Status of Chinook Standardization and Developing Technologies

Coded-wire Tag Workshop Lynnwood, Washington 9 June 2004

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Waypoints

- GAPS Consortium progress report
 - Marker selection/standardization
 - Database construction
 - Remaining challenges
- General considerations
 - International harvest management
 - Funding
 - Agency-specific priorities
 - Marker and baseline sample selection

Project objectives

- Develop a coast-wide DNA baseline that will resolve lineages and stocks within the region for which the CTC is responsible
- 2. Facilitate future expansion of the DNA baseline by the addition of novel genetic markers. This will be accomplished via the exchange of DNA samples
- Develop a data base application to support the dissemination and growth of the baseline

GAPS Collaborators

Agency

NWFSC

OSU

CDFO

SWFSC

CRITFC

UI

IDFG

ADFG

ABL

WDFW

USFWS Abernathy

USFWS Anchorage

P.I.*

Paul Moran

Michael Banks

Terry Beacham

Carlos Garza

Shawn Narum

Madison Powell

Matt Campbell

Lisa Seeb

Richard Wilmot

Sewall Young

Bill Ardren

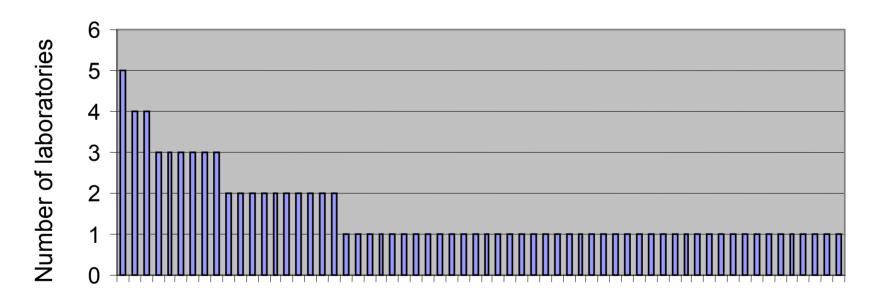
John Wenberg

^{*}Numerous coPIs not listed

Two fundamental challenges to standardization of microsatellite data

- Little overlap in marker sets among labs
- Different DNA fragment sizes among platforms and therefore different allele naming conventions

Only a few markers were widely used among labs



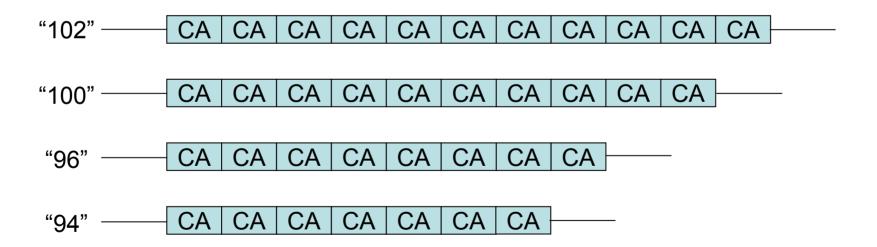
Microsatellite marker (N = 62)

Two fundamental challenges to standardization of microsatellite data

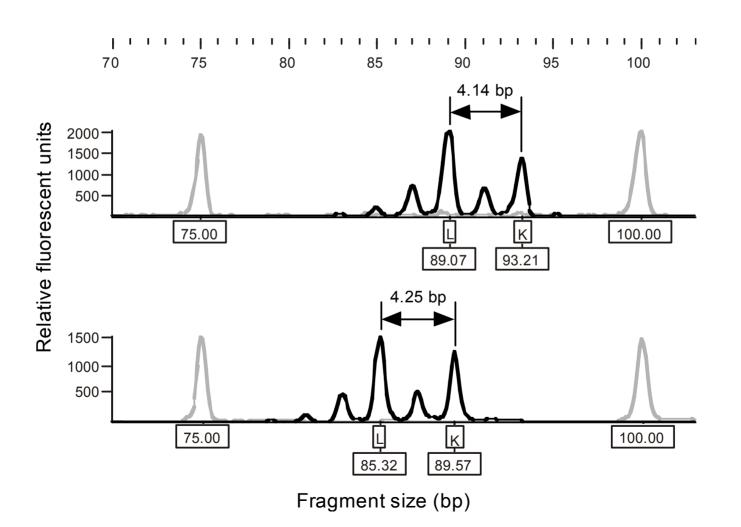
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Microsatellite markers—simple sequence repeats

Allele designations typically related to fragment size



Substantial differences in size estimates among platforms



2-Year GAPS study design

- Year 1: Standardization
 - Selection of a common set of markers
 - Standardization of allele designations
 - Selection of representative populations for year 2
- Year 2: Baseline development
 - Verification via blind samples
 - Sample exchange
 - Data collection and storage
 - Power analysis

"Marker (=locus) sets" and "Sample sets"

Markers*

- 62 sponsored loci
- 25 candidate loci
- 15 finalists for baseline construction

Samples

- 500-sample survey set
 - (10 individuals from each of 47 populations, range-wide)
- 96-sample reference set (8 individuals from each of 12 regions)
- 15,000-sample baseline set (144 individuals from each of 105 populations)

^{*}Current focus on microsatellites, future interest in SNPs

Year 1: Standardization

- Marker selection
 - 500-sample survey set run for 62 loci
 - 25 loci selected for optimization in all labs
 - 96-sample reference set run for all 25 loci
 - 15 loci selected for allele standardization
- Standardization of allele designations
 - Reconciliation among labs
 - "Binning" issues—"lumping" and "splitting"

Two organizational meetings to coordinate standardization

- Workshop associated with the annual meeting of the North Pacific Anadromous Fish Commission, 4-5 Nov 2003
- Ad hoc meeting in Seattle, 10-11 May 2004

Accomplishments at NPAFC

- Established marker selection criteria
- 25 candidate markers (out of 62 examined)
- Range-wide reference sample set to standardize scoring (N = 96, a subset of the 500)
- Putative baseline populations for next year's data collection phase
- Distribution of genotyping effort among labs-regional focus
- Database construction and data sharing

Accomplishments in Seattle

- 25 candidate loci reduced to 15 finalists
- Error-trapping and data reconciliation
- Allele designations, "holotypes," and vouchers
- Locus "curators" volunteered
- Resolution of the data base, lumping, splitting, and imperfect repeats

Ahead....

- 28 May Submit any genotype corrections
- 21 June New alleles to locus curator
- June Data repository—Docushare or FTP server
- 15 July Final standardized allele designations
- July 2004 Final report for year 1
- 1 Aug Baseline and blind samples sent to all labs
- 15 Sept Genotypes for blind samples submitted to NWFSC
- 1 Oct Early evaluation of loci—power analysis, Mendelian segregation, etc.
- 13 Oct Meeting?
- December Distribute samples collected in 2004 field season
- March 2005 Aliquots of DNA from baseline replicate sent to all labs ("Megaswap")
- May 2005 "Beta" version of baseline brought online for collaborators and CTC members
- July 2005 Final report presentation and development of peerreviewed publication

Remaining challenges

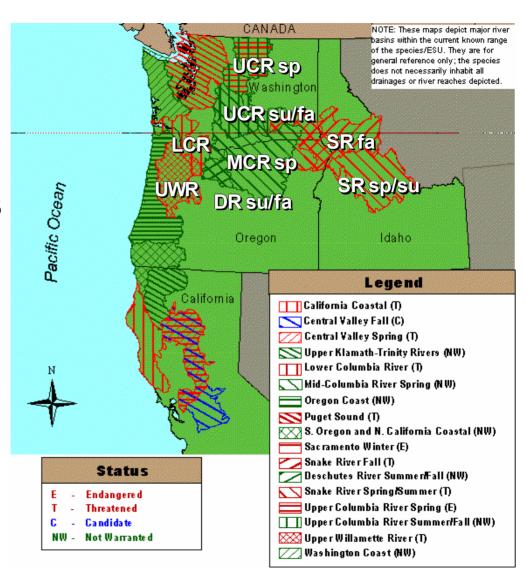
- Get ~15 markers to work in all labs
- Complete standardization of scoring
- Convince ourselves that 15 markers are enough (and not too much)
- Obtain all of the desired samples for the baseline
- Develop safe and secure repository for data

General observations about standardizing genetic data

- Essential to international fishery management
- Doesn't happen without specific funding
- Requires substantial sacrifice by participating labs
- Agency goals diverge from those of the community

NWFSC/WDFW Columbia R. Chinook baseline

- 3,000+ individuals
- 63 populations
- All ESUs
- 8 microsatellites
- 176 alleles
- High assignment accuracy



NWFSC microsatellite baseline for southern coho salmon

12 Loci

Ots103 OtsB3

Ots3 Oki1

P53 One13

Ocl8 Ots213

Oki10 OtsG83b

Oki23 OtsG422

Applications

Mixed Stock Analysis of Ocean-caught juveniles off Oregon and Washington

Recovery Planning for Oregon Coast and Lower Columbia River



D. Van Doornik, unpublished data

Selection of baseline populations

- Emphasis on major contributors to CTC fisheries, balanced against need to fully characterize Chinook population genetic variation range-wide
- Ultimately to provide a robust, state-of-the-art baseline with ample power to identify stock-oforigin of individuals in CTC fisheries
- Provide a standard reference set of Chinook populations that serve as a focal point for a whole range of conservation and recovery research

Marker performance summaries

- Quality of amplification and reliability of scoring
- Number of alleles and other measures of variability

Graphical examples

