Feeding habits of *Paralabrax nebulifer* (Serranidae) during reproductive and non-reproductive seasons in an adjacent area to Magdalena Bay, Baja California Sur, Mexico

Hábitos alimentarios del verdillo *Paralabrax nebulifer* (Serranidae) durante las temporadas reproductiva y no reproductiva en un área adyacente a Bahía Magdalena, Baja California Sur, México

Julio Alejandro Ysla-Guzmán1, Xchel Gabriel Moreno-Sánchez1, Martín Oscar Rosales-Velázquez1, Víctor Carrasco-Chávez1 and José Luis Ortíz-Galindo1*

1Instituto Politécnico Nacional, Centro Interdisciplinario de Ciencias Marinas (CICIMAR-IPN), Av. Instituto Politécnico Nacional s/n, Col. Playa Palo de Santa Rita, C.P. 23096, La Paz, B.C.S., México
*Corresponding author: jortizgalindo@gmail.com

**Resumen.**- El verdillo *Paralabrax nebulifer* es un pez comercialmente importante en la costa occidental de Baja California Sur. Para evaluar la dieta de esta especie y su variación en función del sexo y su condición reproductiva, se capturaron 60 especímenes durante siete muestreos estacionales utilizando trampas de agosto 2016 a agosto 2018 en un área adyacente a Bahía Magdalena, Baja California Sur, México. Se obtuvieron los contenidos estomacales de 50 especímenes (23 machos y 27 hembras). El sexo fue diagnosticado por observación directa de las gónadas. De acuerdo a los índices gonadosomático, hepatosomático y mediante análisis histológico, se corroboró la condición reproductiva del verdillo durante agosto 2016; abril, agosto y septiembre 2017 y agosto 2018; así como la temporada no reproductiva de noviembre 2016 y marzo 2018. El Índice de Importancia Relativa (IRI) se utilizó para clasificar los principales componentes de la dieta, que incluyeron: tres especies de peces, siete especies de crustáceos y una especie de molusco. Según el IRI, la sardina *Sardinops sagax* y la langostilla *Pleuroncodes planipes* fueron las presas que más contribuyeron (55%) a la dieta del verdillo. El análisis ANOSIM mostró que hubo diferencias significativas en la cantidad y tipo de presa consumida por sexo; el análisis SIMPER reveló que la mayoría de las especies que contribuyeron a las diferencias entre sexos, fueron producidas por *S. sagax* (16,58%), *Euphylax dovii* (15,95%), *Stenocionops ovata* (12%) y *P. planipes* (11,82%) para hembras. Hubo diferencias significativas en la cantidad y tipo de presa consumida por tipo de temporada reproductiva; la mayoría de las especies que contribuyeron a las diferencias entre temporadas fueron producidas por *Anchoa* spp. (27,76%) y *P. planipes* (22,67%) para la temporada no reproductiva, y por *S. sagax* (11,08%) para la temporada reproductiva. La estrategia de alimentación del verdillo fue la de un depredador carnívoro especialista, que durante la temporada reproductiva se alimenta principalmente de los peces *Sardinops sagax*, que cubren los requerimientos nutricionales dietéticos de los lípidos AGAI (ácido araquidónico, 20:4n-6; ácido eicosapentaenoico, 20:5n-3; ácido docosahexaenoico 22:6n-3), nutrientes necesarios para lograr el éxito reproductivo.

**Palabras clave:** Estacionalidad reproductiva, ecología alimentaria, Clupeidae, Galatheidae

**Abstract.**- The barred sand bass *Paralabrax nebulifer* is a commercially important fish off the west coast of Baja California Sur. To assess the diet of this species and variations as a function of sex and reproductive condition, 60 specimens were captured using traps during seven seasonal sampling trips from August 2016 to August 2018 in an adjacent area to Magdalena Bay, Baja California Sur, Mexico. The stomach contents of 50 specimens were obtained (23 males and 27 females). Sex was diagnosed by direct observation of the gonads. Based on the gonadosomatic index, hepatosomatic index, and histological analyses, the reproductive season of the barred sand bass was corroborated for August 2016, April, August, and September 2017, and August 2018, and the non-reproductive season was corroborated for November 2016 and March 2018. The Index of Relative Importance (IRI) was used to classify the main diet components, which comprised three fish species, seven crustacean species, and one mollusk species. According to the IRI, the South American pilchard *Sardinops sagax* and the red pelagic crab *Pleuroncodes planipes* were the prey that contributed the most (55%) to the barred sand bass diet. The ANOSIM showed that there were significant differences in the amount and type of prey consumed by sex; the SIMPER analysis revealed that the species contributing the most to differences between the sexes were *S. sagax* (16.58%), *Euphylax dovii* (15.95%), *Stenocionops ovata* (12%), and *P. planipes* (11.82%) for females. There were significant differences in the amount and type of prey consumed between types of reproductive season; the species contributing the most to differences between seasons were *Anchoa* spp. (27.76%), and *P. planipes* for non-reproductive season (22.67%), and *S. sagax* (11.08%) for reproductive season. The feeding strategy of the barred sand bass was that of a specialist carnivorous predator that fed mainly on the fish *Sardinops sagax* during the reproductive season, which supply the dietary nutritional requirements of the lipids HUFA (arachidonic acid, 20:4n-6; eicosapentaenoic acid, 20:5n-3; docosahexaenoic acid, 22:6n-3), nutrients required to achieve reproductive success.

**Key words:** Reproductive seasonality, feeding ecology, Clupeidae, Galatheidae
INTRODUCTION

Changes in the amplitude and phasing (i.e., phenology) of seasonally varying processes can significantly affect the functioning of marine ecosystems, from primary producers to fish stocks (Beare & McKenzie 1999, Bograd et al. 2002). In particular, at mid- to high trophic levels, the phenology of upwelling may be linked to match and mismatch between productivity and the life cycles of predators and prey (Ward et al. 2006, Durant et al. 2007). These types of oceanographic phenomena occur in the coastal zone of Magdalena Bay (Zaytsev et al. 2005).

In fish, reproductive success must be strictly adapted to seasonal events in the environment, whether such fluctuations are the result of cyclical seasonal variations or migrations during the natural life cycle of some species. In those with seasonal reproduction, an important adaptation refers to the spawning time, which is adjusted in such a way that the specific feeding needs of the larvae coincide with the seasonal availability of food (Yamahira 2004, Jalabert 2005).

Seasonal, ontogenetic, and reproductive variations in the diet have been recorded in various species of serranids (Ferry et al. 1997, Mendoza-Carranza & Rosales-Casían 2000, Nakai et al. 2001, Bocanegra-Castillo et al. 2002, Freitas et al. 2015, Vasiliki 2016, Rachedi et al. 2018), but none addresses the relationship of reproductive seasonality and diet composition associated with changes in the oceanic environment. On the other hand, data on the ecobiology of each species of interest in its natural environment may be very useful in creating adequate culture conditions, leading to reproductive maturation and spawning (Mylonas et al. 2010).

The barred sand bass Paralabrax nebulifer (Girard, 1854) belongs to the family Serranidae. Its distribution ranges from Santa Cruz, California (USA), to Magdalena Bay, Baja California Sur, Mexico, including Guadalupe Island (Miller & Lea 1972), where it is captured at depths between 30 and 185 m (Smith-Vaniz et al. 2010, Robertson & Allen 2015). In a study off the California coast, Love et al. (1996) found that P. nebulifer spawns from April to August, with a peak in July and that it tends to form large reproductive aggregations on sandy bottoms during its reproductive season. They estimated that 50% of females mature at 23.9 cm TL at 2 to 5 years of age, while 50% of males mature at 21.9 cm TL at 2 to 4 years of age. The barred sand bass migrates seasonally to the coast during its reproductive season, returning towards rocky bottoms during the non-reproductive season (Jarvis et al. 2010, Smith-Vaniz et al. 2010). The barred sand bass is an important species of high economic value in Mexico, with catches oscillating between 4,000 and 6,000 tons per year. This species is captured year-round, with peaks in catches between March and August in the state of Baja California Sur (SAGARPA 2016), where fishermen take advantage of their reproductive migration to the coast. In a study on reproductive aggregations, Erismann et al. (2017) found that in Punta Abreojos, Baja California Sur, local commercial fisheries do not increase their catch during the months of spawning activity (July and August). In the Gulf of Ulloa on the western coast of Baja California Sur, where 91% of the state fishing resource is captured (SAGARPA 2016), the commercial capture of this species has been affected by the creation of a fishing refuge (published by the Mexican government in June 2016; DOF 2016), which was justified by the interaction with marine turtles. This refuge was extended for five more years in June 2018 (DOF 2018). For this reason, it is necessary to obtain bio-ecological information that contributes to *P. nebulifer* fisheries management plan (DOF 2021).

Despite its economic importance in the state of Baja California Sur, there are no studies on how diet influences on reproductive seasonality of this species. The objective of the present study was to characterize the diet composition of the barred sand bass in an area adjacent to Magdalena Bay, Baja California Sur, during its reproductive and non-reproductive seasons. This will allow us to approximate the nutritional requirements of this species, so that in the future an inert food can be designed to achieve reproduction in captivity and contribute to fisheries management plan.
**Materials and Methods**

**Study area**
All samples were collected with help from artisanal fishermen in August and November 2016; April, August, and September 2017; and March and August 2018, using wire mesh traps with 5 cm mesh width (Erisman et al. 2017). Fishing occurred exclusively in the daytime (08:00 to 13:00 h). One fisherman deployed two traps in the morning at two different sites and retrieved those 40 min later, and repeated this procedure as long as weather conditions allowed. Each trap was placed at an approximately 50 m depth in an adjacent area to Magdalena Bay, Baja California Sur, Mexico (Fig. 1). The sea surface temperature values of the sampling sites were obtained from the satellite image consulted of NOAA’s CoastWatch\(^4\) website, provided by the Aqua satellite of the MODIS sensor, with a resolution of 500 m.

**Sampling**
The barred sand bass specimens were identified (Fischer et al. 1995b), transported to shore alive, and sacrificed with an overdose of anesthetic (40 mg L\(^{-1}\) clove oil). The total length (TL, ± 0.5 mm), standard length (SL, ± 0.5 mm), and weight (W, ± 1 g) of each specimen were recorded. Sex was identified macroscopically: females were identified based on the appearance of the ovaries and the presence of oocytes, and males were identified based on large and firm testes and milt released with gentle pressure on the abdomen (Brown-Peterson et al. 2011). A portion of the gonad (approximately one cubic centimeter) was also preserved with Davidson fixative. The stomachs were extracted by cutting from the esophagus to the pylorus; stomachs were frozen and transported to the laboratory for analysis.

**Reproduction**
To corroborate the reproductive or non-reproductive condition of fish collected in the wild, at least five samples were obtained from each of the catches when possible, and gonad samples were processed histologically. The samples were dehydrated and embedded in paraffin, and 3-mm transverse sections were stained with hematoxylin-eosin (Humason 1979). The histological preparations were analyzed under a microscope and photographs were taken with a digital camera (CoolSNAP-Pro, Media Cybernetics) coupled to an optical microscope (Olympus® BX41). The histological features used to assign the reproductive phases of female and male gonads were based on Brown-Peterson et al. (2011); the most abundant types of oocytes and the presence and abundance of germ cells were considered.

---

\(^4\)http://coastwatch.pfeg.noaa.gov

---

Figure 1. Study area. The dark ellipse indicates the area where barred sand bass *Paralabrax nebulifer* were caught, an adjacent area to Magdalena Bay, Baja California Sur, Mexico / Área de estudio. La ellipse oscura indica el área de captura del verdillo *Paralabrax nebulifer* en un área adyacente a Bahía Magdalena, Baja California Sur, México
The reproductive season was determined through the temporal variation of the frequency of the stages of gonadal maturity, the histological analysis, and the seasonal variation of the gonadosomatic index, hepatosomatic index, and Fulton’s condition factor (García-Díaz et al. 1997).

For each individual, the gonadosomatic index (GSI) was calculated as:

\[ \text{GSI} = \frac{W_{\text{gonad}}}{W_{\text{fish}}} \times 100 \]

The hepatosomatic index (HSI) was calculated as:

\[ \text{HSI} = \frac{W_{\text{liver}}}{W_{\text{fish}}} \times 100 \]

and Fulton’s condition factor (K) was calculated as:

\[ K = \frac{W_{\text{fish}}}{L^3} \]

where \( W_{\text{gonad}} \) is the gonad weight (g), \( W_{\text{liver}} \) is the liver weight (g), \( W_{\text{fish}} \) is the fish body weight (g), and \( L^3 \) is the standard length (mm³).

The mean GSI, HSI, and K were calculated for each seasonal sampling during the study period.

The sex ratio was evaluated with a Chi-square test at a 95% significance level based on the proportion of males and females.

**Diet**

In the laboratory, prey was counted, weighed, and identified to the lowest possible taxonomic level using specialized identification keys (Miller & Lea 1972, Brusca 1980, Wolff 1984, Fischer et al. 1995a, b). Data were grouped by sex (male or female) and season (reproductive or non-reproductive). The bait (Sardinops sagax) was differentiated from consumed prey because the bait did not appear complete but in pieces, as it had been previously crushed.

The quantitative analysis of diet was based on the frequency of occurrence (%F), number (%N), and weight (%W) indices (Hyslop 1980), which were used to calculate the IRI proposed by Pinkas et al. (1971) and modified by Hacunda (1981):

\[ IRI = (%N + %W) \times %F \]

This index is expressed as a percentage (Cortés 1999) and indicates the importance of each prey item in the predator’s diet:

\[ \%IR_i = \left( \frac{100}{\sum_{j=1}^{n} IRI_j} \right) \]

Diet width was calculated with Levin’s standardized index (Hurlbert 1978), based on the absolute number of each prey item. If the index is close to zero it indicates a specialist feeding strategy, and if it is close to one it indicates a generalist strategy (Krebs 1999):

\[ B_i = \frac{1}{n-1} \left( \frac{1}{\sum_{j} p_j} - 1 \right) \]

where \( B_i \) is the niche Breadth, \( \sum p_j \) is the proportion of the \( j \) item in predator \( i \)’s diet, and \( n \) is the total number of prey items.

An analysis of similarities (one-way ANOSIM, with 999 permutations) was performed to test for differences in diet according to sex and reproductive season (Clarke 1993). The resulting R statistic (\(-1 < R < 1\)) describes the similarity between groups defined according to the above factors. Values close to zero indicate no difference, and values close to 1 or -1 indicate a significant separation between groups. P-values were considered significant when \( P < 0.05 \). (Clarke & Gorley 2006). Following the ANOSIM global test, post-hoc tests were done using a Similarity Percentage analysis (SIMPER) (Clarke & Warwick 1994), which identified which prey types were the most responsible for significant differences found between samples (data of each sex or reproductive season).

Costello’s analysis (Costello 1990) modified by Amundsen et al. (1996) was used to interpret the feeding strategy of the barred sand bass and to assess population and individual patterns. The trophic level was calculated based on the number of prey in stomach contents, using the equation proposed by Cortés (1999):

\[ TL_i = 1 + \left( \sum_{j=1}^{n} p_j \times TL_j \right) \]

where \( TL_i \) is the trophic level of each prey category \( j \), and \( p_j \) is the proportion of each prey category in the diet. The trophic level of each prey category was based on Cortés (1999) and López-García et al. (2012).

**Results**

**General biological parameters and reproduction**

The total length shown as mean ± standard error (and range) of *Paralabrax nebulifer* specimens was 30.1 ± 3.4 (29.2-31.0) cm and weight was 377.6 ± 127.9 (345.3-410.0) g (Table 1). The sex ratio (F:M) was 1.17:1 (Chi-square: \( \chi^2 = 0.102; \text{fd}= 1 \)), which is not significantly different from 1:1.
According to the histological analyses, in August 2016, the female gonad was at a very advanced vitellogenesis, since it was even possible to observe the migration of the germinal vesicle, which corresponds to the developing phase (Fig. 2A); whereas the variety and diversity of the spermatogenic cells of the male gonad corresponded to the developing phase (Fig. 2B). In November 2016, the female gonad contained early and late perinucleolar cells, which corresponds to the immature phase (Fig. 2C); in the case of males, some spermatogenic sinuses with sperm were observed, as well as two melanomacrophage centers characteristic of the regenerating phase (Fig. 2D). In April 2017, the female gonad was observed with oocytes in the cortical alveolus stage, which characterizes the developing phase (Fig. 2E); while in males the presence of purple spermatozoa (Sz) was observed, evidence that the gonad was in the developing phase (Fig. 2F).

### Table 1

<table>
<thead>
<tr>
<th>Capture</th>
<th>n</th>
<th>Season</th>
<th>W (g)</th>
<th>TL (cm)</th>
<th>SL (cm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Aug (2016)</td>
<td>16</td>
<td>RS1</td>
<td>438.5 ± 229.3</td>
<td>30.6 ± 6.5</td>
<td>25.1 ± 4.5</td>
</tr>
<tr>
<td>Nov (2016)</td>
<td>3</td>
<td>NRS1</td>
<td>260.7 ± 87.6</td>
<td>27.3 ± 3.1</td>
<td>23.3 ± 3.3</td>
</tr>
<tr>
<td>Apr (2017)</td>
<td>6</td>
<td>RS2</td>
<td>384.3 ± 152.1</td>
<td>30.2 ± 3.8</td>
<td>25.1 ± 3.1</td>
</tr>
<tr>
<td>Aug (2017)</td>
<td>10</td>
<td>RS3A</td>
<td>352.9 ± 72.7</td>
<td>29.9 ± 2.1</td>
<td>24.9 ± 1.8</td>
</tr>
<tr>
<td>Sept (2017)</td>
<td>12</td>
<td>RS3B</td>
<td>376.8 ± 105.1</td>
<td>30.0 ± 2.9</td>
<td>25.9 ± 2.3</td>
</tr>
<tr>
<td>Mar (2018)</td>
<td>9</td>
<td>NRS2</td>
<td>413.2 ± 146.2</td>
<td>31.8 ± 3.4</td>
<td>26.7 ± 3.6</td>
</tr>
<tr>
<td>Aug (2018)</td>
<td>4</td>
<td>RS4</td>
<td>417.0 ± 102.0</td>
<td>30.9 ± 2.2</td>
<td>26.0 ± 2.4</td>
</tr>
</tbody>
</table>

**Figure 2.** Reproductive phases of the gonads of female (left side) and male (right side) specimens of *Paralabrax nebulifer* off an adjacent area to Magdalena Bay, Baja California Sur, Mexico. H & E staining. Developing phase (A-B) (August 2016); regenerating (C) and regressing (D) (November 2016); developing (E-F) (April 2017). Nucleolus (nu), nucleus (n), lipid inclusions (li), chromatin nucleolus (Cn), early perinucleolus (Ep), late perinucleolus (Lp), cortical alveolus (Ca), vitellogenesis 1 (Vtg1), vitellogenesis 2 (Vtg2), vitellogenesis 3 (Vtg3), germ vesicle migration (Gvm), spermatogonia (Sg), spermatocyte (Sc), spermatid (Sd), spermatozoa (Sz), lamella (L), gonadal lumen (GL), and melanomacrophage center (MMC).
In August 2017, oocytes at an advanced stage of vitellogenesis (Vtg 3) were observed in the female gonad, which characterizes the spawning capable phase (Fig. 3A); in males the gonad had well-defined sperm tubules (ST) and sperm, characteristic of the spawning capable phase (Fig. 3B). In September 2017, in the female gonad, post-ovulatory follicles (Pof) were observed in the stroma and cells at the nucleolus chromatin stage (N) were abundant, which indicates the end of the spawning period, however, mature oocytes were observed in the rest of the section, so it was characterized as the spawning capable phase (Fig. 3C); the male gonads showed residual spermatozoa, to the extent that their own seminiferous tubules had lost their shape due to the large amount of sperm inside them; this was considered the spawning capable phase (Fig. 3D).

In March 2018, showed a female gonad with a postovulatory follicle (Pof), in which a strong predominance of immature cells was evident, but not a single one was at the cortical alveolus stage, which indicates that the female gonad corresponded to the regenerating phase (Fig. 3E); the male gonad showed the presence of completely immature sex cells, as well as the presence of well-pigmented melanomacrophage centers, corresponding to the regenerating phase (Fig. 3F).

Figure 3. Reproductive phases of the gonads of female (left side) and male (right side) specimens of Paralabrax nebulifer off an adjacent area to Magdalena Bay, Baja California Sur, Mexico. Spawning capable (A-B) (August 2017); regressing (C-D) (September 2017); regenerating (E-F) (March 2018). Early perinucleolus (Ep), late perinucleolus (Lp), vitellogenesis 1 (Vtg1), vitellogenesis 2 (Vtg2), vitellogenesis 3 (Vtg3), postovulatory follicle (Pof), spermatogonia (Sg), spermatid (Sd), spermatocyte (Sc), spermatozoa (Sz), gonadal lumen (GL), melanomacrophage center (MMC), and seminiferous tubule (ST) / Fases reproductivas de las gónadas de los ejemplares hembras (lado izquierdo) y machos (lado derecho) de Paralabrax nebulifer de un área adyacente a Bahía Magdalena, Baja California Sur, México. Capacidad de desove (A-B) (agosto 2017); regresión (C-D) (septiembre 2017); regeneración (E-F) (marzo 2018). Núcleolo cromatina (Cn), perinucleolo temprano (Ep), perinucleolo tardío (Lp), vitelogénesis 1 (Vtg1), vitelogénesis 2 (Vtg2), vitelogénesis 3 (Vtg3), folículo postovulatorio (Pof), espermatogonia (Sg), espermatocisto (Sc), espermátide (Sd), espermatozoide (Sz), lumen gonadal (GL), centro melanomacrófago (MMC), túbulo seminífero (ST)
The sea surface temperature (SST), gonadosomatic index (GSI), hepatosomatic index (HIS), and Fulton’s Condition Factor (K) are shown in Table 2. The highest SST were found in August/September (2017), and August (2018). The highest GSI were found in August (2016), April/August (2017), and August (2018). The highest HIS were found in August/November (2016) and April (2017). The K values remained high during most of the seasonal samplings, with the exception of April (2017).

Histological results and the gonadosomatic index, hepatosomatic index, Fulton’s condition factor, and sea surface temperature suggest that the reproductive season occurs between April and September.

**Diet**

Three fish species, seven crustacean species, and one mollusk species were identified. A total of 75 prey organisms weighing 256.73 g were counted. The prey with greatest number and weight were the fish *Sardinops sagax* (16 %N, 35.9 %W), the red pelagic crab *Pleuroncodes planipes* (14.7 %N, 15.2 %W), and the fish *Anchoa* spp. (14.7 %N, 6.8 %W). According to the %IRI, the prey with greatest relative importance in the diet were the fish *S. sagax* (35.9%), the crustacean *P. planipes* (18.9%), and the fish *Anchoa* spp. (13.6%), as well as the crustaceans *Stenocionops ovata* (11.9%), *Euphylax dovii* (9.7%), and *Penaeus californiensis* (4.9%). These preys represented 95% of the IRI as a whole (Table 3 and Fig. 4).

Nine prey types were identified in the stomachs of the 23 barred sand bass males. The %IRI indicated that the most important prey were the fish *S. sagax* (41.4%), the crustacean *E. dovii* (24.3%), and the fish *Anchoa* spp. (13.0%). Eleven prey types were identified in the stomachs of the 27 barred sand bass females; the most important were the crustacean *P. planipes* (44.6%), the fish *S. sagax* (21.7%), and the crustacean *S. ovata* (14.2%) (Fig. 4).

### Table 2

<table>
<thead>
<tr>
<th>Capture</th>
<th>n</th>
<th>SST</th>
<th>GSI</th>
<th>HIS</th>
<th>K</th>
<th>Season</th>
</tr>
</thead>
<tbody>
<tr>
<td>Aug (2016)</td>
<td>16</td>
<td>23.7 ± 0.6</td>
<td>3.8 ± 1.3</td>
<td>30.6 ± 6.5</td>
<td>1.5 ± 0.7</td>
<td>RS1</td>
</tr>
<tr>
<td>Nov (2016)</td>
<td>3</td>
<td>23.5 ± 0.3</td>
<td>0.5 ± 0.2</td>
<td>27.3 ± 3.1</td>
<td>1.2 ± 0.0</td>
<td>NRS1</td>
</tr>
<tr>
<td>Apr (2017)</td>
<td>6</td>
<td>17.7 ± 1.0</td>
<td>1.1 ± 0.4</td>
<td>30.2 ± 3.8</td>
<td>0.8 ± 0.1</td>
<td>RS2</td>
</tr>
<tr>
<td>Aug (2017)</td>
<td>10</td>
<td>24.6 ± 0.6</td>
<td>3.9 ± 1.6</td>
<td>1.7 ± 0.5</td>
<td>1.3 ± 0.1</td>
<td>RS3A</td>
</tr>
<tr>
<td>Sept (2017)</td>
<td>12</td>
<td>25.2 ± 0.4</td>
<td>Undetermined</td>
<td>Undetermined</td>
<td>1.3 ± 0.1</td>
<td>RS3B</td>
</tr>
<tr>
<td>Mar (2018)</td>
<td>9</td>
<td>18.0 ± 0.5</td>
<td>0.5 ± 0.3</td>
<td>1.4 ± 0.4</td>
<td>1.3 ± 0.3</td>
<td>NRS2</td>
</tr>
<tr>
<td>Aug (2018)</td>
<td>4</td>
<td>24.2 ± 0.6</td>
<td>1.2 ± 0.9</td>
<td>1.3 ± 0.1</td>
<td>1.4 ± 0.1</td>
<td>RS4</td>
</tr>
</tbody>
</table>

### Table 3

<table>
<thead>
<tr>
<th>Prey</th>
<th>%IRI</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>Euphylax dovii</em></td>
<td>24.3%</td>
</tr>
<tr>
<td><em>Stenocionops ovata</em></td>
<td>11.9%</td>
</tr>
<tr>
<td><em>Penaeus californiensis</em></td>
<td>4.9%</td>
</tr>
</tbody>
</table>

### Table 4

<table>
<thead>
<tr>
<th>Prey</th>
<th>%IRI</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>Sardinops sagax</em></td>
<td>35.9%</td>
</tr>
<tr>
<td><em>Anchoa</em> spp.</td>
<td>13.0%</td>
</tr>
<tr>
<td><em>Caranx</em> spp.</td>
<td>13.6%</td>
</tr>
</tbody>
</table>

Nine prey types were identified in the stomachs of the 23 barred sand bass males. The %IRI indicated that the most important prey were the fish *S. sagax* (41.4%), the crustacean *E. dovii* (24.3%), and the fish *Anchoa* spp. (13.0%). Eleven prey types were identified in the stomachs of the 27 barred sand bass females; the most important were the crustacean *P. planipes* (44.6%), the fish *S. sagax* (21.7%), and the crustacean *S. ovata* (14.2%) (Fig. 4).
There were statistically significant differences between the sexes (Chi-square: $\chi^2 = 19.28$, fd= 10).

A total of 38 barred sand bass stomachs (76%) were analyzed during the reproductive season and the prey that contributed most to the diet, according to the %IRI, were the fish $S$. sagax (52.5%) and the crustaceans $S$. ovata (16.8%) and $E$. dovii (14.7%). A total of 12 stomachs (24%) were analyzed during the non-reproductive season and the most important prey, according to the %IRI, were the crustacean $P$. planipes (57.1%) and the fish Anchoa spp. (40.7%) (Fig. 4).

There were significant differences between the sexes (ANOSIM, R= 0.024; $P = 0.01$). The SIMPER analysis showed that most of the differences between the sexes (56.36%) were produced by $S$. sagax (16.58%), $E$. dovii (15.95%), $S$. ovata (12%), and $P$. planipes (11.82%) for females. There were significant differences between the seasons (ANOSIM, R= 0.113; $P = 0.01$). The SIMPER analysis showed that most of the differences between the seasons (61.51%) were produced by Anchoa spp. (27.76%), and $P$. planipes (22.67%) for non-reproductive season, and $S$. sagax (11.08%) for reproductive season. No significant interactions were found between the sexes and the reproductive season (ANOSIM, R= 0.031; $P = 0.13$).

The feeding strategy of the barred sand bass was that of a specialist predator ($B = 0.38$). This behavior was seen in males ($B = 0.35$) and females ($B = 0.26$), as well as during the reproductive season ($B = 0.20$), whereas during the non-reproductive season ($B = 0.52$) the barred sand bass tended to be a generalist predator. The Costello graph modified by Amundsen et al. (1996) showed that the barred sand bass consumed few preys, and that the abundance and frequency of those prey were similar in the diet, so that there was not a dominant prey (Fig. 5a). Males fed mainly on $S$. sagax, $E$. dovii, and Anchoa spp., whereas females fed mainly on $P$. planipes, $S$. sagax, $S$. ovata, and Anchoa spp. (Fig. 5b, c). The most important prey during the reproductive season were $S$. sagax, $S$. ovata, and $E$. dovii, whereas the most important prey during the non-reproductive season were $P$. planipes, Anchoa spp., and $P$. californiensis (Fig. 5d, e). The trophic level of the barred sand bass was 3.8; it was 3.9 for males and 3.8 for females; 3.8 during the reproductive season, and 3.9 during the non-reproductive season.
**Discussion**

The size of the *P. nebulifer* specimens captured during the present study corresponded to adult specimens, according to the size at first maturation recorded for the California coast (Love *et al.* 1996; Baca-Hovey *et al.* 2002), due to the selectivity of the fishing gear used.

According to results obtained using the gonadosomatic index, hepatosomatic index, Fulton’s condition factor, and macroscopic and histological analyses of the gonads, the reproductive season occurred in August 2016, April, August and September 2017, and August 2018; according to the SST, these months can be characterized as the warm season. The non-reproductive season occurred in November 2016 and March 2018, characterized as the cool season. According to the developmental phases of the gonads, the highest reproductive activity (spawning capable phase) was observed in the warm season and particularly in the month of August 2017. During this reproductive seasonality, also reported from California to Punta Abreojos (Jarvis *et al.* 2010, Erisman *et al.* 2017), *P. nebulifer* migrate from offshore waters to more productive near-shore habitats and spawn at short intervals for a period of two to three months. This same situation was reported for *Sciaenops ocellatus* in the Western Atlantic (Fuiman & Faulk 2013).

According to the results of the diet composition, barred sand bass are carnivores that fed mainly on fish (*Sardinops sagax* and *Anchoa* spp.) and crustaceans (*Pleuroncodes planipes* and *Stenocionops ovata*) in an adjacent area to Magdalena Bay. This contrasts with the study by Roberts *et al.* (1984) in California, where the main prey were brachyuran crabs, mysids, pelecypods, and epibenthic fish, and with the study by Mendoza-Carranza & Rosales-Casián (2002) in Baja California, who reported gammarid amphipods (50.91 %IRI) and fish (16.28 %IRI) as the main prey. This feeding pattern shows that the barred sand bass is a predator that can feed on prey on the benthos as well as in the water column, which places this species as a euryphagous carnivore (Smith 1989). It is important to note that fish and invertebrate species consumed by specimens in this study were not the same as those reported in previous studies and that the proportion of each prey was different. This could be due to characteristics of the study location; Magdalena Bay can be considered the southern boundary of the transition zone between the subarctic and subtropical waters of the northeastern Pacific Ocean, with...
marked seasonality (Zaytsev et al. 2014), which acts as an ecological barrier for several species (Brusca & Wallerstein 1979, Parrish et al. 1981, Norton et al. 1985). Therefore, these oceanographic boundaries could explain differences found in the prey consumed by barred sand bass in its most southern location, compared with the rest of its distribution.

Despite differences in prey consumed by barred sand bass in different geographic locations, it should be noted that fish are important in the diet (up to 50% of diet). This has been confirmed by Roberts et al. (1984), who reported that off southern California, large barred sand bass were mainly piscivores, whereas small specimens preferred crustaceans (Brachyura) and pelecypods, and other organisms associated with the benthos. Serranids in general show high trophic diversity (Gómez et al. 1999), which suggests that the barred sand bass is able to adapt according to the food availability in its distribution area.

There were significant differences by sex, as males fed mainly on fish (54%) and on a lower proportion of crustaceans (24%), whereas females showed the opposite pattern (21% fish and 68% crustaceans). This could suggest that there is resource partitioning between the sexes as a function of habitat.

Seasonal variations in diet were also found, with high consumption of fish (S. sagax 52.5 %IRI) and of the crustaceans S. ovata (16.8 %IRI) and E. dovii (14.7 %IRI) during the reproductive season, and consumption mainly of the crustacean red pelagic crab during the non-reproductive season (P. planipes, 57.1 %IRI). In a study by Mendoza-Carranza & Rosales-Casián (2002), seasonal variations in the diet of the barred sand bass were observed; the species fed mainly on fish and gammarids in the summer (46.2 %IRI, 40.0 %IRI, respectively), while in winter gammarids became the preferred diet (88.4 %IRI) and the proportion of fish consumed was reduced (5.1 %IRI). A similar result was found in the congeneric species spotted sand bass Paralabrax maculofasciatus in Bahía de Los Ángeles, Baja California, in the Gulf of California (Ferry et al. 1997), where crustaceans were the preferred prey during the spring (cool season) and fish remains, ophiuroids, and brachyuran crabs were the main food components during the fall (warm season). These results contrast with what were reported by Mendoza-Carranza & Rosales-Casián (2000) for the spotted sand bass on the Pacific coast in Punta Banda Estuary, Baja California and Ojo de Liebre Lagoon, Baja California Sur (Bocanegra-Castillo et al. 2002), where crustaceans were the most important prey during summer (warm season). In Punta Banda Estuary fishes and decapod crustaceans were the preferred prey in winter (cool season), whereas Callinectes denticulatus and fish, gammarids, and mollusks were the most important food components in November (cool season) in Ojo de Liebre Lagoon.

These similarities and differences in the diet that occur between serranid species or even in the same species in different locations can be explained by prey availability during each season and by the opportunistic behavior of the predator. The prey obtained in the present study showed high %IRI values of S. sagax in the warm season (reproductive season) and P. planipes in the cool season (non-reproductive season), which is when this prey reproduce and reach their greatest abundances (Aurioles-Gamboa et al. 1994, Torres-Villegas et al. 2007).

In the Magdalena Bay zone, during warm season when P. nebulifer undertake their reproductive migration towards the coastal zone, the coastal upwelling events increases off April to June (Zaitsev et al. 2003) and causes the match and mismatch between the reproductive process and the opportunity of the predators to have new prey at their disposal to obtain adequate nutrition for the reproductive process (Yamahira 2004, Ward et al. 2006, Durant et al. 2007). According to McKinzie (2014), the barred sand bass population is associated with different substrates depending on the reproductive condition. During the reproductive season, their migration is associated with sandy bottoms and carries out large movements in the water column, a situation that favors the capture of fish, while during the non-reproductive season it is associated with sandy-reef bottoms, so that it coincides with the abundance of the benthic resource Pleuroncodes planipes.

Thus, it was found that during this season the main food was the sardine Sardinops sagax, a small pelagic fish that presents its highest reproductive peak during winter (Torres-Villegas et al. 2007); according to evidence obtained in other eastern boundary systems, for small pelagic fishes as sardines the warm season corresponds to their so-called “feeding season”, characterized by lipid storage (Silva et al. 2006), which in turn are transferred to predatory fish (Fuiman & Faulk 2013). One of the main contributions of stored lipids are the supply of the dietary nutritional requirements of lipids HUFA (arachidonic acid, 20:4n-6; eicosapentaenoic acid, 20:5n-3; docosahexaenoic acid, 22:6n-3), nutrients required to achieve reproductive success, which are important for fertility, hatching rate, and larval survival (Fernández-Palacios et al. 1995, Cardona et al. 2015). The prey P. planipes, consumed mainly during the non-reproductive season, are rich in carotenoids, particularly astaxanthin, which serves as a source of antioxidants that protect the lipids HUFA (Torrisen & Christiansen 1995), that are stored during the feeding season. Regarding diet variations during the reproductive process, the feeding pattern of barred sand bass is similar to what was reported for the serranid Cephalopholis urodeta in a study carried out in southern Japan, where fish (50% W) were abundant in the diet of mature fish, and the proportion of Galatheidae consumed was higher in February than during the other months (Nakai et al. 2001).
The trophic level found for barred sand bass in this study was 3.8, which places it as a tertiary consumer (Cortés 1999) feeding on prey at the second level of the trophic chain, such as crustaceans (P. planipes, Penaeus californiensis, S. ovata) and teleosts (S. sagax and Anchoa spp.). This is similar to what has been reported for Serranus cabrilla (3.9, Rachedi et al. 2018) and Serranus scriba (3.8, Vasiliki 2016). According to the food group table reported by Vasiliki (2016), trophic level values between 3.7 and 4.0 obtained with TROPH correspond to carnivores with a preference for decapods. The other species in the genus would be considered carnivores with a preference for fish, with trophic level values between 4.0 and 4.5 (Vasiliki 2016), P. maculatofasciatus (4.2) and Paralabrax auroguttatus (4.2) (Aurioles-Gamboa et al. 2013).

In conclusion, the barred sand bass is a third-order opportunistic carnivore with the feeding strategy of a specialist that preys on species forming aggregations in the pelagic and benthic zone of Magdalena Bay, and that fed mainly on the fish Sardinops sagax during the reproductive season, which warranting the nutritional contribution of the lipids HUFA, nutrients required to achieve reproductive success.

ACKNOWLEDGMENTS
The authors thank the Consejo Nacional de Ciencia y Tecnología (CONACyT) for economic support provided and the Instituto Politécnico Nacional for BEIFI, EDI, and COFFA scholarships. The graphic material was custom tailored by Armando Hernández López.

LITERATURE CITED
Feeding habits during reproductive seasonality of *Paralabrax nebulifer*.


Recibido el 15 de mayo 2020 y aceptado el 25 de febrero de 2021

Editor asociado: Mauricio Landaeta D.