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Feeding ecology of elasmobranch fishes in coastal waters of the Colombian Eastern Tropical Pacific

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Abstract

Background: Stomach contents of 131 specimens of five elasmobranch species (*Mustelus lunulatus*, *Dasyatis longa*, *Rhinobatos leucorhynchus*, *Raja velezi* and *Zapteryx xyster*) caught in the central fishing zone in the Pacific Ocean of Colombia were counted and weighed to describe feeding habits and dietary overlaps.

Results: Twenty-one prey items belonging to four major groups (stomatopods, decapods, mollusks and fish) were identified. Decapod crustaceans were the most abundant prey found in stomachs. The mantis shrimp *Squilla panamensis* was the main prey item in the diet of *M. lunulatus*; tiger shrimp *Trachypenaeus* sp. was the main prey item in the diet of *Rhinobatos leucorhynchus* and *Raja velezi*, and Penaeidae shrimp were the main prey items in the diet of *Z. xyster*. Furthermore, fish were important in the diet of *Raja velezi*, *Z. xyster* and *D. longa*. The greatest diet breadth corresponded to *Z. xyster* whereas *M. lunulatus* was the most specialized predator. Finally, four significant diet overlaps between the five species were found, attributable mainly to Squillidae, Penaeidae and Fish.

Conclusion: Shrimps (Penaeidae and stomatopods) and benthic fishes were the most important food types in the diet of the elasmobranch species studied. Diet breadth and overlap were relatively low. Determination of food resource partitioning among the batoid species studied was not possible. However, we identified partitions in other niche axes (time of feeding activity and habitat utilization). It is possible to assume that diffuse competition could be exceeding the biunivocal competition among the studied species. Therefore, this assemblage would have a strong tendency to trophic guild formation.

Background

Elasmobranch fishes are among the top predators in the marine environment and thus play an important role in marine ecosystems, potentially regulating, through predation, the size and dynamics of their prey populations [1]. An understanding of competitive and predatory processes

is thus necessary to gain insight into the role of predators in influencing niche, community and food web structure, and ultimately ecosystem dynamics [2]. In this sense, the theory about resource partitioning is frequently attributed to competitive or cooperative interactions. This approximation predicts that spatial or temporal partitioning may

increase the tolerance of niche overlap, and may reduce the pressure of competition among coexisting species [3,4].

Several studies have evaluated the influence of resource partitioning in teleost fishes to explain reduction in competition potential [5-7]. These authors reported a positive relationship between habitat partitioning and the magnitude of diet overlap or competitive interactions, when the interacting species had the same diet preferences. Furthermore, Ross [8] found that the first resource that is fractioned is food, followed by habitat. Despite the ecological importance of elasmobranch fishes for the marine ecosystem, resource partitioning and competitive exclusions in this taxonomic group are poorly understood.

A substantial amount of data on the diet of different elasmobranch species has been reported to date. Elasmobranch fishes are often typified as opportunistic predators, with a wide trophic spectrum that includes plankton to marine mammals. In general, oceanic elasmobranch species feed on squid and big fishes [9,10], whereas the coastal and benthic species feed on crustaceans, mollusks and small or juvenile fishes [11-13]. A few species feed on other elasmobranchs, birds, reptiles or marine mammals [14,15]. Ontogenetic variation in diet is well known [16,17], with a strong tendency to ingest larger and more mobile animals with increasing size.

Research on trophic relationships among sympatric species of elasmobranchs is scarce and results vary among studies [18-20]. Varying overlap values between coexisting species, successive sizes of a same species, or even between sexes have been reported. Moreover, the effect of habitat and feeding time on the diet of sympatric species is very poorly known [21-23]. In the Eastern Pacific Ocean of Colombia (EPOC) the study of elasmobranch diet is in its infancy [24-26]. The existing studies have only considered three of 87 elasmobranch species reported for the area [27]. Furthermore, no research has addressed trophic interactions among sympatric fish species in coastal or oceanic environments.

Therefore, the object of this study was to quantify and compare the diet and trophic interactions of coastal elasmobranch species from the Eastern Tropical Pacific of Colombia, and to suggest possible mechanisms for their coexistence.

Results

Feeding activity

A total of 131 specimens of five species (*Mustelus lunulatus*, *Dasyatis longa*, *Rhinobatos leucorhynchus*, *Raja velezi* and *Zapteryx xyster*) were analyzed. All specimens were captured in shallow waters (between 15 and 60 m depth). Significant differences in the bathymetric distribution of the species were found (KW-H (4,162) = 105.16, p << 0.0001), with *Zapteryx xyster* and *Raja velezi* found in narrower depth ranges and under 35 and 40 m, respectively. *Dasyatis longa* and *Rhinobatos leucorhynchus* were associated with the shallowest depths (Figure 1). *Mustelus lunulatus*, *D. longa* and *Rhinobatos leucorhynchus* were captured mainly during diurnal bottom trawling, while *Z. xyster* were captured mainly during nocturnal bottom trawling. *Raja velezi* was captured only at night. Only *R. leucorhynchus* and *Z. xyster* showed significant differences in hourly feeding activity (Table 1). We did not carry out analyses of ontogenetic shifts in diet because we did not capture individuals from all sizes intervals.

Diet composition

The proportion of empty stomachs varied widely among species, with a minimum vacuity index of 9.5% in *M. lunulatus* and a maximum vacuity index of 47.6% in *D. longa* (Table 2). The main prey item of *Mustelus lunulatus* were stomatopods from the family Squillidae (%N = 73.2%, %W = 36.1%, and occurrence 78.6%) (Table 3), with *Squilla panamensis* being the main prey item (92.8 %IRI). An additional twelve prey categories were identified in the stomachs of *M. lunulatus*, which feed opportunistically on shrimps, crabs and gasteropods in low percentages.

In *D. longa* stomatopods were the most frequent prey item (14.3%) and had the highest percent by number (58.3%),

Table 1: Sampling data of elasmobranchs caught

Species	n	Capture (n)		p	Size (cm)	
		Diurnal	Nocturnal		Min	Max
<i>Mustelus lunulatus</i>	42	15	16	0.8202	55	125
<i>Dasyatis longa</i>	21	19	5	0.8906	106	210
<i>Rhinobatos leucorhynchus</i>	24	10	18	0.0017	37	67
<i>Raja velezi</i>	13	0	15	-	48	80
<i>Zapteryx xyster</i>	31	3	51	0.0353	27	66

Sampling data of elasmobranchs caught in the Eastern Tropical Pacific Ocean of Colombia and range size of these species. n: number of individuals captured. p: p-value of K-W test.

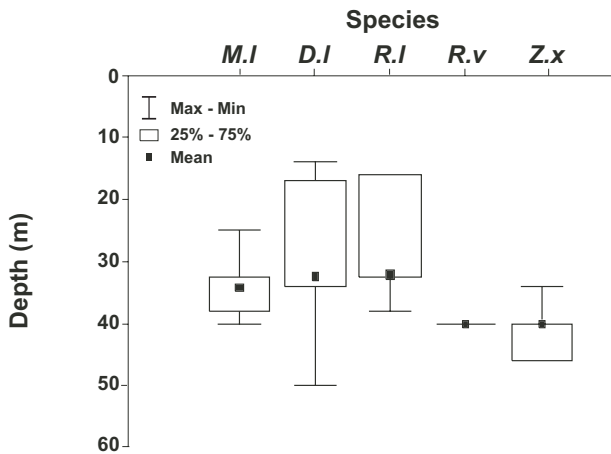


Figure 1
Bathymetric distribution of the elasmobranch species studied. (M.I) is *Mustelus lunulatus*, (D.I) is *Dasyatis longa*, (R.I) is *Rhinobatos leucorhynchus*, (R.v) is *Raja velezi* and (Z.x) is *Zapteryx xyster*.

whereas fishes had the highest percent by weight (36.8%) (Table 3). Digested material was 35.1% and vacuity index was 47.6%. The main prey items were Squillidae (54.9% %IRI) and fishes (31.1%IRI). The diet of this species also included decapods and gastropods.

In *Rhinobatos leucorhynchus* the most frequent (66.6%), abundant (85.6%) and with highest percent by weight (41.7%) prey item was decapods, following by fishes (12.5, 11.4 and 30.9% respectively) (Table 3). The main prey item was *Trachypena* sp. (60.4%IRI). However, stomatopods and crabs (Portunidae) were also found in the diet of this species. The vacuity index was low (20.8%).

Only two prey items were identified in *R. velezi*, fishes and decapods. Fishes were the most frequent item (53.8%), and also the most abundant by number (53.9%) and

weight (50.8%), and collectively (70.4% IRI). Among the fishes, only Batrachoididae could be identified (Table 3).

Six prey categories were identified in stomach contents of *Z. xyster* (Table 3). Fishes and penaid shrimp occurred most frequently (> 16%) and were the most abundant by number (27.3% each), however shrimp were the most abundant by weight (31.4%). For this species decapods were the most important prey item with 43.2 %IRI.

Diet breadth and overlap

Use of a three-dimensional graphical representation of the diet [28] facilitated the description of feeding styles of the five species studied (Figure 2). The diet of *M. lunulatus* was specialized, with most individuals consuming *Squilla pan-amensis*, although this species also feed on twelve other prey items. In contrast, the diet of *Z. xyster* was more heterogeneous and generalized because fishes, decapods and stomatopods were consumed by the majority of individuals. These results were confirmed by Levin's breadth index, which yielded the greatest diet breadth for *Z. xyster*, while *M. lunulatus* was the most specialized predator (Table 4).

Following Zares & Rand's [29] criteria, four biologically significant diet overlaps were established among the studied species (Table 5). A significantly high overlap was found between *M. lunulatus* and *D. longa*, which shared shrimps of the family Squillidae. Significant overlaps were also found between *Rhinobatos leucorhynchus* and *Raja velezi*, and *R. leucorhynchus* and *Z. xyster*, which shared Penaeidae, and between *Raja velezi* and *Z. xyster* which shared fish and Penaeidae. The random overlap hypothesis was rejected ($p = 0.010$), suggesting low overlap and low partitioning of resources [30].

Discussion

The composition of diets suggests that all elasmobranchs studied are species that feed on epibenthic invertebrates (mainly stomatopods and decapods) and fish. This result is highly associated with the source of the samples analyzed, i.e., elasmobranchs captured in bottom trawl fish-

Table 2: Feeding activity of captured elasmobranchs

Specie	n	Es	VI	χ_n	CI $_{\chi_n}$	χ_w	CI $_{\chi_w}$
<i>Mustelus lunulatus</i>	42	3	9.5	2.5	1.9 – 3.1	8.9	7.8 – 11
<i>Dasyatis longa</i>	21	10	47.6	0.9	0.3 – 1.2	1.1	0.3 – 1.9
<i>Rhinobatos leucorhynchus</i>	24	5	20.8	1.8	1.1 – 2.5	1.3	0.6 – 2.0
<i>Raja velezi</i>	13	4	38.8	1.0	0.3 – 1.7	6.3	0.2 – 12.4
<i>Zapteryx xyster</i>	31	11	35.5	0.9	0.5 – 1.3	1.1	0.6 – 1.6

Feeding activity of elasmobranch captured in the Eastern Tropical Pacific Ocean of Colombia. Number of stomachs analyzed (n), Empty stomachs (Es), Vacuity Index (VI), mean number of prey per stomach (χ_n), 95% Confidence Interval for χ_n (CI $_{\chi_n}$), mean weight of prey per stomach (χ_w), 95% Confidence Interval for χ_w (CI $_{\chi_w}$).

Table 3: Diet composition of captured elasmobranchs

	<i>M. lunulatus</i>				<i>D. longa</i>				<i>R. leucorhynchus</i>				<i>R. velezi</i>				<i>Z. xyster</i>			
	% O	% N	%W	% IRI	% O	% N	%W	% IRI	% O	% N	%W	% IRI	% O	% N	%W	% IRI	% O	% N	%W	% IRI
CRUSTACEA																				
STOMATOPODA																				
DA																				
Squillidae					14.28	58.30	4.48	54.87	4.16	2.85	0.31	0.31								
<i>Squilla panamensis</i>	61.90	62.90	28.10	92.77													12.90	18.20	26.00	20.83
<i>Squilla parva</i>	14.28	9.27	5.95	4.01																
<i>Pseudosquilla similis</i>	2.38	1.03	2.07	0.12																
Total Stomatopoda	78.56	73.20	36.10	96.90	14.28	58.30	4.48	54.87	4.16	2.85	0.31	0.31					12.90	18.20	26.00	20.83
DECAPODA																				
Portunidae									8.33	8.57	0.47	1.79								
<i>Euphyllax</i> sp.	2.38	1.03	0.40	0.05																
<i>Portunus asper</i>	4.76	3.09	7.32	0.78																
<i>Callinectes toxotes</i>	2.38	1.03	0.40	0.05																
<i>Hepatus</i> sp.	4.76	2.06	0.75	0.21																
Brachiura																	3.20	4.54	0.59	0.56
Penaeidae									25.00	31.30	10.30	24.82					19.40	27.3	31.4	41.89
<i>Pennaeus occidentales</i>	7.14	2.06	0.75	0.23																
<i>Trachypenaeus</i> sp.	4.76	2.06	0.40	0.49					33.30	45.70	30.30	60.44	15.3	46.2	45.7	29.57				
									8	0	0		8	0	0					
Palaemonidae	2.38	9.27	4.11	0.49																
Hippidae	7.14	4.12	2.20	0.71	4.76	8.33	15.70	6.98									3.20	4.54	2.09	0.74
Total Decapoda	35.70	24.70	16.50	3.01	4.76	8.33	15.70	6.98	66.63	85.60	41.70	87.05	15.38	46.20	45.70	29.57	25.80	36.40	34.10	43.19
MOLLUSCA																				
GASTEROPODA																				
A																				
Buccinidae	2.38	1.03	0.03	0.04																
<i>Agaronia testacea</i>					4.76	8.33	7.62	4.59												
<i>Distorsio decussata</i>					4.76	8.33	7.62	2.71												
BIVALVIA																				
Terebridae	2.38	1.03	0.16	0.04																
Total Mollusca	4.76	2.06	0.19	0.08	9.52	16.70	7.92	7.30												
FISHES																				
Fish					9.50	16.70	36.80	31.05	12.50	11.40	30.90	12.64	38.4	38.5	37.4	61.17	16.10	27.30	14.60	24.81
													6	0	0					
Batrachoididae													15.3	15.4	13.4	9.26				
													8	0	0					
Cynoglossidae																	9.70	18.20	13.70	11.16
Total Fishes					9.50	16.70	36.80	31.05	12.50	11.40	30.90	12.64	53.84	53.90	50.80	70.43	25.80	45.50	28.30	35.97
Digested material			47.20				35.10				27.10				3.52				11.40	

Prey items observed in stomachs of five species of elasmobranchs captured in the Eastern Tropical Pacific Ocean of Colombia. %O is frequency of occurrence, %N is percentage by number, %W is percentage by weight

Table 4: Diet breadth of elasmobranchs caught

Species	N	Percentage by number (%N)	
		B_i	B_A
<i>Mustelus lunulatus</i>	13	2.40	0.12
<i>Dasyatis longa</i>	5	2.57	0.39
<i>Rhinobatos leucorhynchus</i>	5	3.05	0.51
<i>Raja velezi</i>	3	2.60	0.53
<i>Zapteryx xyster</i>	6	4.57	0.71

Levins' measure of niche breadth (B_i) and standardized B_i (B_A) calculated with percentage by number (%N) data. N: total number of prey categories used.

ery. Predators found during this activity, feed mainly on benthic infauna (prey species living in the sediment), epibenthic fauna (prey species living on the surface of the sediment), benthic prey (prey species living on the bottom) or demersal prey (prey species living near the bottom but not linked to it). Therefore, the results of this research are restricted to elasmobranchs that feed on prey species living on the sediment surface or inside the sediment. However, our results contribute significantly to increase the knowledge on the feeding ecology of the other 138 species of fishes coexisting in the same habitat [31], with similar food and space requirements.

Mustelus species have been reported to feed mainly on crustaceans and fish [18,25,26,32], although cephalopods would also be important in their diets [33]. Despite the large number of taxa found in the stomach contents of *M. lunulatus*, one species of stomatopod accounted for most of the prey consumed (stomatopoda = %IRI = 96.9). Therefore, *M. lunulatus* in coastal waters of the Colombian Eastern Tropical Pacific have a specific dietary preference. Besides Ellis et al. [18], this is the only other report of feeding specialization in *Mustelus*, a genus that has been classified as opportunistic polyphagous [34,35].

The diet of *Raja velezi* in the neritic zone of the Eastern Pacific Ocean of Colombia showed a low number of prey

items in compared with other studies [20,36,37] that found polychaetes, copepods, amphipods, Myscridacea, stomatopods, cephalopods and bivalves. However, the preference of *R. velezi* to feed on decapods and fishes detected in this study was similar to that reported for *R. naevus* [18], *R. clavata* [37], *R. radiata* [38], and *R. brachyura*, *R. montagui* and *R. eglanteria* [39], which also feed upon decapods or fish. Although *R. velezi* is a deep-water ray with a widespread distribution, our results are the first description of the diet of this species in the Tropical Pacific Ocean. *Rhinobatos leucorhynchus* has been identified as a crustacean predator, with preference for penaeid shrimp [22,34,40,41], although portunid crabs have also been reported as alternative prey [42].

Very little is known of the diet of *Dasyatis longa* and *Zapteryx xyster* and this study is the first record on the diet of these species. Similar to other benthic species *Dasyatis longa* feed on crustacean decapods (stomatopods) and fishes. The preferences for these dietary items have also been reported for *Dasyatis americana* in the Caribbean Sea [43]. *Zapteryx xyster* showed the widest feeding spectrum of all elasmobranch studied. This species included stomatopods, decapods and fishes in its diet and, was the most generalist species. Bornatowski et al. [44] reported similar results for *Z. brevirostris* from South Brazil.

Macpherson [45] and Cortés [46] suggested that the relatively big body size of elasmobranch makes it easy for them to expand their feeding spectrum, preying simultaneously on the pelagic and benthic communities. Our study shows that diet breadth was reduced, and prey item distribution by number and weight were similar. Moreover, all studied elasmobranch fed on three main items: Squillidae, Penaeidae and fish. This can be explained by the benthic habits and similarity of sizes and dental structure between *M. lunulatus* and *D. longa*, as well as between *Rhinobatos leucorhynchus*, *Raja velezi* and *Z. xyster*.

The complete dietary overlap detected between *M. lunulatus* and *D. longa*, their similar geographical distribution in the study area, and their diurnal activity, suggest that these

Table 5: Niche overlap of elasmobranchs caught

Species A	Species B				
	<i>M. lunulatus</i>	<i>D. longa</i>	<i>R. leucorhynchus</i>	<i>R. velezi</i>	<i>Z. xyster</i>
<i>M. lunulatus</i>	1.000	-	-	-	-
<i>D. longa</i>	0.930	1.000	-	-	-
<i>R. leucorhynchus</i>	0.097	0.072	1.000	-	-
<i>R. velezi</i>	0.035	0.208	0.729	1.000	-
<i>Z. xyster</i>	0.343	0.515	0.603	0.935	1.000

Niche overlap between five species of elasmobranch caught in the Eastern Tropical Pacific Ocean of Colombia. Values were calculated using the Pianka measure, indicating the extent to which the diet of Species A overlaps with the diet of Species B. Significant diet overlap (>0.6).

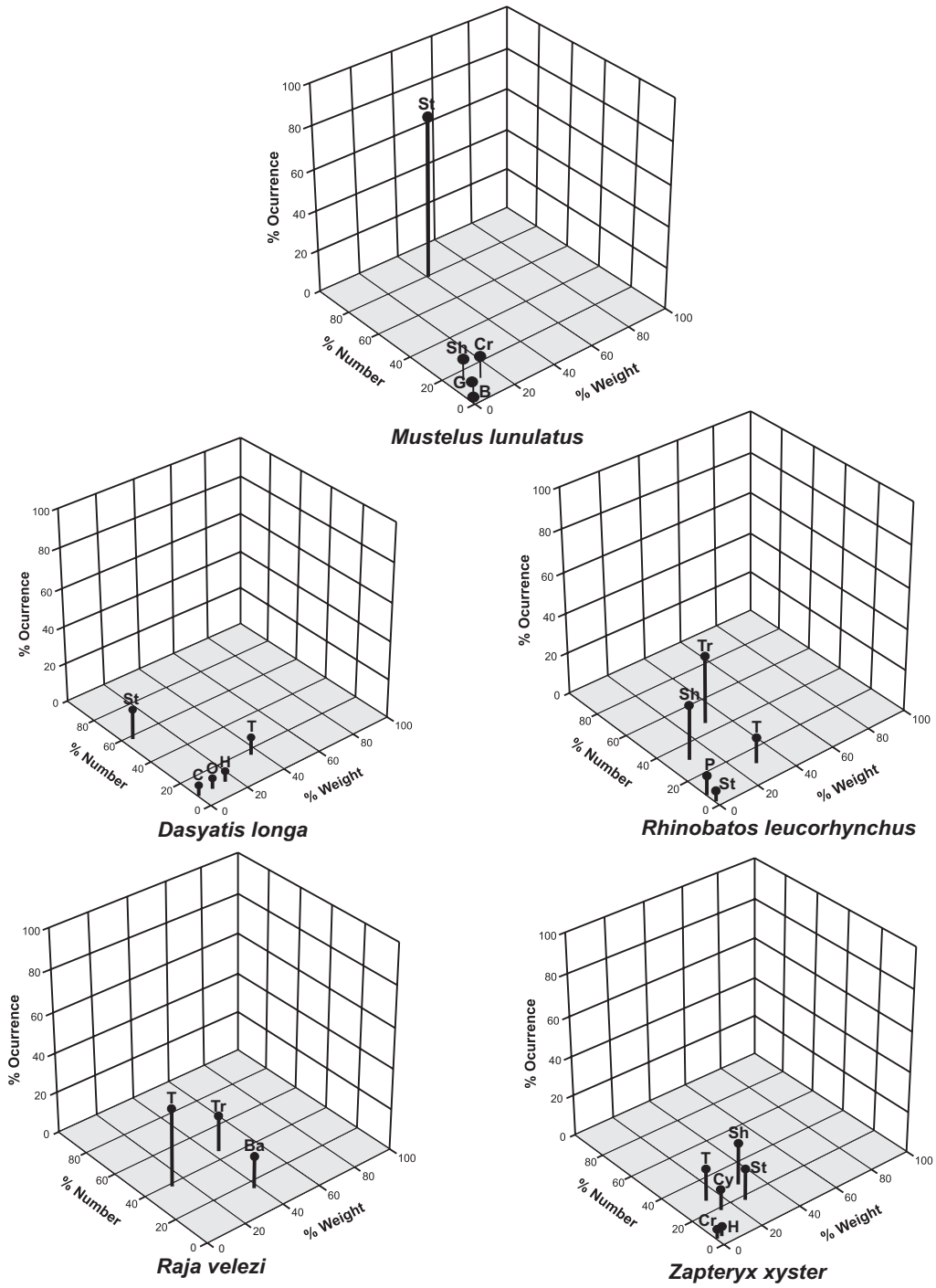


Figure 2

Three dimensional graphical representation of the relative importance of prey in the diet of five elasmobranch species: percentage by number (%N), percentage by weight (%W) and frequency of occurrence (%O). Prey items are: St) Stomatopods, Cr) Crabs, Sh) Shrimps, B) Bivalvia, G) Gasteropoda, T) Teleost, O) Olividae, H) Hippidae, C) Cymatidae, Tr) *Trachypenaeus*, P) Portunidae, Ba) Batrachoididae, Cy) Cynoglossidae.

species could show a competitive exclusion trend for food resources [47].

However, the dietary overlap would be reduced when the deep distribution is considered because *M. lunulatus* occupies deep waters and *D. longa* shallow waters. In contrast the dietary overlap detected between *Rhinobatos leucorhynchus*, *Raja velezi* and *Z. xyster* would be significant, since these species have morphological (size and position mouth) and behavioral similarities.

The dietary overlap between *Rhinobatos leucorhynchus* – *Raja velezi* and *R. leucorhynchus* – *Z. xyster*, based on the presence of *Trachypenaeus* shrimp and fishes, could be compensated by differential diel feeding activity. *Rhinobatos leucorhynchus* feed mainly during the day whereas *R. velezi* and *Z. xyster* feed mainly at night. Furthermore, 89% of *Rhinobatos* whose stomachs were analyzed were caught at a deep of 15–30 m, whereas all *Raja* and *Zapteryx* specimens were caught at a depth of 40 m. In contrast, the dietary overlap between *R. velezi* and *Z. xyster*, species with similar feeding activity and bathymetric distribution, suggests that they share only a fraction of their feeding niche, protecting a portion as an uncontested space refuge [47]. In this sense, mantis shrimp would be the feeding refuge of *Z. xyster*. Dietary overlap between *Z. xyster* and *R. velezi* was fish unidentified and decapods; however this food category was so indeterminate that the evidence of effective dietary overlap is weak. Furthermore, this wide category of prey item overestimate dietary overlap and underestimate niche breadth.

Values of overlap based on simulations suggest that the species studied showed low dietary overlap (mean = 0.44). In this study only five species of elasmobranchs were examined. In the study area at least another 12 species of elasmobranchs [31,48] and six species of fishes [31] that share morphological and ecological characteristics with the studied species have been reported. However, all these species and the elasmobranchs that we studied would conform apackaging throughout a feeding axis, and the temporal dynamics of this packaging could be determined by: (i) variety and productivity of available resources, (ii) short-term and long-term environmental variability, and (iii) niche overlap [49,50]. Therefore, if the packaging is modulated by resource availability [50] then the shrimp trawl fishery in the neritic zone of Eastern Pacific of Colombia has severely affected the natural availability of prey for elasmobranch communities [51,52], affecting their structure.

Conclusion

Shrimps (penaeid and stomatopods) and benthic fishes were the main component of the diet of the elasmobranchs studied. Diet breadth and overlap were relatively

low. Determination of food resource partitioning among the batoid species studied was not possible. However, we identified partitions in other niche axes (time of feeding activity and habitat used). It is possible to assume that diffuse competition could be exceeding the biunivocal competition among the studied species. Therefore, this assemblage would have a strong tendency to trophic guild formation.

Methods

Study area and sampling

Stomachs were collected from elasmobranchs caught by bottom trawling with commercial shrimp nets (2 m mouth, 50 mm mesh size), during three cruises in the central fishing area of Eastern Pacific of Colombia (Fig. 3). Twenty four trawls were made over the continental shelf (5–60 m depth) from the bottom prawn trawler vessel *Arraijan* from June to November 2001.

The total length of captured elasmobranch was recorded and the stomachs of all specimens were extracted on board, fixed in 10% formalin, labeled, bagged and transported to the laboratory. Once in the laboratory an incision was made along the longitudinal axis and the stomach contents were emptied into a 1000 ml plastic bucket for rinsing and sorting. Food items were identified to the lowest taxonomic level possible. Numbers and weights (to the nearest 0.1 g) of food items were recorded after the items were dried with blotting paper to remove surface moisture.

Numerical analysis

The quantitative importance of each prey group in the diet of elasmobranch species was estimated by using the frequency of occurrence (%O), percentage by number (%N) and percentage by weight (%W) of prey items in stomachs [28] as:

$$\%O = \frac{n}{N} \times 100 \quad \%N = \frac{N_n}{N_p} \times 100 \quad \%W = \frac{P_p}{P_t} \times 100$$

where *n* = number of stomachs that have the prey *i*, *N* = total number of analyzed stomachs; *N_n* = number of prey items of prey group *n* observed, *N_t* = total number of prey items of all prey groups, *P_p* = weight of prey items of prey group *p* observed and *P_t* = total weight of prey items of all prey groups. The contribution of each prey to the diet was also estimated with the Index of Relative Importance (IRI) and its standardized value (%IRI) [28] as:

$$IRI = (\%N + \%W) \times \%O \quad \%IRI = \frac{100 \times IRI_i}{\sum IRI_i}$$

where IRI_{*i*} is the IRI value for each prey category of prey *i*.

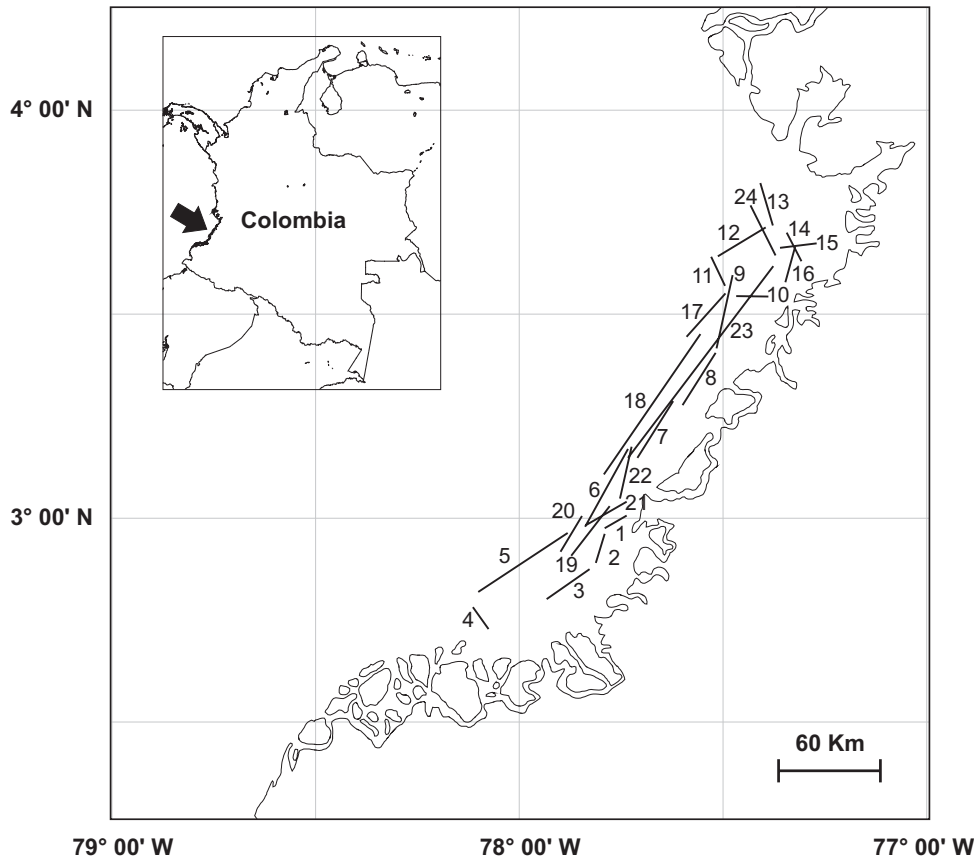


Figure 3
Eastern Pacific Ocean of Colombia (EPOC) showing the central fishing area. Each number represents one trawling episode.

The %O, %N and %W values of each prey were plotted following the method proposed by Cortés [28], which allows for an easy and adequate interpretation of prey importance in the diet of predators.

Differences in bathymetric distribution and diel (day-night) feeding activity were analyzed with a Kruskal-Wallis test [53]. For the analysis of feeding activity, the percentage by weight data of each stomach were grouped into four daily activity periods (morning, afternoon, night and dawn).

Diet breadth and overlap were estimated from %IRI values. Breadth of diet was calculated by using both Levins' index and its standardized form [54] as follows:

$$B_i = \left(\frac{1}{\sum P_j^2} \right) \text{ (iv) and } B_A = \left(\frac{B-1}{n-1} \right) \text{ (v)}$$

where B_i = Levins' measure of niche breadth, P_j = proportion of diet of predator that is made up of prey j , B_A = Levins' standardized niche breadth and n = number of prey categories. Levins' index of niche breadth (B_i) ranges from 1 to n , whereas, Levins' standardized niche breadth (B_A) ranges from 0 to 1; low values indicate diets dominated by few prey items (specialist predators) while higher values indicate generalist diets [54].

A numerical abundance dietary matrix of prey items was constructed to calculate diet overlap between elasmobranch species by using the Pianka index [54] as follows:

$$O_{jk} = \frac{\sum_i P_{ij} P_{ik}}{\sqrt{\sum_i P_{ij}^2 \sum_i P_{ik}^2}}$$

where O_{jk} = Pianka measure of niche overlap between species j and species k ; and P_{ij} and P_{ik} = proportions of pred-

ator j and k with prey i in their stomachs. Diet overlap increases as the Pianka index increases and overlap is generally considered to be biologically significant when the value exceeds 0.60 [29].

To evaluate the statistical significance of estimated overlaps, observed overlap values were compared with a distribution of expected values based on simulations of a null model. The distribution of this model was realized with 1000 repetitions using the random algorithm R3. The observed values were considered statistically different from the null distribution values if they were higher or lower than 95% of the simulated indices [30].

Authors' contributions

AFN conceived the study, conducted the *in situ* work and measurements, analyzed and interpreted the data and wrote the first draft of manuscript. PAM participated in the *in situ* work, analyzed the data and contributed to the critical review of the draft. AG analyzed the data and contributed to the critical review of the draft. All authors assisted in writing the manuscript and approved the final version.

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References

1. Wetherbee B, Cortés E: **Food consumption and feeding habits.** In *Biology of sharks and their relatives* Edited by: Carrier JF, Musick JA, Heithaus M. CRC press. USA; 2004.
2. Krebs CJ: **Ecology.** Benjamin Cummings, San Francisco, CA 2001.
3. Root RB: **The niche exploitation pattern of the blue-gray gnat-catcher.** *Ecol Mon* 1967, **37**:317-350.
4. Schoener TW: **Trophic structure, reproductive success, and growth rate of fishes in natural and modified head-water stream.** *Can J Fish Aquat Sci* 1974, **39**:968-978.
5. Zaret TM, Rand AS: **Competition in tropical stream fishes: support for the competitive exclusion principle.** *Ecology* 1971, **52**(2):336-342.
6. Hixon MA: **Competitive interactions between California reef fishes of the genus *Embiotoca*.** *Ecology* 1980, **61**(4):918-931.
7. Platell ME, Potter IC: **Partitioning of food resources among 18 abundant benthic carnivorous fish species in marine waters of the lower west coast of Australia.** *J Exp Mar Biol Eco* 2001, **261**:31-54.
8. Ross ST: **Resource partitioning in fish assemblages: a review of field study.** *Copeia* 1986:352-388.
9. Smale MJ: **Occurrence and feeding of three shark species, *Carcharhinus brachyurus*, *C. obscurus* and *sphyrna zygaena*, on the Eastern Cape coast of South Africa.** *S Afr J Mar Sci* 1991, **11**:31-42.
10. Vaske-Junior T, Rincón-Filho G: **Conteúdo estomacal dos tubarões azul (*Prionace glauca*) e anequim (*Isurus oxyrinchus*) en águas oceanicas no sul do Brasil.** *Rev Brasil Biol* 1998, **58**(3):445-452.
11. Carrasón M, Stefanescu C, Cartes JE: **Diets and bathymetric distributions of two bathyal sharks of the Catalan deep sea (Western Mediterranean).** *Mar Ecol Prog Ser* 1992, **82**:21-30.
12. Cortés E, Gruber SH: **Diet, feeding habits, and estimates of daily ration of young lemon shark, *Negaprion brevirostris* (Poey).** *Copeia* 1990:204-218.
13. Simpfendorfer CA: **Diet of the Australian Sharpnose shark *Rhizoprionodon taylori*, from northern Queensland.** *Mar Fresh Res* 1992, **49**:757-761.
14. Wetherbee BM, Lowe CG, Crowe GL: **Biology of the Galapagos shark, *Carcharhinus galapagensis*, in Hawaii.** *Environ Biol Fish* 1996, **45**:299-310.
15. Heithaus MR: **The biology of tiger sharks, *Galeocerdo cuvier*, in Shark Bay, Western Australia: sex ratio, size distribution, diet, and seasonal changes in catch rates.** *Environ Biol Fish* 2001, **61**:25-36.
16. Bethea D, Carlson JK, Buckel JA, Satterwhite M: **Ontogenetic and site-related trends in the diet of the atlantic sharpnose shark *Rhizoprionodon terraenovae* from the northeast gulf of Mexico.** *Bull Mar Sci* 2006, **78**(2):287-307.
17. Brickle P, Laptikhovsky V, Pompert J, Bishop A: **Ontogenetic changes in the feeding habits and dietary overlap between three abundant rajoids species on the Falkland Island's shelf.** *J Mar Biol Ass UK* 2003, **83**:1119-1125.
18. Ellis JR, Pawson MG, Shackley SE: **The comparative feeding ecology of six species of shark and four species of ray (Elasmobranchii) in the North-East Atlantic.** *J Mar Biol Ass UK* 1996, **76**:89-106.
19. Bethea D, Buckel JA, Carlson JK: **Foraging ecology of the early life stages of four sympatric shark species.** *Mar Ecol Prog Ser* 2004, **268**:245-264.
20. Orlov AM: **The diet and feeding habits of some Deep-Water benthic skates (Rajidae) in the pacific waters off the Northern Kuril islands southeastern Kamchatka.** *Alas Fish Res Bull* 1998, **5**(1):1-17.
21. Platell ME, Potter IC, Clarke KR: **Resource partitioning by four species of elasmobranch (Batoidea:Urolophidae) in coastal waters of temperate Australia.** *Mar Biol* 1998, **131**:719-734.
22. White WT, Platell ME, Potter IC: **Comparison between the diets of four abundant species of elasmobranch in a subtropical embayment: implications for resource partitioning.** *Mar Biol* 2004, **144**:439-448.
23. Papastamatiou YP, Wetherbee BM, Lowe CG, Crow GL: **Distribution and diet of four species of carcharhinid shark in the Hawaiian Islands: evidence for resource partitioning and competitive exclusion.** *Mar Ecol Prog Ser* 2006, **320**:239-251.
24. Mejía-Falla PA, Navia AF, Giraldo A: **Notas biológicas de la raya ocelada (*Zapteryx xyster*) en la zona central de pesca del Pacífico colombiano.** *Invest Mar* 2006, **34**(2):181-185.
25. Navia AF, Giraldo A, Mejía-Falla PA: **Notas sobre la biología y dieta del toyo vieja (*Mustelus lunulatus*) de la zona central de pesca del Pacífico colombiano.** *Invest Mar* 2006, **34**(2):217-222.
26. Gómez G, Zapata LA, Franke R, Ramos GE: **Hábitos alimentarios de *Mustelus lunulatus* y *M. henlei* (Pisces: Triakidae) colectados en el Parque Nacional Natural Gorgona, Pacífico colombiano.** *Bol Inv Mar Cost* 2003, **32**:219-231.
27. Mejía-Falla PA, Navia AF, Mejía-Ladino LM, Acero A, Rubio E: **Tiburones y rayas de Colombia (Pisces: Elasmobranchii). Lista revisada, actualizada y comentada.** *Bol Inv Mar Cost* 2007 in press.
28. Cortés E: **A critical review of methods of studying fish feeding base on analysis of stomach contents: application to elasmobranch fishes.** *Can J Fish Aquat Sci* 1997, **54**:726-738.
29. Zares TM, Rand AS: **Competition in tropical stream fishes: support for the competitive exclusion principle.** *Ecology* 1971, **52**:336-342.
30. Gotelli NJ, Graves GR: **Null Models in ecology.** Smithsonian Institution Press. Washington. DC; 1996.
31. Navia AF, Mejía-Falla PA: **Aspectos biológicos, ecológicos y pesqueros de la fauna capturada incidentalmente en la pesca de camarón de aguas someras en el Pacífico colombiano: informe final y propuesta metodológica para monitoreos a bordo de embarcaciones arrastreas.** *Doc tec No FS0206* 2006:42p.
32. Rojas JR: **Reproducción y alimentación del tiburón enano *Mustelus dorsalis* (Pisces: Triakidae) en el Golfo de Nicoya, Costa Rica: Elementos para un manejo sostenible.** *Rev Biol Trop* 2006, **54**(3):861-871.

33. Capapé C: **Observations sur le régime alimentaire de 29 Selaciens pleurotêrmes des côtes tunisiennes.** *Arch Inst Pasteur Tunis* 1975, **52**:395-414.
34. Talent L: **Food habits of the gray smoothhound *Mustelus californicus*, the Brown-smoothhound, *Mustelus henlei*, the shovelnose guitarfish *Rhinobatos productus* and the bat ray *Myliobatis californica* in Elkhorn slough, California.** *Cal Fish Game* 1982, **68**:224-234.
35. Chiaramonte G, Pettovello P: **The biology of *Mustelus schmitti* in southern Patagonia, Argentina.** *J Fish Biol* 2000, **57**:930-942.
36. Galvan-Magaña F, Nienhuis HJ, Klimley AP: **Seasonal abundance and feeding habits of sharks of the lower gulf of California, México.** *Calif Fish and Game* 1989, **75**(2):74-84.
37. Morato T, Solá E, Grós MP, Menezes G: **Diets of thornyback ray (*Raja clavata*) and tope shark (*Galeorhinus galeus*) in the bottom longline fishery of the Azores, northeastern Atlantic.** *Fish Bull* 2003, **101**:590-602.
38. Pedersen SA: **Feeding habits of starry ray (*Raja radiata*) in West Greenland waters.** *ICES J Mar Sci* 1995, **52**:43-53.
39. Schwartz FJ: **Biology of the clearnose skate *Raja eglanteria* from North Carolina.** *Florida Scientist* 1996, **59**(2):82-95.
40. Ismen A, Yigin C, Ismen P: **Age, growth, reproductive biology and feed of the common guitarfish (*Rhinobatos rhinobatos* Linnaeus, 1758) in Iskenderun Bay, the eastern Mediterranean sea.** *Fish Res* 2007, **84**(2):263-269.
41. Shibuya A, Rosa RS, Soares MC: **Note on the diet of the guitarfish *Rhinobatos percellens* (Walbaum, 1792) (Elasmobranchii: Rhinobatidae) from the coast of Paraíba, Brazil.** *Acta Biologica Leopoldensia* 2005, **27**(1):63-34.
42. Capapé C, Zaouali J: **Diet of two selachians common to the Gulf of Gabes (Tunisia): *Rhinobatos rhinobatos* (Linné, 1758) and *Rhinobatos cemiculus* (Geoffroy Sainte-Hilarie, 1817).** *Arch Inst Pasteur Tunis* 1979, **56**(3):285-306.
43. Stokes MD, Holland ND: **Southern Stingray (*Dasyatis americana*) feeding on lancelets (*Branchiostoma floridae*).** *J Fish Biol* 1992, **41**:1043-1044.
44. Bornatowski H, Abilhoa V, Freitas MO: **Alimentacao da raia-viola *Zapteryx brevirostris* na baía de ubatuba-enseada, Sao franciso do sul, Santa Catarina, Brasil.** *Estud Biol* 2005, **27**(61):31-36.
45. Macpherson E: **Resource partitioning in a Mediterranean demersal fish community.** *Mar Ecol Progr Ser* 1981, **4**:183-193.
46. Cortés E: **Standardized diet compositions and trophic levels of sharks.** *ICES J Mar Sci* 1999, **56**:707-717.
47. Pianka ER: *Evolutionary Ecology* 6th edition. Addison Wesley; 2000.
48. Navia AF: **Aspectos de la biología de los elasmobranquios capturados como fauna acompañante del camarón en aguas someras del Pacífico colombiano.** B.Sc. Thesis, Universidad del Valle, Cali, Colombia 2002.
49. MacArthur RH: **Species packing and competitive equilibrium among many species.** *Theo Pop Biol* 1970, **1**:1-11.
50. May RM, MacArthur RH: **Niche overlap as a function of environmental variability.** *Proc Nat Acad Sci* 1972, **69**:1109-1113.
51. Invenmar: **Informe del estado de los ambientes marinos y costeros en Colombia: 2000.** Serie documentos generales No. 3, Santa Marta, Colombia 2001.
52. Zapata LA, Rodríguez G, Beltrán B: **Prospección de los principales bancos de pesca en el Pacífico colombiano, durante noviembre de 1998.** *Boletín científico INPA* 1999, **6**:111-175.
53. Zar J: *Biostatistical Análisis* 4th edition. Pearson education. India; 1999.
54. Krebs CJ: *Ecological methodology* 2nd edition. Benjamin Cummings press, Menlo Park, California; 1999.

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