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Benoit Lebreton, Pierre Richard, Emmanuel Parlier, Gaël Guillou, Gérard  
Blanchard

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1 Trophic ecology of mullets during their spring migration in a  
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8 Benoit Lebreton<sup>a,1\*</sup>, Pierre Richard<sup>a</sup>, Emmanuel P. Parlier<sup>a,2</sup>, Gaël Guillou<sup>a</sup>, Gérard F.  
9 Blanchard<sup>a</sup>  
10  
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15 <sup>a</sup>: UMR Littoral, Environnement et Sociétés (LIENSs), UMR 6250 CNRS-Université de La  
16 Rochelle, Institut du Littoral et de l'Environnement, 2 rue Olympe de Gouges, 17000 La  
17 Rochelle, France  
18  
19  
20  
21

22 \* : Corresponding author: benoit.lebreton@univ-lr.fr, phone: + 33 (0)5 46 50 76 61, fax: + 33  
23 (0)5 46 50 76 63  
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25  
26

27 <sup>1</sup>: Present address: Harte Research Institute for Gulf of Mexico Studies, Texas A & M  
28 University – Corpus Christi, 6300 Ocean Drive, Corpus Christi, TX 78412- 5869, USA  
29  
30  
31

32 <sup>2</sup>: Present address: Parlier environnement SAS, Pôle technologique, 40 rue Chef de Baie,  
33 17000 La Rochelle, France  
34  
35  
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37 E-mail :

38 Pierre Richard: pierre.richard01@univ-lr.fr;  
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40 Emmanuel P. Parlier: emmanuel.parlier@yahoo.fr;  
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44 Abbreviations:

45 CPUE: Catch per unit effort

46 CPUE<sub>n</sub>: Number CPUE

47 CPUE<sub>m</sub>: Biomass CPUE

48 DTW: Digestive tract weight

49 G0: Young-of-the-year

50 G1: 1-year olds

51 G2: 2-year olds

52 G3+: 3-year-old and older

53 IR: Instantaneous ration

54 SPOM: Suspended particulate organic matter

55 SSOM: Surface sediment organic matter

56 TW: Total fish weight

57 VI: Vacuity index

58 Gaël Guillou: gael.guillou@univ-lr.fr;  
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## ABSTRACT

Mullet populations are abundant in littoral waters throughout the world and play a significant role in organic matter fluxes. Mulletts are opportunistic feeders: adults have frequently been shown to feed on primary producers (e. g. fresh or detrital plant material, microphytobenthos) but they may also feed on meiofauna. The population structure and stomach contents of mulletts that colonize salt marsh creeks in Aiguillon Bay (French Atlantic coast) were studied to determine if they use salt marshes as a feeding ground in spring. Stable isotope analyses were carried out on mulletts sampled to assess their diet during their spring migration. The mullet population was primarily composed of young-of-the-year (G0), 1-year old (G1) of both *Liza ramada* and *Liza aurata* species and 3-year-old or older (G3+) *L. ramada* individuals. G0 and G3+ population densities increased during the spring period: catch per unit effort (CPUE) increased from 0.22 to 1.49 ind.min<sup>-1</sup> for the G0 age group; but stomach content analyses revealed that only G1 and G3+ feed in the salt marsh. Isotopic signatures of G1 (spring:  $\delta^{13}\text{C}$ : -14.8‰,  $\delta^{15}\text{N}$ : 14.1‰) and G3+ mulletts (spring:  $\delta^{13}\text{C}$ : -16.9‰,  $\delta^{15}\text{N}$ : 13.8‰) indicate that mullet growth is supported largely by primary consumers, such as benthic meiofauna or small macrofauna. Mulletts are thus positioned at a much higher trophic level than true primary consumers.

## KEY WORDS

Mugilidae; Salt marsh; Diet; Stable isotope analysis; Stomach content; Food web; European Atlantic coasts

## 1. INTRODUCTION

Mulletts are important members of fish communities in estuarine and littoral waters throughout the world. The study of their feeding activity is of particular interest for a better understanding of ecosystem functioning because they are among the few large fish able to feed directly on the lowest trophic levels (Hickling, 1970; Odum, 1970; Laffaille et al., 2002; Gautier and Hussenot, 2005). They inhabit numerous habitats in the littoral zone: mudflats (Almeida et al., 1993), salt marshes (Lefeuvre et al., 1999; Laffaille et al., 2002) as well as estuaries and even rivers (Lam Ho , 1969; Keith and Allardi, 2001; Cardona, 2006). Depending on their habitat, mulletts can have many feeding strategies (e. g. water filtering,

1 sediment scraping...) (Bruslé, 1981; Almeida et al., 1993; Cardona, 1995) to exploit the most  
2 accessible food resources, in terms both of quantity and quality (Odum, 1968; Almeida et al.,  
3 1993). Due to their high biomass in European salt marshes, (Laffaille et al., 2000; Parlier et  
4 al., 2006), mullets play a major role in coastal food webs as biotic vectors of organic matter  
5 between littoral habitats and the open sea (Lefeuvre et al., 1999; Laffaille et al., 2002) and  
6 also as competitors with other consumers for food resources (Lasserre et al., 1976).  
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10 On European Atlantic coasts, mullets spend their reproduction period off-shore, on the  
11 continental shelf. Spawning occurs at different periods depending on location and species:  
12 from August to February for *L. aurata* and from fall to winter for *L. ramada* (Keith, 2001).  
13 Recruitment also varies in coastal areas depending on species and generally lasts from the end  
14 of winter through summer (Hickling, 1970; Gautier and Hussenot, 2005). During that period,  
15 young-of-the-year shift from a planktonic to a benthic diet (Albertini-Berhaut, 1973, 1974;  
16 Ceccherelli et al., 1981; Ferrari and Chieregato, 1981).  
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23 Adult mullets also migrate in early spring to the very littoral areas where they perform  
24 daily trophic migrations. They can exploit the full extent of the mudflats, including the lower  
25 part of salt marsh, they can reach via the numerous creeks and channels that fill up at every  
26 high tide (Laffaille et al., 2000; Parlier et al., 2006). Stomach content analyses tend to show  
27 that mullets have a limno-benthophagous feeding mode in these areas: when foraging, they  
28 scrap the superficial sediment layers and ingest a mix of sediment and organic matter. Food  
29 selection is achieved by the mechanical (gill-rakers) and gustatory elements (Bruslé, 1981).  
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36 Trophic ecology of mullets has been most often studied through stomach content analysis.  
37 Although it provides considerable information on fish food resources, stomach content  
38 analysis only reveals what has been recently ingested. Stable isotope analyses of tissues are  
39 complementary to stomach content studies since they allow the determination of what is  
40 actually assimilated (Pinnegar and Polunin, 1999). Stable isotope analyses can also be used to  
41 assess changes of diet during migrations (Fry, 2006). The distinction between ingested and  
42 assimilated resources is important for mullets because their limno-benthophagous feeding  
43 strategy leads them to ingest a large variety of available food items of different degrees of  
44 digestibility, including plants or detrital matter (Odum, 1970; Bruslé, 1981), microalgae  
45 (Almeida et al., 1993; Laffaille et al., 2002), but also meiofauna (Ezzat, 1963; Lasserre et al.,  
46 1976; Laffaille et al., 1998) and small macrofauna (Riera et al., 1999; Bouchard and Lefeuvre,  
47 2000; Quan et al., 2007). Assimilation processes are related to the quality, and thus the  
48 digestibility of ingested food items, with better assimilation rates for fresh tissues or tissues  
49 with higher nitrogen or nutrient content (Tenore, 1983; Mann, 1988; Cebrián, 1999).  
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1 This study was conducted to determine the origin of food sources and the trophic level of  
2 mullets. It has been carried out in the salt marshes of the Aiguillon bay, an intertidal bay on  
3 the French Atlantic coast, in which salt marshes are in the same type of these which have  
4 previously been examined using stomach content analyses (Laffaille et al., 1998; Laffaille et  
5 al., 2002). Mullet abundance is high in this bay during the spring (Parlier et al., 2006), when  
6 mullets colonize salt marsh creeks after a winter stay in off-shore waters (Laffaille et al.,  
7 1998). In this study, the population structure of mullets was determined to define which age  
8 groups colonize the salt marsh creeks. The feeding activity of each age group entering and  
9 leaving the creeks was analyzed by comparing stomach content weight between flood and ebb  
10 tides. The instantaneous and long-term diets were determined using stable isotope analyses of  
11 stomach content and muscle tissues, respectively.  
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## 22 **2. MATERIALS AND METHODS**

### 23 **2.1 Study site**

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25 Aiguillon Bay, located on the French Atlantic coast, is an bay of 56 km<sup>2</sup> in area and  
26 includes 38 km<sup>2</sup> of bare mudflats and 18 km<sup>2</sup> of salt marshes (Fig. 1) (Verger, 2005). This bay  
27 is a semi-diurnal macrotidal system, which causes relatively strong tidal currents (average  
28 0.2-0.6 m.s<sup>-1</sup>) (SHOM, 2001). The bay receives most of effluents of the Marais Poitevin, the  
29 second largest wetland in France. Freshwater inputs come from the Sèvre Niortaise River, the  
30 Lay River and many channels, including the largest, the Curé Channel. Watershed inputs  
31 occur mainly from autumn to spring (Meunier and Joyeux, unpublished results). Salinity  
32 ranged from 4.5 to 35 during our spring sampling period.  
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41 Mulletts were collected in a tributary creek of the Curé Channel (Fig. 1). This creek drains  
42 a 10 ha watershed on the southern part of Aiguillon Bay (Parlier et al., 2006). The whole  
43 creek is integrated in the salt marsh part of the bay, dominated by halophytic plants, such as  
44 *Halimione portulacoides* (Meunier and Joyeux, unpublished results). No vegetation was  
45 present on creek banks, which were covered with wide patches of microphytobenthos. The  
46 sampling site (46°15'49 N, 01°07'09 W) was located 50 m upstream from the creek mouth  
47 and had a cross-section of 15 m wide and 5 m deep.  
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### 56 **2.2 Fish sampling**

57 Mulletts were regularly sampled during three non-consecutive weeks from the beginning  
58 of March to mid-April 2005 (Spring 1: March 8 to 14; Spring 2: March 24 to 30; Spring 3:  
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April 6 to 12). Sampling was performed during spring tides, with tidal ranges from 5.70 to 6.25 m as in the study of Parlier et al. (2006). A total of 22 fish samplings were carried out during the three sampling weeks to determine the population structure with precision and to obtain enough samples for stomach content and stable isotope analyses (muscle tissues and stomach contents). In order to compare stomach contents of mullets before and after their visit to the salt marsh creek, 16 fish samplings were performed at ebb tide and 6 at flood tide (Laffaille et al., 2002).

The whole mullet community was sampled by using a series of four successive nets in the creek (Laffaille et al., 1998; Parlier et al., 2006): three trammel nets (70 to 30 mm mesh size, 2 m high and 30 m long) to capture large individuals and a fyke net (4 mm mesh size, 5 m deep, 1.80 m high, 20 m long) to catch the smallest individuals. To ensure maximum adult capture rates, trammel nets were set out diagonally across the creek and spaced 10 m apart because adult mullets can avoid nets and thereby escape (Laffaille et al., 2002). For ebb tide sampling, nets were set out across the creek at high tide until the creek was totally drained. Nets were set up on the creek for periods of 20 minutes separated by intervals of 5 minutes; sampling was therefore efficient for 80% of the tide. Using this kind of protocol, the number and biomass of fish which migrate into the creek during one tidal cycle can be estimated (Laffaille et al., 1998). Moreover, with this regular sampling, mullets can be frozen quickly, thereby arresting digestion and avoiding regurgitation. For flood tide sampling, nets were set across the empty creek and were left until high tide. The sampling protocol was the same as for the ebb tide. All samples were stored in iceboxes in the field and deep frozen (-20°C) at the laboratory until analysis.

### 2.3 Population structure

All sampled mullets were measured (fork length) to the nearest mm. Mulletts of 50 mm length or more were identified to the species level using the identification keys of Farrugio (1977), Cambrony (1984), synthesized in Gautier and Hussenot (2005), and Keith and Allardi (2001). Mulletts were individually weighed to the nearest 0.01 g. Mulletts less than 50 mm long, for which identification is very difficult (Cambrony, 1984; Keith and Allardi, 2001), were not identified. These mullets were pooled into size classes to the nearest mm and mean individual weights were calculated for each size class ( $\pm 0.01$  g). Within species, age was defined based on growth curves from Le Dantec (1955) and Lam Hoï (1969), tables in Gautier and Hussenot (2005) and young-of-the-year growth curves for *Liza aurata* and *Liza ramada* from Aiguillon Bay (Parlier, unpublished results).

1 For all ebb samples (one sample corresponds to a period of 20 minutes), the number ( $N_i$ )  
2 and biomass ( $M_i$ ) of mullets per sample were defined for each species and each age group  
3 (Laffaille et al., 1998). Numbers of non-caught mullets ( $N_j$ ) during the 5 minute intervals  
4 between samples  $i$  and  $i+1$  were estimated by extrapolation using the moving average of  
5 samples ( $N_i$  and  $N_{i+1}$ ) with the formula:  $N_j = [[(N_i/t_i)+(N_{i+1}/t_{i+1})]/2]*t_j$  where  $t_i$  is the sampling  
6 time for sample  $i$  (min),  $t_{i+1}$ : sampling time for sample  $i+1$  (min) and  $t_j$  is the time between  
7 sample  $i$  and  $i+1$  (min). Biomass of non-caught mullets per sample ( $M_j$ ) was determined using  
8  $M_i$  and the same calculations. Total number ( $\sum N_i + \sum N_j$ ) and biomass ( $\sum M_i + \sum M_j$ ) of  
9 mullets were calculated per species and per age group for each ebb fish sampling. For each  
10 ebb fish sampling, catch per unit effort (CPUE), i.e. the number (CPUE<sub>n</sub>) or biomass  
11 (CPUE<sub>m</sub>) of mullets caught per minute of sampling, was calculated for each species and  
12 cohort using the formulas (Laffaille et al., 1998): CPUE<sub>n</sub> = ( $\sum N_i + \sum N_j$ )/( $\sum t_i + \sum t_j$ ) and  
13 CPUE<sub>m</sub> = ( $\sum M_i + \sum M_j$ )/( $\sum t_i + \sum t_j$ ).  
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## 25 2.4 Stomach content analysis

26 A total of 123 stomach contents of mullets less than 50 mm long were sampled among the  
27 three sampling weeks (flood tides,  $n = 55$ ; ebb tides,  $n = 68$ ). The stomach contents of these  
28 fish were too small to be weighted with enough accuracy, so whole digestive tracts were  
29 dissected under a stereomicroscope ( $\times 20$ ) and weighed. The ratio between digestive tract  
30 weight (DTW,  $\pm 0.0001$  g) and total fish weight (TW,  $\pm 0.0001$  g) was calculated to compare  
31 mullets sampled at flood tide and at ebb tide. Non-parametric unilateral Wilcoxon tests were  
32 used to test the differences in these ratios between flood tide and ebb tide ( $H_1$ : DTW/TW at  
33 flood tide < DTW/TW at ebb tide).  
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42 Digestive tracts of mullets of 50 mm long or more were regularly collected on mullets  
43 sampled at flood tide ( $n = 123$ ) and ebb tides ( $n = 131$ ). Percentages of empty stomachs  
44 (vacuity index, VI) were determined and compared between flood tide and ebb tide with Chi-  
45 square tests. The whole stomach contents were weighed ( $\pm 0.01$ g) and ratios between stomach  
46 content fresh weight and total fish fresh weight (instantaneous ration, IR) were determined.  
47 Differences in IR of mullets per cohort and species between flood tide and ebb tide were  
48 tested using non-parametric Wilcoxon tests.  
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## 56 2.5 Stable isotope ratio analysis

57 Samples of each age group and of each species for mullets >50 mm long were collected  
58 during the three sampling weeks. Stomach contents and white dorsal muscle tissues were  
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collected to determine the instantaneous and the long-term diets of mullets, respectively. Muscles were dissected with the highest care to avoid the presence of bone pieces. For smaller mullets (<50 mm long), the mass of the stomach contents was too small for the stable isotope analysis and fish analyses were performed on whole eviscerated and topped fish. All samples were freeze-dried and ground using a ball mill. Fish samples from mullets <50 mm and all stomach contents were acidified with 1 mol.L<sup>-1</sup> HCl to remove carbonates then dried in a dry bath at 60°C and ground again. Since acidification may have an effect on δ<sup>15</sup>N values, these samples were analyzed separately for δ<sup>13</sup>C (acidified samples) and δ<sup>15</sup>N (raw samples). No delipidation was performed on samples since no significant effect on isotope values was noticed during preliminary tests on the different fish class sizes.

Fish isotopic signatures were compared with signatures of food sources from Riera et al. (1999) and from Richard (Unpublished results). Other food sources were hand-picked directly from fresh stomach contents with a stereomicroscope and Dumont #55 forceps: nematodes (sample of 200 individuals), annelids (sample of 50 individuals) and plant detritus and treated as above.

Samples were analyzed using an EA-IRMS (Isoprime, Micromass, UK). Isotopic values were expressed in the δ unit notation as deviations from standards (Vienna Pee Dee Belemnite for δ<sup>13</sup>C and atmospheric N<sub>2</sub> for δ<sup>15</sup>N) following the formula: δ<sup>13</sup>C or δ<sup>15</sup>N = [(R<sub>sample</sub>/ R<sub>standard</sub>) - 1] × 10<sup>3</sup>, where R is <sup>13</sup>C/<sup>12</sup>C or <sup>15</sup>N/<sup>14</sup>N. The analytical precision for the measurement was 0.15‰ for carbon and 0.2‰ for nitrogen. Stable isotope signatures were compared with non-parametric Wilcoxon tests.

### 3. RESULTS

#### 3.1 Population structure

From the 16 ebb and the 6 flood fish samplings carried out, 1748 mullets were captured. Among individuals with fork length ≥50 mm, 131 *Liza ramada* and 59 *Liza aurata* were identified. Four age-groups were distinguished (Fig. 2, Table 1): young-of-the-year (G0), 1 year old mullets (G1), 2 year-old mullets (G2) and 3 year-old or older mullets (G3+), the G3+ group being exclusively composed of *L. ramada*.

G0 represented the main percentage (50.9 to 96.9%) on the population at each sampling and their mean CPUE<sub>n</sub> increased from 0.22 to 1.49 ind.min<sup>-1</sup> between Spring 1 and Spring 3 sampling weeks (Table 1). The G1 age group was dominated by *L. ramada* and represented the second most abundant age group. G2 individuals from either species were very rare. *L.*



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*ramada* G3+ represented a very small proportion of the population, with the highest G3+ CPUE<sub>n</sub> being 0.01 ind.min<sup>-1</sup>, corresponding to 0.9 to 1.2% of total densities. Nevertheless, their CPUE<sub>m</sub> accounted for the largest proportion of total population biomass, representing between 56.2 and 96.6%. Biomass proportions of other groups were extremely low except during Spring 1 sampling, when the sum of G1 accounted for 32.9% of the population biomass (Table 1).

### 3.2 Stomach content analyses

For <50 mm mullets, DTW/TW ratios were 8.6 ± 3.5 (n = 55) at flood tide and 8.7 ± 5.5 (n = 68) at ebb tide and there was no difference in whole digestive tract weights between mullets sampled at flood and ebb tides (p = 0.955). All of the 153 studied digestive tracts were orange-colored and only four presented light traces of sediment.

For ≥ 50 mm mullets, *L. ramada* G1 had lower VI at ebb tide, when they left the salt marsh creek, than at flood tide (Table 2). No significant differences in VI were observed between ebb and flood tides for *L. aurata* G1 and *L. ramada* G2 and G3+ groups. Numerous G1 individuals sampled at flood tide were indeed characterized by light food traces in the cardiac part of their stomach, explaining the lack of differences in VI between flood and ebb tides. Except *L. ramada* G2, age groups of both species presented much higher IR at ebb than at flood tide and very low IR values (ranging from 0 to 1%) were much more frequently observed at flood tides than at ebb tide (Table 2). At flood tide, the highest IR values were measured in *L. ramada* G3+. The number of *L. aurata* G2 stomachs collected was too low for VI and IR comparisons. All of the 202 full stomach contents were composed by a greenish-black mixture of sediment and organic matter.

### 3.3 Stable isotope ratios of stomach contents and muscle tissue

Food sources directly sorted from stomach contents had the following signatures: nematodes: δ<sup>13</sup>C: -15.9‰, δ<sup>15</sup>N: 11.7‰; annelids: δ<sup>13</sup>C: -15.3‰, δ<sup>15</sup>N: 12.2‰; plant detritus: δ<sup>13</sup>C: -15.6‰, δ<sup>15</sup>N: 9.4‰ (Fig. 3 and 4). Values reported in Riera et al. (1999) presented lower δ<sup>15</sup>N, particularly those of nematodes and annelids with a difference of up to 4.2‰. The mean isotope ratios for microphytobenthos from Richard (Unpublished results) and Riera et al. (1999) ranged from -16.6 to -14.4‰ for δ<sup>13</sup>C and from 4.5 to 6.3‰ for δ<sup>15</sup>N.

Isotope ratios of G0 ranged from -22.1 to -18.4‰ for δ<sup>13</sup>C and from 8.0 to 12.8‰ for δ<sup>15</sup>N (Fig. 3) and show a very large increase in δ<sup>13</sup>C and δ<sup>15</sup>N values between individuals of 20 to

30 mm in length. Mulletts longer than 30 mm in length presented  $\delta^{13}\text{C}$  and  $\delta^{15}\text{N}$  values approaching those of G1.

In spring,  $\delta^{13}\text{C}$  and  $\delta^{15}\text{N}$  values of *L. ramada* and *L. aurata* G1 ranged from -17.4 to -12.7‰ for  $\delta^{13}\text{C}$  and from 12.4 to 18.7‰ for  $\delta^{15}\text{N}$  (Fig. 3); they were not significantly different between species (Wilcoxon-tests,  $\delta^{13}\text{C}$ :  $p = 0.156$ ,  $\delta^{15}\text{N}$ :  $p = 0.734$ ).  $\delta^{13}\text{C}$  and  $\delta^{15}\text{N}$  values of G1 individuals were, on average, higher than those of G0 individuals (Wilcoxon-tests,  $\delta^{13}\text{C}$ :  $p < 0.001$ ,  $\delta^{15}\text{N}$ :  $p < 0.001$ ). Mean differences in  $\delta^{15}\text{N}$  between muscle and stomach content signatures were 2.7‰ for both species (Table 3). Stomach content signatures were similar between both species (Wilcoxon tests,  $\delta^{13}\text{C}$ :  $p = 0.607$ ,  $\delta^{15}\text{N}$ :  $p = 1.000$ ), ranging from -24.2 to -14.5‰ for  $\delta^{13}\text{C}$  and from 9.6 to 14.1‰ for  $\delta^{15}\text{N}$  (Fig. 4). Range of stomach content  $\delta^{13}\text{C}$  was wider and more depleted than the one of muscle tissue; nevertheless most of stomach content  $\delta^{13}\text{C}$  ranged from -18.0 to -14.0‰.

Signatures of G2 muscle tissue were highly variable in spring, particularly for  $\delta^{13}\text{C}$ , with values ranging from -25.4 to -10.5‰, and not significantly different between species (Wilcoxon tests,  $\delta^{13}\text{C}$ :  $p = 0.373$ ,  $\delta^{15}\text{N}$ :  $p = 1.000$ ).  $\delta^{15}\text{N}$  values (both species taken together) were similar to the ones of G1 individuals (Wilcoxon test,  $p = 0.905$ ) whereas G2  $\delta^{13}\text{C}$  values were much lower than G1  $\delta^{13}\text{C}$  (Wilcoxon test,  $p = 0.001$ ) (Table 3). The few stomach content samples from *L. ramada* G2 individuals showed stable isotope ratios similar to those of *L. ramada* G1 (Wilcoxon-tests,  $\delta^{13}\text{C}$ :  $p = 0.582$ ,  $\delta^{15}\text{N}$ :  $p = 0.727$ ). Differences between muscle and stomach content signatures were high, particularly for  $\delta^{15}\text{N}$  (3.2‰ for *L. ramada*). Only one stomach content from *L. aurata* was sampled in spring. There were no G2 individuals in the summer sample.

$\delta^{13}\text{C}$  of G3+ muscle tissues were similar to those of G2 individuals but lower than those of G1 (Wilcoxon tests, G2:  $p = 0.150$ , G1:  $p = 0.001$ ).  $\delta^{15}\text{N}$  of G3+ muscle tissues were similar to those of G2 and G1 individuals (Wilcoxon tests, G2:  $p = 0.632$ , G1:  $p = 0.507$ ). Almost all values ranged from -17.7 to -12.9‰ for  $\delta^{13}\text{C}$  and from 9.7 to 15.1‰ for  $\delta^{15}\text{N}$ . Only three individuals were outside of this range, with lower  $\delta^{13}\text{C}$  values (-19.3, -24.7 and -26.1‰) and very high  $\delta^{15}\text{N}$  values (17.2 and 18.0‰) for two of them (Fig. 3). G3+ stomach contents showed a very large range of signatures ( $\delta^{13}\text{C}$ : -22.6 to -15.3‰;  $\delta^{15}\text{N}$ : 9.3 to 13.5‰). No differences were observed between G3+ and G1 age groups (Wilcoxon-tests,  $\delta^{13}\text{C}$ :  $p = 0.063$ ,  $\delta^{15}\text{N}$ :  $p = 0.568$ ). Differences between G3+ muscle tissue and stomach contents were on average 0.4‰ for  $\delta^{13}\text{C}$  and 2.7‰ for  $\delta^{15}\text{N}$  (Table 3).

## 4. DISCUSSION

### 4.1 Which groups of mullet colonize salt marsh creeks?

Only two species of mullet, *Liza ramada* and *Liza aurata*, colonize the salt marsh. No specimens of *Chelon labrosus* were caught during our sampling, such as in a previous study in the same site (Parlier et al., 2006). *C. labrosus* has however been observed in similar embayments along European Atlantic coasts (Labourg et al., 1985).

G0 individuals, not identified to the species level during this study, were probably specimens of both species (Parlier et al., 2006). The large increase in G0 densities that occurred between March and April reflects the coastal recruitment of these species (Lam Hoï, 1969; Hickling, 1970; Labourg et al., 1985). The G0 age group was the most frequent age group of the mullet population. However, due to very low individual biomass, the G0 group only represented a very small proportion of the total biomass, which was largely dominated by G3+. High densities of the G1 age group in both *L. ramada* and *L. aurata* were also observed, but very few G2 mullets were observed. This population structure is similar to that observed by Parlier et al. (2006) in the same bay and of Laffaille et al. (2000, 2002) in the Mont Saint-Michel Bay in spring.

This pattern of population structure may have several explanations. First, mullets are known to be highly euryhaline at the G0 and G1 stages of development (Lam Hoï, 1969; Lasserre and Gallis, 1975; Shusmin, 1990). Thus, these individuals were able to withstand salinity variations that occurred in the salt marsh, particularly in spring. Second, G0 and G1 migration into creeks is probably related to the typical high tidal currents in Aiguillon Bay (SHOM, 2001). These currents may have facilitated the migration of these age groups in particular (Dame and Allen, 1996). This hypothesis is strengthened by the observation that the catches of G0 and G1 mullets in the creek were the greatest during the highest tidal current periods, at mid ebb tide.

The absence of *L. aurata* adults (G3+) is probably linked to the relatively low salinities sometimes observed in salt marsh creeks. Adults from this species are known to tolerate only very short periods of low salinity (Lam Hoï, 1969; Keith and Allardi, 2001; Gautier and Hussenot, 2005; Cardona, 2006). On the contrary, *L. ramada* adults are highly euryhaline, well-known for their migrations into freshwater habitats (Keith and Allardi, 2001), and are thus able to stay in salt marsh creeks during these low salinity periods. The colonization of *L. ramada* adults in these creeks can be considered opportunistic and probably related to their high dietary demand. Mulletts show an increase of feeding activity in spring (Hickling, 1970;

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Odum, 1970; Bruslé, 1981) that may be needed to improve overall condition after winter spawning.

Among the four age groups of mullets present in coastal areas, only some of them regularly colonized the salt marsh creeks of Aiguillon Bay in spring: G0, *L. ramada* and *L. aurata* G1 and *L. ramada* G3+. These observations support the view that this bay is an important habitat for juvenile fish, as are other European salt marshes (Mathieson et al., 2000; Cattrijsse and Hampel, 2006).

#### 4.2 Stable isotope ratios of food sources

$\delta^{13}\text{C}$  values of food sources collected in this work are generally in the range of those of previous studies carried out in the Aiguillon bay (Richard, unpublished results; Riera et al., 1999). Three groups are clearly distinguished: 1. food sources originating from neritic waters, i.e. suspended particulate organic matter (SPOM) and phytoplankton (with  $\delta^{13}\text{C}$  values ranging from -23 to -20‰), 2.  $^{13}\text{C}$ -enriched salt marsh sources, i.e. microphytobenthos and C4 halophyte plants, such as *Spartina maritima* (with  $\delta^{13}\text{C}$  from -17 to -12‰), and 3.  $^{13}\text{C}$ -depleted salt marsh sources, i.e. C3 halophyte plants, such as *Plantago maritima* (with  $\delta^{13}\text{C}$  lower than -24‰). Sediment surface organic matter (SSOM)  $\delta^{13}\text{C}$  value (-20.6‰ : Riera et al., 1999) indicates a mixing of depleted (C3) and enriched (C4) salt marsh plant detritus and of microphytobenthos.

Some of these primary producers are not ingested as fresh matter, but as partially degraded matter that may have different isotopic values. Plant detritus sampled in stomach contents had lower  $\delta^{13}\text{C}$  values than fresh plant material characterized in previous studies, likely because degraded material is mainly composed of less labile compounds, particularly lignin which is known to have lower  $\delta^{13}\text{C}$  than fresh material (Benner et al., 1987). High  $\delta^{15}\text{N}$  values observed in plant detritus were probably related to bacterial film on detritus surfaces, which increases  $^{15}\text{N}$  concentration due to trophic isotopic fractionation (Dijkstra et al., 2008). A similar trend has been observed between fresh and detrital material of *S. maritima* (Riera et al., 1999).

Nematodes and annelids sampled from stomach contents during this study and nematodes from Riera et al. (1999) had  $\delta^{13}\text{C}$  values in the range of those of microphytobenthos and C4 plant detritus, suggesting that these consumers exploit these salt marsh resources. However, nematodes and annelids from stomach contents showed relatively high  $\delta^{15}\text{N}$  values compared with values from Riera et al. (1999), with a difference of 4.2‰ for nematodes. This discrepancy suggests that ingested nematodes may not be representative of the *in situ*

1 population. A selection of prey occurs in fact when mullets forage in the superficial sediment  
2 layer because they use gill-rakers and taste buds to select the most appetizing food items  
3 (Bruslé, 1981).  
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#### 7 4.3 Differential role of the salt marsh as a feeding ground for mullet groups 8

9 The absence of increase of DTW/TW ratios for G0 between flood tide and ebb tide  
10 showed that G0 individuals probably feed continuously (Albertini-Berhaut, 1974), within and  
11 outside the salt marsh creeks. The very few traces of sediment noticed in stomach contents  
12 confirmed that these mullets had not yet shifted to a limno-benthophagous diet, excluding  
13 SSOM as a food resource.  $\delta^{13}\text{C}$  values of muscle tissue, which were intermediate to those of  
14 SPOM and phytoplankton, indicated that G0 individuals exploit food resources from neritic  
15 waters. The orange coloration of stomach content suggested that their diet was based on  
16 crustaceans, probably zooplankton, which was confirmed by the observation of five stomach  
17 content samples that showed zooplankton fragments (swim appendages, etc.) and small  
18 copepods.  $\delta^{15}\text{N}$  values of these mullets corresponded to a mean trophic isotopic fractionation  
19 of 2.3‰ for each of the two trophic levels from phytoplankton to mullets, via zooplankton,  
20 which is in the range of isotopic fractionations generally reported for sources and consumers  
21 at the base of the food web (Vander Zanden and Rasmussen, 2001). This confirms the trophic  
22 level of the G0 age group and that the diet of this age group is based on zooplankton  
23 (Albertini-Berhaut, 1973, 1974; Ceccherelli et al., 1981; Ferrari and Chierogato, 1981). So G0  
24 individuals do not depend on salt marsh food resources just after their migration from  
25 spawning areas to the salt marsh. These migrations are probably not active due to the weak  
26 swimming capacities of G0 compared to the strong tidal currents (SHOM, 2001).  
27 Nevertheless, the large ranges of  $\delta^{13}\text{C}$  and  $\delta^{15}\text{N}$  suggest that a diet shift occurs in G0 diet,  
28 from planktonic feeding to a limno-benthophagous diet, as previously demonstrated by  
29 Albertini-Berhaut (1973, 1974) and Ceccherelli et al. (1981). More generally, this trophic  
30 shift is also well-known in numerous species of mullets (Bruslé, 1981; Koussoropolis et al.,  
31 2010).  
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51 The very few G2 individuals observed in salt marsh creeks and the absence of differences  
52 for IR between flow and ebb tides demonstrated that these areas are not preferential feeding  
53 grounds for these fish, as also shown by Laffaille et al. (2000). The larger range of  $\delta^{13}\text{C}$   
54 values in muscle tissue confirms that these mullets have a large range of trophic behaviors  
55 and/or habitats, probably from terrestrial to oceanic areas. However, the few samples  
56 collected do not allow a clear understanding of G2 diet.  
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On the contrary, salt marsh resources are heavily used by G1 and G3 individuals. For G1 individuals, the strong increase in their IR when they were present in the salt marsh creeks indicates that they were feeding there. The quantity of food ingested before mullets migrate to the creeks is extremely low, demonstrating that salt marsh creeks are preferential feeding grounds for these mullets when they colonize these areas.

As for G1, the strong increase of *L. ramada* G3+ IRs between flood and ebb tides showed that these mullets migrate to salt marsh creeks for feeding purposes during high tides, as in Mont Saint-Michel Bay (Laffaille et al., 1998; Laffaille et al., 2002). However, contrary to G1, mean IR values at flood tide showed that numerous G3 individuals enter the creek with a partially full stomach. Retention time of food in the digestive tract of mullets ranges from 2 to 6 h (Bruslé, 1981), leading to the conclusion that adult mullets feed on the adjacent mudflat before they migrate into salt marsh creeks. The same behavior has been observed in Mont Saint-Michel salt marsh creeks (Laffaille et al., 2002). In this bay, mean observed IR values (close to 3% in March and to 7% in April) are clearly higher than those observed in spring in Aiguillon Bay (only 1.4%), indicating that mullets feed less intensively on Aiguillon Bay mudflats before migrating to the salt marsh.

Muscle and stomach content  $\delta^{13}\text{C}$  of G1 and G3+ individuals demonstrated the salt marsh origin of food resources (meiofauna, halophyte plant detritus, microphytobenthos...). The large quantities of sediment in stomach content confirmed the limno-benthophagous diet of G1 (*L. ramada* and *L. aurata*) and of G3+ (*L. ramada*), as in many other ecosystems (Hickling, 1970; Albertini-Berhaut, 1974; Bruslé, 1981; Ferrari and Chieragato, 1981; Almeida et al., 1993; Shapiro, 1998; Laffaille et al., 2002) and such as many other species (Bruslé, 1981; Koussoropolis et al., 2010). The numerous foraging traces on the sediment at ebb tides corroborated these observations (Laffaille et al., 2002). The large range of G1 and G3+ stomach content  $\delta^{13}\text{C}$  showed that they contained a mixture of  $^{13}\text{C}$ -enriched salt marsh sources and of  $^{13}\text{C}$ -depleted salt marsh plants, such as *Aster tripolium*, *Halimione portulacoides* or *Plantago maritima* which grow on this area (Verger, 2005) and generally have low  $\delta^{13}\text{C}$  values (Riera et al., 1999). Nevertheless the range of G1 and G3+ muscle, narrower and more enriched than these of stomach content, demonstrated that C3 halophyte plant material, even if it is ingested, is thus hardly assimilated by mullets. G3+ showed sometimes some variations of individual behavior. In spring, two mullets showed very low  $\delta^{13}\text{C}$  values (-24.7 and -26.1‰). These  $\delta^{13}\text{C}$  values, similar to typical terrestrial sources, indicated that these mullets probably spent the winter in river waters and have not carried out their catadromous migration for breeding. One other individual showed  $\delta^{13}\text{C}$  values typical of

1 oceanic food resources, indicating that this individual was sampled probably just after its  
2 migration to coastal areas after the spawning season.

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4 Thus G1 and G3+ clearly colonize intertidal salt marsh creeks for the purpose of feeding,  
5 as previously demonstrated for mullets in other salt marshes or even in rivers (Lam Hoï, 1969;  
6 Sauriau, 1990; Keith and Allardi, 2001; Laffaille et al., 2002; Almeida, 2003; Koussoropolis  
7 et al., 2010) and as for numerous other nektonic species (Deegan et al., 1990; Laffaille et al.,  
8 2001; Lefeuvre et al., 1999). Salt marshes are in fact well known for their role as feeding  
9 areas (Kneib, 1997). However, the macrotidal range of the Aiguillon and Mont Saint Michel  
10 Bays salt marshes gives them strong particularities because fishes migrate daily to these  
11 feeding areas (Lefeuvre et al., 1999). Colonizing salt marsh creeks is riskier for mullets  
12 (stranding, bird predation for G1) and represents higher energy loss than feeding on bare  
13 mudflats (Wolff et al., 1981; Gibson, 2003), where they are not likely to be food-limited.  
14 These risks and energetic costs of migrations may be counteracted by a maximization of G1  
15 and G3+ food intake in salt marsh creeks (Gibson, 2003).  
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#### 27 4.4 Mullet trophic level

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29 High  $\delta^{15}\text{N}$  values of muscles indicated a high trophic level, at least secondary, for G1 and  
30 G3+ individuals. As for the G0 age group, there was a good relationship between  $\delta^{15}\text{N}$  values  
31 of muscle tissue and stomach contents, with a mean trophic fractionation ranging from 2.7.  
32 This value matches Vander Zanden and Rasmussen's (2001) observations for high-level  
33 consumers. Stomach content signatures, close to those of primary consumers (nematodes,  
34 annelids), highlight that diet of G1 and G3+ individuals were partially based on these prey,  
35 which were observed in large quantities in stomach contents along with benthic diatoms and  
36 detrital matter. The high muscle  $\delta^{15}\text{N}$  values indicate that among the large diversity of food  
37 sources they ingest, from primary producers to animals, mullets incorporate more N from the  
38 most  $^{15}\text{N}$ -enriched food sources (e. g. meiofauna, small macrofauna). This can be due to a  
39 better assimilation of animal prey than of primary producers (e. g. C3 and C4 salt marsh  
40 plants) which have lower nutritional quality, and thus lower digestibility (Cebrián, 1999). It  
41 can also be due to the higher protein content of muscle tissue, which N isotope signature  
42 reflects thus more the isotope composition of animal preys than of plants, much less proteic.  
43 Primary producers have a high content of carbohydrates, mainly and readily used for animal  
44 metabolism, the fish muscle  $\delta^{13}\text{C}$  may therefore not completely reflect their actual  
45 assimilation.  
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1 Our study gives information on assimilated food sources and thus on more profitable salt  
2 marsh resources for mullets. It well complete works previously made on stomach contents  
3 (Almeida et al., 1993; Laffaille et al., 2002), which determined the food sources ingested by  
4 mullets, and on which animal preys have been regularly observed in stomach contents (Ezzat,  
5 1963; Lasserre et al., 1976; Laffaille et al., 1998), without necessarily determining their role  
6 in mullet resources. Although detritus and benthic diatoms constitute a large proportion of  
7 stomach contents, stable isotopes demonstrate that animal prey represent a very high  
8 contribution of resources incorporated into fish tissues when they migrate in the salt marshes.  
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10 High  $\delta^{15}\text{N}$  values have already been observed in mullets from a bare mudflat (Richard,  
11 unpublished results), but are slightly lower ( $13.2 \pm 0.7\%$  on average) than those observed in  
12 this study. Bacterivorous nematodes, that are  $^{15}\text{N}$ -enriched relative to herbivorous species  
13 (Rzeznik-Orignac et al., 2008), are likely to be present in higher proportions than herbivorous  
14 ones when detrital organic matter and thus bacteria are very abundant, as in salt marshes  
15 (Moens et al., 1999). This difference is also probably due to a larger part of meiofauna in the  
16 diet of mullets feeding in creeks. Nematode abundance can be several times higher in salt  
17 marshes covered with halophytic plants ( Heip et al., 1985; Giere, 1993) than in bare mudflats  
18 (Rzeznik-Orignac et al., 2003). The higher availability of more protein-rich food sources,  
19 such as meiofauna in the salt marsh creeks may explain why mullets migrate in the salt marsh  
20 creeks, when they can find enough food resources on the bare mudflat they pass through. For  
21 G3+ individuals, this feeding behavior may be induced by the strong increase in their  
22 nutritional requirements since they have a poor overall condition after the spawning period.  
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## 40 **5. CONCLUSION**

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42 Five groups of mullets migrate to salt marsh creeks for feeding purposes: *L. ramada* and  
43 *L. aurata* G0 after their trophic shift, *L. ramada* and *L. aurata* G1 and *L. ramada* adults. All  
44 fishes from these five groups are well known for their euryhaline behavior (Keith and Allardi,  
45 2001; Gautier and Hussenot, 2005; Cardona, 2006). These migrations for purpose of feeding,  
46 already well known in adults, are also realized by young-of-the-year, after their trophic shift,  
47 and by G1 but not by G2. During these periods of salt marsh colonization, mullets have a  
48 limno-benthophagous diet behavior and a diet mostly based on animal prey, particularly  
49 meiofauna, positioning mullets at a high trophic level. This assertion contrasts with the  
50 general knowledge on mullets, which are generally considered as primary consumers (Bruslé,  
51 1981; Almeida et al., 1993; Shapiro, 1998; Laffaille et al., 2002).  
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## FIGURE CAPTIONS

Fig. 1. Location of the Aiguillon Bay and sampling site.

Fig. 2. Size-frequency distributions and age groups of the two studied mullet species, *Liza aurata* (fork length >50 mm) and *Liza ramada* (fork length >50 mm) in spring 2005.

Fig. 3.  $\delta^{13}\text{C}$  and  $\delta^{15}\text{N}$  of mullet muscle tissue, potential food sources and prey. For mullet muscle tissues, each point represents one individual. Potential food sources and prey: Ann: annelids, Mpb: microphytobenthos, Nem: nematodes, Phy: phytoplankton, Pma: *Plantago maritima*, Sma: fresh *Spartina maritima*, Sma det: detritic *S. maritima*, SPOM: suspended particulate organic matter, SSOM: surface sediment organic matter, Pla det: plant detritus. Food source data taken from Richard (1998, unpublished results) are italicized, from Richard (2004, unpublished results) underlined, from Riera et al. (1999) are in normal face type, and those from the present study are in bold face type. Plotted G0 points are circled.

1 Fig. 4.  $\delta^{13}\text{C}$  and  $\delta^{15}\text{N}$  of mullet stomach contents at ebb tide, potential food sources and prey.  
2 For mullets, each point represents one individual. Potential food sources and prey: Ann:  
3 annelids, Mpb: microphytobenthos, Nem: nematodes, Phy: phytoplankton, Pma: *Plantago*  
4 *maritima*, Sma: fresh *Spartina maritima*. Sma det: detritic *S. maritima*, SPOM: suspended  
5 particulate organic matter, SSOM: sediment surface organic matter, Pla det: plant detritus.  
6 Food source data taken from Richard (1998, unpublished results) are italicized, from Richard  
7 (2004, unpublished results) underlined, from Riera et al. (1999) are in normal face type, and  
8 those from the present study are in bold face type.  
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## TABLES

Table 1. Fork lengths (FL, mean  $\pm$  standard deviation), catch per unit effort by number (CPUE<sub>n</sub>, mean  $\pm$  standard deviation) and by biomass (CPUE<sub>m</sub>, mean  $\pm$  standard deviation) of mullets sampled at ebb tide in spring 2005. Age groups: G0: young-of-the-year, G1: 1 year-old mullets, G2: 2 year-old mullets, G3+: 3 year-old and older mullets.

Species	Age group	Spring sampling week	n	Fork length mm	CPUE <sub>n</sub>		CPUE <sub>m</sub>	
					ind.min <sup>-1</sup>	Mean % per week	g.min <sup>-1</sup>	Mean % per week
<i>Liza sp.</i> (FL <50 mm)	G0	1	106	21 $\pm$ 1	0.22 $\pm$ 0.39	50.9	0.02 $\pm$ 0.02	0.5
		2	562	20 $\pm$ 2	1.12 $\pm$ 1.31	94.5	0.09 $\pm$ 0.10	0.7
		3	890	20 $\pm$ 2	1.49 $\pm$ 2.76	96.9	0.10 $\pm$ 0.19	1.0
<i>L. aurata</i> (FL >50 mm)	G1	1	34	80 $\pm$ 15	0.07 $\pm$ 0.09	17.4	0.42 $\pm$ 0.50	14.0
		2	10	81 $\pm$ 20	0.01 $\pm$ 0.01	0.8	0.04 $\pm$ 0.04	0.3
		3	13	78 $\pm$ 13	0.02 $\pm$ 0.02	1.0	0.06 $\pm$ 0.08	0.6
	G2	1	1	164	< 0.01 $\pm$ 0.01	0.7	0.18 $\pm$ 0.36	6.0
		2	1	150	< 0.01 $\pm$ < 0.01	0.2	0.10 $\pm$ 0.22	0.7
		3	0	-	0	0	0	0
<i>L. ramada</i> (FL >50 mm)	G1	1	63	77 $\pm$ 14	0.13 $\pm$ 0.09	29.7	0.57 $\pm$ 0.45	18.9
		2	21	77 $\pm$ 13	0.04 $\pm$ 0.02	3.4	0.23 $\pm$ 0.04	1.8
		3	11	81 $\pm$ 18	0.02 $\pm$ 0.03	1.0	0.11 $\pm$ 0.19	1.0
	G2	1	1	195	< 0.01 $\pm$ < 0.01	0.4	0.14 $\pm$ 0.27	4.5
		2	0	-	0	0	0	0
		3	2	173 $\pm$ 21	< 0.01 $\pm$ 0.01	0.3	0.20 $\pm$ 0.46	1.9
G3+	1	2	335 $\pm$ 56	< 0.01 $\pm$ 0.01	0.9	1.70 $\pm$ 3.40	56.2	
	2	15	401 $\pm$ 27	0.01 $\pm$ 0.01	1.2	12.61 $\pm$ 10.02	96.6	
	3	14	391 $\pm$ 40	0.01 $\pm$ 0.01	0.9	10.08 $\pm$ 10.75	95.5	



Table 2. Comparisons of vacuity indexes (%) and instantaneous feeding rations (IR, % mean  $\pm$  standard deviation) of *Liza aurata* and *L. ramada* at flood tide and at ebb tide in spring 2005. Age groups: G0: Young-of-the-year, G1: 1 year-old mullets, G2: 2 year-old mullets, G3+: 3 year-old and older mullets.

Species	Age group	Vacuity indexes			Instantaneous feeding rations			IR values < 1 %		n	
		Flood tide %	Ebb tide %	Khi <sup>2</sup> test p	Flood tide %	Ebb tide %	Wilcoxon test p	Flood tide %	Ebb tide %	Flood tide	Ebb tide
<i>L. aurata</i>	G1	34.0	27.8	0.537	0.75 $\pm$ 1.81	2.58 $\pm$ 2.30	< 0.001	81	36	53	36
	G2	-	-	-	-	-	-	-	-	2	1
<i>L. ramada</i>	G1	32.7	13.0	0.009	0.63 $\pm$ 1.08	3.41 $\pm$ 3.54	< 0.001	83	30	52	69
	G2	20.0	0.0	0.673	0.36 $\pm$ 0.33	3.02 $\pm$ 4.73	0.760	100	67	5	4
	G3+	0.0	4.6	0.528	1.39 $\pm$ 1.06	4.23 $\pm$ 2.90	0.002	45	14	11	22

Table 3.  $\delta^{13}\text{C}$  and  $\delta^{15}\text{N}$  (mean  $\pm$  standard deviation) of muscle tissue and stomach content of mullets in spring 2005 and difference ( $\Delta$ ) between muscle and stomach content signatures. Number of analyzed samples is shown in parentheses. Age groups: G0: young-of-the-year, G1: 1 year-old mullets, G2: 2 year-old mullets, G3+: 3 year-old and older mullets.

	Age group	Species	Muscle tissue ‰	Stomach contents ‰	$\Delta$ ‰
$\delta^{13}\text{C}$					
	G0	<i>Liza sp.</i>	-20.3 $\pm$ 0.9 (49)	-	-
	G1	<i>L. ramada</i>	-14.9 $\pm$ 1.3 (14)	-16.1 $\pm$ 1.6 (16)	1.2
		<i>L. aurata</i>	-14.1 $\pm$ 1.0 (9)	-17.0 $\pm$ 2.6 (15)	2.9
	G2	<i>L. ramada</i>	-17.6 $\pm$ 4.1 (9)	-16.2 $\pm$ 1.4 (3)	1.4
		<i>L. aurata</i>	-20.1 $\pm$ 3.4 (3)	-18.0 (1)	2.1
	G3+	<i>L. ramada</i>	-16.9 $\pm$ 3.3 (20)	-17.3 $\pm$ 3.9 (13)	0.4
$\delta^{15}\text{N}$					
	G0	<i>Liza sp</i>	9.8 $\pm$ 1.2 (27)	-	-
	G1	<i>L. ramada</i>	14.1 $\pm$ 1.5 (14)	11.4 $\pm$ 1.2 (9)	2.7
		<i>L. aurata</i>	14.1 $\pm$ 1.3 (9)	11.4 $\pm$ 0.5 (6)	2.7
	G2	<i>L. ramada</i>	13.9 $\pm$ 2.8 (9)	10.7 $\pm$ 1.5 (2)	3.2
		<i>L. aurata</i>	13.7 $\pm$ 2.8 (3)	10.9 (1)	2.8
	G3+	<i>L. ramada</i>	13.8 $\pm$ 2.2 (20)	11.1 $\pm$ 1.2 (11)	2.7

Figure 1

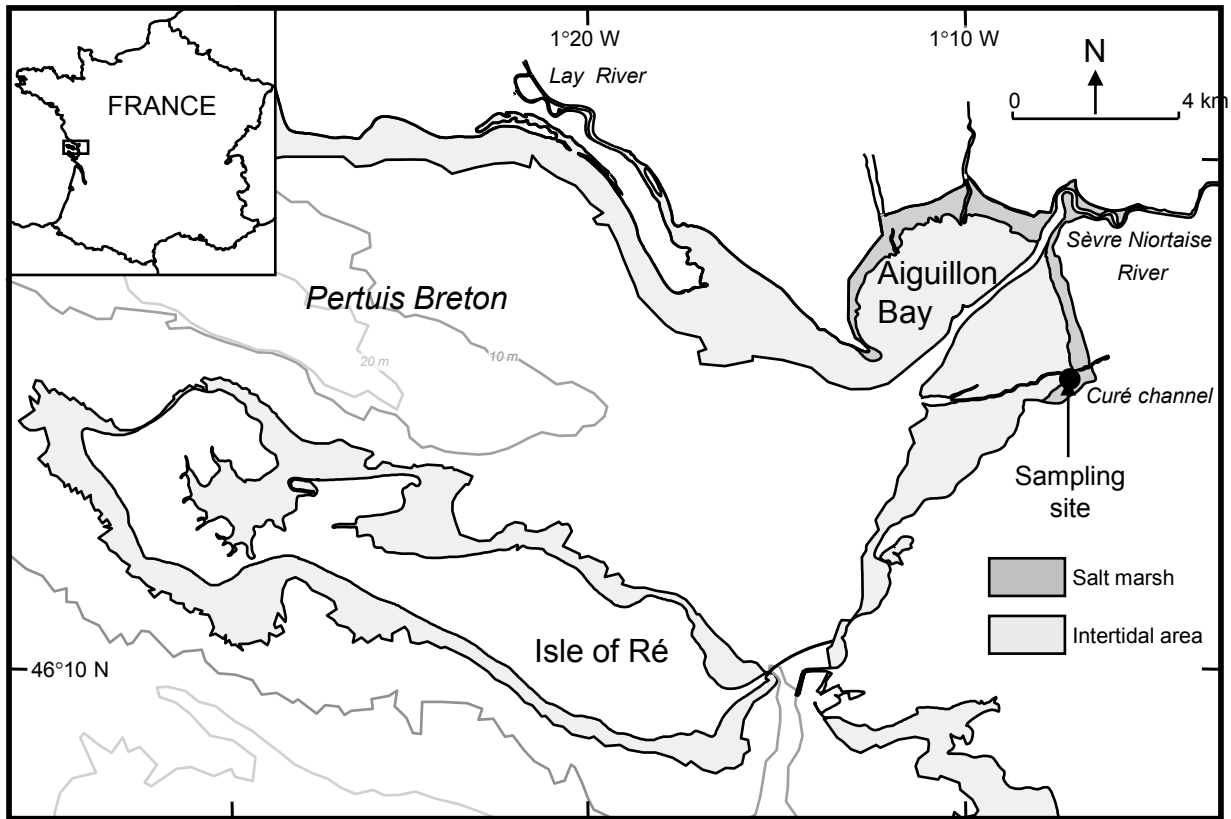


Figure 2

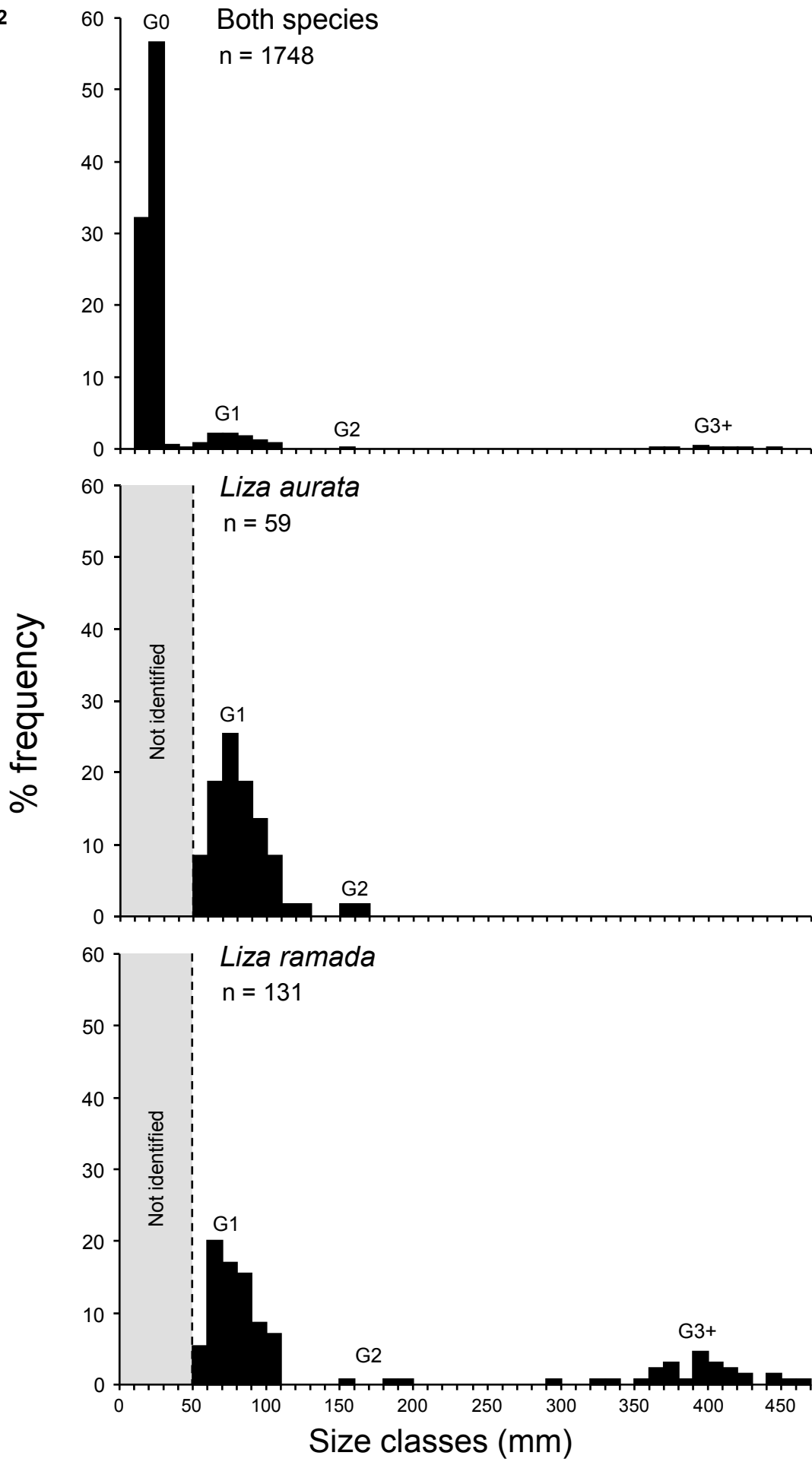


Figure 3

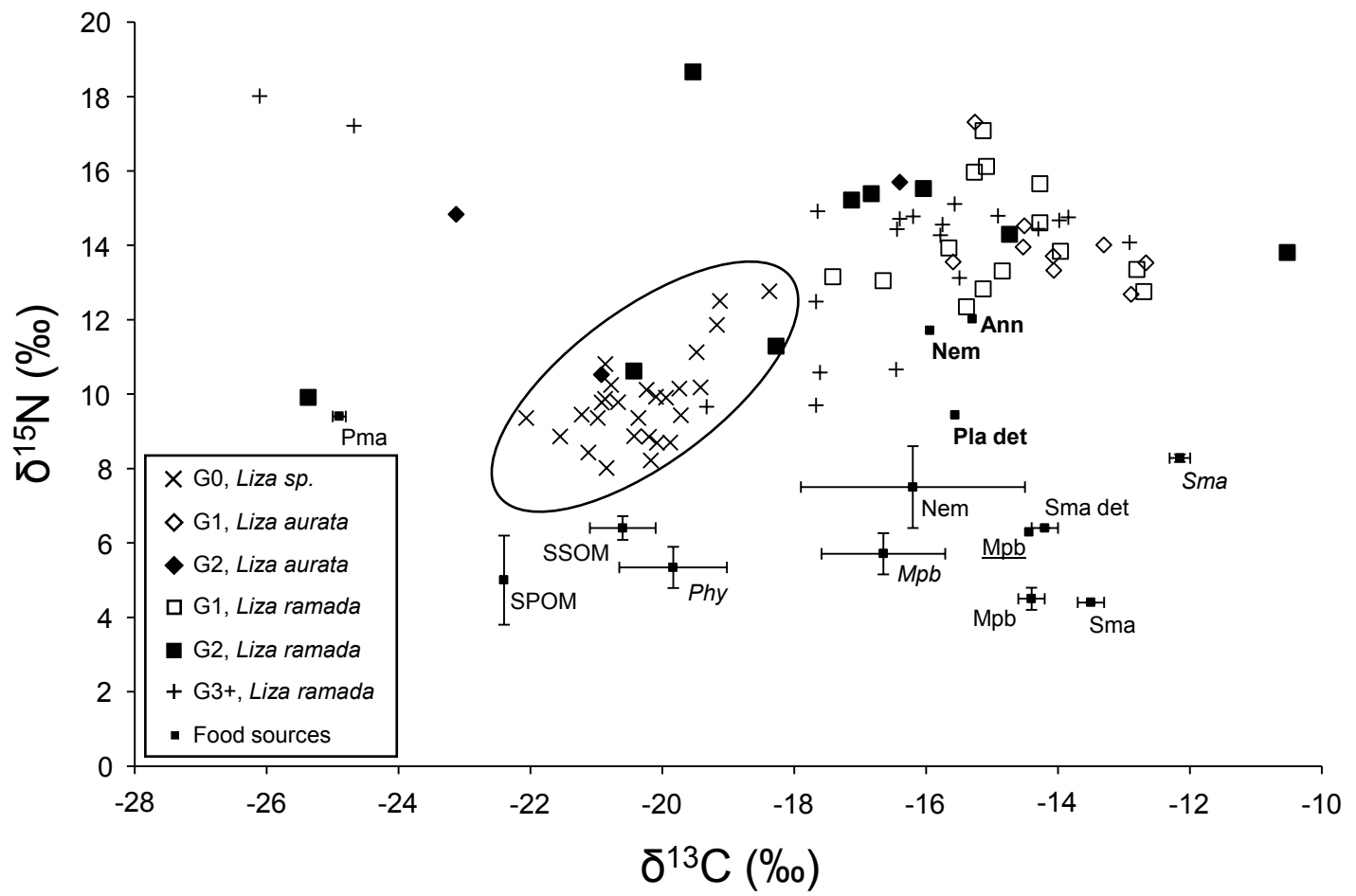


Figure 4

