Parasitism, space constraints, and gonad asymmetry in the pollan (Coregonus autumnalis)

Chris Harrod and David Griffiths

Abstract: Both sexes of Lough Neagh pollan (Coregonus autumnalis) show directional asymmetry in gonad mass, the left being larger than the right in 70% of individuals. The frequency and degree of asymmetry are size independent in males but become progressively more marked in females of >120 g somatic mass. There is little evidence to support the hypothesis that the size of the body cavity constrains gonad size. Pollan are heavily and size-dependently infected by metacercariae of the trematode Ichthyocotylurus erraticus. Total gonad mass in both sexes of pollan is negatively correlated with infection intensity of this parasite. In females, mean egg size shows no trend but fecundity declines and the degree of asymmetry in ovary mass increases with infection intensity up to a threshold, beyond which it remains constant. While directional asymmetry in gonad size is strongly developmentally controlled in many fish species, variation in the degree of gonad asymmetry in pollan provides a potentially sensitive indicator of stress.

Résumé : Les deux sexes du pollan (Coregonus autumnalis) au Lough Neagh présentent une asymétrie directionnelle de la masse de leurs gonades, la gauche étant plus grande que la droite chez 70 % des individus. La fréquence et l’importance de l’asymétrie sont indépendantes de la taille chez les mâles, mais elles deviennent progressivement plus marquées chez les femelles de masse somatique >120 g. Il y a peu d’indications pour appuyer l’hypothèse selon laquelle la dimension de la cavité corporelle restreint la taille des gonades. Les pollans sont fortement infectés par les métacercaires du trématode Ichthyocotylurus erraticus et l’infection varie en fonction de la taille. Il y a une corrélation négative entre la masse totale des gonades chez les pollans et l’intensité de l’infection par ce parasite. Chez les femelles, la taille moyenne des œufs ne suit aucune tendance particulière; en revanche, la fécondité décroît et l’importance de l’asymétrie de la masse ovarienne augmente en fonction de la gravité de l’infection jusqu’à une limite supérieure au-delà de laquelle elles demeurent constantes. Bien que l’asymétrie directionnelle de la taille des gonades soit fortement contrôlée par le développement chez plusieurs espèces de poissons, la variation de l’importance de l’asymétrie des gonades chez les pollans fournit un indicateur potentiellement sensible du stress.

Introduction

Asymmetries in paired structures are subject to both genetic and environmental influences. In some species, the asymmetry is largely under developmental or genetic control and extreme. For example, there is no left testis in most Oryzias species (Hamaguchi 1996), the left ovary has more or less disappeared in mbuna, the rock-frequenting group of cichlid species in Lake Malawi (Fryer and Iles 1972), and only the right ovary is functional in six genera of cartilaginous fishes and only the left in two genera (Dodd 1983). Other asymmetries are less extreme and have attracted much attention recently because they have been shown to be influenced by environmental factors (Clarke 1995; Møller 1997). Fluctuating asymmetry has most frequently been examined as a measure of environmental stress, but variation in the degree of directional asymmetry has also been linked to stress (Sasal and Pampoulie 2000). However, directional asymmetry in internal structures such as gonads, in a body cavity of finite size, could also be a consequence of space constraints (Brown and Kamp 1942; MacGregor 1968). The distended abdomens characteristic of gravid female fish close to spawning time suggest that abdominal space might limit ovary size and affect symmetry.

Directional gonad asymmetry has been observed in whitefish (Coregonus lavaretus) (Heese 1990), salmonids (Oncorhynchus spp.) (Rounsefell 1957; A. Hendry, Department of Biology, McGill University, Montréal, QC H3A 2K6, Canada, unpublished data), Atlantic salmon (Salmo salar) (C. Harrod, personal observation), brown trout (Salmo trutta) (Brown and Kamp 1942), and other fishes, e.g., anguillid eels (Tesch 2003), anchovies (Anchoa naso, Engraulis mordax, and Engraulis encrasius) (MacGregor 1968; Vallisneri and Scapolatempo 2000), capelin (Mallotus villosus) (Winters 1971), and albacore (Thunnus alalunga) (Ramón and Bailey 1996). In most species, the left gonad is larger than the right.

Asymmetry in paired fins has been correlated with reproductive effects (Hecht et al. 2000; Wedekind and Müller 2004) and with parasitism (Reimchen 1997; Sasal and Pampoulie 2000). Parasitism can affect fish physiology and...
reproductive potential (e.g., Arme and Owen 1968; Pampoulie et al. 1999; Tolonen et al. 2002), but parasitism has not been associated with gonad asymmetry.

Pollan (Arctic cisco in North America) (Coregonus autumnalis) occur as glacial relicts in Ireland, far outside their main high Arctic range, and are subject to a variety of environmental stressors, both external and internal (Harrod et al. 2001, 2002; Harrod and Griffiths 2005). Almost all adult fish in Lough Neagh (99.9% of fish older than 1+, n = 2126) are infected with metacercariae of a trematode, Ichthyocotylurus erraticus, at densities (median infection intensity, i.e., the number of parasites per fish, 536 in fish 2+ and older) considerably greater than those shown to cause death or reduced condition in other fish species (Harrod and Griffiths 2005). The pericardial aggregations of the parasite probably affect heart function, and this, coupled with low oxygen concentrations in hypertrophic Lough Neagh, could reduce fitness by affecting survival, condition, or reproduction. Females are more heavily infected than males. 

Ichthyocotylurus erraticus affects somatic condition and liver size in pollan (Harrod and Griffiths 2005) and condition, liver, and reproductive indices and heart mass in C. lavaretus in Loch Lomond (Dolezel and Crompton 2000).

Harrod and Griffiths (2004) described the reproductive biology of pollan in Lough Neagh, the only stock to have been examined in detail. This paper describes patterns of gonad asymmetry in pollan and investigates to what extent space constraints and parasitism are correlated with these patterns. We find no support for the space constraints hypothesis but demonstrate differences in the pattern of asymmetry between male and female fish and show that asymmetry in females is linked to infection intensity.

Materials and methods

Pollan were sampled from Lough Neagh (6°25′W, 54°35′N) using surface- and bottom-set multipanel gill nets on five occasions from September to early December 1999, when fish normally spawn. Fish were aged from scales following the procedure of Wilson and Pitcher (1984). The contents of the body cavity were divided into gonads, liver, and alimentary canal and each weighed separately to the nearest 0.01 g. Sex was assessed through visual inspection of the gonads. Left and right gonads of 136 female and 94 male fish were weighed individually. Ripe ovaries (maturity stage 5; Lagler 1978) were stored individually in Bouin’s fixative prior to measurement of egg size and fecundity. Pollan hearts and any encysted Ichthyocotylurus metacercariae found in the pericardial cavity were stored in 80% alcohol. To measure infection intensity, cysts were teased from the heart and host reactive tissue using mounted needles and this material squeezed between microscope slides before counting cysts under a low-power microscope.

Asymmetry was measured as the signed difference between left and right gonad mass, the signed difference expressed as a percentage of the left gonad mass or the ratio of left to right gonad mass. Most analyses were restricted to fish with ripe gonads (stage 5). When investigating the effect of parasitism, the analysis was restricted to fish collected shortly before spawning (November) for which all three measures of reproductive potential (i.e., ovary size, egg size, and fecundity) were available.

Asymmetry might vary with fish size (Somarakis et al. 1997). Two measures of fish size were used to detect effects on gonads and on the body cavity contents. When examining relationships between body cavity components, gutted mass (i.e., total body mass minus body cavity content mass) was used as the independent variable, while in investigations of possible parasite effects on gonads, somatic mass (i.e., total body mass minus gonad mass) was used. Regressions of right versus left gonad mass were carried out: this tests for a fish size effect, since gonad size is correlated with fish size (see below). Slopes of log-transformed gonad mass regressions significantly different from 1 indicate that there is a body size effect, while intercepts significantly different from 1 indicate asymmetry. To show nonlinear trends, locally weighted scatterplot smoothing (LOWESS) was used. To test the possibility, suggested by the data, that the degree of asymmetry (Y) changes above a threshold body size or infection intensity (X), linear and piecewise (Toms and Lesperance 2003) models

\[ Y = a + bX \]

\[ Y = a + b_1X + b_2(X - T) \]

were fitted to the data using nonlinear least squares regression. The terms in parentheses in the piecewise model control where the change in slope (T) occurs. The small-sample Akaike’s Information Criterion (AICc) was calculated for each model because the ratio of number of observations to number of estimated parameters was appreciably below 40 for most data sets. Model fits were then compared with AICc, Akaike weights, and evidence (Akaike weight) ratios (Burnham and Anderson 1998). Since gonad size, infection intensity, and degree of asymmetry varied with body size and age, the effects of the latter were controlled for by partial correlation. All errors are standard errors.

Results

Paired t tests showed left gonad masses to be significantly larger than right in both sexes (Table 1): the percent difference was much greater in females. Note, however, that the right gonads were larger than the left in about one third of the fish (Fig. 1). In the absence of estimates of measurement error, we assumed that gonads were asymmetric if they differed in mass by more than 10%: about 70% of fish showed differences of this magnitude or greater. The distribution of differences of left and right gonad masses in female or male fish in gonad state 5 showed no significant skewness or kurtosis.

The pattern of asymmetry differs between the sexes (Table 2): the degree of gonad asymmetry was size dependent in females but not in males. Note that this conclusion does not depend on whether ordinary least squares or reduced major axis regression is used. The frequency of ovarian asymmetry was also size dependent. Only half of the 20 smallest fish examined (maximum somatic mass 129 g) had larger left ovaries, but this had risen to 85% for the 20 largest fish (minimum somatic mass 177 g) and one fish had symmetrically ovaries.
Size-linked asymmetry was much more marked in ripe fish but was still significant in fish with late-developing ovaries (Fig. 2). However, the degree of asymmetry, when expressed as a percentage of left gonad mass, did not differ between fish of a given somatic mass in the two gonad states (analysis of covariance: slopes, $F_{1,190} = 0.44$; intercepts, $F_{1,191} = 2.52$). For fish with ripe ovaries, piecewise regression gave a better fit than the linear model (AICc = 3.90). The Akaike weights (0.87 and 0.12) show that the piecewise model is seven times more likely to be the better model.

Ovarian asymmetry did not appear until fish reached a somatic mass of 120 ± 12 g, equivalent to a fork length of 235 mm (age 2+ fish).

The ratio of left to right gonad size varied with somatic mass, but the slope and intercept of this relationship did not depend on sampling date (slopes, $F_{4,100} = 1.23$; intercepts, $F_{4,191} = 2.52$). For fish with ripe ovaries, piecewise regression gave a better fit than the linear model (AICc = 3.90). The Akaike weights (0.87 and 0.12) show that the piecewise model is seven times more likely to be the better model. Ovarian asymmetry did not appear until fish reached a somatic mass of 120 ± 12 g, equivalent to a fork length of 235 mm (age 2+ fish).

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Similarly, there were no significant partial correlations between right or left ovary masses or the size difference in ovary mass and combined gut and liver mass when gutted mass was controlled (i.e., there was no evidence of a body cavity size constraint).

Ichthyocotylurus numbers increased with gonad size in both sexes: the slopes and intercepts did not differ between the sexes. Partial correlations were calculated to control for the effect of somatic mass. Gonad mass declined with increasing Ichthyocotylurus numbers in both sexes (male partial $r = -0.38$, 26 df, $P < 0.05$; female partial $r = -0.72$, 17 df, $P < 0.001$) (Fig. 3). Ichthyocotylurus numbers were not correlated with gonad asymmetry in males, but females were affected: the difference between right and left gonad masses expressed in absolute (or percentage) terms increased with the number of parasites below the breakpoint of 855 (±213) parasites per fish ($r = 0.58$, 26 df, $P < 0.01$, $n = 25$) and the intercept was not significantly different from 0 ($a = 0.90 ± 0.88$) (i.e., uninfected fish had symmetrical ovaries). Increasing ovarian asymmetry with infection intensity in fish with less than 855 Ichthyocotylurus was not related to fish size or age. There was no

### Table 1. Comparison of left and right gonads of pollan (*Coregonus autumnalis*) in maturity stage 5.

<table>
<thead>
<tr>
<th>Sex</th>
<th>Mean left gonad mass (g)</th>
<th>Mean right gonad mass (g)</th>
<th>% difference</th>
<th>n</th>
<th>t</th>
<th>P</th>
<th>% fish with left &gt; right</th>
</tr>
</thead>
<tbody>
<tr>
<td>Female</td>
<td>12.85</td>
<td>11.43</td>
<td>11.1</td>
<td>76</td>
<td>4.10</td>
<td>&lt;0.001</td>
<td>68</td>
</tr>
<tr>
<td>Male</td>
<td>1.08</td>
<td>1.01</td>
<td>6.5</td>
<td>60</td>
<td>2.51</td>
<td>0.02</td>
<td>68</td>
</tr>
</tbody>
</table>

*Note:* The paired $t$ tests were performed on log-transformed data but the reported masses are the back-transformed values.

### Table 2. Summary of regression statistics of log right pollan gonad mass against log left gonad mass.

<table>
<thead>
<tr>
<th>Sex</th>
<th>$a \pm SE$</th>
<th>$t$ ($a = 0$)</th>
<th>$P$</th>
<th>OLS $b \pm SE$</th>
<th>$t$ ($b = 1$)</th>
<th>$P$</th>
<th>RMA $b$</th>
<th>$n$</th>
</tr>
</thead>
<tbody>
<tr>
<td>Female</td>
<td>0.184±0.057</td>
<td>3.23</td>
<td>&lt;0.01</td>
<td>0.788±0.051</td>
<td>4.10</td>
<td>&lt;0.001</td>
<td>0.901</td>
<td>76</td>
</tr>
<tr>
<td>Male</td>
<td>–0.025±0.012</td>
<td>2.12</td>
<td>&lt;0.05</td>
<td>0.871±0.079</td>
<td>1.64</td>
<td>&gt;0.10</td>
<td>1.058</td>
<td>60</td>
</tr>
</tbody>
</table>

*Note:* OLS, ordinary least squares; RMA, reduced major axis.

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**Fig. 1.** Histograms of left to right gonad mass ratios for (a) male and (b) female pollan (*Coregonus autumnalis*).
change in the degree of asymmetry for fish with larger infection intensities (mean difference in ovary mass = 18.7% ± 3.8%).

Mean egg size (diameter) did not differ between left and right ovaries (paired t test: t = 0.52, 61 df). Egg size increased with fish mass and infection intensity (r = 0.58, 35 df, P < 0.001, and r = 0.43, 35 df, P < 0.01, respectively), but after controlling for mass, there was no parasite effect (partial r = 0.17). Egg size was independent of fish size for any age of fish (i.e., egg size was not related to growth rate). There was no correlation between skewness in egg size of left and right ovaries (r = 0.16, P > 0.05, n = 116). Skewness increased as ovaries developed (r = 0.48, P < 0.001, n = 116): skewness was correlated with gonad mass when somatic mass was controlled for (partial r = 0.52, 88 df, P < 0.001). However, skewness was not related to infection intensity after controlling for somatic mass (partial r = -0.06, 66 df).

Fecundity declined slowly and egg size increased rapidly as spawning date approached (log fecundity = 2.50 – 0.0011 ± 0.0004 × day + 0.79 ± 0.10 × log somatic mass (R² = 0.43, n = 84), log egg diameter = -1.18 – 0.0030 ± 0.0002 × day + 0.24 ± 0.05 × log somatic mass (R² = 0.77, n = 84), where day is the number of days since 1 January 1999). There was a significant negative correlation between egg size and fecundity after controlling for body size and day (partial r = -0.38, 80 df, P < 0.001). Fecundity declined with increasing infection intensity (Fig. 5) when the effect of somatic mass was controlled (partial r = -0.59, 34 df, P < 0.001).

Discussion

The literature contains a number of vague references to the effect that gonad asymmetry is not uncommon in fishes (e.g., MacGregor 1968; Sanwal and Khanna 1972; Ramon and Bailey 1996), but there is insufficient information to determine its frequency. In species in which both gonads develop, the ovary size ratio ranges from 8.8 in capelin (Winters 1971) to around 1.3 in trout, anchovies, and pollan (Brown and Kamp 1942; MacGregor 1968; Vallisneri and Scapolatempo 2000). The left gonad is usually the larger (11/14 species), suggesting some general constraint (whether physical or phylogenetic; Harvey and Pagel 1993). However, geographic variation in the degree (capelin, Winters 1971; albacore, Ramon and Bailey 1996) and even direction (sockeye salmon (Oncorhynchus nerka), Rounsefell 1957) of asymmetry has been reported, consistent with an environmental influence.

Pollan provide only very limited support for the space constraint hypothesis. The testes were more symmetrical than the much larger ovaries as predicted by this hypothesis, but the ovaries did not become more asymmetrical as they developed and spawning date approached (although absolute size differences did increase). In November, the ovaries constituted a median of 69.5% (n = 50) of the mass of the body cavity contents. MacGregor (1968) argued that since gonads are the last organs to develop in the body cavity, they will occupy whatever space is available. Brown and Kamp (1942) noted that gonad asymmetry in female S. trutta was associated with the anatomical configuration of the intestine, which, in its posterior region, bends to one side (usually the right). Pollan might be similarly affected, but we have no information on the orientation of the anterior portion of the intestine. While the left gonad was larger in most pollan, consistent with this explanation, the converse was true in about 30% of fish. There is probably little selective pressure on internal (gonad) asymmetry but strong pressure on external morphology, since this is likely to affect swimming ability. If this is correct, internal asymmetry in one organ would be expected to necessitate a compensatory asymmetry in other internal structures. However, there was no evidence of such compensation in pollan. If the other body cavity structures differ in their compressibility, one might expect the degree of asymmetry to change as ovaries develop, but we found no support for a temporal change in relative asymmetry. Furthermore, if, as our results suggest, egg number is largely determined early in the reproductive cycle and well before egg size (Kinnison et al. 2001), space constraints are unlikely to be important. The negative correlation between fecundity and egg size after controlling for body size is consistent with a space constraint. However, this seems unlikely to be the cause, since the correlation occurred both early (September) and late (November) in the reproductive cycle despite a threelfold increase in gonad mass over this period: the correlation seems more likely to reflect an energetic trade-off.

This study shows that variation in the degree of directional asymmetry is associated with parasite infection intensity in females but not in males. While asymmetry was independent of fish size and not affected by infection intensity in male pollan, total testis mass declined with increasing levels of parasitism. Total ovary mass was also correlated...
with parasitism, and asymmetry increased above a fish size threshold because infection intensity was itself fish size dependent. Fecundity declined strongly with increasing parasite numbers, but there was no asymmetry in egg size or evidence that infection intensity affected egg size. The extremely high infection intensities observed (median intensity 536 and range 5–3941 in fish 2 or more years old and with 10% of fish more than twice the median) suggest that *Ichthyocotylurus* has a minor effect on survival, although there are clear energetic effects, with both somatic and liver condition being reduced (Harrod and Griffiths 2005). However, the mechanism by which parasitism generates the gonad asymmetry is unknown. While it is tempting to conclude that parasitism by *Ichthyocotylurus* reduces fitness by affecting gonad size, we have insufficient information to estimate lifetime reproductive success and the correlations do not exclude the possibility that fish that are stressed for other reasons are also more susceptible to parasitism.

Pollan were also attacked, less frequently, by the small intestinal cestode *Proteocephalus exiguus* (45% of fish) and the ectoparasitic copepod *Argulus foliaceus* (0.3%). Infection intensity by *Proteocephalus* was far less than by *Ichthyocotylurus*, with no sign of gut distortion, and consequently, we have assumed that the latter had the major impacts. Lough Neagh supports large numbers of waterbirds, which are the final host of *Ichthyocotylurus*, and one would expect greater gonad symmetry in populations with lower infection intensities. Unfortunately, there is no information on relative gonad sizes in the other three, endangered, Irish populations or on the species in its normal, high arctic range.

One might expect reproductive traits to be more sensitive indicators of biotic stress than meristic traits. The latter are subject to effects throughout the life of the fish, whereas reproductive traits might well be determined in the early stages of development. However, gonad development only occurs in larger individuals and is probably determined on an annual, rather than a whole life, basis. The asymmetry in pollan gonads is consistent with this possibility: the majority of small fish showed relatively low levels of infection by *Ichthyocotylurus* (Harrod and Griffiths 2005) and symmetric gonads, but the intensity of infection increased dramatically for fish larger than 200 mm fork length. However, as the earlier discussion makes clear, gonad asymmetry is not always an indicator of stress.

We have demonstrated that variation in the degree of gonad asymmetry and in fecundity is associated with levels of biotic stress (parasite infection intensity) in one species. We suggest that such measures have the potential to act as sensitive indicators of stress. The study highlights the need for more systematic reporting of inter- and intra-specific varia-

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**Fig. 3.** (a) Pollan (*Coregonus autumnalis*) testis and (b) ovary masses decline with increasing numbers of *Ichthyocotylurus* after controlling for the effects of body size (log somatic mass).

**Fig. 4.** Change in absolute asymmetry in pollan (*Coregonus autumnalis*) ovary mass as a function of infection intensity.

**Fig. 5.** Pollan (*Coregonus autumnalis*) fecundity declines with increasing numbers of *Ichthyocotylurus* after controlling for the effects of body size (log somatic mass).
tion in relative gonad sizes and examination of the factors that influence that asymmetry.

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**References**


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