## NEWS AND VIEWS

## PERSPECTIVE

## Puny males punch above their weight to preserve genetic diversity in a declining Atlantic salmon population

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Many salmonid fish populations have anadromous (i.e. migratory) and nonanadromous individuals co-existing in sympatry. The nonanadromous individuals, frequently males, mature at a much smaller size in freshwater without undergoing marine migrations and often successfully fertilize many eggs laid by anadromous females. Because these small males do not recruit to fisheries, they are often not regarded in high esteem by fishers. In this issue of Molecular Ecology, Johnstone et al. (2013) demonstrate that by substantially contributing to reproduction, such males help maintain genetic diversity in a declining population of Atlantic salmon (Salmo salar). Their results show that estimates of effective population size $\left(N_{\mathrm{e}}\right)$, obtained by counting the number of anadromous adults returning from sea and correcting for unequal sex ratios, are lower than estimates generated from genetic markers. Many mechanisms are expected to reduce $N_{\mathrm{e}}$ below the adult census population size ( $N$ ); the opposite pattern of $N_{\mathrm{e}}>N$ observed by Johnstone et al. (2013) is difficult to explain unless the reproductive effort of nonanadromous males is accounted for. The results have important implications for the conservation of small populations and highlight the challenges of relating $N_{\mathrm{e}}$ to $N$ in organisms with complex life histories.

Keywords: animal mating/breeding systems, conservation genetics, ecological genetics, fish, fisheries management, life history evolution

Received 30 November 2012; revised 17 December 2012; accepted 19 December 2012

It is not easy being an anadromous Atlantic salmon (Salmo salar). Born in freshwater, you are exposed to an array of predators as you migrate to marine feeding grounds in the North Atlantic. Once you attain a large enough size,

[^0]typically after one or more winters at sea, you return to natal spawning beds following an arduous upstream migration. It is therefore not surprising that in many populations, a substantial proportion of males - whose reproductive output is less tightly linked to body size than that of females - opt instead for the much easier option of foregoing marine migration and maturing at a small size in freshwater. These males then 'sneak' in between their larger anadromous brethren while they are fighting over access to anadromous females, and often successfully fertilize a significant portion of the eggs (Fleming 1996). Much recent work has strongly suggested that the variable life history of Atlantic salmon constitutes a conditional strategy wherein juveniles displaying certain phenotypes (often a larger size at age or growth rate) forego anadromous migrations in favour of freshwater residence (e.g. Hutchings \& Myers 1994; Piché et al. 2008). In some populations, males adopting freshwater residence far outnumber the larger anadromous males (Fleming 1996). Because fishers do not care much for those small males, their presence has often been seen as having a negative effect on recruitment to the fishery and on the harvestable biomass (Myers 1984). Indeed, nonanadromous males can eventually grow to become anadromous males, but the energy invested in early reproduction often results in increased mortality. For instance, Myers (1984) estimated that precocious male maturation reduced the anadromous male population by $60 \%$ in a Newfoundland stream. The study by Johnstone et al. (2013) published in this issue of Molecular Ecology suggests that, while small mature males do reduce the number of anadromous males, they contribute significantly to the preservation of genetic diversity in a declining population. They could thus contribute to the long-term persistence of populations - and thus to fisheries - that might otherwise suffer from a number of genetic consequences associated with small population size.
Johnstone et al. (2013) took advantage of a long-term data set of adult census population size ( $N$ ) estimates collected between 1985 and 2011 from a small salmon stream in southeast Newfoundland (Fig. 1), which they supplemented with microsatellite data from samples collected over the same period. The authors predicted that, if the nonanadromous males contributed significantly to genetic diversity in the population, effective population size ( $N_{\mathrm{e}}$ ) estimates based on genetic markers should be higher than those obtained from counting the returning number of anadromous adults (after correcting for unequal sex ratios). Few other mechanisms would be expected to lead to such a pattern. Indeed, a variety of well-known mechanisms reduce $N_{\mathrm{e}}$ compared with $N$ (e.g. variance in reproductive success, biased sex ratios, fluctuating population sizes), but the opposite pattern is difficult to explain


Fig. 1 The counting fence set up on Northeast Brook, Trepassey in southeast Newfoundland, Canada, used by Johnstone et al. (2013) to obtain estimates of the anadromous adult Atlantic salmon run sizes between 1985 and 2011. The inset shows two nonanadromous small mature male parr Atlantic salmon swimming next to a much larger anadromous male. Photo credits: top: D. Ruzzante, bottom: L. Weir \& J. Hutchings.
unless one considers the contribution of nonanadromous males. Because of the prevalence of early maturation in males, the sex ratio of returning anadromous adults was female-biased; $N_{\mathrm{e}}$ was therefore estimated based on theoretical predictions of the effect of biased sex ratio on $N_{\mathrm{e}}$. Note that, many other phenomena not taken into account here (e.g. high variance in reproductive success) would also reduce effective population size, but this only renders the conclusions of the study more conservative. Estimating $N_{\mathrm{e}}$ using genetic markers can be challenging and estimates often vary between methods or can have large confidence intervals (Luikart et al. 2010; Palstra \& Fraser 2012). The authors mitigated this potential problem by generating many estimates from a variety of estimation methods (for a recent review of available genetic methods, see Luikart et al. 2010). The data set allowed the use of several different temporal methods on samples separated by roughly five generations. These temporal estimates were supplemented with those generated from a single year of sampling. In this case again, multiple methods were used, but only the linkage disequilibrium method was found to give reliable results. Variation among estimates generated with different methods was considerable (between 290 and 584). Nevertheless, in all cases, the $N_{\mathrm{e}}$ estimates obtained from microsatellites were $34-420 \%$ higher than the demographic estimate of $N_{\mathrm{e}}$ obtained using the information on the sex ratio of the anadromous run.

Conditional mating strategies are widespread among salmonids and other taxa (Gross 1996), and the results of Johnstone et al. (2013) have implications for the conserva-
tion of small populations of other species of salmonids. It remains to be seen whether the contribution of these puny males to the preservation of genetic diversity is a widespread phenomenon, but in the population studied by Johnstone et al. (2013), they certainly punch above their weight class. The study also highlights some of the challenges associated with calculating $N_{\mathrm{e}} / N$ ratios in organisms with complex life histories. Estimates of $N_{e}$ are easily obtained from molecular markers and offer the promise of a simple and cost-effective tool to estimate $N$ in conservation programs strapped for cash (Luikart et al. 2010). But a simple conversion factor based on $N_{\mathrm{e}} / N$ ratios remains elusive (Palstra \& Fraser 2012), and such ratios are likely to vary among species, among populations within species and even over time within populations. More studies that provide reliable estimates of both $N_{\mathrm{e}}$ and $N$ are needed to understand the relationship between the two parameters. Reliable estimates of census population size are particularly hard to come by (Palstra \& Fraser 2012), and the study by Johnstone et al. (2013) suggest that they may be particularly unreliable in species where one component of the population is more difficult to catch or survey than the remainder.

## References

Fleming IA (1996) Reproductive strategies of Atlantic salmon: ecology and evolution. Reviews in Fish Biology and Fisheries, 6, 379-416.
Gross MR (1996) Alternative reproductive strategies and tactics: diversity within sexes. Trends in Ecology \& Evolution, 11, 92-98.
Hutchings JA, Myers RA (1994) The evolution of mating strategies in variable environments. Evolutionary Ecology, 8, 256-268.
Johnstone DL, O'Connell MF, Palstra FP, Ruzzante DE (2013) Mature male parr contribution to the effective size of an anadromous Atlantic salmon (Salmo salar) population over 30 years. Molecular Ecology, 22, 2394-2407.
Luikart G, Ryman N, Tallmon DA, Schwartz MK, Allendorf FW (2010) Estimation of census and effective population sizes: the increasing usefulness of DNA-based approaches. Conservation Genetics, 11, 355-373.
Myers RA (1984) Demographic consequences of precocious maturation of Atlantic salmon (Salmo salar). Canadian Journal of Fisheries and Aquatic Sciences, 41, 1349-1353.
Palstra FP, Fraser DJ (2012) Effective/census population size ratio estimation: a compendium and appraisal. Ecology and Evolution, 2, 2357-2365.
Piché J, Hutchings JA, Blanchard W (2008) Genetic variation in threshold reaction norms for alternative reproductive tactics in male Atlantic salmon, Salmo salar. Proceedings of the Royal Society B: Biological Sciences, 275, 1571-1575.
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doi: $10.1111 / \mathrm{mec} .12222$


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