

Faunal Remains Associated with Late Saladoïd and Post-Saladoïd Occupations at Anse à la Gourde, Guadeloupe, West Indies: Preliminary Results

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ABSTRACT: The study of the large faunal assemblage recovered from the Saladoïd (400-600 A.D.) and Troumassoïd (800-1400 A.D.) occupation levels in a trench at the site of Anse à la Gourde, Guadeloupe, permitted an evaluation not only of the richness and diversity of the faunal spectrum, but also of the ecosystems exploited and the techniques of capture, based on the taxonomic list and the size estimates for snappers (Lutjanidae), grunts (Haemulidae), and parrotfishes (Scaridae). The large sample size also permitted a statistical analysis of the data obtained from each archaeological layer. These analyses indicated significant changes in the exploitation of the animal resources through time.

KEY WORDS: ZOOARCHAEOLOGY, GUADELOUPE, WEST INDIES, FISH, GRUNT, SNAPPER, PARROTFISH, HAEMULIDAE, LUTJANIDAE, SCARIDAE

RESUMEN: La capas saladoïdes tardías (400 – 600 A.C.) y troumassoïdes (800 – 1 400 A.C.) del sondeo realizado en el sitio Anse à la Gourde, Guadeloupe, permitió caracterizar tanto el espectro de fauna (gracias a los índices de riqueza y de diversidad), así como los ecosistemas explotados y las técnicas de captura utilizadas (gracias al conjunto faunístico específico y a la talla de ciertos peces: Haemulidae, Lutjanidae y Scaridae). La gran cantidad de huesos permitió también testar estadísticamente las informaciones obtenidas sobre cada capa arqueológica. Gracias a estos análisis se pudo observar que ciertas mutaciones eran significativas.

PALABRAS CLAVE: ARQUEOZOOLOGÍA, GUADELOUPE, PEQUEÑAS ANTILLAS, PEZ, HAEMULIDAE, LUTJANIDAE, SCARIDAE

RÉSUMÉ: Les couches saladoïdes tardives (400 – 600 ap. J.-C.) et troumassoïdes (800 - 1400 ap. J.-C.) d'un sondage du site l'Anse à la Gourde ont livré un abondant matériel faunique. L'étude de ces restes a permis d'une part, de caractériser la richesse et la diversité du spectre de faune, et d'autre part, de reconstituer les écosystèmes exploités et les techniques de capture (au travers de la liste taxonomique et par l'estimation de la taille de trois familles de poissons: Haemulidae, Lutjanidae et Scaridae). La grande quantité d'ossements a permis d'étayer statistiquement les informations obtenues sur chaque couche archéologique. Ces approches indiquent quelques variations significatives de l'exploitation des ressources animales dans le temps.

MOTS CLEFS: ARCHEOZOOLOGIE, GUADELOUPE, PETITES ANTILLES, POISSON, GORETTE, VIVANEAU, POISSON-PERROQUET, HAEMULIDAE, LUTJANIDAE, SCARIDAE

INTRODUCTION

The importance of marine resources in the pre-Columbian Caribbean is suggested by the island

environment and substantiated by several zooarchaeological studies (Wing, 1977, 1989, 1994, 1995; Wing & Scudder, 1983; deFrance, 1988; Carlson, 1995; Keegan, 1997). Analyses of faunal

assemblages from various sites in the Lesser Antilles provide data on the subsistence patterns and on the environments that were exploited by the pre-Columbian Amerindians (Rouse, 1989a; Watters & Rouse, 1989; Watters, 1998). Guadeloupe is an island for which we have a small amount of data. The site of Anse à la Gourde in Grande-Terre provides a long chronological sequence with large refuse areas. Based on one refuse area, this zooarchaeological study was undertaken to investigate the nature of both marine and terrestrial exploitation in a chronological perspective (molluscs have been excluded from this study). The large faunal samples from successive levels of the site provide an understanding of the prehistoric economy, the prehistoric fisheries and the long term effects of fishing on the natural resources.

This article deals with the preliminary results from the material recovered during the 1997-1998 excavations. Although we are still in the process of studying the new data recovered during the 1999-2000 excavations, the initial results confirm the main interpretations discussed here.

MATERIAL AND METHODS

Anse à la Gourde is one of the major village sites located along the eastern coast of Grande-Terre in Guadeloupe (Figures 1, 2). It occupies 4.5 hectares (Figure 3). The coastal strip is a refuse area and the remains of house posts, burials and fireplaces are located inland from the shore. A trench through the large refuse area indicated a deeply stratified unit (Z64S93C01), corresponding to an Amerindian Ceramic period occupation from 400 to 1400 A.D. (Rouse, 1989b, 1992, 1995; Delpeuch *et al.*, 1997). This last unit was selected for analysis because it is very rich in well preserved faunal remains (vertebrates, crabs, and urchins) and it covers all the occupation periods.

The trench measuring 1 x 1 square meter was excavated in 10 cm arbitrary levels. The material from this unit was water sieved through a 2.8 mm gauge screen. The stratigraphy reveals nine successive occupations (Figure 4). The analysis of the ceramics reveals that occupation levels III, IV, VI, IX, X, and XVIII are of the late Saladoïd period, ranging from 400 to 600 A.D. Occupation levels XX, XXI, XXIII, and XXIV are different phases of the Troumassoïd period, ranging from 800 to 1400 A.D. (Delpeuch *et al.*, 1997). In order to

point out differences between samples, the data is grouped into five phases: Saladoïd 1 (S1 = levels III, IV and VI), Saladoïd 2 (S2 = IX, X, XVIII), Post-Saladoïd 1 (PS1 = XX and XXI), Post-Saladoïd 2 (PS2 = XXIII), and Post-Saladoïd 3 (PS3 = XXIV).

Spectrum of taxa

The faunal remains were sorted and identified to their lowest taxonomic levels using comparative collections at the following institutions: the Florida Museum of Natural History in Gainesville; The *Laboratoire d'Anatomie Comparée* of the *Muséum National d'Histoire Naturelle* in Paris; The Laboratory of Zooarchaeology of the *Centre de Recherches Archéologiques* in Valbonne; and the specimens collected by the author in 1997, which are now at the *Service Régional de l'Archéologie de Guadeloupe*.

The number of identified specimens (NISP), the minimum number of individuals calculated by paired elements (MNI), and the weight of the remains (in grams) were quantified for each taxon in each level. Each of these methods of quantification has its own advantage (see Chaplin, 1971; Ducos, 1975; Poplin, 1976a, 1976b, 1977; Grayson, 1984; Lyman, 1994). Comparisons were made between samples using both NISP and MNI.

An understanding of subsistence during antiquity, at least concerning the flesh meat portion of the diet, is based on the identified taxa and their relative abundances in each period. One dietary aspect is diversity, which can be measured by calculating the taxonomic richness, according to the size of samples. In order to check the reliability of samples in relation to their size, a rarefaction curve was constructed by plotting MNI on the *x* axis and the number of taxa (*S*) on the *y* axis. Richness was also estimated by applying the Margaleff index (*dI*) (1958, cited by Bobrowski & Ball, 1989) to the samples from each level. This index was calculated using the formula $dI = (S-1)/(\log_e N)$ where *S* is the number of species and *N* is the total NISP for each level. Diversity and homogeneity were estimated using the Simpson Reciprocity index (*H'*), calculated using the formula $H' = 1/\sum p_i^2$, where $p_i = n_i/N$, $n_i = \text{NISP per taxon}$, and *N* = total NISP for the sample (Grayson, 1984). In order to evaluate the differences between the samples, Chi² square tests of NISP across species were carried out (the MNI of some species was too small to be used in a Chi² test).

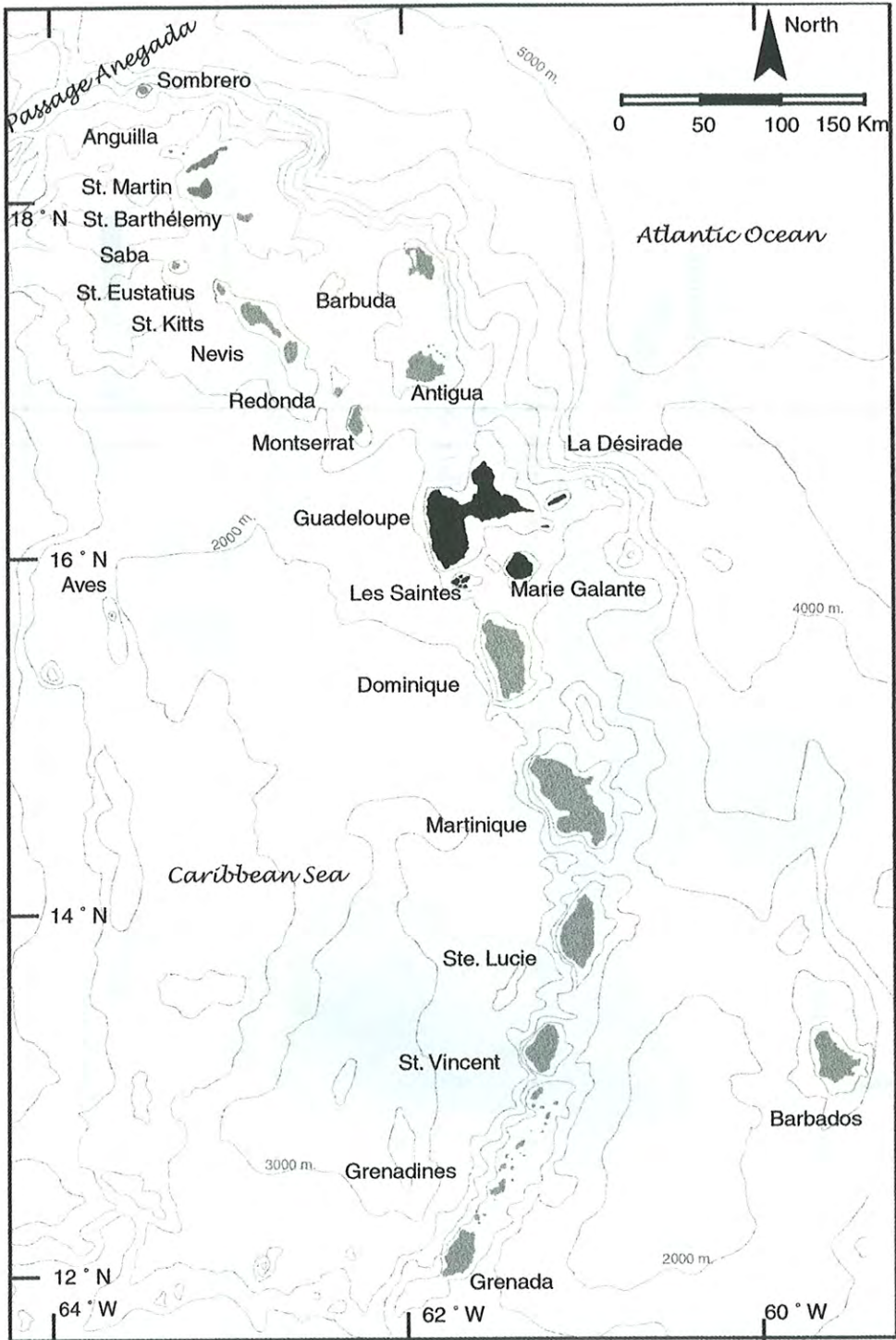


FIGURE 1
Map of the West Indies.

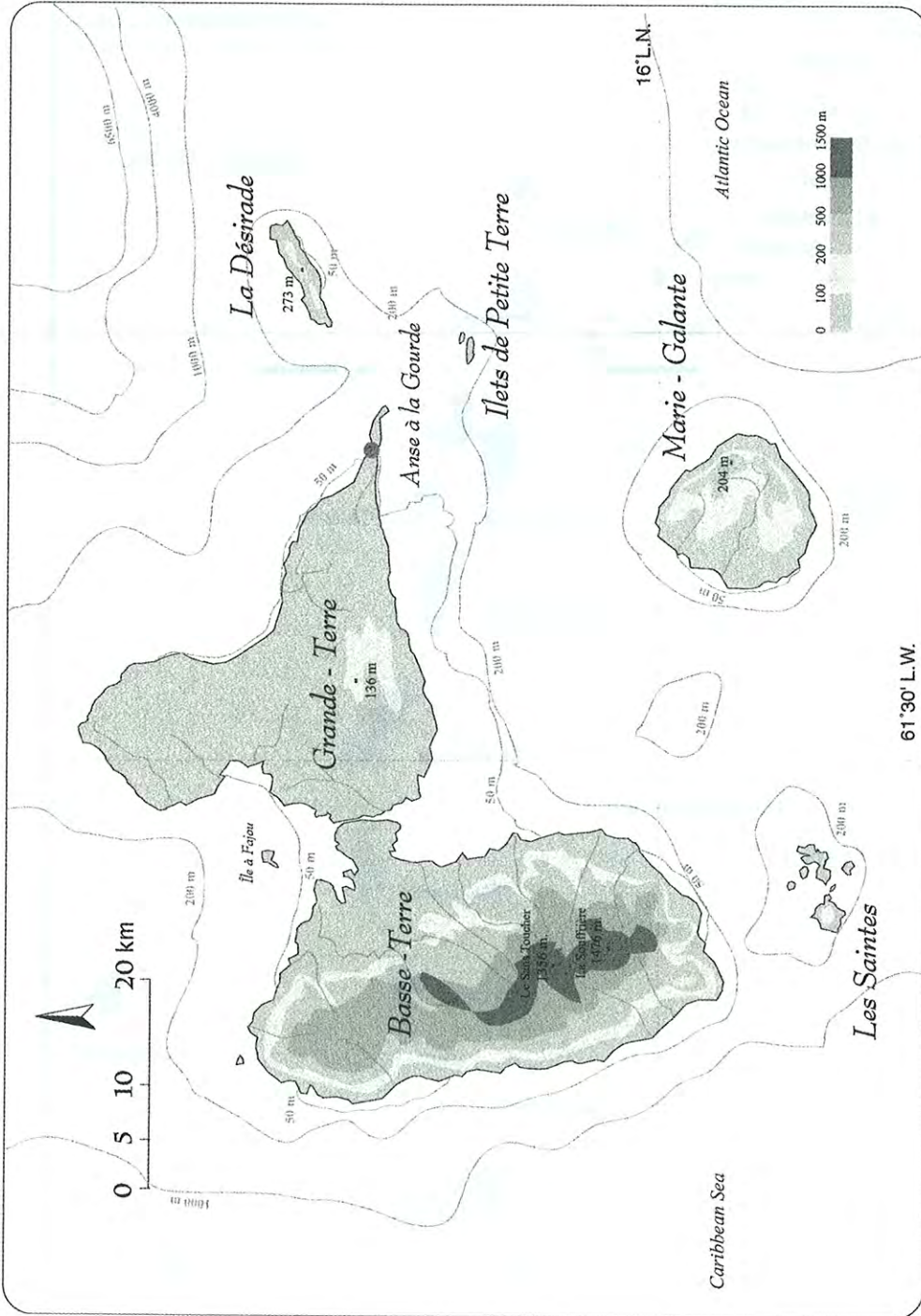
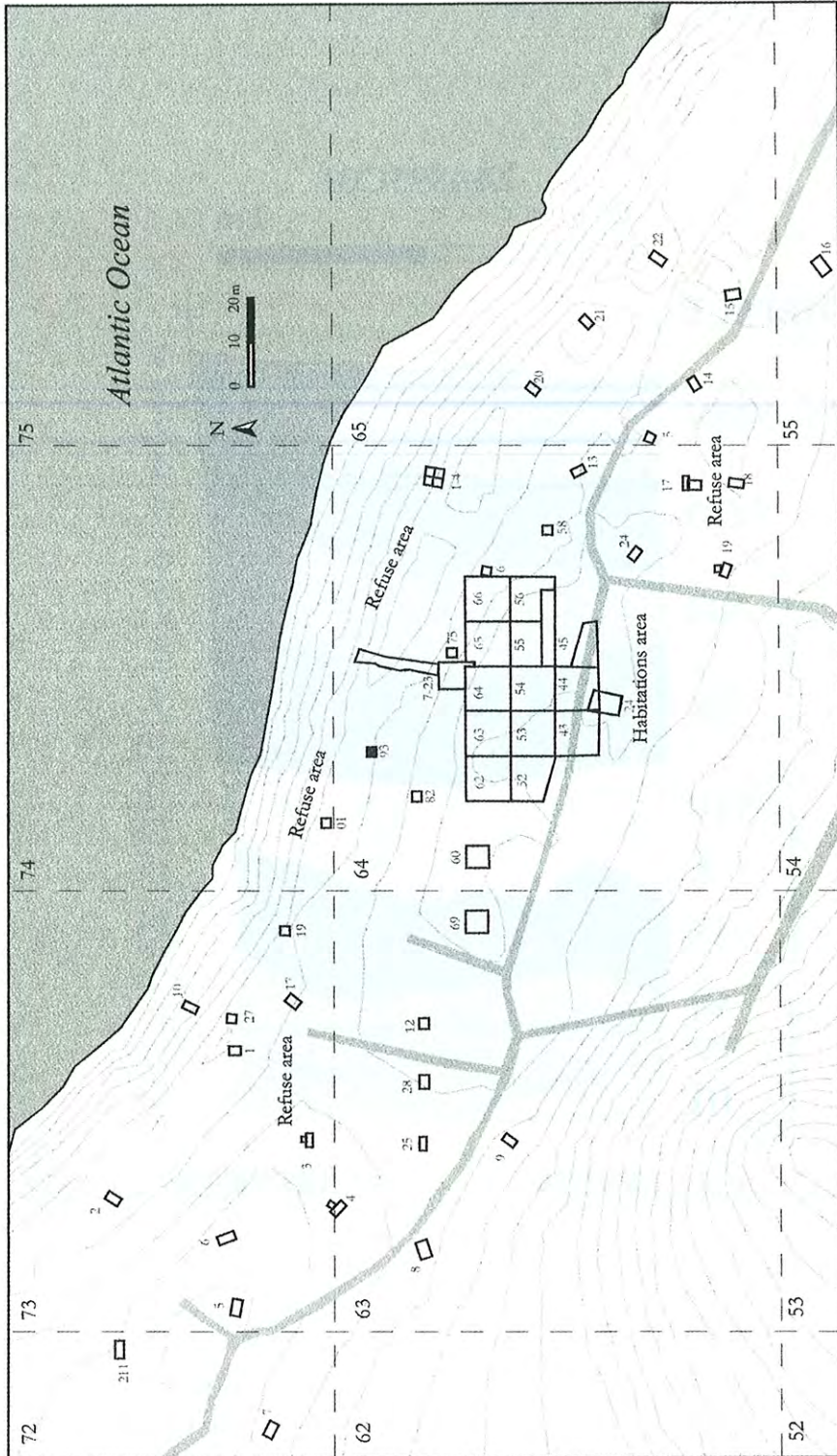


FIGURE 2
Location of Anse à la Gourde in Guadeloupe.



units
 Z64S93C01 in 1997
 1 m
 roads

Anse à la Gourde - Guadeloupe - French West Indies

FIGURE 3
Location of unit Z64S93C01 at Anse à la Gourde.

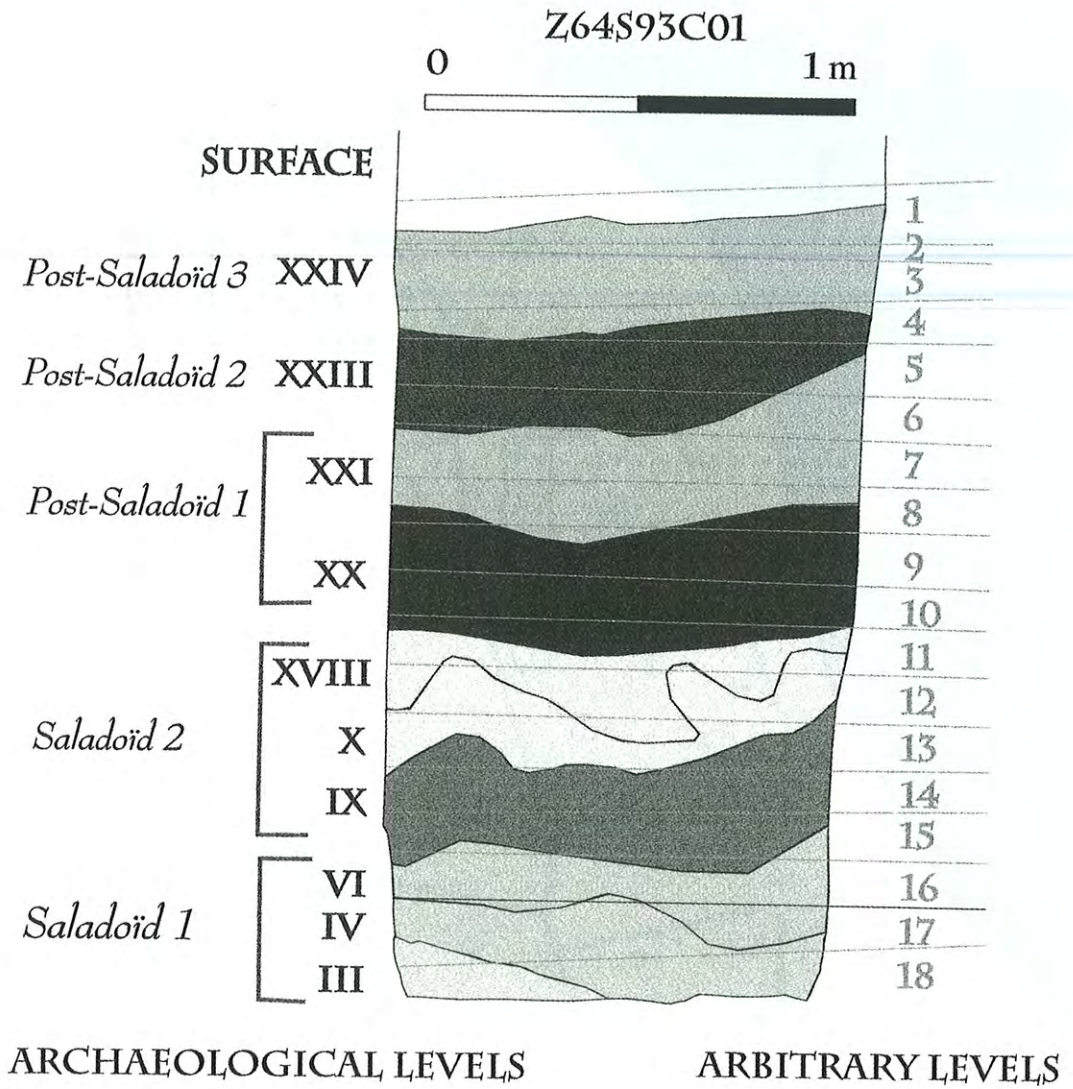


FIGURE 4
Arbitrary levels and archaeological level of unit Z64S93C01 at Anse à la Gourde.

Body size reconstruction

The sizes of fish from archaeological sites illustrate both the different ecosystems where the animals were probably caught, as well as the techniques used to catch them. There are many ways of estimating body size (live weight or standard length) based upon different measurements of fresh fish skeletal parts (Casteel, 1974, 1978; Desse, Desse-Berset & Rocheteau, 1987, 1989, 1996; Wheeler & Jones, 1989; Leach & Boocock, 1993, 1995; Béarez, 1995; Leach *et al.*, 1996a, 1996b, 1996c). For some Caribbean taxa, body weights are estimated using the first vertebrae width or otolith length (Reitz & Cordier, 1983; Adams, 1985; Reitz *et al.*, 1987; Reitz & Wing, 1999). Correlations between the body weight or length and otolith length or vertebral width did not seem to provide accurate estimates of weight for some taxa in this study (correlation coefficients were too small). As a result, various regression lines were fitted and equations were calculated for the different skeletal parts of the most common taxa from the samples, i.e., grunts (Haemulidae), snappers (Lutjanidae) and parrot-fishes (Scaridae). The comparative specimens used in producing the equations belong to the Greater and Lesser Antilles, and Florida. The "least squares method" was applied to determine the metrical relationship between live standard lengths and various measurements of skull elements (examples in Figure 5). As illustrated by the example of the *Sparisoma* standard length for all the taxa and skeletal parts (Figure 6), the standard error of the estimates was smaller for the power curve, than for the linear, the exponential, and the logarithmic curves. Consequently, the power curve was chosen to estimate the length of the archaeological specimens.

Equations were calculated at the level of species, genus and family when correlations were significant, as recommended by Desse & Desse-Berset (1996a, 1996b) (examples at the level of genus in Appendix 1).

Before choosing measurements for analysis, the random nature of the processes of destruction indicated by the distribution of body parts was verified for each sample of unit Z93S64C01, as suggested by Leach & Boocock (1994:73; 1995:27). All measurements are in millimetres and all weights are in grams. Details of the basic statistics of these formulas will be published elsewhere.

RESULTS AND INTERPRETATIONS

The taphonomic study concluded that all bones were dissociated and partly broken, and that all parts of the skeleton were present. The majority of bones were well-preserved, although some of them showed evidence of erosion possibly due to exposure to acidic conditions or weathering. None of the bones exhibited evidence of cut marks, gnaw marks or burning. Because their bones were also dissociated and broken, mammals such as dogs (*Canis familiaris*), agoutis (*Dasyprocta cf. leporina*), and rice rats (*Oryzomyini*) were probably eaten, though no evidence of butchery or cooking was present on their bones. These taxa are common in Lesser Antillean sites and they were undoubtedly consumed (Wing, 1995; Wing & Wing, 1995). The animal bones regularly thrown into a refuse area near the habitation during the occupations, were not disturbed by animals and survived the attacks from sand, sea, and weather elements, etc.

Richness, diversity and faunal assemblage

A total of 191,978 fragments were identified, at least by class, with 22,287 crab and sea urchin fragments, 27,864 fishes, 1,414 reptiles, 273 birds, and 4,199 mammal remains. These represent a minimum of 3,842 individuals (MNI). A total of 46 families (89 species) of vertebrates, crustacean, and sea urchins were identified (Table 1, Appendix 2).

Crustaceans and sea urchins were represented by seven families and 20 species (Table 1, Appendix 2). The land crabs (Gecarcinidae) and the land hermit crabs (Coenobitidae) were the most abundant. The fishes belong to 26 families and 55 species, of which the doctor-fishes (Acanthuridae), jacks (Carangidae), trigger-fishes (Balistidae), and parrot-fishes (Scaridae) were the most abundant. Among the reptiles, sea turtles (Cheloniidae) and iguanas (Iguanidae) were the most frequent taxa. Pigeons and doves (Columbidae) predominated among the birds. Rice rats (Cricetidae tribe *Oryzomyini*) and agoutis (*Dasyproctidae*) were the most abundant mammals.

From the bottom to the top of the stratigraphy, we observed a strong increase in the relative abundance of fish remains until the late Saladöid level (S2), and a slight decline after the Post-Saladöid

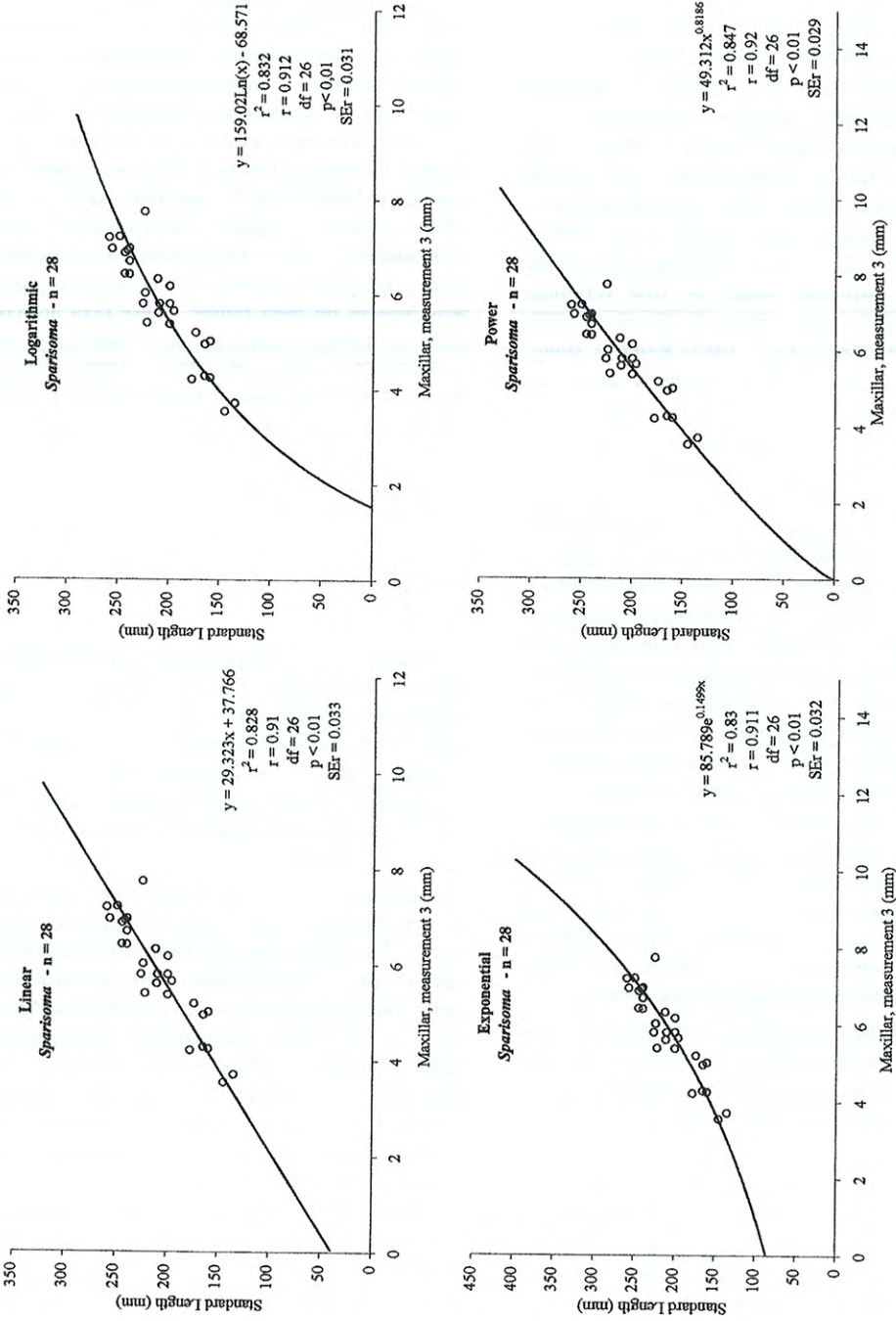


FIGURE 5
Examples of the best-fit curves (for *Sparisoma*).

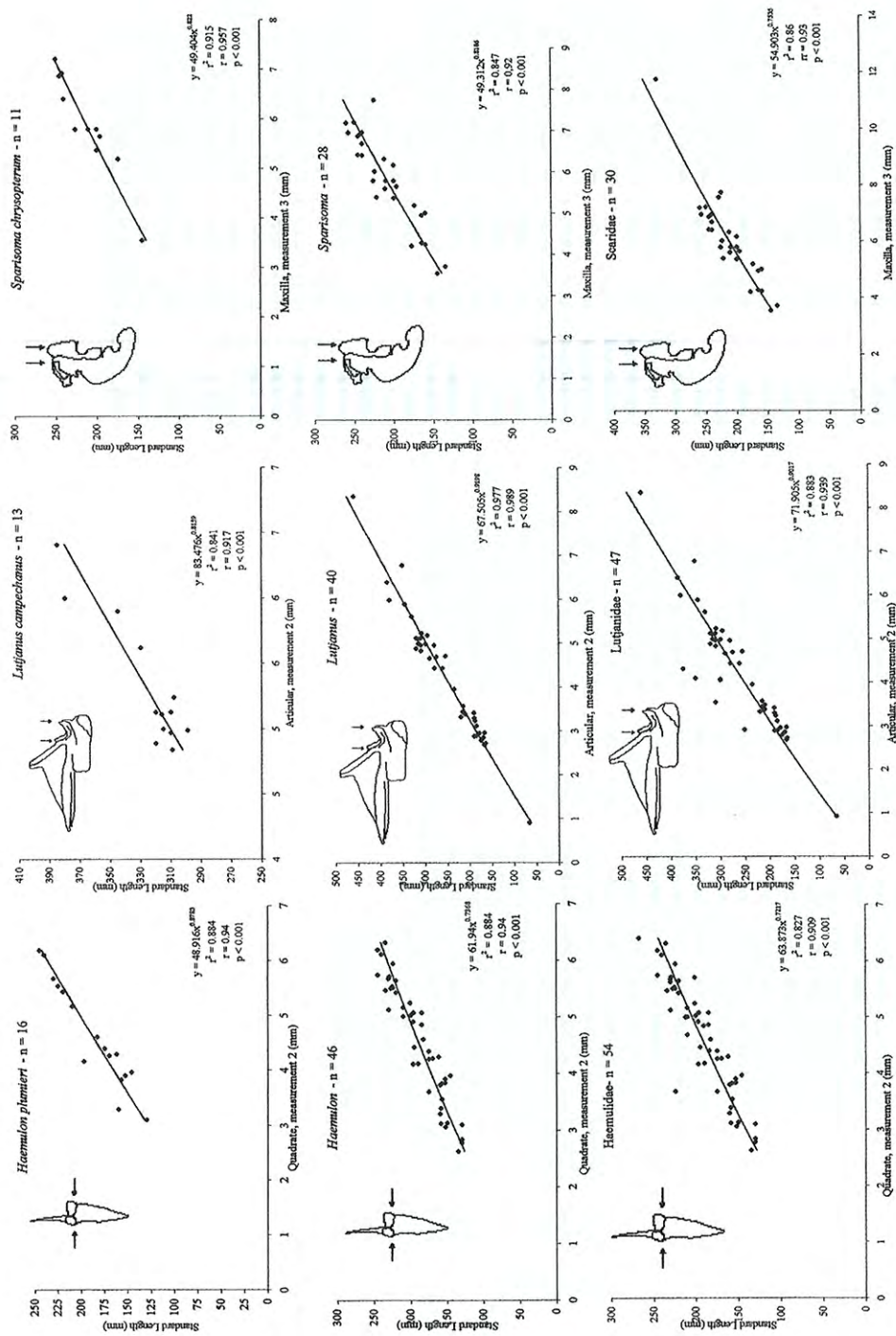


FIGURE 6
 Examples of allometric regressions for the estimation of the Standard Length at the rank of species, genus and family from the same measurement.

Class	Family	NISP	Weight (g.)	MNI	% NISP	% Weight	% MNI
Invertebrata	Palinuridae	187	12.5	8	0.8	1.3	0.3
	Coenobitidae	9 947	367.8	1 756	44.6	38.3	72.7
	Portunidae	33	1.0	2	0.2	0.1	0.1
	Xanthidae	998	88.2	16	4.5	9.2	0.7
	Gecarcinidae	5 032	325.0	573	22.6	33.8	23.7
	Ocypodidae	27	1.2	7	0.1	0.1	0.3
	Echinoidea	6 063	165.6	52	27.2	17.2	2.2
		7	22 287	961.3	2 414	100	100
Pisces	Carcharhinidae	8	0.6	2	0	0	0.2
	Dasyatidae	12	0.6	3	0	0	0.2
	Elopidae	6	0.6	2	0	0	0.2
	Albulidae	23	1.0	4	0.1	0.1	0.3
	Muraenidae	27	1.0	3	0.1	0.1	0.2
	Clupeidae	2 282	12.1	95	8.2	1.1	7.1
	Belontiidae	759	24.0	26	2.7	2.2	1.9
	Holocentridae	679	18.3	76	2.4	1.7	5.6
	Centropomidae	69	5.1	5	0.3	0.5	0.4
	Serranidae	620	76.1	31	2.2	6.9	2.3
	Priacanthidae	1	0	1	0	0	0.1
	Carangidae	5 467	172.6	204	19.6	15.6	15.1
	Lutjanidae	1 334	95.0	53	4.8	8.6	3.9
	Haemulidae	1 915	58.3	200	6.9	5.3	14.8
	Sparidae	26	3.7	2	0.1	0.3	0.2
	Sciaenidae	62	2.1	11	0.2	0.2	0.8
	Kyphosidae	13	0.2	2	0.1	0	0.2
	Pomacanthidae	1	0	1	0	0	0
	Sphyrenidae	41	1.7	4	0.2	0.1	0.3
	Labridae	1 123	39.0	41	4.0	3.5	3.0
	Scaridae	2 934	279.4	221	10.5	25.3	16.4
	Acanthuridae	6 153	211.1	297	22.1	19.1	22.0
	Scombridae	406	54.5	9	1.5	4.9	0.7
	Balistidae	3 842	48.2	50	13.8	4.4	3.7
	Ostraciidae	24	0.5	3	0.1	0	0.2
	Diodontidae	37	1.1	3	0.1	0.1	0.2
	26	27 864	1 106.8	1 349	100	100	100
Amphibia	Anura	2	0.1	1	0.2	0	5.3
Reptilia	Cheloniidae	902	125.5	2	63.8	88.4	10.5
	Iguanidae	388	14.9	8	27.4	10.5	42.1
	Lacertilia	122	1.5	8	8.6	1.1	42.1
		4	1 414	142.0	19	100	100
Aves	cf. Procellariidae	15	0.8	2	5.5	11.6	13.3
	cf. Anatidae	8	0.6	1	2.9	9.8	6.7
	cf. Laridae	1	0.1	1	0.4	1.2	6.7
	cf. Columbidae	246	5.1	10	90.1	77.1	66.6
	cf. Mimidae	3	0	1	1.1	0.3	6.7
		5	273	6.6	15	100	100
Mammalia	Chiroptères	74	1.3	9	1.8	0.7	20.0
	Canidae	34	3.3	2	0.8	1.9	4.5
	Oryzomyine	2 798	92.6	28	66.6	53.7	62.2
	Dasyproctidae	1 293	75.4	6	30.8	43.7	13.3
		4	4 199	172.6	45	100	100
Total	46	56 037	2389.3	3 842			

TABLE 1

NISP, Weight, MNI, and S by family and class for the unit as a whole.

Class	Family	Species	English name	French name	Ecosystems
Crustacea	Palinuridae	<i>cf. Panulirus sp.</i>	lobster	langouste épineuse	reef
	Coenobitidae	<i>Coenobita cybeatus</i>	hermit crab	bernard-l'hermite terrestre	terrestrial
	Portunidae	<i>cf. Callinectes sp.</i>	blue crab	crabe carcaïde bleu	reef
	Xanthidae	<i>cf. Carpillus corallinus</i>	coral crab	crabe corallien	reef
		<i>sp. 1</i>	sea crab	crabe de mer	reef
		<i>sp. 2</i>	sea crab	crabe de mer	reef
		<i>sp. 3</i>	sea crab	crabe de mer	reef
		<i>sp. 4</i>	sea crab	crabe de mer	reef
	Gecarcinidae	<i>cf. Cardisoma guanhumi</i>	great land crab	crabe terrestre	terrestrial
		<i>cf. Gecarcinus ruricola</i>	black, blue or mountain crab	crabe terrestre	terrestrial
	Ocypodidae	<i>cf. Gecarcinus lateralis</i>	black land crab	crabe terrestre	terrestrial
		<i>Ocypode quadrata</i>	sand crab	crabe blanc	terrestrial
		<i>Uca sp. 1</i>	fiddler crab	crabe violoniste	terrestrial
		<i>Uca sp. 2</i>	fiddler crab	crabe violoniste	terrestrial
		<i>Uca sp. 3</i>	fiddler crab	crabe violoniste	terrestrial
		<i>Uca sp. 4</i>	fiddler crab	crabe violoniste	terrestrial
		<i>Uca sp. 5</i>	fiddler crab	crabe violoniste	terrestrial
Echinoida	Echinometridae	<i>Echinometra sp.</i>	rock-boring urchin	oursin lie-de-vin	reef
	Echinidae		variegated urchin	oursin commun	reef
Pisces	Carcharhinidae		sharks	requins	reef
	Dasyatidae	<i>cf. Dasyatis americana</i>	southern stingray	penélague américaine	estuarine
		<i>cf. Aetobatis narinari</i>	spotted eagle ray	ange de mer	estuarine
	Elopidae	<i>Elops saurus</i>	mullet	mulet	estuarine
	Albulidae	<i>Albula vulpes</i>	bonefish	baranme	estuarine
	Muraenidae	<i>cf. Echnida catenata</i>	chain moray	murène à gueule pavée	reef
		<i>Cymnothorax sp.</i>	moray	murène	reef
	<i>cf. Clupeidae</i>	<i>cf. Engraulidae, or cf. Atherinidae</i>	herring/sardine/anchovy	pisquette	estuarine
	Belontiidae	<i>Tylosurus crocodilus</i>	houndfish	orpie de terre	estuarine
		<i>cf. Platybelone argalus</i>	keeled needlefish	orpie de canal	estuarine
	Holocentridae	<i>cf. Strongilura timucu</i>	Timucu needlefish	aiguillette timucu	estuarine
		<i>cf. Holocentrus sp.</i>	squirrelfish	cardinal	reef
	Centropomidae	<i>cf. Sargocentron sp.</i>	groupers	brochet de mer	reef
	Serranidae	<i>Centropomus cf. undecimalis</i>	groupers	mérou, vieille	reef
		<i>Epinephelus</i>	groupers	mérou, vieille	reef
		<i>cf. Epinephelus</i>	rock hind	waliwa	reef
		<i>cf. Epinephelus adscensionis</i>	coney	tanche	reef
	Priacanthidae	<i>cf. Cephalopholis fiva</i>	glass-eye	soleil batard	reef
	Carangidae	<i>cf. Priacanthus cruentatus</i>	jack	coulirou de canal	reef
		<i>Decapterus sp.</i>	mackerel scad	carangue	reef
		<i>Caranx sp.</i>	jack	carangue franche	reef
		<i>Caranx ruber</i>	bar jack	carangue grasse	reef
		<i>Caranx hippos</i>	crevalle jack	carangue grasse	reef
	Lutjanidae	<i>Lutjanus sp.</i>	snapper	vivaneau	reef
		<i>Lutjanus buccanella</i>	blackfin snapper	vivaneau oreilles noires	reef
		<i>Lutjanus purpuraceus</i>	red snapper	vivaneau rouge	reef
		<i>Lutjanus vivamus</i>	silk snapper	vivaneau franc	reef
		<i>Ocyurus chrysurus</i>	yellowtail snapper	colas	reef
	Haemulidae	<i>Anisostrems sp.</i>	margate	lippu	reef
		<i>Haemulon sp.</i>	grunt	gorette	reef
	Sparidae	<i>cf. Calamus pennatulata</i>	pluna	daubenet plume	estuarine
		<i>cf. Calamus sp.</i>	porgy	guelle ferrée	estuarine
	Sciaenidae	<i>cf. Archosargus rhomboidalis</i>	sea bream	rondeau des pâturages	estuarine
		<i>cf. Equetus acuminatus</i>	cubbyu	monsieur l'abbé	estuarine
		<i>cf. Equetus lanceolatus</i>	jackknifefish	monsieur l'abbé	estuarine
	Kyphosidae	<i>Equetus sp.</i>	dem	monsieur l'abbé	estuarine
		<i>Kyphosus sectatrix</i>	bermuda chub	sgouti	reef
	Pomacanthidae	<i>cf. Holocanthus sp.</i>	rock beauty	ange des Caraïbes	reef
	Sphyrnidae	<i>Sphyrna barracuda</i>	great barracuda	barracuda	reef
	Labridae	<i>Sphyrna sp.</i>	southern sennet	bambokoï	reef
		<i>Bodianus rufus</i>	spanish rufus	capitaine de roche	reef
	Scaridae	<i>Halihoeres sp.</i>	wrasse	girelle	reef
<i>Scarus sp.</i>		parrotfish	chat	reef	
	<i>Sparisoma sp.</i>	parrotfish	chat	reef	
	<i>Sparisoma chrysopterygum</i>	redtail parrotfish	perroquet queue rouge	reef	
	<i>Sparisoma viride</i>	stoplight parrotfish	chat	reef	
	<i>Sparisoma aurofrenatum</i>	yellow parrotfish	perroquet à bandes rouges	reef	
	<i>Sparisoma rubripinne</i>	yellowtail parrotfish	perroquet queue jaune	reef	
Acanthuridae	<i>Acanthurus sp.</i>	dosterfish	surgén	reef	
Scombridae	<i>Euthynnus sp.</i>	little tunny	bonite	pelagic	
	<i>Thunnus cf. albacares</i>	yellowfin tuna	thon ailes jaunes	pelagic	
Ballistidae	<i>cf. Balistes vetula</i>	queen triggerfish	bourse blanche	reef	
Ostraciidae	<i>cf. Lactophrys sp.</i>	smooth trunkfish	coffre mouton	estuarine	
Diodontidae	<i>Chilomycterus sp.</i>	burrfish	diodon	reef	
	<i>Diodon cf. hystrix</i>	porcupinefish	diodon de roche	reef	
Amphibia	<i>cf. Leptodactylidae</i>	<i>cf. Leptodactylus fulvix</i>	frogs	grenouilles	terrestrial
Reptiles	Cheloniidae		sea turtle	tortue marine	pelagic, terrestrial
	Iguanidae	<i>cf. Chelonia mydas</i>	green turtle	chélone franche	pelagic, terrestrial
	Laertilia <i>cf. Teiidae</i>	<i>Iguana sp.</i>	West Indian iguana or green iguana	iguane des Petites Antilles ou iguane vert	terrestrial
	<i>cf. Ameiva cinerea</i>	ameiva	ameive de Guadeloupe	terrestrial	
Aves	<i>cf. Procellariidae</i>	<i>Puffinus sp.</i>	shearwater	puffin	terrestrial
	<i>cf. Anatidae</i>	<i>Anas sp.</i>	duck, teal	canard, sarcelle	reef
	<i>cf. Laridae</i>	<i>cf. Larus sp.</i>	gull	mouette	terrestrial
	Columbidae	<i>Zenaidra cf. aurita</i>	dove	tourterelle	terrestrial
	<i>cf. Mimidae</i>	<i>cf. Margarops sp.</i>	trashier	grive fine, grosse grive	terrestrial
Mammalia	Chiroptera		bat	chauve-souris	terrestrial
	Canidae	<i>Canis familiaris</i>	domestic dog	chien domestique	terrestrial
	Oryzomyine	<i>Megalomys cf. desmarestii</i>	rice rat	rat pilori	terrestrial
	Dasyproctidae	<i>cf. Dasyprocta leporina</i>	sgouti	sgouti	terrestrial

Number of taxa

89

APPENDIX 2

Faunal spectrum and ecosystems of the whole assemblage Z64S93C01.

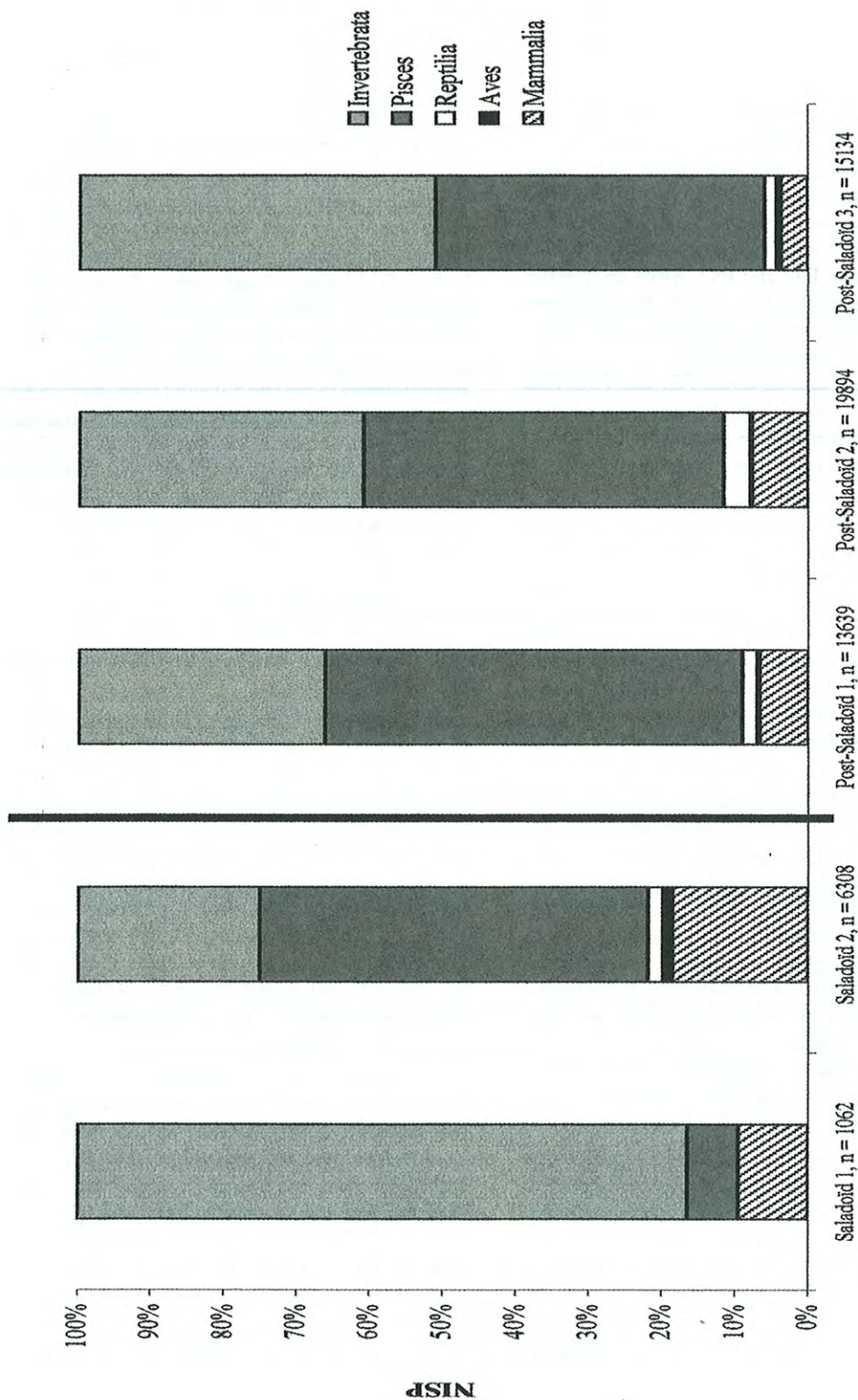


FIGURE 7
Faunal classes by archaeological level (percentage of the NISP).

level (PS1) (Figure 7). Conversely, crustaceans and sea urchins showed a decrease between levels S1 and S2, then a slight increase in level PS3. Mammals were most abundant in the late Saladoïd level (S2). Dogs were absent from the Saladoïd and the third Post-Saladoïd levels (Appendix 3). The NISP of agoutis increased progressively from S1, but decreased between PS2 and PS3. Remains of rice rats increased between S1 and S2, and between PS1 and PS2, but decreased between S2 and PS1 and between PS2 and PS3. Birds and reptiles were scarce throughout the levels. In conclusion, the global spectrum changed among the levels: mammals were abundant towards level S2 and well represented among levels PS1 and PS2. Fish decreased, while crabs and urchins increased from level S2 until level PS3. It seems that each occupation had a distinctive faunal selection.

The rarefaction curve (Figure 8) indicates that the sample from level S1 was not large enough to reflect species richness, nor to reflect the proportions of the rarest species. The samples from levels S2, PS1, PS2 and PS3 were close to the asymptote of this curve and included common as well as rare taxa. They were considered to be an adequate representation of the population being studied. The samples from levels S1, S2, and PS1 included respectively 25, 53, and 72 taxa, but PS2 and PS3 contained 67 and 66 taxa, although they are the largest samples.

The richness index for the whole sequence was 8.1, but differences can be seen between levels in Figure 9. The richness for S2, PS2, and PS3 reached values between 5.9 and 6.8, while PS1 had a higher index of 7.5. These results indicate that Saladoïd 2, Post-Saladoïd 2 and 3 lacked of some of the rarer taxa present in the first Post-Saladoïd deposit. According to Grayson (1984), the higher the value of the richness index, the richer the spectrum, thus the broader the exploitation of resources. Consequently, the inhabitants of Post-Saladoïd 1 had a broader exploitation subsistence than those of Saladoïd 2, Post-Saladoïd 2, or 3.

The diversity index for the whole sequence was 11.3, but differences were also observed between levels. The S2 and PS1 had high diversity indices (12.3 and 12.9, respectively), while PS2 and PS3 had small indices (11.4 and 7.8, respectively) (Figure 9). The diversity index represents the number of equally common species; the higher the values, the more evenly distributed the NISP is across species (Grayson, 1984: 160). Thus, the more the acquisition of the resources is diversified,

the more equal the distribution between the samples. A faunal sample with both high richness and diversity suggests a generalised subsistence (Leonard & Jones, 1989).

Figure 10 showing the 20 most abundant families, based on NISP, illustrates this phenomenon. According to the decreasing relative frequencies, the most important taxa of the whole assemblage were land hermit crabs (Coenobitidae), doctor-fishes (Acanthuridae), sea urchins (Echinoidea), jacks (Carangidae), land crabs (Gecarcinidae), trigger-fishes (Balistidae), parrot-fishes (Scaridae), and rice rats (Oryzomyini), anchovies (Clupeidae) and grunts (Haemulidae). The rice rats predominated in the late Saladoïd level, with jacks, land hermit crabs, and land crabs. A terrestrial animal (either a rice rat, a land hermit crab, or a land crab) appeared among the three most important taxa of each level. Sea urchins were among the three most important taxa of the most recent three levels. Fish, especially jacks and doctor-fishes were abundant, jacks particularly in the earlier levels and doctor-fishes in the later deposits. These lists illustrate the natural importance of animals that are typically found on land and in the sea around Grande-Terre.

In conclusion, a strategy of selection based on a small number of dominant species was found in Post-Saladoïd 3. However, Post-Saladoïd 1 had both high richness and diversity indices, and its spectrum of taxa illustrated a generalised subsistence, with a large spectrum and equally distributed species. Saladoïd 2 was apparently impoverished, but had a high diversity index and its spectrum illustrated equally distributed species. Post-Saladoïd 2 had a high richness index, but a low diversity index, and its spectrum illustrated a specialisation with few taxa, but accompanied by a large quantity of species.

Comparisons of the distributions were undertaken in chronocultural stages, and statistically tested pairwise (χ^2). Differences between the assemblages were all significant (Table 2). Hermit crabs, rice rats, urchins, jacks, and doctor-fishes were the taxa that created the differences between levels S2, and PS1, PS2 and PS3. Land crabs, hermit crabs, rice rats, jacks, sea turtles, herrings, and trigger-fishes created the differences between levels PS1, PS2 and PS3. Land crabs, hermit crabs, rice rats, sea turtles, and agoutis created the differences between levels PS2 and PS3. Without those taxa, the differences between the levels would

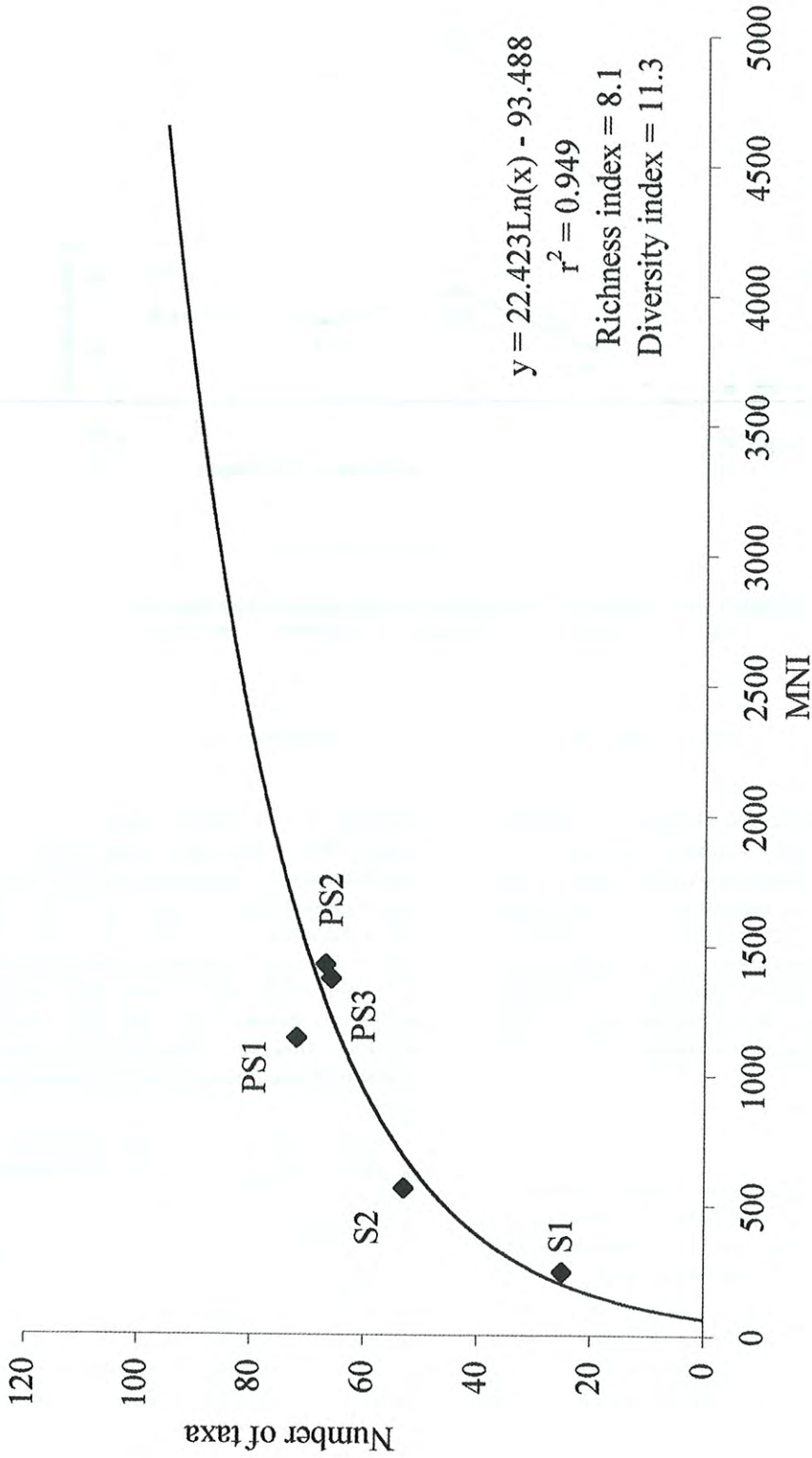


FIGURE 8
Richness curve (number of taxa and MNI) by archaeological level (PS = Post-Saladoïd, S = Saladoïd).

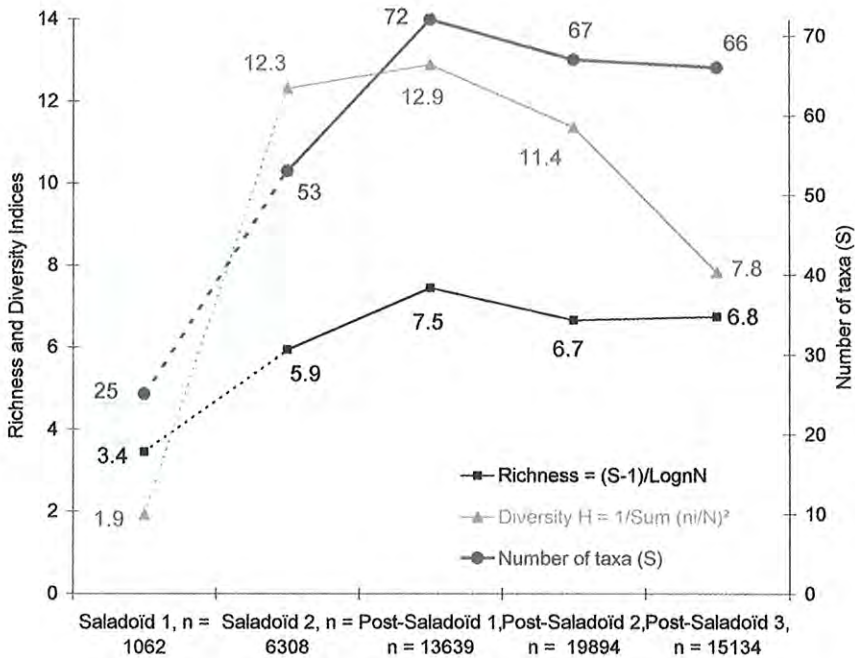


FIGURE 9

Richness and diversity indices, and number of taxa by archaeological level.

have been rendered insignificant. In conclusion, terrestrial crabs and mammals, sea turtles and some reef fishes (like doctor-fishes, jacks or triggerfishes) were not exploited in the same proportions throughout each period. The Post-Saladoïd 3 seems to have had a specialised economy, and although Saladoïd 2, Post-Saladoïd 1 and 2 had generalised economies, the inhabitants selected different species during each occupation.

Fish sizes

The results of the length estimates for fish are presented in Table 3 and Figure 11. Details of the basic statistics of the 3,951 measurements will be presented the author's PhD dissertation.

According to the Agostino test (Chenorkian, 1996), none of the size class distributions of grunts (Haemulidae) were normal (Table 4). Most individuals were between 120 and 160 mm long and few small individuals gave a slight skew toward the small size classes (Figure 11). The smallest indi-

vidual was 52 mm, and the largest was 317 mm length (Table 3). The general mean length was 147 mm. However, the maximum, minimum and mean were slightly different in each level. For example, maximal lengths varied between 251 mm (PS1) and 317 mm (PS3); minimal lengths varied between 52 mm (PS1) and 73 mm (PS3); and mean lengths vary between 145.7 mm (PS1) and 148.9 mm (PS3). However, shapes of the size class distributions of grunts seem generally similar through time.

The PS1 and PS2 size class distributions of snappers (Lutjanidae) had normal distributions (Table 4), while S1+S2 and PS3 did not have a normal distribution, although they resembled normal curves for size with a few large individuals giving the curve a slight skew toward the large size (Figure 11). The smallest individual was 59 mm and the largest 824 mm long (Table 3). The mean was 207 mm long. Maximum lengths varied from 401 mm (PS2) to 824 mm (PS3); minimal lengths varied from 59 mm (PS1) to 82 mm (PS2); and mean lengths varied from 201 mm (PS2) to 215

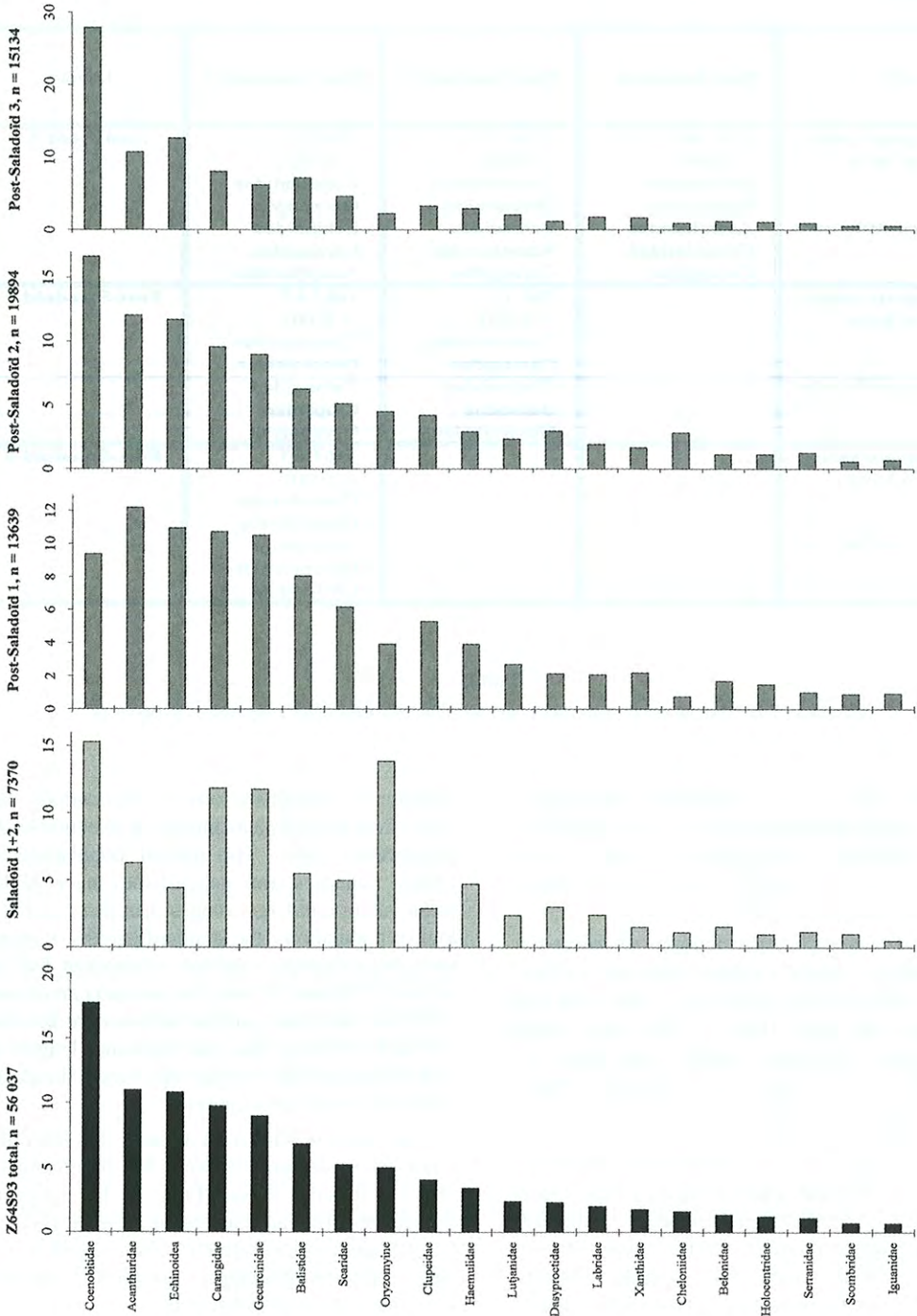


FIGURE 10
Diversity distribution by archaeological level (percentage of NISP).

Chi² tests

df = 26	Post-Saladoïd 1	Post-Saladoïd 2	Post-Saladoïd 3	Levels
chi square value	1251.89	1856.73	3097.32	Saladoïd 2
risk of error	< 0.001	< 0.001	< 0.001	
main contributions	Oryzomyini Echinoidea Acanthuridae Coenobitidae Carangidae	Coenobitidae Oryzomyini Echinoidea Acanthuridae Carangidae	Coenobitidae Oryzomyini Echinoidea Carangidae Acanthuridae	
chi square value		784.12	1883.67	Post-Saladoïd 1
risk of error		< 0.001	< 0.001	
main contributions		Coenobitidae Carangidae Cheloniidae Balistidae Gecarcinidae	Coenobitidae Gecarcinidae Carangidae Clupeidae Oryzomyini	
chi square value			1161.00	Post-Saladoïd 2
risk of error			< 0.001	
main contributions			Coenobitidae Gecarcinidae Oryzomyini Dasyproctidae Cheloniidae	

TABLE 2

Chi² values on the NISP by taxon, significance, and taxa providing the highest contributions to the value.

mm (PS3). The size class distributions of snappers slightly changed through time. In S1+S2 and PS3, most of individuals were between 150 and 275 mm long, while in PS1 and PS2, most of individuals were between 125 and 250 mm long.

All of the size class distributions of parrot-fishes (Scaridae) formed a normal distribution (Table 4). The smallest individual was 65 mm long and the largest 567 mm (Table 3). The mean length was 218 mm. Maximum length varied from 463 mm (PS1) to 567 mm (S1+2); minimal lengths varied from 65 mm (PS1) to 105 mm (S1+2); and mean lengths varied from 212 mm (S1+2) to 222 mm (PS1). The size class distributions of parrot-fishes also changed slightly through time. Most individuals of S1+S2 were between 150 and 200 mm long. Most individuals of PS1 were between 200 and 250 mm long. PS2 showed relatively equal proportions in each size class between 125 and 275 mm long, while PS3 resembled normal curves for size with a few large individuals.

Most fish were intermediate in size. Few juveniles and few large adults were caught, and a large

quantity of individuals were of intermediate size. This effect suggests a technique of selection of the intermediate sizes. The natural populations of grunts, snappers and parrot-fishes have higher mean values and larger ranges than these archaeological populations: the standard lengths of grunts from the reference collection of modern fish are between 130 and 267 mm, the snappers are between 66 and 460 mm, and parrotfishes are between 135 and 260 mm; thus the estimated lengths of archaeological fish beyond this range should be interpreted with great caution.

The pairwise Chi² tests showed that the differences between samples were not significant for grunts and parrot-fishes (Table 5). For snappers, significant differences appeared between PS1 and S2 and between PS2 and PS3 (Table 5). However, the distribution of snappers during PS1 and PS2 were not significantly different.

The sizes of the fish did not decrease nor increase through time, and if the curves are globally homogeneous throughout the levels, selections must have been made on the size of the fishes

Haemulon	number of individuals	mean	maximum	minimum	standard deviation
<i>Natural population (Fisher, 1978)</i>		198.00	171.42	229.63	14.32
<i>Reference collection</i>	54	192.30	267.00	130.00	34.41
Saladoïd 1 + 2	330	145.82	282.53	60.73	30.78
Post-Saladoïd 1	519	145.70	251.34	52.62	30.89
Post-Saladoïd 2	625	146.76	299.09	67.15	30.03
Post-Saladoïd 3	431	148.91	317.51	73.27	32.59
Total	1905	146.80	317.51	52.62	31.01
Lutjanus	number of individuals	mean	maximum	minimum	standard deviation
<i>Natural population (Fisher, 1978)</i>		385.99	224.39	797.01	73.65
<i>Reference collection</i>	47	264.17	460.00	66.00	75.83
Saladoïd 1 + 2	155	209.87	544.74	78.59	64.51
Post-Saladoïd 1	340	205.30	562.05	62.26	71.65
Post-Saladoïd 2	376	201.38	401.83	82.03	57.55
Post-Saladoïd 3	278	215.42	824.12	59.76	66.11
Total	1149	207.08	824.12	59.76	65.20
Sparisoma	number of individuals	mean	maximum	minimum	standard deviation
<i>Natural population (Fisher, 1978)</i>		471.49	1628.49	231.41	157.6
<i>Reference collection</i>	28	208.11	260.00	135.00	35.32
Saladoïd 1 + 2	144	212.26	566.77	105.94	58.88
Post-Saladoïd 1	252	222.00	463.92	65.33	63.31
Post-Saladoïd 2	288	215.87	489.55	104.61	63.82
Post-Saladoïd 3	213	218.55	467.81	69.10	66.38
Total	897	217.65	566.77	65.33	63.63

TABLE 3

Basic statistics of the estimated standard length by archaeological level for three fish taxa.

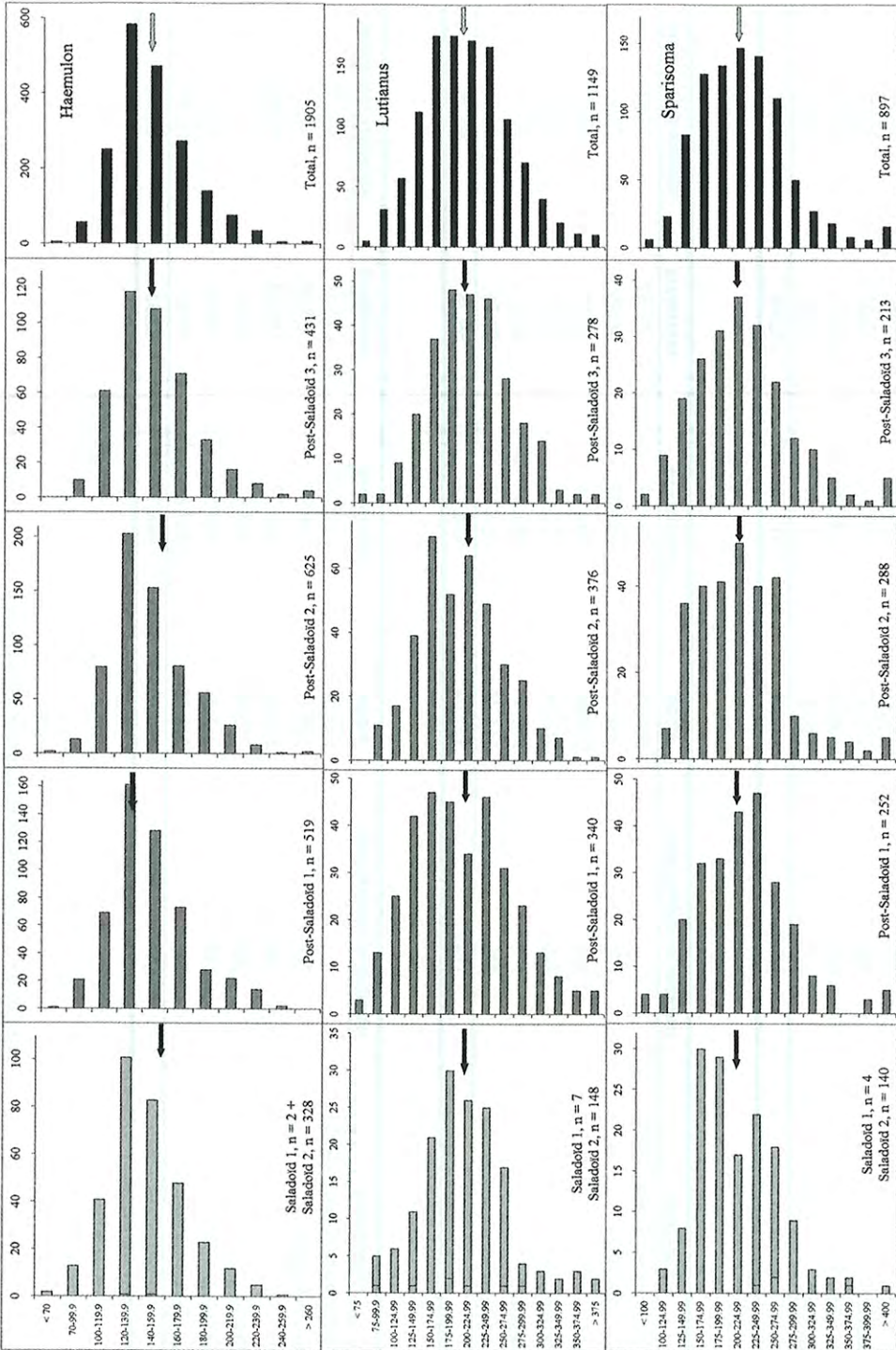


FIGURE 11

Curves of estimated standard length for *Haemulon*, *Lutjanus*, and *Sparisoma* by archaeological level.

Agostino tests of Normality

Ho = normal distribution at the confidence level of 0.05

	Saladoïd	Post-Saladoïd 1	Post-Saladoïd 2	Post-Saladoïd 3
Haemulon	Statistic U	922434.00	2292547.09	1645544.81
	Statistic D	0,28	0,276	0,27
	Statistic Q	-4,23	-5,16	-7,11
	Number of Measurements	330	519	431
	Standard error	30,84	30,92	32,63
	Signification	Ho rejected	Ho rejected	Ho rejected
Lutjanus	Statistic U	407977,55	2308092,51	1277765,85
	Statistic D	0,26	0,28	.25
	Statistic Q	-7,84	-2,11	-17,79
	Number of Measurements	155	340	278
	Standard error	64,72	71,76	66,22
	Signification	Ho rejected	Ho accepted	Ho rejected
Sparisoma	Statistic U	317631,24	1097306,83	827623,80
	Statistic D	0,26	0,27	0,27
	Statistic Q	-8,78	-4,86	-7,73
	Number of Measurements	144	252	213
	Standard error	59,08	63,44	66,53
	Signification	Ho rejected	Ho rejected	Ho rejected

TABLE 4

Agostino test values and significance of the normality of the estimated length distributions.

Chi² tests**Haemulon**

df = 8	Post-Saladoïd 1	Post-Saladoïd 2	Post-Saladoïd 3	Levels
Chi square value	2.71	5.21	5.70	Saladoïd
risk of error, p	0.95	0.74	0.68	
difference	ns	ns	ns	
Chi square value		11.40	10.27	Post-Saladoïd 1
risk of error, p		0.18	0.25	
difference		ns	ns	
Chi square value			8.46	Post-Saladoïd 2
risk of error, p			0.39	
difference			ns	

Lutjanus

df = 11	Post-Saladoïd 1	Post-Saladoïd 2	Post-Saladoïd 3	Levels
Chi square value	18.12	16.89	8.95	Saladoïd
risk of error, p	0.08	0.11	0.63	
difference	ns	ns	ns	
Chi square value		20.99	25.59	Post-Saladoïd 1
risk of error, p		0.03	0.007	
difference		significant	significant	
Chi square value			14.67	Post-Saladoïd 2
risk of error, p			0.20	
difference			ns	

Sparisoma

df = 10	Post-Saladoïd 1	Post-Saladoïd 2	Post-Saladoïd 3	Levels
Chi square value	11.78	14.51	14.01	Saladoïd
risk of error, p	0.30	0.15	0.17	
difference	ns	ns	ns	
Chi square value		11.58	3.85	Post-Saladoïd 1
risk of error, p		0.31	0.95	
difference		ns	ns	
Chi square value			10.24	Post-Saladoïd 2
risk of error, p			0.42	
difference			ns	

TABLE 5

Chi² values on the estimated lengths and significance for the main fish taxa.

during the Saladoïd and the first two Post-Saladoïd levels. The absence of the smallest specimens could be linked to the use of pots, tramails (large pots), nets, bottom trawls, or hooks and lines, which are good techniques for selecting the biggest fish. Only the specimens in the last Post-Saladoïd level resembled a normal curve, suggesting a selection over the natural range of the species. The large range of size of the samples suggests the simultaneous use of diverse fishing techniques, with nets, pots, hooks and lines, spears, bows, poisons, trawls, etc. This result corroborates our observation concerning the richness and diversity indices, that is, a generalised subsistence during the last Post-Saladoïd occupation.

Selected ecosystems and fishing techniques

The reconstruction of the selected ecosystems have been based on the ecological preference of the species and the size profiles of the faunal sample (Appendix 2).

The species represented on this site could all have been procured from adjacent environments (Figure 12). Most of the fish are primarily reef fish, such as trunkfish, porcupinefish, doctor-fish, parrotfish, triggerfish, grunt, squirrelfish, and small grouper. Throughout the Post-Saladoïd levels (from PS1 until PS3), there was a decline in the use of animals typically found on land, such as rice rats and agoutis. Moreover, beginning in the first Post-Saladoïd period, there was also a decrease of the exploitation of fish from the rocky substrate in inshore water and a corresponding increase in the use of coral reef species, particularly doctor-fishes, parrotfishes, and triggerfishes. As suggested by the insignificant number of small and juvenile fishes, there was no overexploitation of the natural ecosystems.

The geological, geographic, edaphic, and climatic characteristics of the Grande-Terre area produce a mosaic of conditions which sustains a diverse animal life. In particular, the varied marine ecosystems are determined by the nature of the lagoon, the island shelf, and the banks. The present location of the site is on a white-sand beach one kilometre long on the Atlantic Ocean. It is separated from the island of La Désirade by swift ocean currents. A coral barrier reef protects the lagoon, and the beach vegetation is xerophytic. A salt pond is located around La Pointe des Châteaux to the

east and a mangrove forest is situated in the Grand-Cul-de-Sac Marin to the west.

The fishing techniques in use today in Guadeloupe are adapted to the sea bottoms, period of the year, species and size of fish. Reef fish are caught from a small boat with hoop nets, pots, or traps, during the whole year, but especially from May to December, because of the rain and hurricane season. Small fish and juveniles are caught with cast nets and lines from the beach in shallow waters, lagoons, or mangrove canals, where they are used to feeding and sheltering. For example, grunts less than 122 mm long (standard length) are juveniles that hide in coral reefs during the day and feed in the shallow waters near the shore, and on the seagrass beds at night (Wing, pers. comm.). The bottom fish, such as large grouper or snapper, are caught with tramails all year round, except from December to February, when the currents are unfavorable. The permanent pelagic fishes, such as anchovies, needlefish, houndfish, little tunny, mackerel scad, and barracuda, are caught during the whole year with surface hooks and lines, seines, or casting nets. The seasonal pelagic species, such as mackerel and tuna, are caught by dragnet fishing from January to June in inshore waters, or in the swift currents flowing between Guadeloupe and La Désirade, from large boats (Aubin-Roy, 1968; Odum, 1971; Hurault, 1972; Wing, 1977, 1994; Bonniol, 1979, 1981; Wing & Reitz, 1982; Béarez, 1995). Although these fishing techniques are employed today, they could have not been used by the pre-Columbian inhabitants of the Caribbean, because some of these techniques were introduced by Europeans, Africans or Asians (seines, cast nets, tramails and pots). However, similar techniques such as reed curtains could have been used. In fact, Amerindians from French Guyana (Galibis) used to close the estuary with a reed curtains during high tide. When the tide ebbed, fishes and crabs that tried to reach the sea remained prisoners. At least, at the beginning of European colonisation, Amerindians had a very high level of navigation technology, using large pirogues to travel from the Guyanas to the Caribbean islands.

Thus, one of the explanations for the diverse fauna is that the wide variety of ecosystems near the site were exploited and that numerous and diverse fishing techniques were probably used.

CONCLUSIONS

The archaeological faunal assemblage from Anse à la Gourde provided information about the

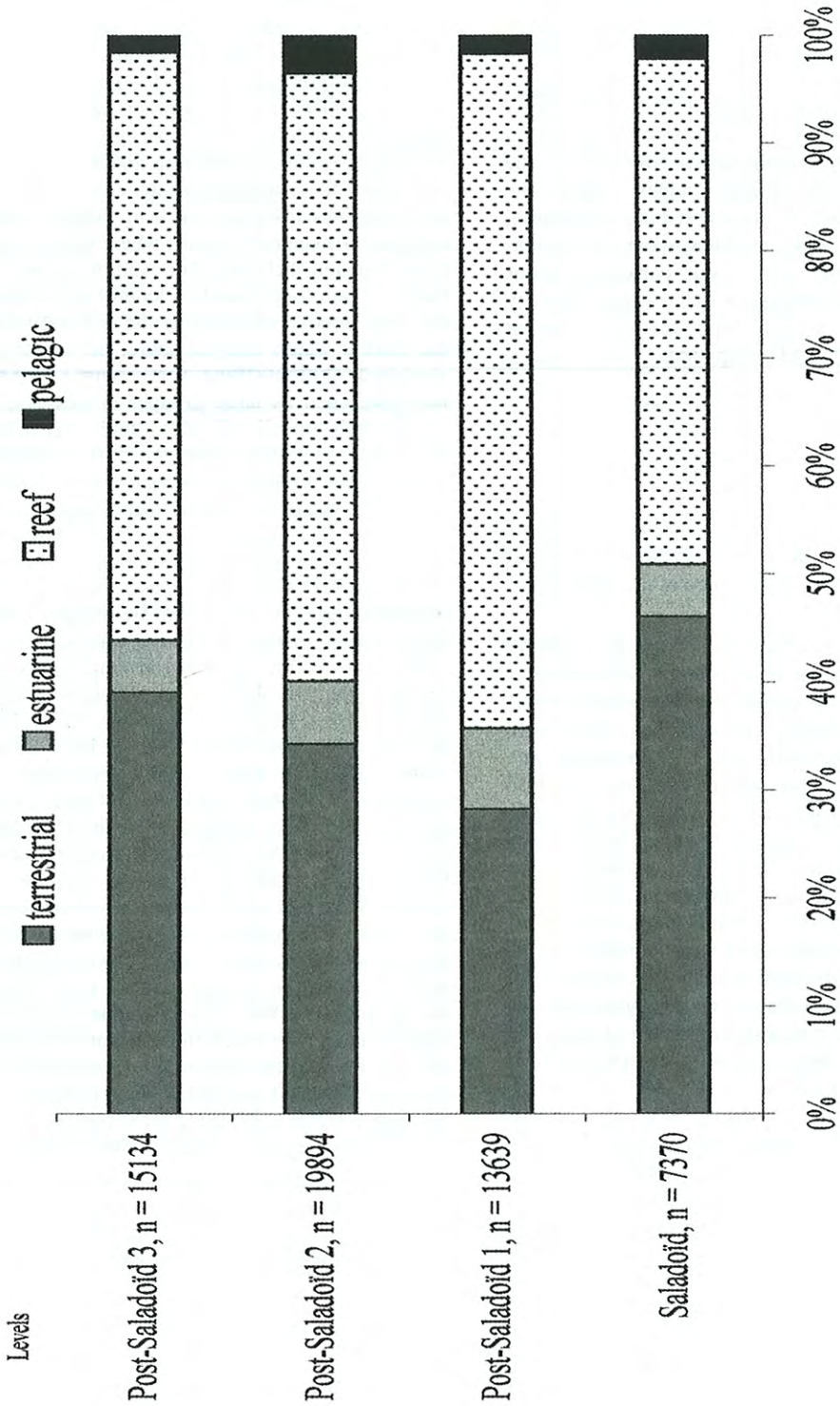


FIGURE 12
Selected ecosystems by archaeological level.

Families	Saladoïd 1		Saladoïd 2		Post-Saladoïd 1		Post-Saladoïd 2		Post-Saladoïd 3		TOTAL	
	NR	NMI	NR	Weight	NR	NMI	NR	Weight	NR	NMI	NR	Weight
Palaeobirdidae	5	0.15	21	1.08	4	7	27	1.95	82	5.3	187	12.49
Columbidae	748	23.38	380	13.32	67	245	3312	122.69	4226	145.84	9947	367.69
Poronidae											33	1.01
Xanthidae			113	11.41	10	3	335	30.88	20	19.38	998	88.22
Oecouridae	131	5.14	738	49.85	84	216	1787	105.25	216	53.28	5032	325.04
Oxyopidae			4	0.11	1	8	2327	58.62	23	19.16	27	1.19
Echinidae			4	0.29	4	2	2	0.12	2	0.2	8	0.61
Carcharhinidae			3	0.17	2	2	4	0.44	2	0.01	12	0.56
Dasyptidae			3	0.2	2	2	4	0.44	2	0.01	6	0.64
Elopidae			1	0.03	1	1	11	0.47	4	0.35	23	1.00
Albulidae			1	0.01	1	1	10	0.43	2	0.26	27	0.96
Muraenidae			215	1.36	21	42	839	4.56	504	2.97	2282	12.13
Clupeidae			111	4.51	14	232	646	7.74	16	182	759	23.99
Belontiidae	3	0.06	74	2.16	14	205	232	6.53	22	4.47	679	18.34
Holocentridae			15	0.68	6	10	0.66	4	5	1.73	69	5.08
Centropomidae			83	5.8	14	141	253	18.87	25	35.28	620	76.05
Serranidae			4	0.64	1	26	1	0.01	1	0.01	1	0.01
Pisacanthidae			857	25.62	56	1465	1905	41.74	97	1222	5872	172.55
Carangidae	18	0.67	2	0.88	2	370	473	27.56	29	313	1334	95.01
Lutjanidae	7	0.04	1	0.04	1	60	539	10.65	60	1915	58.25	
Haemulidae	2		2	0.11	2	15	2.22	0.5	2	6	26	3.71
Sparidae			5	0.07	4	27	0.41	9	9	0.26	62	2.06
Sciaenidae			1	0.02	1	1	0.12	2	2	0.1	13	0.24
Kyphosidae			4	0.2	1	1	0.02	1	6	0.42	1	0.02
Pomacanthidae			4	0.2	1	14	0.49	5	9	0.42	41	1.73
Sphyraenidae	3	0.09	176	5.56	19	285	387	12.48	26	272	1123	38.99
Labridae	7	0.69	2	0.26	362	844	844	86.22	83	60	2934	279.48
Scorpaenidae	5	0.26	1	0.26	1	1663	61.44	96	1626	57.13	6153	211.2
Acanthuridae	9	0.76	67	13.43	7	125	13.81	8.91	7	85	406	54.48
Scombridae	17	0.25	390	5.83	11	1099	13.25	18	14.97	13.85	3842	48.15
Balistidae			1	0.1	1	8	0.29	3	0.04	1	24	0.49
Omnacidae			1	0.1	1	19	0.29	5	9	0.52	37	1.11
Dysommidae			83	34.77	5	105	19.62	6	568	38.93	902	125.53
Atherinidae			37	1.71	6	132	4.63	7	80	4.14	388	14.91
Chenidae			14	0.34	4	47	0.36	10	27	0.51	122	1.54
Iguanidae			9	0.34	4	1	0.36	10	6	0.42	15	0.76
Lacertillidae			3	0.41	1	1	0.08	1	5	0.23	8	0.64
Procelariidae			68	1.45	6	44	1.01	6	75	1.11	1	0.08
Anartidae			17	0.19	2	29	0.56	3	1	0.01	246	5.05
Laridae	3	0.09	1	0.19	2	29	0.56	3	2	0.11	3	0.02
Columbidae			938	34.35	17	538	16.71	12	343	15.29	74	3.28
Mimidae			211	13.05	6	294	35.36	5	182	12.65	34	3.28
Chiroptera			6308	310.92	562	13639	619.94	1139	19894	731.19	2798	92.67
Carnivora			1892	48.91	562	13639	619.94	1444	15134	691.2	1293	75.42
Oryzomyini			337	5.56	8325	234.62	4774	142.3	2814	66.48	18142	497.87
Dasyproctidae			201	3.39	35468	517.54	41691	668.3	25650	453.32	116548	1854.45
Total identified	1062	35.84	244	6308	310.92	562	13639	619.94	1139	19894	731.19	2389.29
Unidentified												
Crustaceans												
Osteichthyes												
Mammalia												
Total unidentified	538	0.00	15746	0.00	44228	0.00	46776	0.00	28653	0.00	135941	0.00
Total	1600	44.79	244	72054	577.19	562	57867	1380.65	1139	66670	1547.74	43787
												4767.78
												3842

APPENDIX 3

NISP, Weight, NMI, and S by family for each archaeological level.

animals that were used for food during the prehistoric occupation of the site.

Although the same complex of common species appeared throughout the excavation of unit Z64S93C01, a shift in the predominant species was observed. Hermit crabs, rice rats, urchins, jacks, and doctor-fishes predominated in the Saladoïd deposits. Land crabs, hermit crabs, rice rats, jacks, sea turtles, herrings, and trigger-fishes predominated in the Post-Saladoïd levels. An analysis of the remains did not indicate overexploitation of these animals but rather a shift toward a greater dependence upon marine resources during Post-Saladoïd periods.

In general, grunts, snappers and parrotfishes had adults lengths, with few small and large individuals. Selections must have been made on the size of fish during both Saladoïd levels and the first two Post-Saladoïd levels. Only the last Post-Saladoïd level suggested a selection over the natural range of the species.

Based on the ecological context in which the species are usually found, it is clear that a great variety of different ecosystems was exploited. These included the tropical forests, dry lands, sand beaches, salt ponds, shallow inshore waters, rocky substrates, coral reefs, estuaries, mangroves, and pelagic waters. Catching animals from such varied habitats would naturally result in a diverse faunal assemblage.

The fauna from unit Z64S93C01 as a whole is rich and diverse, particularly in the second Saladoïd and the first Post-Saladoïd deposits. This suggests a more generalised subsistence during these periods. The second and third Post-Saladoïd periods were still rich, but less diversified. This suggests a more specialised subsistence economy, mainly during the Post-Saladoïd 3. The decrease in diversity could be linked to a less extensive use of the wider ecosystems and a more intensive use of the immediate vicinity, namely coral reefs and the lagoon.

The size characteristics, species characteristics, and the natural environment of the encountered species suggest that a variety of different techniques were used to catch them.

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REFERENCES

- ADAMS, W.H. 1985: *Aboriginal subsistence and settlement archaeology of the Kings Bay locality*. University of Florida, Department of Anthropology, Gainesville.
- AUBIN-ROY, J. 1968: *Vieux-Fort (Guadeloupe). Techniques et coopération dans un village de pêcheurs*. Département d'Anthropologie-Université de Montréal, juin 1968.
- BÉAREZ, P. 1995: *Comparaison des ichthyofaunes marines actuelles et Holocène et reconstitution de l'activité halieutique dans les civilisations précolombiennes de la côte du Manabi sud (Equateur)*. Diplôme de Doctorat: Ichtyologie Générale et Appliquée, Muséum National d'Histoire Naturelle.
- BOBROWSKI, P.T. & BALL, B.F. 1989: The theory and mechanics of ecological diversity in archaeology. In: Leonard, R.D. & Jones, G.T. (eds): *Quantifying diversity in archaeology*: 4-12. Cambridge University Press, Cambridge.
- BONNIOL, J.L. 1979: La perception de l'environnement dans une petite île de la Caraïbe: Terre de Haut des Saintes. In: Renard, Y. (ed.): *Perceptions of the environment. A selection of interpretative essays*: 51-68.

- Association Caraïbe pour l'Environnement, Caribbean Environment, Environmental Studies, St. Michael, Barbados.
- BONNIOL, J.L. 1981: La pêche en mer: l'exemple des Saintes. In: Bonniol, J.L. (ed.): *L'histoire Antillais*: 382-406. Dajani Editions, Pointe-à-Pître.
- CARLSON, B. 1995: The Coralie site (GT 3): Preliminary excavations at an Ostionan Ostionoïd site on Grand Turk and Caïcos. *Actes du 16è Congrès International d'Archéologie de la Caraïbe*: 88-94. Conseil Régional de la Guadeloupe, Mission Archéologique et Patrimoine, Basse-Terre, Guadeloupe.
- CASTEEL, R.W. 1974: On the number and sizes of animals in archaeological faunal assemblages. *Archaeometry* 16: 238-243.
- CASTEEL, R.W. 1978: Faunal assemblages and the "Wiegemethods" or "Weight method". *Journal of Field Archaeology* 5 (1): 71-77.
- CHAPLIN, R.E. 1971: *The study of animal bones from archaeological sites*. Seminar Press, London.
- CHENORKIAN, R. 1996: *Pratique archéologique statistique et graphique*. Editions Errance et Adam, Paris.
- DEFRANCE, S.D. 1988: *Zooarchaeological investigations of subsistence strategies at the Maisabel site, Puerto Rico*. Master of Arts Research Project. Department of Anthropology, University of Florida, Gainesville.
- DELPUECH, A.; HOFMAN, C. & HOOGLAND, M. 1997: Fouilles sur le site précolombien de l'Anse à la Gourde (Saint-François, Guadeloupe). *Journal de la Société des Américanistes* 83: 279-282.
- DESSE, J. & DESSE-BERSET, N. 1996a: Archaeozoology of groupers (Epinephelinae). Identification, osteometry and keys to interpretation. *Archaeofauna* 5: 121-127.
- DESSE, J. & DESSE-BERSET, N. 1996b: On the boundaries of osteometry applied to fish. *Archaeofauna* 5: 171-179.
- DESSE, J.; DESSE-BERSET, N. & ROCHETEAU, M. 1987: Contribution à l'ostéométrie du Mulet *Liza (Liza) ramada* Risso, 1826 (= *Mugil capito*, 1829). C.R.A. du C.N.R.S., APDCA, Juan-les-Pins.
- DESSE, J.; DESSE-BERSET, N. & ROCHETEAU, M. 1989: Les profils rachidiens globaux. Reconstitution de la taille des poissons et appréciation du nombre minimal d'individus à partir des pièces rachidiennes. *Revue de Paléobiologie* 8 (1): 89-94.
- DESSE, J.; DESSE-BERSET, N. & ROCHETEAU, M. 1996: Ostéométrie et archéologie de la Daurade Royale (*Sparus aurata*, Linné 1758). C.R.A. du C.N.R.S., APDCA, Juan-les-Pins.
- DUCOS, P. 1975: Analyse statistique des collections d'ossements d'animaux. In: Clason, A.T. (ed.): *Archaeozoological Studies*: 35-44. North-Holland Publ. Co. & American Elsevier, Amsterdam.
- GRAYSON, D.K. 1984: *Quantitative zooarchaeology. Topics in the analysis of archaeological faunas*. Academic Press, Orlando.
- HURAUULT, J. 1972: *Français et Indiens en Guyane*, collection 10-18, Paris.
- KEEGAN, W.F. 1997: *Bahamian Archaeology: Life in the Bahamas and Turks and Caicos before Columbus*. Media Publishing, Nassau.
- LEACH, B.F. & BOOCKOCK, A. 1993: *Prehistoric fish catches in New Zealand*. Tempus Reparatum, Archaeological and Historical Associates Limited, Oxford.
- LEACH, B.F. & BOOCKOCK, A. 1994: The impact of pre-European Maori fishermen on the New Zealand snapper *Pagrus auratus*, in the vicinity of Rotokura, Tasman Bay. *New Zealand Journal of Archaeology* 16: 69-84.
- LEACH, B.F. & BOOCKOCK, A. 1995: Estimating live fish catches from archaeological bone fragments of snapper, *Pagrus auratus*. *Tuhinga: Records of the Museum of New Zealand Te Papa Tongarewa* 3: 1-28.
- LEACH, B.F.; DAVIDSON, J.M.; HORWOOD, L.M. & ANDERSON, A.J. 1996a: The estimation of live fish size from archaeological cranial bones of the New Zealand barracouta *Thyrstites atun*. *Tuhinga: Records of the Museum of New Zealand Te Papa Tongarewa* 6: 1-25.
- LEACH, B.F.; DAVIDSON, J.M.; HORWOOD, L.M. & MALLON, S. 1996b: The estimation of live fish size from archaeological cranial bones of the New Zealand kahawai *Arripis trutta*. *Tuhinga: Records of the Museum of New Zealand Te Papa Tongarewa* 7: 1-20.
- LEACH, B.F.; DAVIDSON, J. & ATHENS, J.S. 1996c: Mass harvesting of fish in the waterways of Nan Madol, Pohnpei, Micronesia. In: Davidson, J.M.; Irwin, G.; Leach, B.F.; Pawley, A. & Brown, D. (eds.): *Oceanic Culture History: essays in honour of Roger Green*: 319-341. *New Zealand Journal of Archaeology Special Publication*.
- LEONARD, R.D. & JONES, G.T. 1989: *Quantifying diversity in archaeology*. Cambridge University Press, Cambridge.
- LYMAN, L.R. 1994: Quantitative units and terminology in zooarchaeology. *American Antiquity* 59 (1): 36-71.
- ODUM, E.D. 1971: *Fundamentals of ecology*. Saunders, Toronto.
- POPLIN, F. 1976a: A propos du Nombre de Restes et du Nombre d'Individus dans les échantillons d'ossements. *Cahier du Centre de Recherche Préhistorique de l'Université de Paris I* 5: 61-74.
- POPLIN, F. 1976b: Essai d'ostéologie quantitative sur l'estimation du Nombre d'Individus. In: Schwabedissen, F.H. (ed.): *Kölner Jahrbach für Ur- und Frühgeschichte*: 153-164.
- POPLIN, F. 1977: Problèmes d'ostéologie quantitative relatifs à l'étude de l'écologie des hommes fossiles. *Approche écologique de l'Homme fossile* 47: 63-68.
- REITZ, E.J. & CORDIER, D. 1983: Use of allometry in zooarchaeological analysis. In: Grigson, C. & Clutton-Brock, J. (eds.): *Animals and Archaeology 2: Shell Middens, Fishes and Birds*: 237-252. B.A.R. (International Series) 183, Oxford.
- REITZ, E.J. & WING, E.S. 1999: *Zooarchaeology*. Cambridge University Press, Cambridge.
- REITZ, E.J.; QUITMYER, I.R.; HALE, H.S.; SCUDDER, S.J. & WING, E.S. 1987: Application of allometry to zooarchaeology. *American Antiquity* 52 (2): 304-317.

- ROUSE, I. 1989a: Peoples and cultures of the Saladoïd frontier in the Greater Antilles. In: Siegel, P.E. (ed.): *Early ceramic population lifeways and adaptive strategies in the Caribbean*: 383-403. B.A.R. (International Series) 506, Oxford.
- ROUSE, I. 1989b: Peopling and re-peopling of the West Indies. In: Woods, C.A. (ed.): *Biogeography of the West Indies*: 119-135. Sandhill Crane Press, Gainesville.
- ROUSE, I. 1992: *The Taïnos. Rise and decline of the people who greeted Columbus*, Yale University Press, New Haven.
- ROUSE, I. 1995: Ceramic chronology of the Leeward Islands. Paper presented at the XVI International Congress for Caribbean Archaeology, Guadeloupe, July 1995.
- WATTERS, D.R. & ROUSE, I. 1989: Environmental diversity and maritime adaptations in the Caribbean area. In: Siegel, P.E. (ed.): *Early ceramic population lifeways and adaptive strategies in the Caribbean*: 129-144. B.A.R. (International Series) 506, Oxford.
- WATTERS, D.R. 1998: Maritime adaptive strategies in the Caribbean archipelago. *Revista de Arqueología Americana* (15): 7-31.
- WHEELER, A. & JONES, A.K.G. 1989: *Fishes*. Cambridge University Press, Cambridge.
- WING, E.S. 1977: Factors influencing exploitation of marine resources. In: Benson, E.P. (ed.): *The sea in the pre-columbian world*: 47-64. Dumbarton Oaks, Washington, D.C.
- WING, E.S. 1989: Human exploitation of animal resources in the Caribbean. In: Woods, C.A. (ed.): *Biogeography of the West Indies: past, present, and future*: 137-152. Sandhill Crane Press, Gainesville.
- WING, E.S. 1994: Patterns of prehistoric fishing in the West Indies. *Archaeofauna* 3 : 99-107.
- WING, E.S. 1995: Rice rats and Saladoïd people as seen at Hope Estate. In: Alegría, R.E. & Rodríguez, M. (eds.): *Comptes-Rendus du Xvème Congrès International d'Archéologie de la Caraïbe*: 219-231. Centro de Estudios Avanzados de Puerto Rico y el Caribe. Fundación puertorriqueña de las humanidades. Universidad de Turabo.
- WING, E.S. & REITZ, E.J. 1982: Prehistoric fishing economies of the Caribbean. *Journal of New World Archaeology* 5 (2): 13-32.
- WING, E.S. & SCUDDER, S.J. 1983: Animal exploitation by prehistoric people living on a tropical marine edge. In: Grigson, C. & Clutton-Brock, J. (eds.): *Animals and Archaeology 2: Shell Middens, Fishes, and Birds*: 197-210. B.A.R. (International Series) 183, Oxford.
- WING, E.S. & WING, S.R. 1995: Prehistoric ceramic age adaptation to varying diversity of animal resources along the West Indian archipelago. *Journal of Ethnobiology* 15 (1): 119-148.