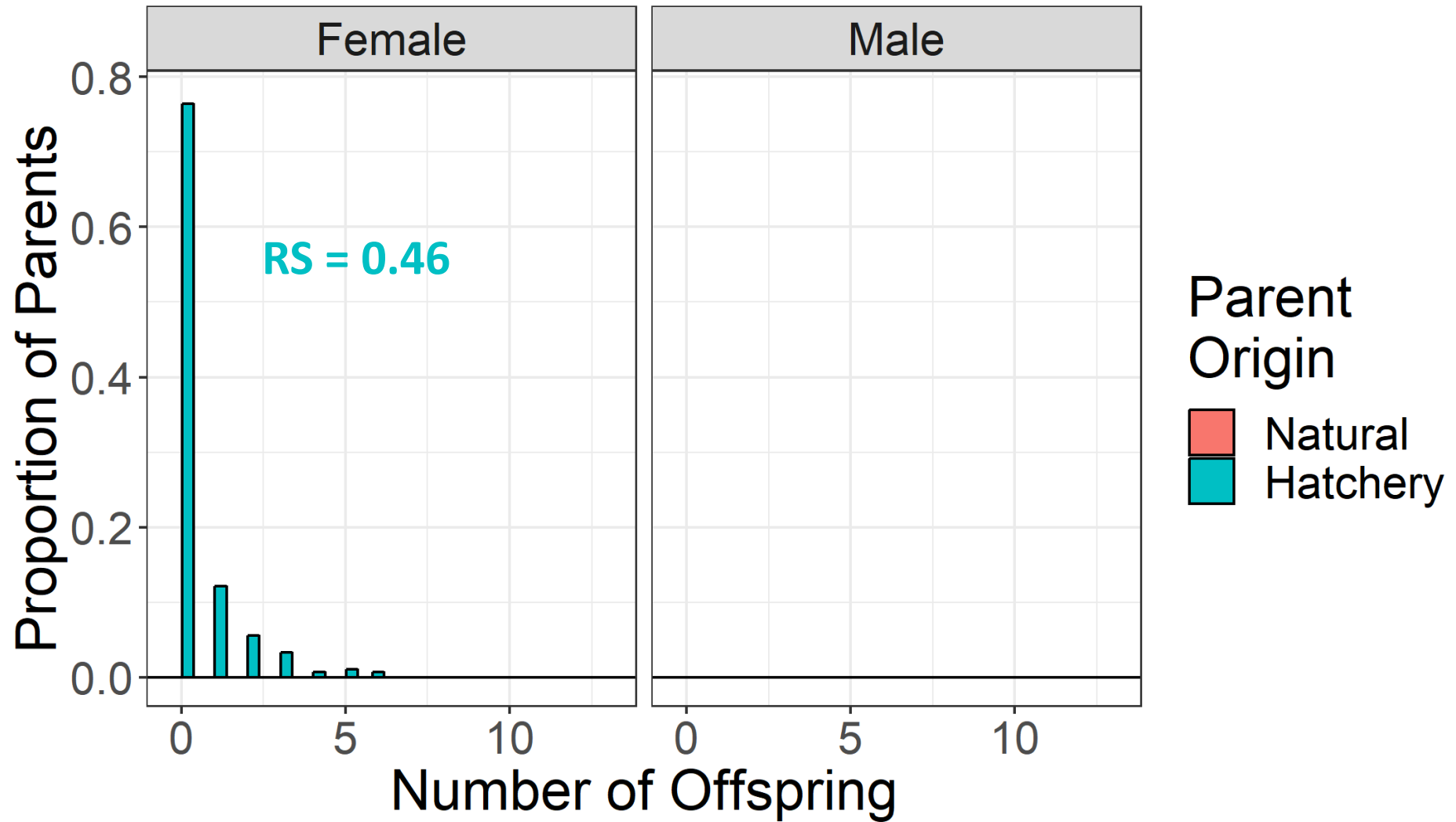


Figure 3b – Lescak et al. *in prep*

RS Distribution: Even-Lineage

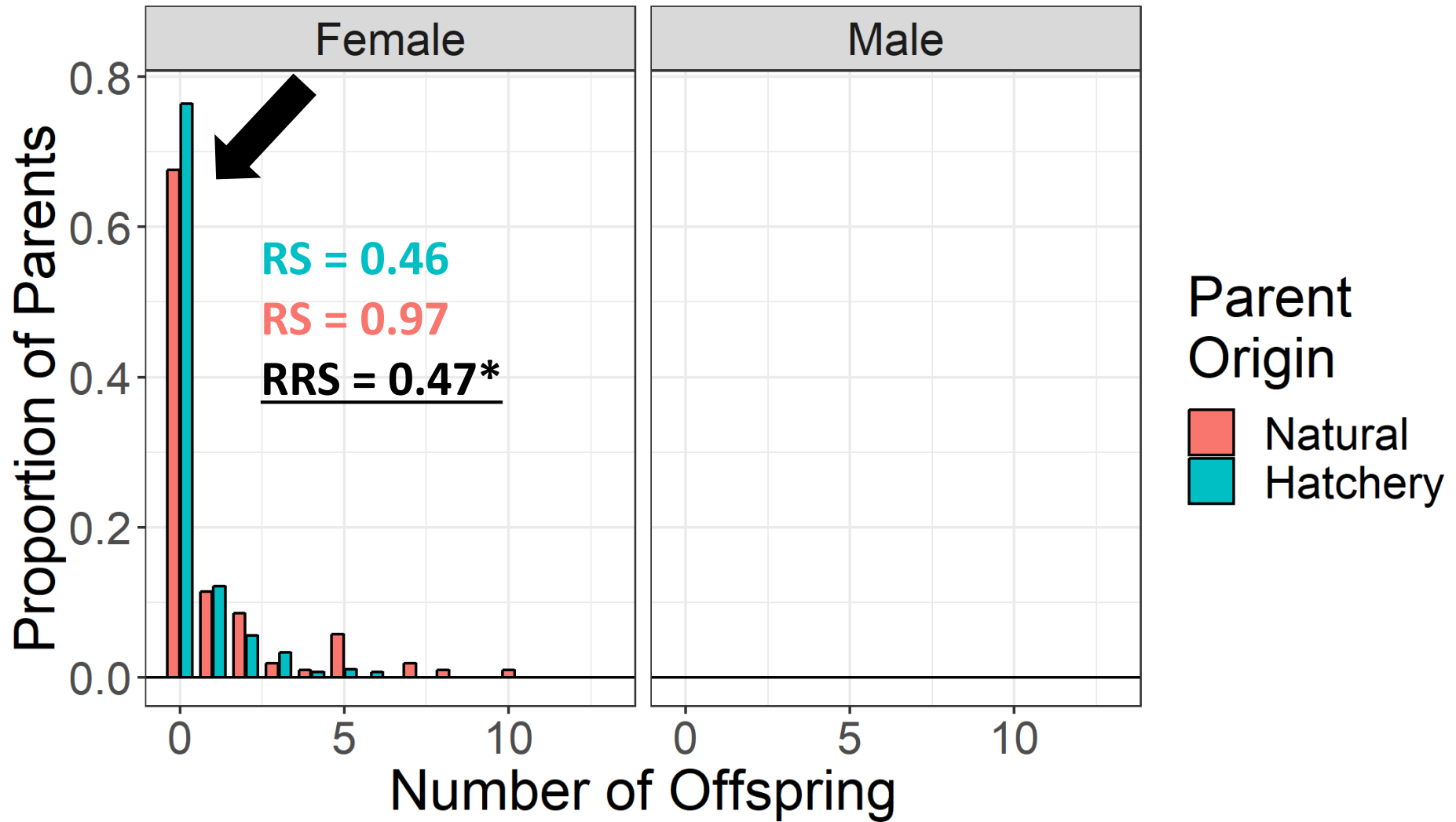


Figure 3b – Lescak et al. *in prep*

RS Distribution: Even-Lineage

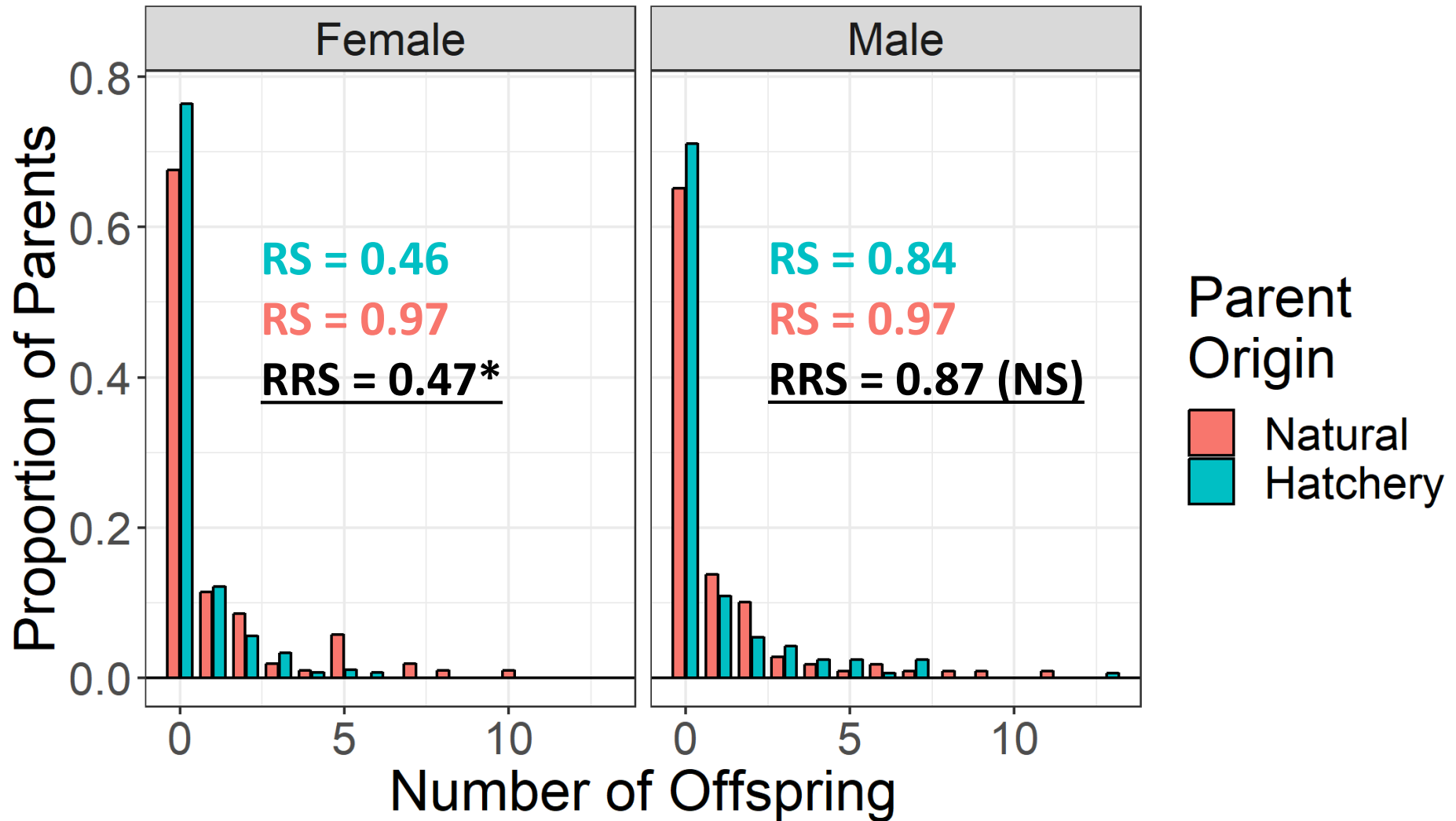
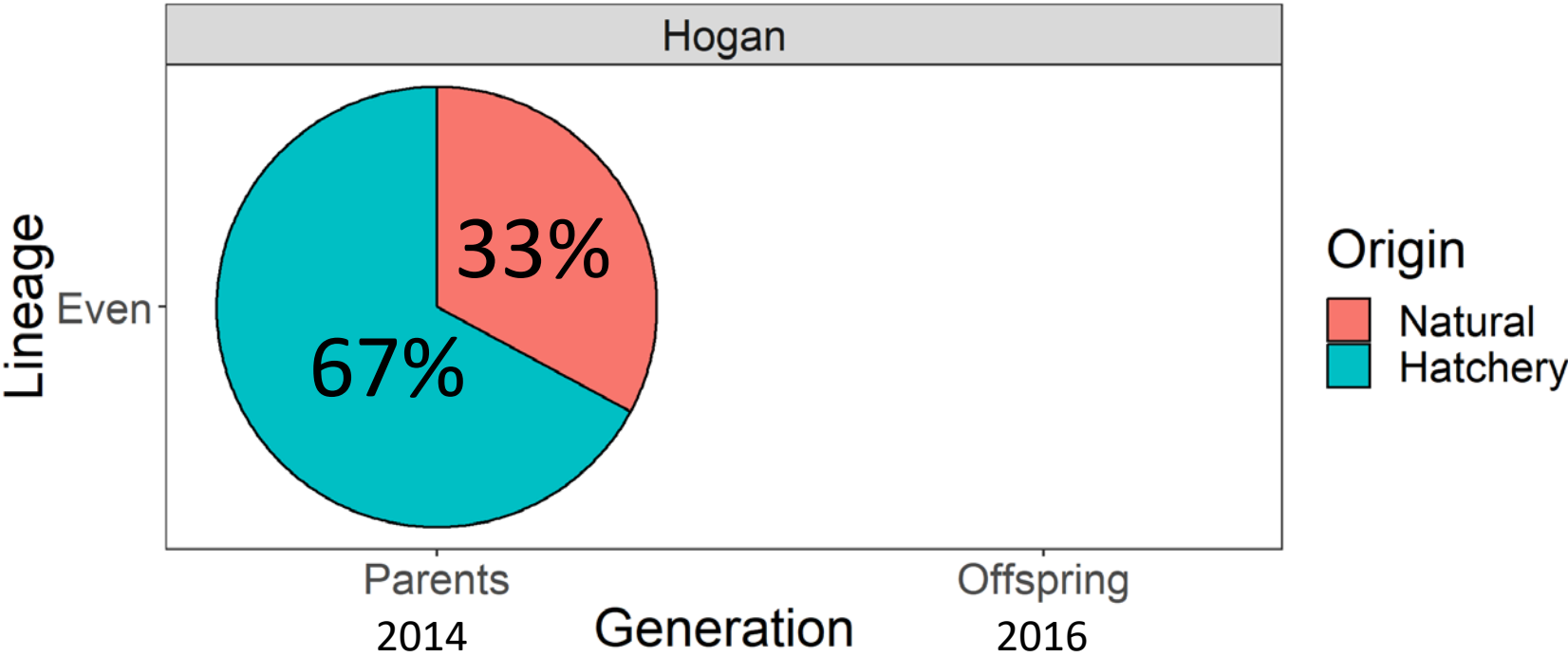
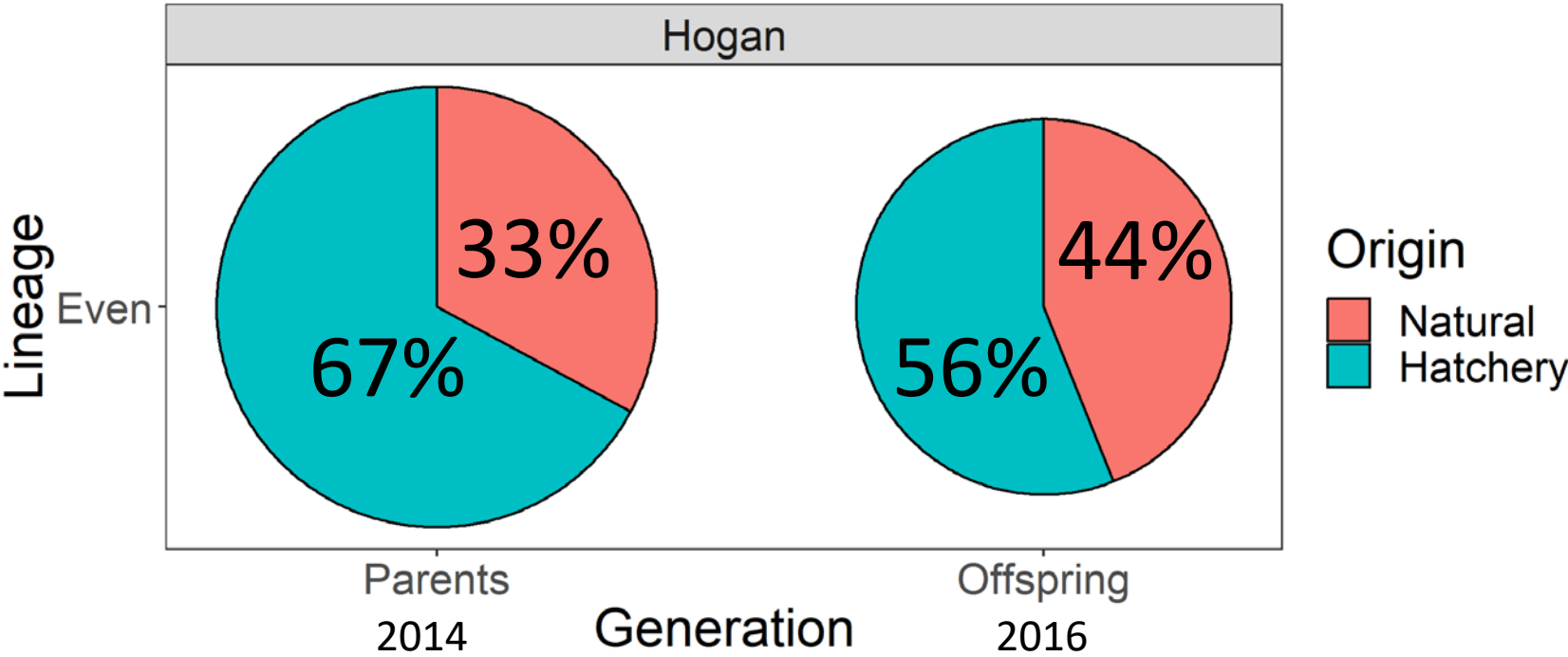


Figure 3b – Lescak et al. *in prep*

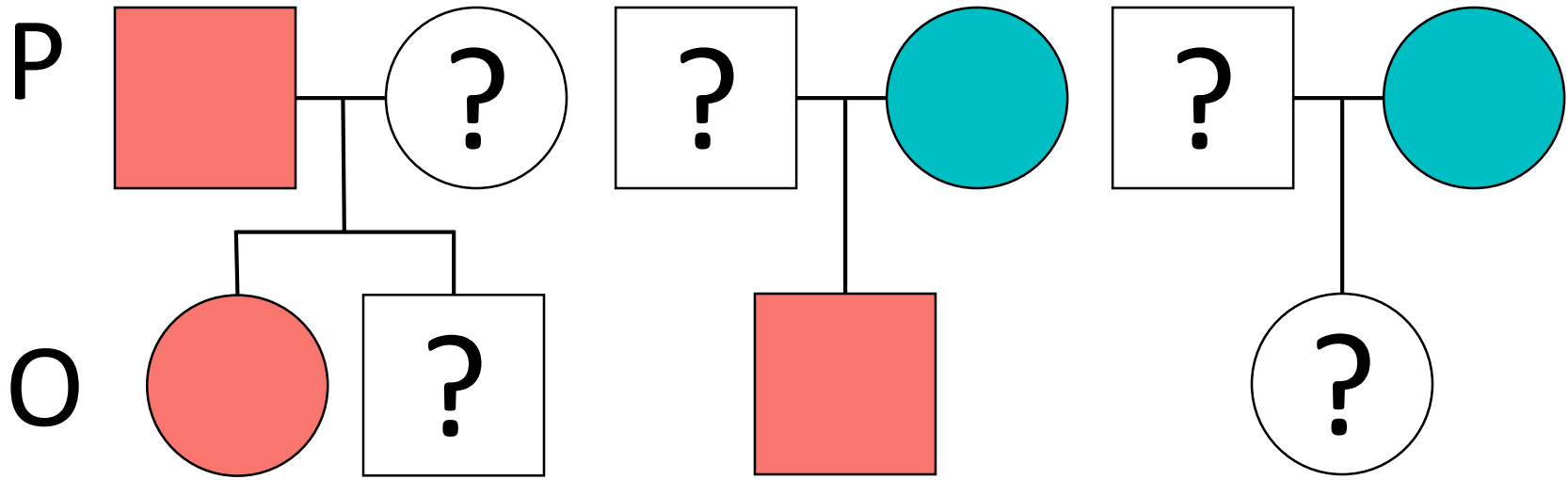
Proportion Test: Even-Lineage



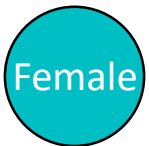
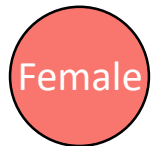
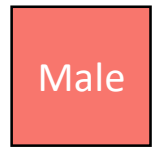
Proportion Test: Even-Lineage



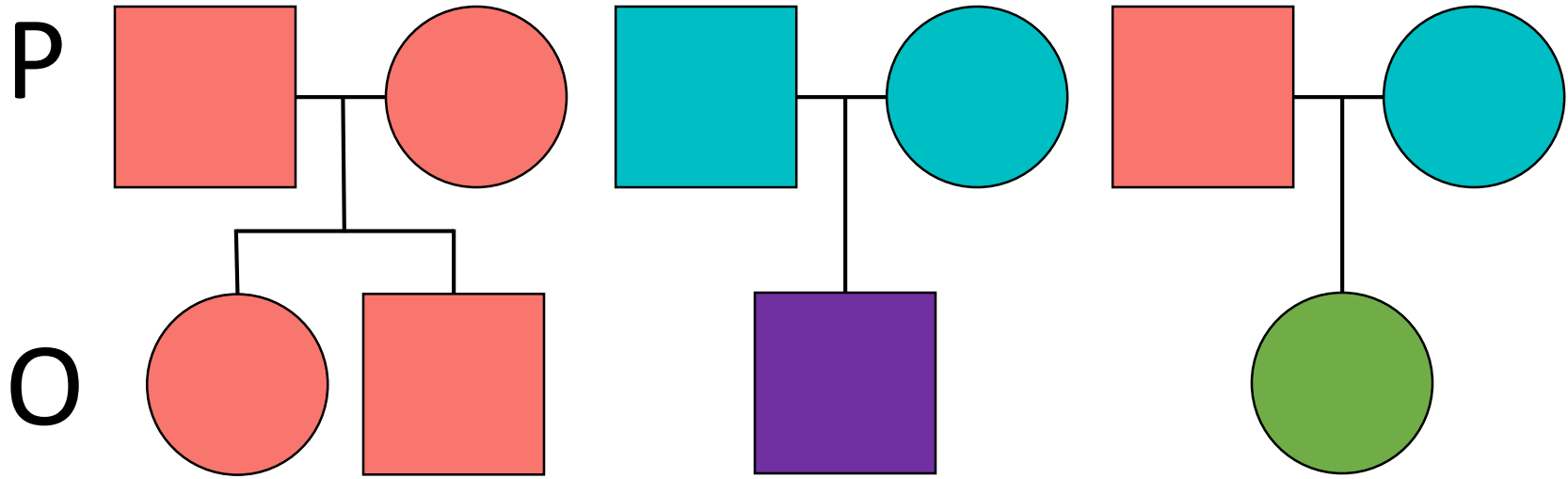
Parent-Offspring Duos



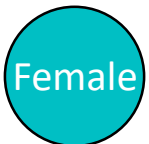
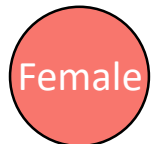
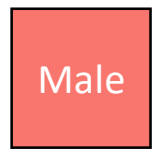
Natural Hatchery



Parent-Offspring Trios



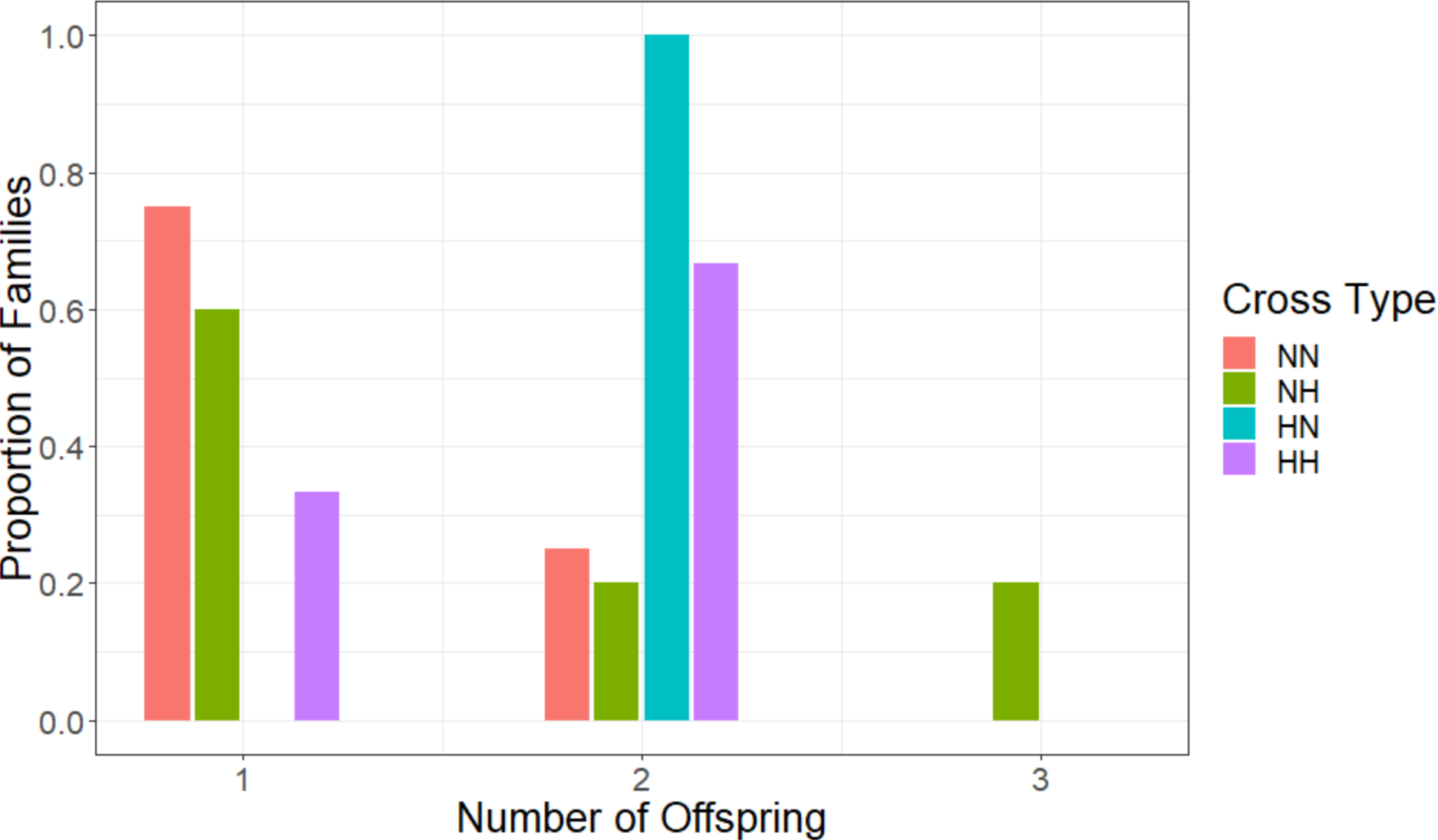
Natural **Hatchery**



Distribution of Reproductive Success by Cross Type



Distribution of Reproductive Success by Cross Type



Analyzed Samples: Odd-Lineage

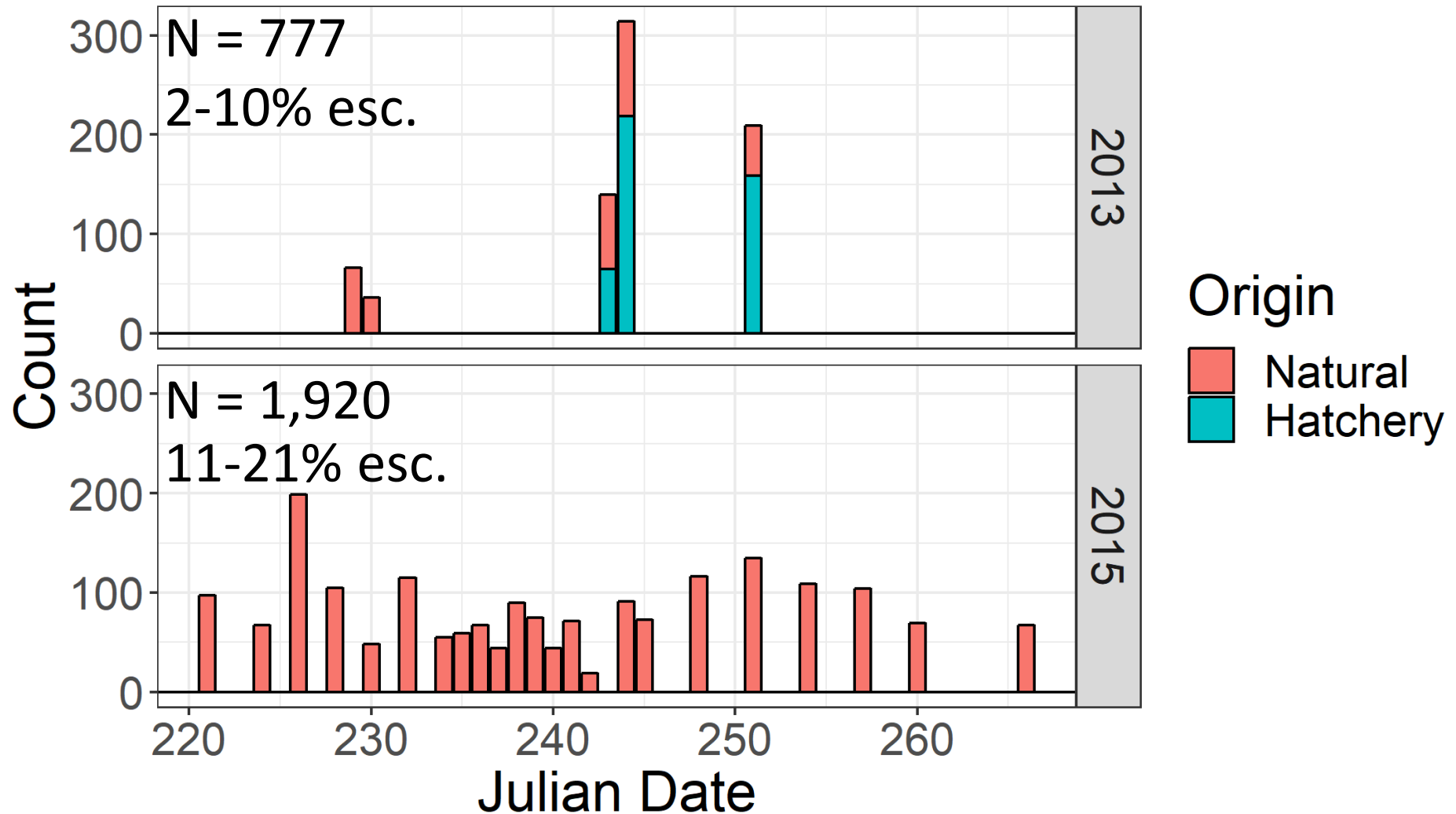


Figure 2a – Lescak et al. *in prep*

Pedigree Results: Odd-Lineage

- 48 offspring (2.3%) assigned to 20 parents
 - 45 → natural-origin parents
 - 3 → hatchery-origin parents
 - 2 – AFK
 - 1 – WNH
 - 0 – CCH
 - 0 – SGH

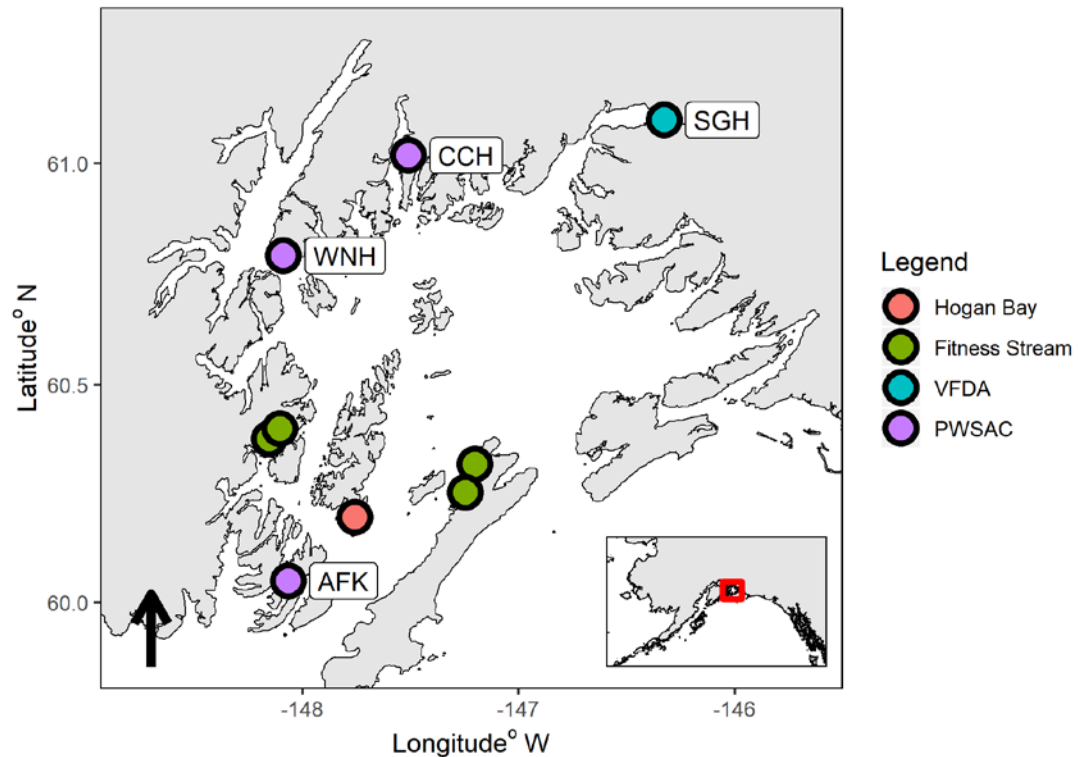


Figure 1 – Lescak et al. *in prep*

RS Distribution: Odd-Lineage

Reproductive Success for Odd-Lineage

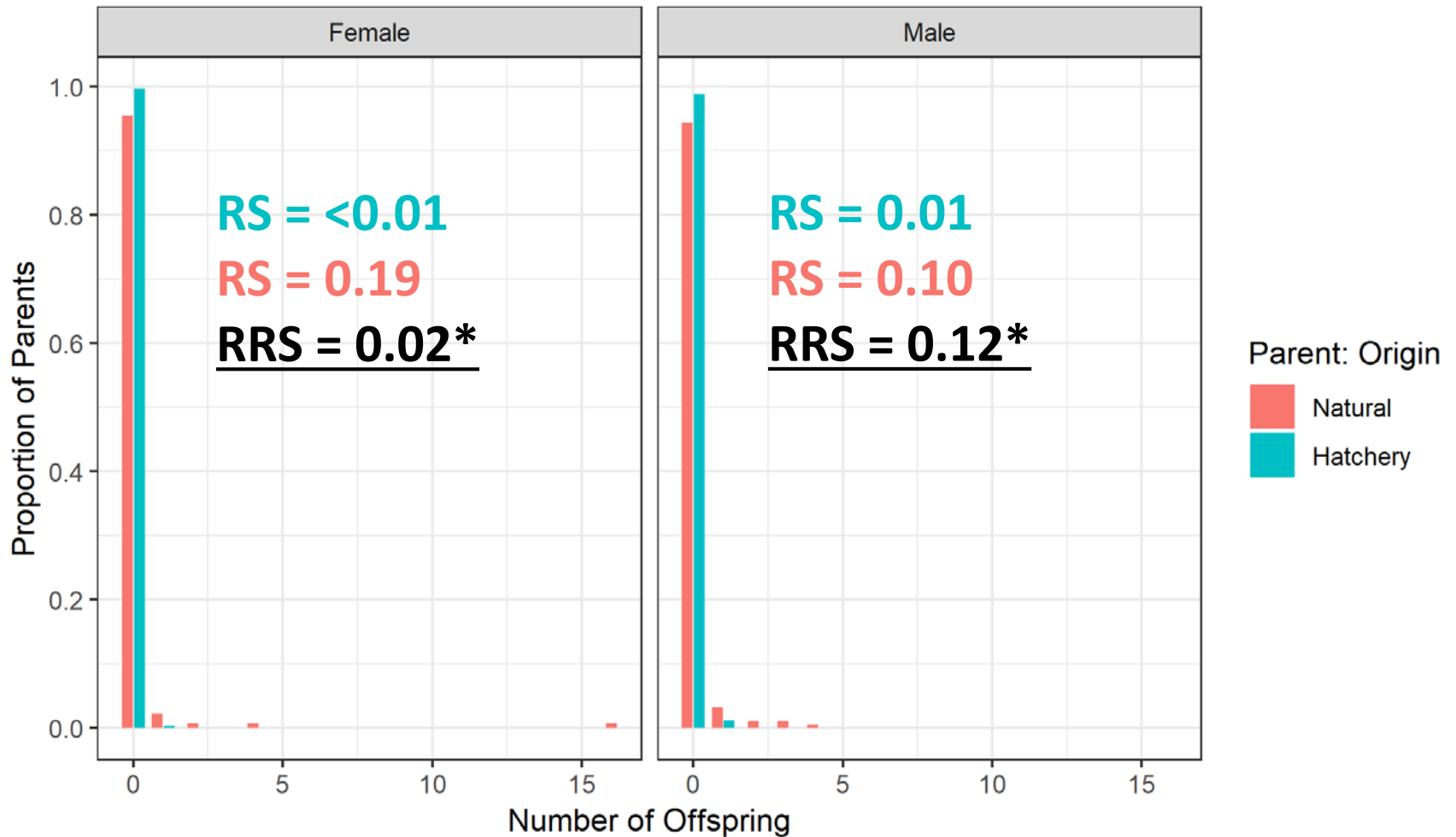
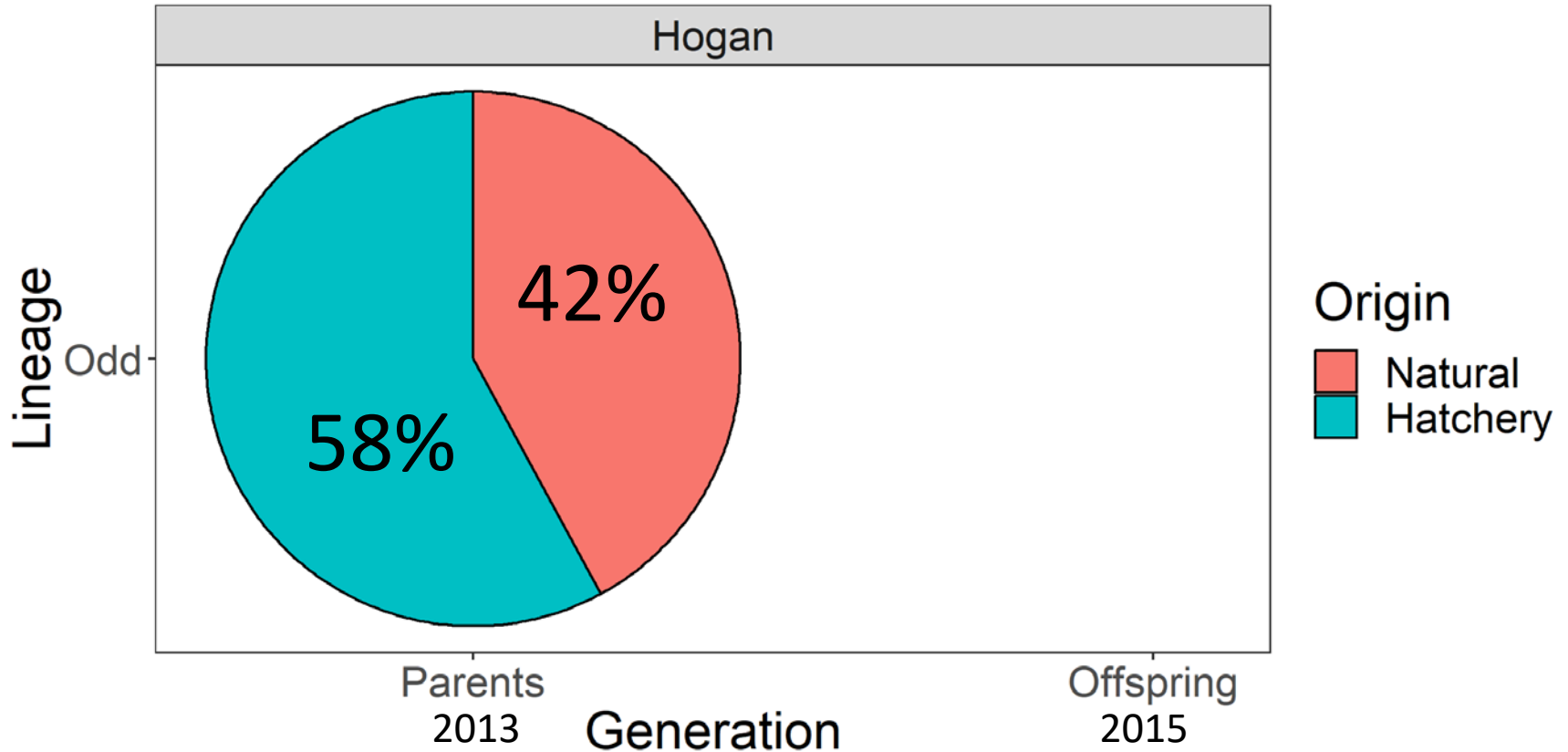
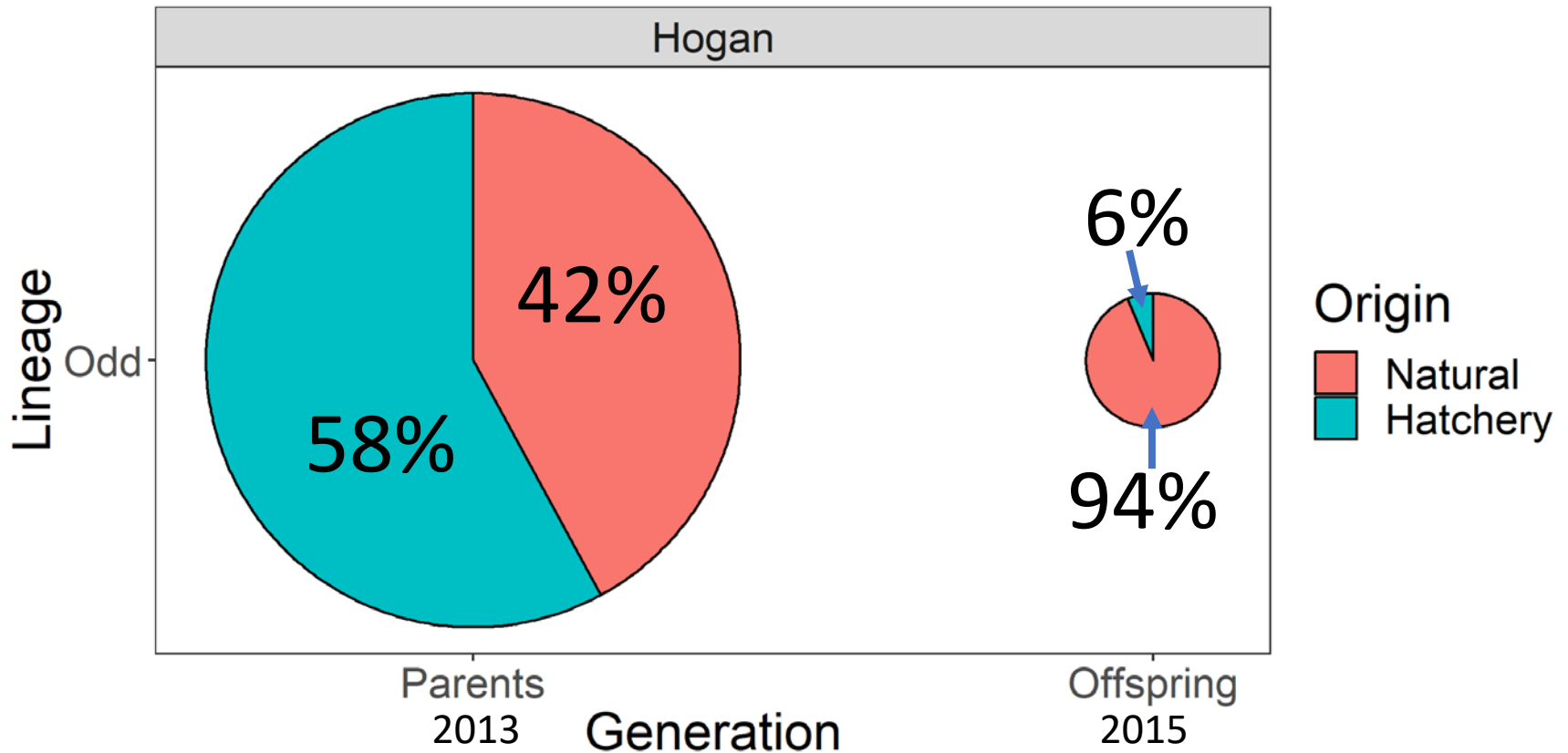


Figure 3b – Lescak et al. *in prep*

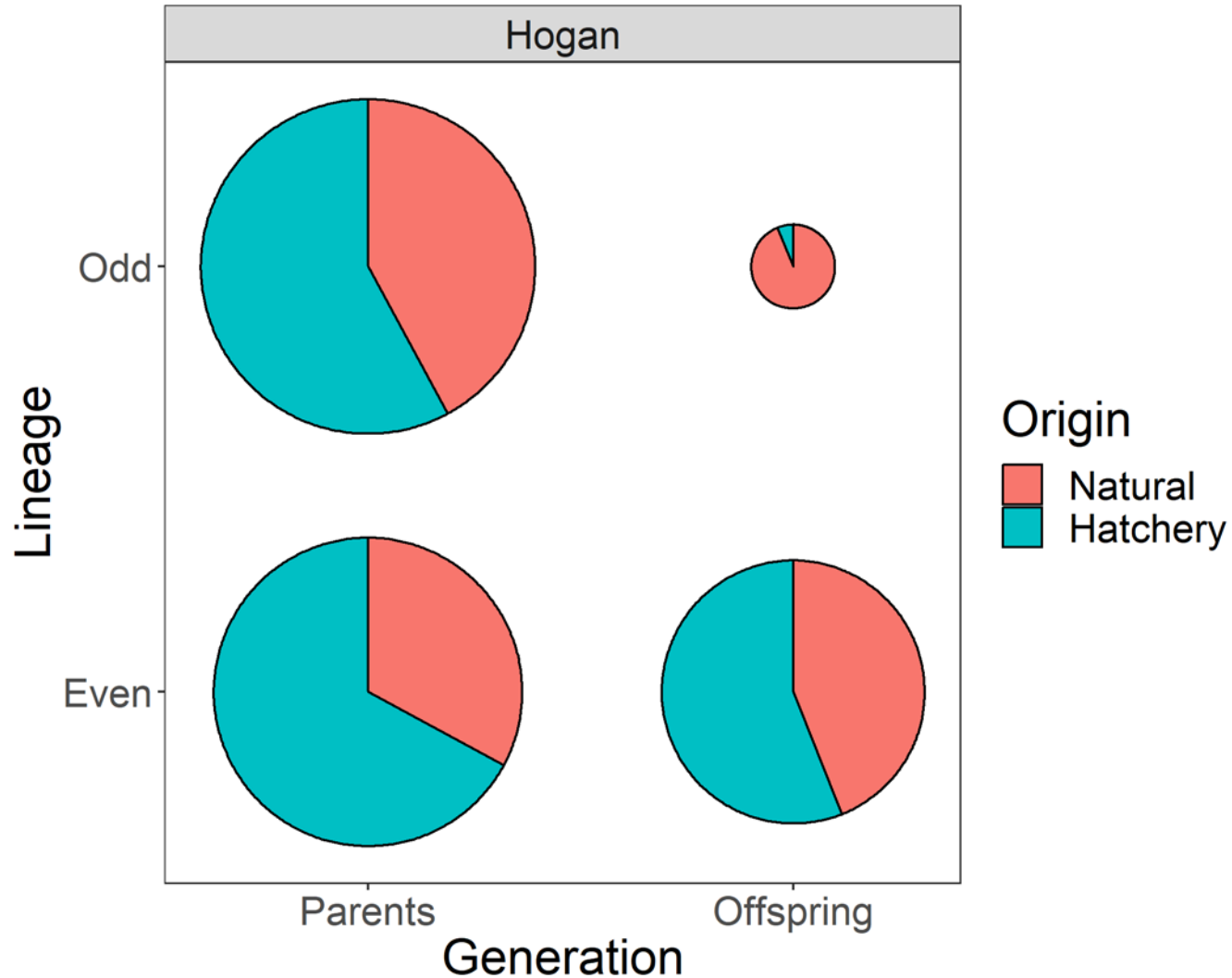
Proportion Test: Odd-Lineage



Proportion Test: Odd-Lineage



Proportions for Both Lineages



How robust are our pedigrees?

- Simulations
 - No incorrect or missed assignments
- Sensitivity analysis for *FRANz* parameters
 - Results robust to changes in genotyping error rates and maximum numbers of potential parents
- All parentage assignments unequivocal
 - No split pedigrees

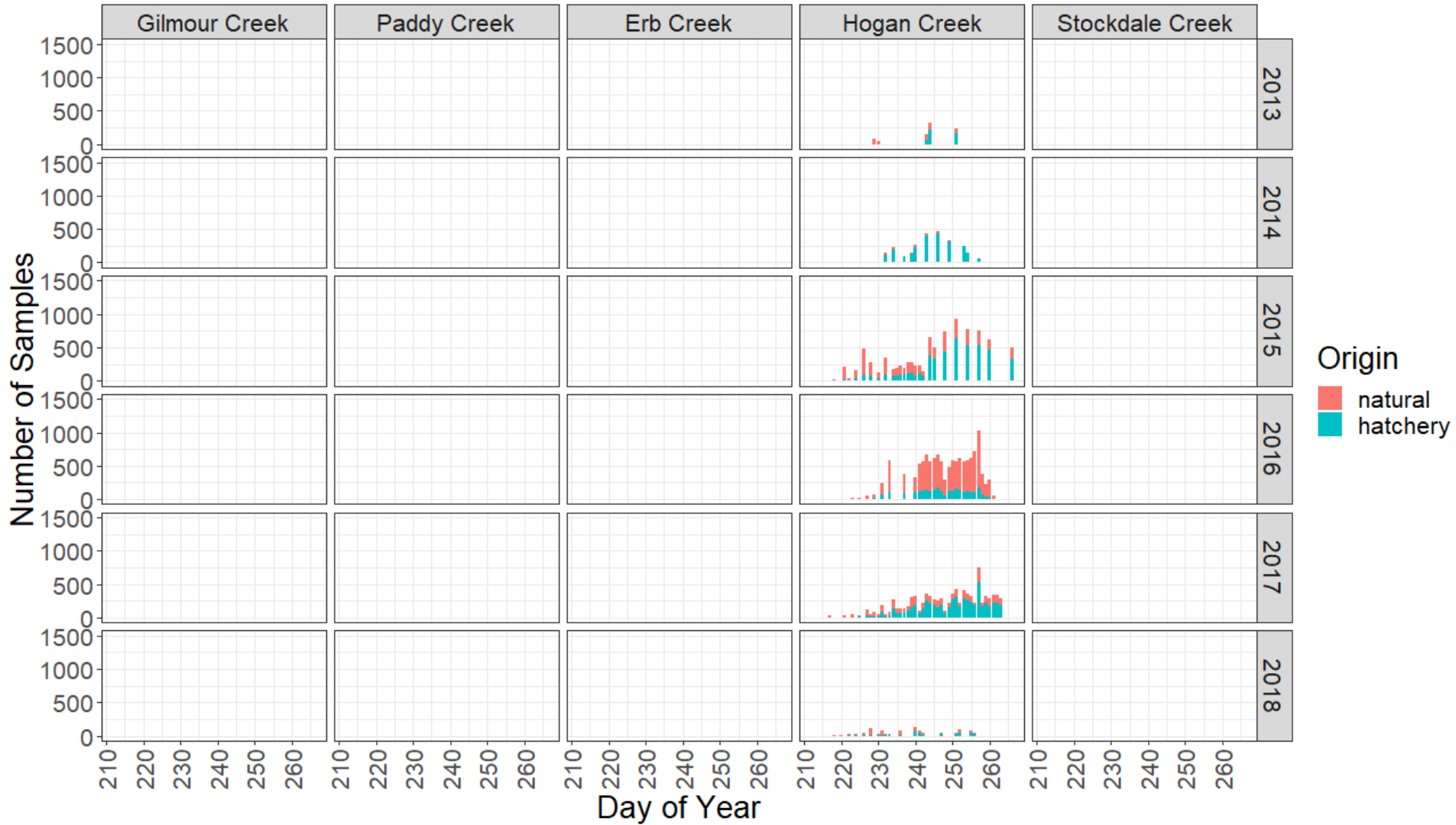
Results from 1 generation of Hogan

- Pedigree in natural system possible
- Even-lineage
 - 451 offspring to 184 parents
 - Offspring assignment rate 11.0%
 - RRS = 0.47 (significant) for females
 - RRS = 0.87 (not significant) for males
- Odd-lineage
 - 48 offspring to 20 parents
 - Offspring assignment rate 2.5%
- Under-representation of offspring assigned to hatchery-origin parents in both lineages

Conclusions from Hogan Bay

- Hatchery-origin fish spawned and produced adult offspring that were sampled
- Hatchery-origin fish spawned with both other hatchery-origin fish as well as natural-origin fish
- On average, hatchery-origin fish produced fewer adult offspring that returned to Hogan Bay and were sampled than their natural-origin conspecifics
- There are potentially important differences in RS between male and female hatchery-origin fish

Future Analyses



Future Analyses



Acknowledgements

- Alaska Hatchery Research Program
 - State of Alaska
 - Seafood industry
 - Private non-profit hatcheries
- North Pacific Research Board (Project #1619)
 - Funding for Hogan Bay analyses
- Prince William Sound Science Center
 - Field collection
- ADF&G Cordova Otolith Lab
- University of Washington - Seeb Lab
- ADF&G Gene Conservation Laboratory



A large group of salmon are captured in mid-air, jumping out of the water onto a rocky shore. The fish are in various stages of their jump, with some fully extended and others just beginning to leave the water. The water is a deep blue-green color, and the rocks on the shore are dark and wet. The overall scene is dynamic and energetic.

Questions?

AHRP Fitness Study: SEAK Chum Salmon



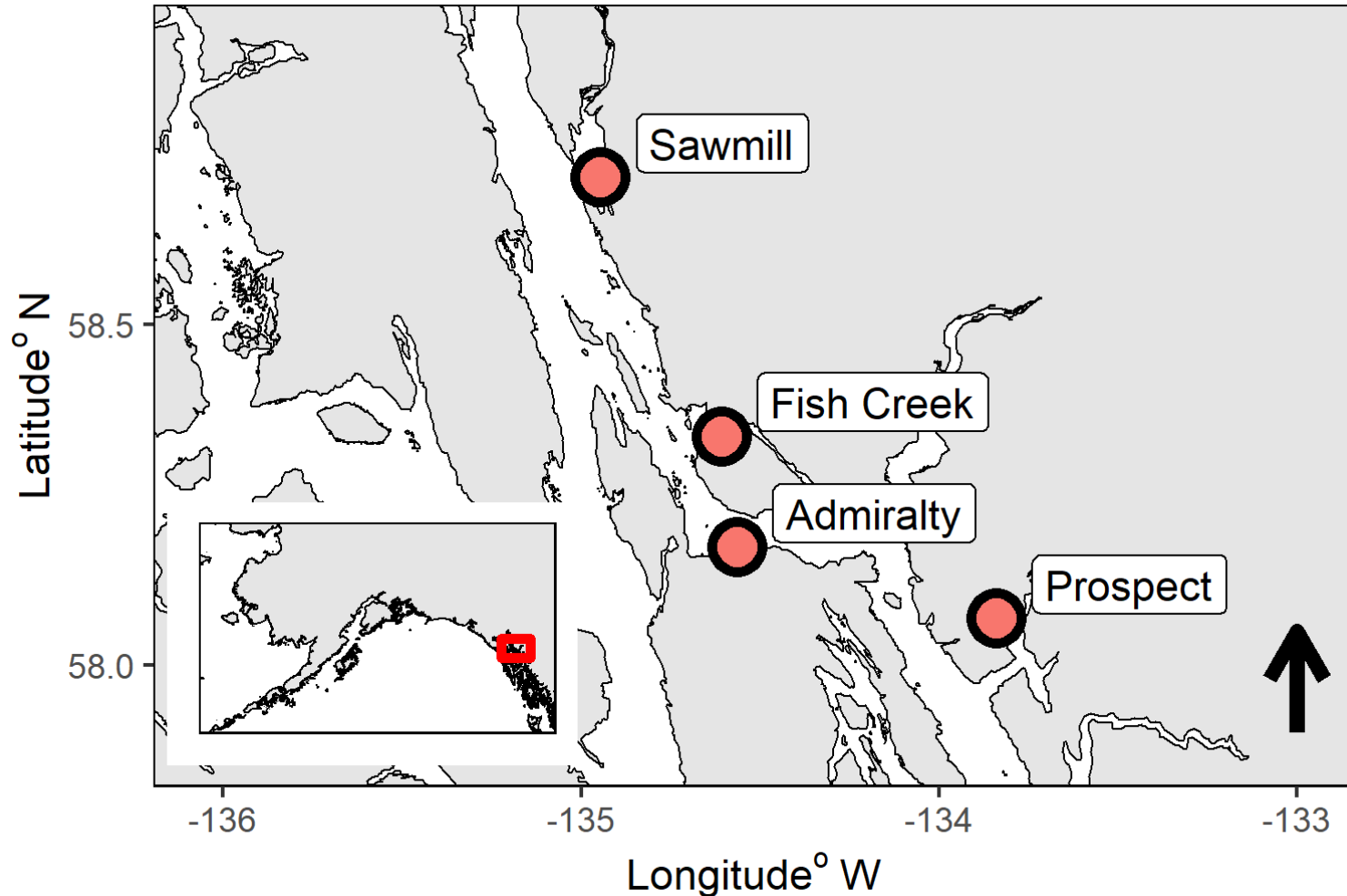
Kyle Shedd, E. Lescak, H. Hoyt, T. Dann, C. Habicht

Alaska Department of Fish and Game Gene Conservation Lab

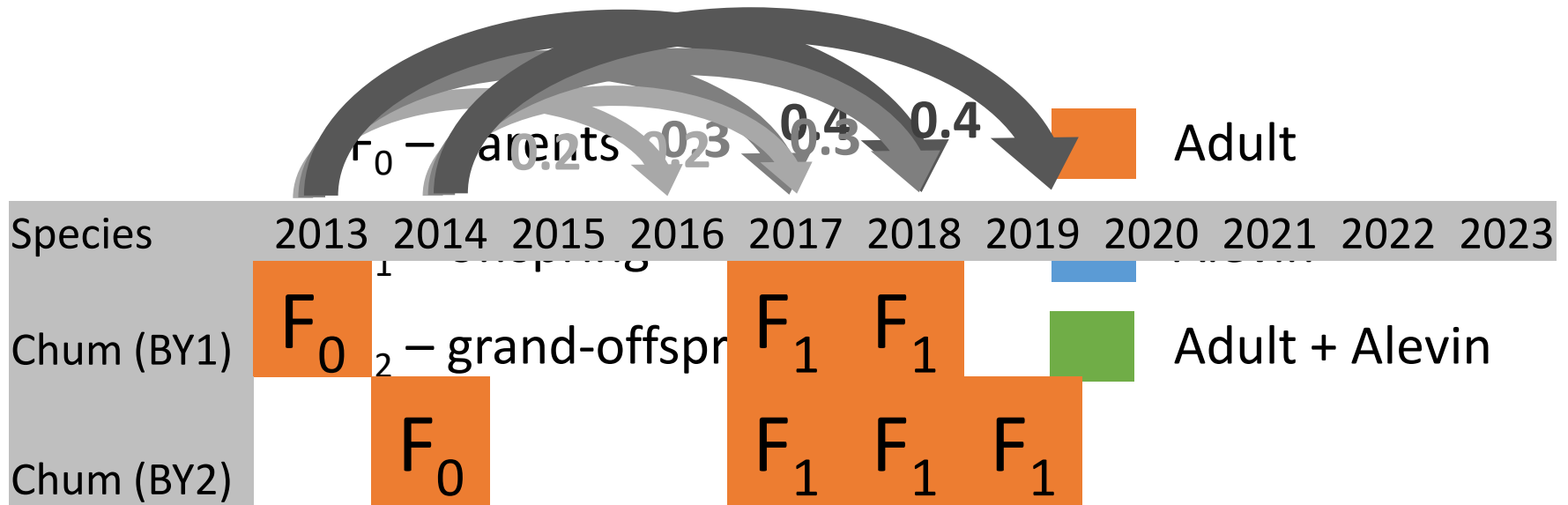
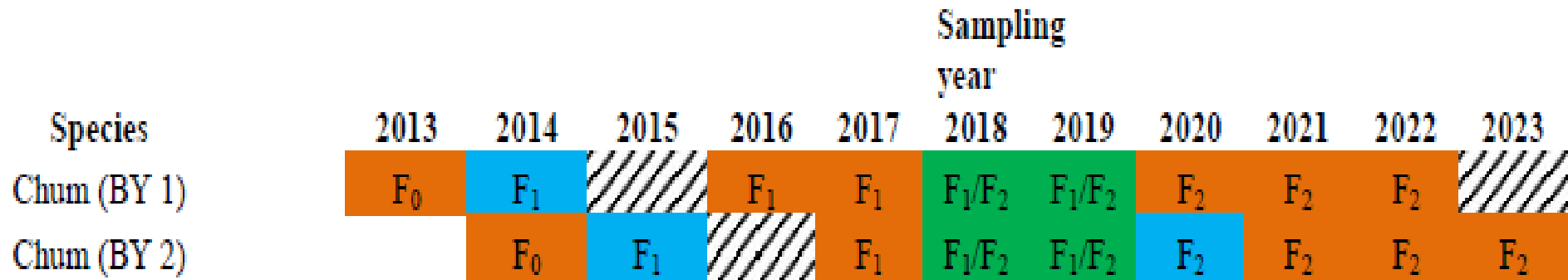
Alaska Board of Fisheries Hatchery Committee

March 8, 2019

Map of SEAK Chum fitness streams



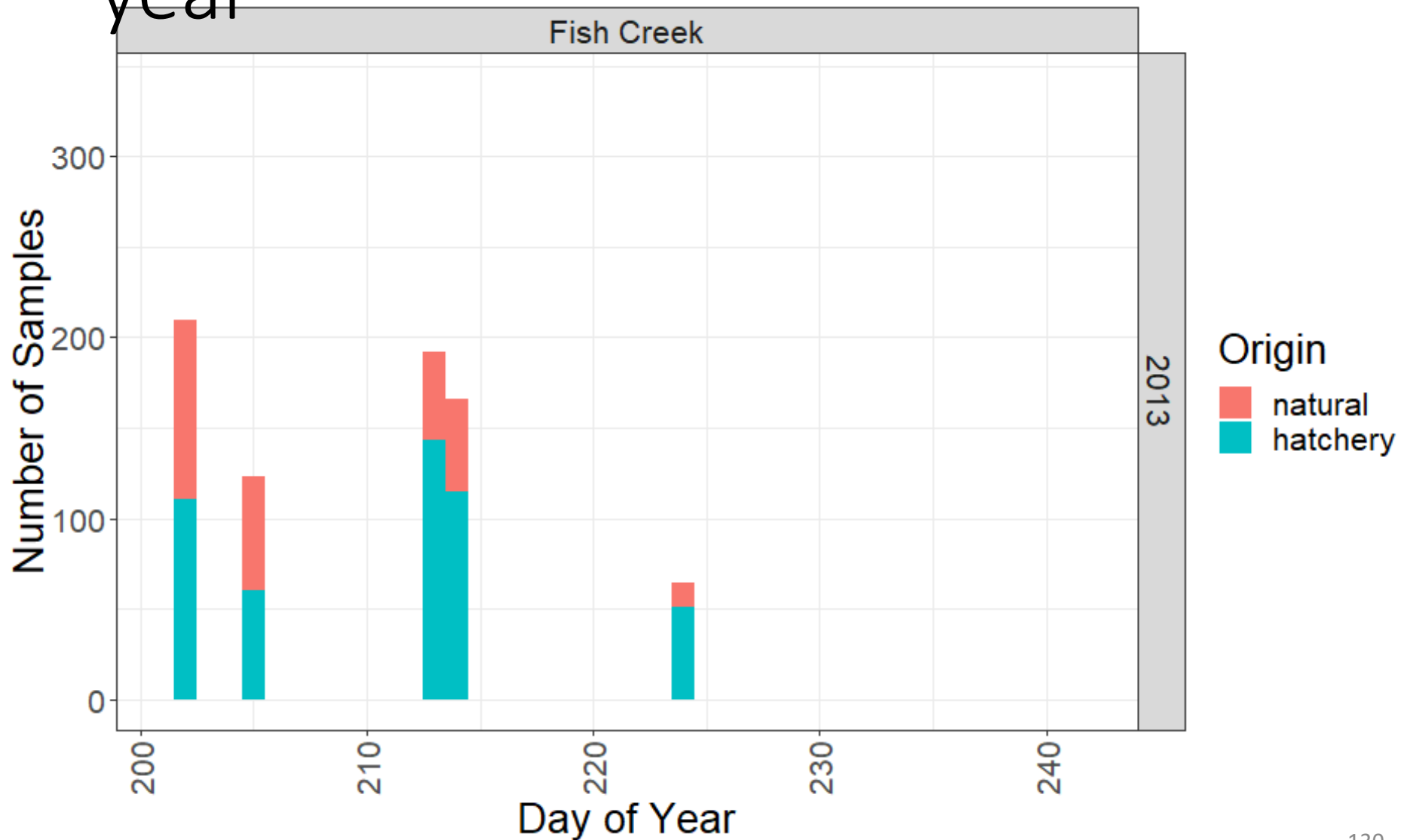
Study plan



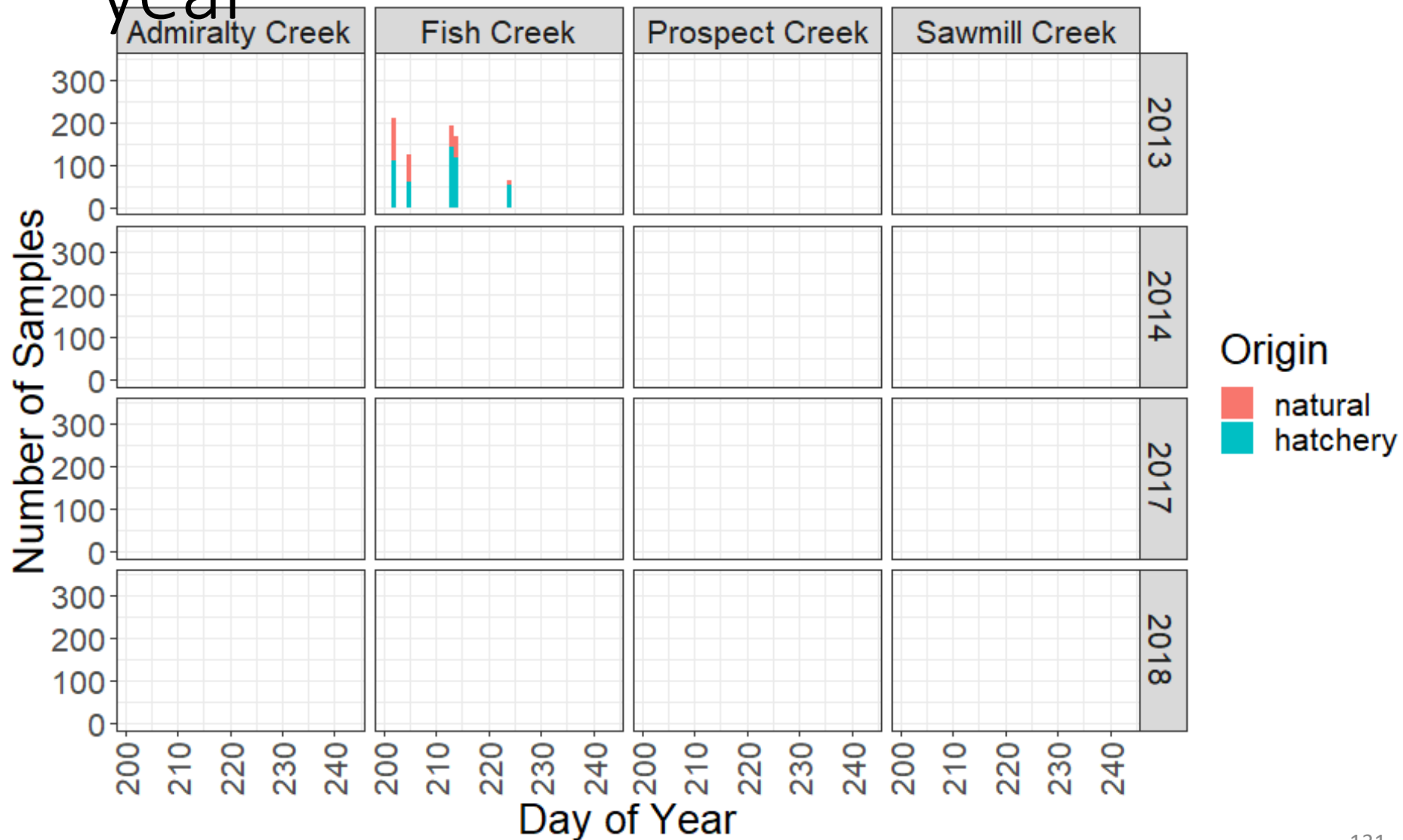
Statistical power of study plan

- Need minimum ~100 parents of each sex/origin
- Ideally a high proportion of parents
 - Hogan Bay 2013/2015
 - Low sampling rate = few parent-offspring assignments
- Sample high proportion of offspring
 - Consistent proportion for all return years
 - Differences in age at return?

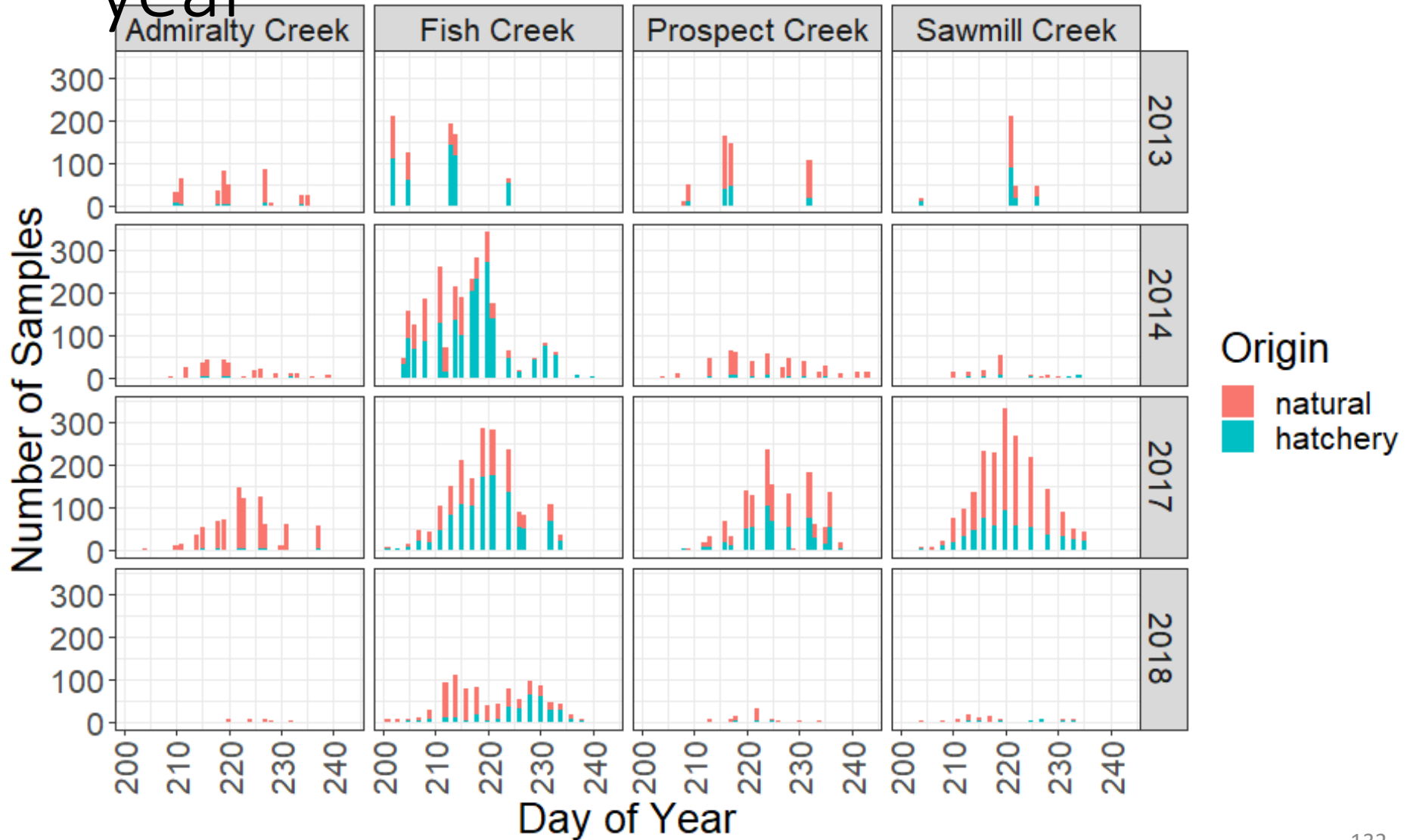
Samples by origin, stream, and year



Samples by origin, stream, and year



Samples by origin, stream, and year



Acknowledgements

- Alaska Hatchery Research Program
 - State of Alaska
 - Seafood industry
 - Private non-profit hatcheries
- Sitka Sound Science Center
 - Field collection
- ADF&G Mark, Tag and Age Lab
- ADF&G Gene Conservation Laboratory



Questions?

