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Natal Homing and Connectivity in Atlantic Bluefin Tuna Populations

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Atlantic bluefin tuna populations are in steep decline, and an improved understanding of connectivity between individuals from eastern (Mediterranean Sea) and western (Gulf of Mexico) spawning areas is needed to manage remaining fisheries. Chemical signatures in the otoliths of yearlings collected from 1999 to 2004 in the eastern Atlantic Ocean (Cantabrian Sea) and western Atlantic Ocean (Gulf of Mexico) prove that the populations are not distinct entities, and that transatlantic mixing is occurring. Adults repeatedly to the same spawning region (natal homing) has been documented, and management activities, the International Commission for the Conservation of Atlantic Tunas (ICCAT) has assumed that Atlantic bluefin tuna occur as two discrete populations that originate in the Mediterranean Sea or the Gulf of Mexico; and these populations are relatively small in number, justifying the use of two broad management regions east and west of 45° W longitude. Despite four decades of regulation by ICCAT, bluefin tuna populations remain severely depressed, causing many to question the effectiveness of the current management regime. Several approaches have been used to examine the population structure of Atlantic bluefin tuna, of which chemical traces in otoliths (ear stones) have considerable potential for quantifying natal homing and connectivity because otolith material deposited during the first year of life serves as a natural tag of the individual’s place of birth. Otolith δ13C and δ18O values for yearling Atlantic bluefin tuna collected from 1999 to 2004 in the eastern Atlantic Ocean/Mediterranean Sea (blue triangles) and western Atlantic Ocean (red triangles) show that bivariate ellipses (one standard deviation of the mean) and normal distribution curves are shown. Yearlings ranged in age from 12 to 18 months. Two regions of the eastern Atlantic Ocean/Mediterranean Sea were sampled over the 6 years: the eastern Atlantic Ocean (Cantabrian Sea; 2000, 2001, and 2002) and the western/central Mediterranean Sea (Ligurian Sea to Adriatic Sea; 1999, 2000, 2002, 2003, and 2004) (n = 113). In the continental shelf waters of the U.S. Atlantic Ocean, yearlings were collected from Maryland to Massachusetts over a 6-year period (n = 81) [see S8 in (8)].

Fig. 1. Otolith δ13C and δ18O values for yearling Atlantic bluefin tuna collected from 1999 to 2004 in the eastern Atlantic Ocean/Mediterranean Sea (blue triangles) and western Atlantic Ocean (red triangles). Gaussian bivariate ellipses (one standard deviation of the mean) and normal distribution curves are shown. Yearlings ranged in age from 12 to 18 months. Two regions of the eastern Atlantic Ocean/Mediterranean Sea were sampled over the 6 years: the eastern Atlantic Ocean (Cantabrian Sea; 2000, 2001, and 2002) and the western/central Mediterranean Sea (Ligurian Sea to Adriatic Sea; 1999, 2000, 2002, 2003, and 2004) (n = 113). In the continental shelf waters of the U.S. Atlantic Ocean, yearlings were collected from Maryland to Massachusetts over a 6-year period (n = 81) [see S8 in (8)].
origin or nursery habitat (6, 7). We determined the origin of bluefin tuna by means of carbon and oxygen stable isotope ratios ($\delta^{13}C$ and $\delta^{18}O$) in otoliths [see S1 in (8)]. Stable isotopes in otoliths and other biogenic carbonates represent a class of chemical tags for which associations with water mass properties are well understood (9). Global records of surface water stable carbon and oxygen isotopes indicate these natural markers vary regionally (10), and in otoliths these isotopes often serve as natal tags because they reflect water composition differences in nurseries, although fractionation due to kinetic and metabolic effects, particularly for $\delta^{13}C$, can influence isotopic composition (11).

The isotopic composition of otoliths from yearling (12 to 18 months of age) bluefin tuna was measured for individuals collected over 6 years (1999 to 2004) from both eastern (Mediterranean Sea/eastern Atlantic Ocean) and western (Gulf of Mexico/U.S. Atlantic Ocean) nurseries (Fig. 1). Otolith composition was distinct between yearlings from eastern and western nurseries [multivariate analysis of variance (MANOVA), $P < 0.01$; based on pooled years], and otolith $\delta^{18}O$ values were significantly higher for yearlings from the eastern nursery in five of the years (all except 2001). Mean (SD) otolith $\delta^{18}O$ values for the eastern and western nurseries were $-0.89\%$ (0.23) and $-1.66\%$ (0.37), respectively. No significant differences in otolith $\delta^{13}C$ values were observed between eastern and western bluefin tuna in five of the six years sampled (exception 2002, ANOVA, $P < 0.05$) (Fig. 1). Using quadratically discriminant function analysis (QDFA) parameterized with otolith $\delta^{13}C$ and $\delta^{18}O$ values from all year classes, cross-validated classification success to eastern and western nurseries was high (87%), indicating that stable isotopes were useful markers of natal origin for bluefin tuna [see S2 in (8)].

Milled cores of otoliths were used to represent the yearling period of school (<60 kg), medium (60 to 140 kg), and giant (>140 kg) category bluefin tuna [see S3 in (8)], and maximum likelihood estimates were generated using a mixed-stock algorithm [see S4 in (8)] (12) to assign individuals to eastern and western nurseries. Although the temporal stability of the primary marker, $\delta^{18}O$, justified the pooling of years for establishing a baseline, temporal variability in otolith core $\delta^{18}O$ or $\delta^{13}C$ values was investigated because our calibration set of juveniles (baseline) did not match our adults. No differences were detected in otolith core $\delta^{18}O$ values between bluefin tuna with birthdates before or during our baseline period in either region, suggesting that the otolith $\delta^{18}O$ composition remained relatively constant over the years investigated [see S5 in (8)]. Changes in $\delta^{13}C$ values in otolith cores were detected, and depletion levels for $\delta^{13}C$ were significant and similar to reductions reported in the oceans due to release of $CO_2$ from the burning of fossil fuels (Suess Effect) (13). Otolith core $\delta^{13}C$ values were adjusted to account for such temporal variation, but estimates of origin were comparable to unadjusted values [see S6 in (8)].

Natal homing, defined as the return of spawning adults to their region of origin, was high and remarkably similar for both eastern and western spawning regions: 95.8% for the Mediterranean Sea and 99.3% for the Gulf of Mexico [see S7 in (8)]. This work corroborates earlier indications of spawning fidelity from electronic tagging data that showed return migrations of both Mediterranean (5) and Gulf of Mexico (5, 14) spawners over multiple years. Additionally, genetic differences have been observed between juveniles collected from the two regions (15). Documentation of natal homing in marine vertebrates is rare, and the size of natal areas for bluefin tuna are much larger than in systems where homing is better known (e.g., streams, lakes, and estuaries). Nevertheless, rates of return to eastern and western spawning areas by Atlantic bluefin tuna populations are at the upper end of ranges reported for teleosts, rivaling those of Pacific salmon (16). Although we could not document repeat spawning in this particular application, long-term archival tag deployments provide evidence of bluefin returning to the same spawning sites over consecutive years (2, 5).

Estimates of natal origin from otolith chemistry indicate that mixing occurs in North American waters, and our data confirm that U.S. waters...
commercial and recreational fisheries are composed of both populations of bluefin tuna (Fig. 2). A large fraction of the school (57.4%) and medium (44.3%) category bluefin tuna present in the U.S. waters of the Mid Atlantic Bight were from the eastern population, and we observed that the occurrence of eastern bluefin tuna in the Mid Atlantic Bight decreased with increasing size (age) (Fig. 3). Our estimates of trans-Atlantic exchange were significantly higher than previous reports from conventional tags (3) and demonstrated substantial intermingling of individuals from eastern and western populations in U.S. waters, a finding supported with recent evidence from electronic tags (5). In contrast, giant category bluefin tuna collected from northern U.S. (Gulf of Maine) and Canadian (Gulf of St. Lawrence) fisheries were almost entirely of western origin (94.8% and 100%, respectively). The mechanism(s) responsible for differences in stock composition of bluefin tuna samples from Mid Atlantic Bight (mixed populations) and Gulf of Maine/Gulf of St. Lawrence (western population) waters appears related to size (age) or reproductive state. The majority of our sample from the Mid Atlantic Bight was composed of adolescent bluefin tuna (<5 years of age), and tagging studies have demonstrated that young bluefin tuna are more likely to display trans-Atlantic movements that are linked to foraging than are adults (2). Ontogenetic shifts in dispersive behaviors often occur for marine vertebrates displaying natal homing, with exploratory movements associated with foraging decreasing at the onset of breeding (17, 18). Similarly, our finding of stock homogeneity of giants (>140 kg, >10 years of age) in the Gulf of Maine and Gulf of St. Lawrence, and increasing contributions from the western population with age in the Mid Atlantic Bight, suggests that movement becomes more limited and structured after bluefin tuna become sexually mature.

Significant trans-Atlantic mixing of eastern adolescents on western foraging areas emphasizes the connectivity of Atlantic bluefin tuna populations. Under the current assessment framework that assumes limited mixing, a high degree of exchange evident from chemical signatures in otoliths indicates that past abundances of western Atlantic bluefin tuna may have been overestimated, particularly at younger age classes. In addition, exchange rates reported here show that U.S. fisheries for bluefin tuna appear dependent, to some extent, on recruits from the Mediterranean Sea. Because the eastern population is at least an order of magnitude higher in abundance than the western population (19), it is unlikely that west-to-east movement of adolescents from the western population contribute significantly to Mediterranean and other eastern Atlantic fisheries. Of greater concern is that adolescents from the western population show similar eastward dispersive behaviors across the 45°W management boundary. If this occurs at rates observed here for eastern adolescents, the smaller, less productive western population will be disproportionately affected by higher fishing rates in the eastern management zone.

The disparity between the eastern and western population sizes and the continued decline of the western stock suggests that some added level of protection is needed to ensure the sustainability of the smaller western component. Natal homing rates reported here were remarkably high to both regions and clearly show that the contribution of eastern adults to the western spawning area is inconsequential. Thus, spawning adults in the Gulf of Mexico appear to be entirely of western origin, and this region should be given high priority for conservation. High connectivity between foraging areas in the Gulf of Maine/Gulf of St. Lawrence and the Gulf of Mexico was also observed, signifying that this region of the northern Atlantic represents critical refugia for western giants. Due to the condition of the western population, a more conservative rate of exploitation of bluefin tuna, inclusive of eliminating bycatch in the Gulf of Mexico, will be required for the recovery of this population.

References and Notes
8. Materials and methods are available as supporting material on Science Online.
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Supporting Online Material
www.sciencemag.org/cgi/content/full/3161473/DC1 Materials and Methods
Fig. S1
References
SOM Data
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Glia Are Essential for Sensory Organ Function in C. elegans
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Sensory organs are composed of neurons, which convert environmental stimuli to electrical signals, and glia-like cells, whose functions are not well understood. To decipher glial roles in sensory organs, we ablated the sheath glial cell of the major sensory organ of Caenorhabditis elegans. We found that glia-ablated animals exhibit profound sensory deficits and that glia provide activities that affect neuronal morphology, behavior generation, and neuronal uptake of lipophilic dyes. To understand the molecular bases of these activities, we identified 298 genes whose messenger RNAs are often associated with glia, such as retinal pig-

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