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CUBA'S MESOPHOTIC CORAL REEFS AND ASSOCIATED FISH COMMUNITIES

Arrecifes de coral mesofóticos de Cuba y comunidades de peces asociadas

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ABSTRACT

A joint Cuba-U.S. expedition was conducted May 14-June 12, 2017 to characterize for the first time the extent and health of mesophotic coral ecosystems (MCEs) along the entire coastline of Cuba. Remotely Operated Vehicle (ROV) dives at 36 sites confirmed the presence of MCE habitat along all coasts of Cuba. ROV dives covered 27 km, at depths of 25-188 m, and documented habitat and species with 103 hours of high-definition video and 21,146 digital images. A total of 477 taxa of benthic macrobiota and 178 fish taxa were identified, and 343 specimens of benthic invertebrates and algae were collected to verify taxonomy and assess population genetic structure. The primary geomorphological features were the Deep Island Slope (125->150 m), Deep Fore-Reef Escarpment (the 'Wall', 50-125 m), and Deep Fore-Reef Slope (30-50 m). Most vertical surfaces of the Wall were covered with dense sponges, algae, octocorals, and black corals. Agaricia was the most abundant scleractinian genus on the Wall at depths of 50-75 m, and was observed to 122 m. Of the 2,240 scleractinian colonies that were counted in this study, only 12 corals (0.53%, mainly Agaricia spp.) showed signs of bleaching, and one Agaricia had black band disease, comprising remarkably low disease prevalence. The most frequently recorded sponge genera were *Xestospon*gia, Aplysina, and Agelas. At least 10 previously unknown sponge species were collected during the expedition. Sites outside of marine protected areas generally had lower fish abundances, a possible indicator of historical overfishing. Lionfish were observed at most sites but abundances were low compared to other Caribbean regions.

KEY WORDS: Cuba, mesophotic reef, geomorphology, biozonation, biodiversity.

RESUMEN

Entre el 14 de mayo y el 12 de junio de 2017 se llevó a cabo una expedición conjunta entre Cuba y EEUU. El objetivo fue mapear y caracterizar, por primera vez, la extensión y salud de los arrecifes de coral mesofóticos (MCE. por sus siglas en inglés) alrededor de las costas de Cuba. 43 inmersiones con un Vehículo Operado por Remoto (ROV, por sus siglas en inglés) en 36 sitios, confirmaron la presencia de arrecifes mesofóticos a lo largo de toda la línea costera cubana. Las inmersiones del ROV desde 25 a 188 m de profundidad abarcaron, en total, 27 km, y se documentaron hábitats y especies con 103 horas de videos de alta definición y 21,146 imágenes digitales. Un total de 477 taxa de macrobiota bentónica y 178 taxa de peces fueron identificados. Se recolectaron 343 especímenes de invertebrados bentónicos y algas para verificaciones taxonómicas y evaluación de estructura de poblaciones. Los zonas geomorfológicas más prominentes fueron: Pendiente profunda (125- >150 m), Escarpe profundo del arrecife frontal (la "pared", 50-125 m), y arrecife de franja profundo (30-50 m). La pared presentó la mayor diversidad y densidad de la macrobiota; todas las superficies verticales estuvieron cubiertas por diversidad de esponjas, algas, gorgonias y coral negro. Agaricia fue el género más abundante de escleractineos y dominó a profundidades entre 40-75 m. Solo 12 colonias de corales escleractíneos (principalmente Agaricia spp.) de 2,240 colonias (0,53%) registradas con el ROV mostraron signos de blanqueamiento, y una Agaricia presentó enfermedad de banda negra, evidenciando la muy baja prevalencia de enfermedades. Los géneros más frecuentes de Porifera fueron Xestospongia, Aplysina, y Agelas. Al menos una decena de especies de esponjas desconocidas para la ciencia fueron colectadas. Los sitios que se encontraban fuera de áreas marinas protegidas, generalmente poseían abundancia de peces baja, lo cual pudiera ser un posible indicador de sobrepesca histórica. Individuos de pez león fueron observados en la mayoría de los sitios pero con abundancias bajas en comparación con otras regiones del Caribe.

PALABRAS CLAVES: Cuba, arrecife mesofótico, geomorfología, biozonación, biodiversidad

INTRODUCTION

Mesophotic coral reef ecosystems (MCEs) are light-dependent benthic communities that occur deeper than shallow reefs and typically range from depths of 30 m to the bottom of the photic zone, which may extend to >150 m in some regions (Lesser et al., 2009; Hinderstein et al., 2010; Baker et al., 2016). MCEs represent in part an extension of shallow-water coral reef ecosystems and support a diverse assemblage of habitat-building taxa, including corals, sponges and algae, and associated commercial and recreational fisheries. However, many of the MCE species are depth specialists not found on shallow reefs. Studies of MCEs worldwide are expanding our understanding of their distribution, health, biodiversity, and ecology (Hinderstein et al., 2010; Baker et al., 2016; Loya et al., 2016, 2018; Roman, 2018). Many MCEs worldwide appear to be thriving compared to shallow reefs. The deep reef refugia hypothesis suggests that MCEs may be less impacted from natural and anthropogenic impacts than shallow coral reefs, and may be more stable and resilient than shallow reefs (Bongaerts et al., 2010; Bridge and Guinotte, 2012). MCEs may also act as refugia for shallow reef species through the export of fish and coral larvae (Serrano et al., 2014; Vaz et al., 2016). However, it is now apparent that MCEs are also vulnerable to disturbances from all facets of perturbations including climate change, bottom trawling, invasive species, and pollution (Bak et al., 2005; Andradi-Brown et al., 2016; Appledoorn et al., 2016a). Smith and Holstein (2016) further state that the main premise of the deep reef refugia hypotheses is incorrect and that it now appears that increases in temperatures above the local mean warmest conditions can lead to thermal stress and bleaching. Also there is growing evidence that the hypothesis may only be applicable to depth generalists on MCEs, and their contribution to replenishment of shallow reefs may not be as consequential as once hoped (Schlesinger *et al.*, 2018).

MCEs have been studied extensively throughout the Western Atlantic regions of the Caribbean, Bahamas, and the Gulf of Mexico (Hinderstein et al., 2010; Baker et al., 2016; Loya et al., 2016, 2018). Whereas considerable data have been reported regarding the distribution, ecology, and health of Cuba's shallow reefs (Alcolado et al., 2001a; Alcolado et al., 2001b; Alcolado et al., 2003; Aguilar et al., 2004; Aguilar Betancourt and González-Sansón, 2007; Alcolado and García, 2007; Martínez-Daranas et al., 2008; Alcolado et al., 2009; González-Díaz et al., 2010; Perera Valderrama et al., 2013; González-Díaz et al., 2014; Alcolado et al., 2015; González-Díaz et al., 2015; Suárez et al., 2015; Caballero Aragon et al., 2016; Ferrer Rodríguez et al., 2016; González-Díaz et al., 2018; Roman, 2018), relatively little is known about the distribution, community structure and health of Cuba's deep mesophotic reefs. The book "The Ecology of the Marine Fishes of Cuba" (Claro et al., 2001) provides a comprehensive description of physical attributes of coastal Cuba including geology, geomorphology, hydrology, and major habitats from intertidal depths out to the shelf break at approximately 30 m. The fish fauna of what Claro et al. (2001) termed as the slope and bathyal zone (100-600 m) was described from dives with Harbor Branch Oceanographic Johnson-Sea-Link Institute's submersible in 1997. In the 1970s, Vassil Zlatarski and Nereida Martinez Estalella were the

pioneers of research on Cuba's mesophotic reefs. Zlatarski and Estalella's (1982) book "The Scleractinians of Cuba" (first published in Russian in 1980, 1982 in French, 2008 in Spanish) provides detailed reef profiles and coral species distributions from 44 sites around the island, some of which extend into the >40-70 m depth range; however, the eastern tip and northeast coast lack data. Scuba dives (some to depths of 70 m) for this study resulted in an extraordinary collection of corals (5,924) specimens) which are now archived in the National Aquarium of Cuba (in Havana). González Ferrer's (2004) book "Corales Petreos Jardines Sumergidos de Cuba" also provides invaluable data on the stony corals of Cuba as well as the history of Cuban reef research since the 16th century. This book provides beautiful color plates for all the hard corals found in Cuba, as well as discussions of reef types, coral disease and bleaching. Kühlmann (1974, 1983) also provided data on the depth distribution of scleractinian corals from scuba dives to 70 m in Cuba.

In 2015, a Joint Statement was developed between the United States and the Republic of Cuba on Cooperation on Environmental Protection (November 24, 2015), and a Memorandum of Understanding (MOU) was signed by the U.S. National Oceanic and Atmospheric Administration (NOAA), the U.S. National Park Service and Cuba's National Center for Protected Areas. This MOU establishes a 'Sister-Sanctuary' relationship between Guanahacabibes National Park and Banco de San Antonio Marine Sanctuary in Cuba, and the Florida Keys National Marine Sanctuary (FKNMS) and Flower Garden Banks National Marine Sanctuary (FGBNMS) in the United States. It is recognized that these regions are all

inextricably linked through ocean currents. In support of these agreements, a joint research expedition was conducted from May 14 to June 12, 2017 on the University of Miami's ship F.G. Walton Smith to survey the mesophotic reefs of Cuba. This project was jointly planned and in collaboration with organizations in Cuba and the U.S., including Centro de Investigaciones Marinas at University of Havana (CIM-UH), Centro Nacional de Áreas Protegidas (CNAP), Instituto de Ciencias del Mar (ICIMAR), Geocuba Estudios Marinos, Guanahacabibes National Park-Sistema Nacional de Areas Protegidas (PNG-SNAP), Acuario Nacional de Cuba (ANC); and for the U.S., two NOAA Cooperative Institutes: the Cooperative Institute for Ocean Exploration, Research, and Technology (CIOERT) at Harbor Branch Oceanographic Institute/ Florida Atlantic University (HBOI-FAU), and the Cooperative Institute for Marine and Atmospheric Studies (CIMAS) at the University of Miami.

The primary goal of this research expedition was to survey the extent of mesophotic reefs along the entire coastline of Cuba using a remotely operated vehicle (ROV). The objectives of the expedition were to: 1) characterize Cuba's MCE benthic habitats, benthic macroinvertebrate and algal communities, and fish populations (emphasis on commercially important grouper/snapper species, spawning grounds, and Lionfish); 2) understand the genetic connectivity of Cuba's MCEs with mesophotic and shallow reefs upstream and cross-stream (including the Sister-Sanctuaries in U.S. and throughout Cuba); 3) characterize the health of Cuba's MCEs; 4) conduct ROV collections of benthic species for taxonomy and molecular analyses; 5) collect CTD, ADCP (Acoustic Doppler Current Profiler) oceanographic data, and water column data for benthic carbonate chemistry including aragonite saturation state. Data collected during this expedition and on planned future cruises will be used to compare the health and connectivity of Cuba's MCEs to reef systems elsewhere in the Caribbean, Gulf of Mexico, and southeastern United States.

This paper provides a preliminary overview of the results of this expedition to date, including general descriptions of the oceanographic data around Cuba, MCE habitats, geomorphology, biozonation, and biodiversity of these reefs. Future taxonomic and genetic analyses of the specimens, along with quantitative analyses of the video and photographic data, will allow a more precise characterization of the diversity, zonation, health, and regional differences of the mesophotic communities of Cuba, as well as a better understanding of the connectivity of Cuban reefs with the Sister Sanctuaries in the U.S. and elsewhere in the Caribbean. These data will also be useful for decision making for new and existing marine protected areas (MPAs) into these deeper waters of Cuba. Cuba's strong marine policies and legislation has already resulted in 105 MPAs, covering nearly 25% of its insular shelf, yet overfishing, poaching, pollution and global warming are threats to these vulnerable ecosystems (Perera Valderrama et al., 2018; Roman, 2018).

METHODS AND MATERIALS

Locations of proposed dive sites were compiled prior to the cruise, based on data from various sources: 1) recommendations from CIM-UH, CNAP, and ICIMAR; 2) NOAA and regional coastal charts; and, 3) site data from HBOI's *Johnson-Sea-Link* manned submersible 1997 expedition (Discovery

Channel "Forbidden Depths" documentary). Some sites were selected directly offshore of well-characterized shallow-water reef sites or within Marine Protected Areas (MPAs). Except for Banco de San Antonio, there are no high-resolution maps of the Cuba's deepwater shelf and slope. Specific sites were selected based on the above sources and plotted in ArcGIS 10.3 using coastal charts at depths of 30-150 m. In order to circumnavigate the island in the time available, transit between sites at night was generally from 50-150 km. Additional dives were made within the Guanahacabibes National Park which is of special interest. Two 3-4 hour dives were made each day with a *Mohawk* ROV. At each site a fathometer profile was made prior to the ROV dive to document the depth and location of the shelf-edge break and to select the exact target site for the ROV. Sketches of reef profiles were drawn to illustrate the various dive sites. The specific depths and slopes were drawn to scale from observations made during the ROV dives. The slope angles, inflections of the slopes at different depths, depths of zones, and depths of shelf-edge reef crest were made from the ROV dive notes.

Video was recorded throughout each ROV dive with an Insite Pacific Mini Zeus high-definition CMOS color zoom camera. Digital still images were taken with a Kongsberg Maritime OE14-408. Both cameras had 10-cm parallel lasers for scale (green-still; red-video). At each site a continuous video/photo transect commenced at a maximum depth of ~150 m and continued upslope to the upper mesophotic reef zone or about 30 m. The photo transects were used for habitat characterization and species identifications. When a second dive was made at the same dive site, it

was usually for sample collections and fish video transects. Photo transects were conducted using the digital still camera oriented perpendicular to the substrate (~1-2 m off bottom), and images were haphazardly taken 2-3 per minute. In addition, at most sites, at least one horizontal transect was made along a constant depth contour wherever Agaricia spp. and Montastraea cavernosa corals first became common (usually between 50 and 70 m depths). These horizontal transects were typically along the face of the vertical rock wall or overhanging rock buttresses, so the camera was pointed straight forward, perpendicular toward the wall. Time and distance of the horizontal transects varied depending on currents and topography, but generally were about 15 minutes, and transited ~100 m, for a total of 30 images. Fish were identified throughout the dive using the video camera pointed forward and down ~20° to view from the horizon to close up. In addition, fish video transects of approximately 30 minutes were made at each site along the upper wall and deep fore-reef crest. Direction of transects were haphazard, but generally headed along slope, but also depended on the ship's maneuverability with the wind and current.

The *Mohawk* ROV was equipped with a collection skid, which consists of a small 5-function manipulator, five suction buckets, and a Plexiglas box. Specimens of corals, sponges, and algae were collected at most sites and will be used for museum specimens and taxonomic identification, genetic analysis, and coral health studies. Specimens were photographed in situ, and in the ship's lab, and then preserved in 95% ethanol, 5% formalin, or TRIzol. ROV sensors included: Seabird Fastcat 49, CO₂ and pH. Shipboard oceanographic data



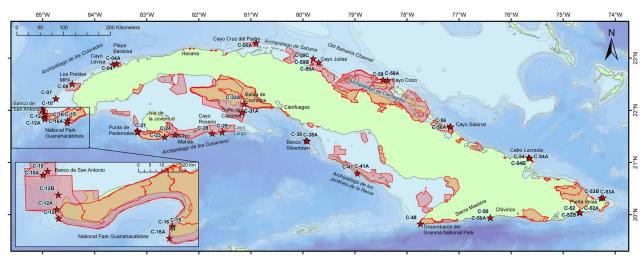


Fig. 1. Map of Cuba showing ROV dive sites (red stars) on mesophotic reef sites (30-150 m depths) during the *F.G. Walton Smith* cruise, May 14 to June 12, 2017. Pink polygons- marine protected areas (UNEP-WCMC, 2017).

continuously recorded surface water data including temperature, salinity, florescence, and dissolved oxygen, and ADCP data. A shipboard CTD and water sample rosette were cast daily to profile the dive sites.

Metadata and habitat dive notes were recorded during the dive into a Microsoft Access database which links to the ROV navigation data. Teams of U.S. and Cuban scientists specializing in coral, sponge, algae, and fish taxonomy recorded data in

separate Access databases which were later compiled with the benthic habitat database. Duplicate sets of the video/photo data and specimens are curated in the U.S. at CIOERT (HBOI-FAU), and in Cuba at CIM-UH/ANC/ICIMAR/CNAP.

RESULTS

Thirty-six mesophotic reef sites around the coast of Cuba were surveyed during 43 ROV dives (Fig. 1, Table 1). Total ship

Table 1. ROV dive sites on Cuba's mesophotic reefs during the F.G. Walton Smith cruise, May 14 to June 12, 2017.

Station	Dive No.	Location	Latitude	Longitude	Depth Range (m)
C-04	1	Cuba, NW coast, Cayo Levisa MPA	22°53.483'N	83°34.969'W	25-188
C-04A	2,3	Cuba, NW coast, 2 nmi north of Cayo Arenas, Cayo Levisa MPA	22°52.638'N	83°39.103'W	49-172
C-06	4,5	Cuba, NW coast, Norte de Los Pretiles MPA	22°30.487'N	84°25.907'W	43-170
C-07	6,7	Cuba, NW coast, Archipiélago de Los Colorados, No MPA	22°13.405′N	84°44.837'W	45-178
C-10	8	Cuba, W coast, Banco de San Antonio MPA, east wall	22°00.901'N	84°59.947'W	25-183
C-10A	9	Cuba, W coast, Banco de San Antonio MPA, south wall	22°00.017'N	85°00.977'W	30-150
C-12	10	Cuba, W coast, Los Cayuelos, Guanahacabibes National Park	21°50.005′N	84°57.426'W	149
C-12A	11	Cuba, W coast, Faro Roncali, off lighthouse, Guanahacabibes National Park	21°52.111'N	84°57.824'W	45-156
C-12B	12	Cuba, W coast, Las Tumbas, Guanahacabibes National Park	21°55.477'N	84°57.494'W	38-148
C-15	13	Cuba, W coast, El Almirante, Punto de Buceo, Cabo Corrientes, Guanahacabibes National Park	21°48.224'N	84°31.137'W	25-145

C-16A	14	Cuba, W coast, Cabo Corrientes, Guanahacabibes National Park	21°45.512'N	84°31.874'W	28-150
C-21	15	Cuba, S coast, SW tip of Isla de la Juventud, Punta de	21°35.263'N	83°10.325'W	25-158
0-21	10	Pedernales MPA	21 33.203 N	03 10.323 W	20-100
C-21A	16	Cuba, S coast, SW Isla de la Juventud, bay on west side of point, Punta Francés MPA	21°36.010'N	83°10.943'W	28-150
C-23	17	Cuba, S coast, SE Isla de la Juventud, Punta Brava MPA	21°29.419'N	82°39.228'W	37-150
C-24	18	Cuba, S coast, E of Isla de la Juventud, Archipiélago de Los Canarreos, Cayo Matías MPA	21°31.924'N	82°27.480'W	40-150
C-28	19	Cuba, S coast, Archipiélago de Los Canarreos, Cayo Rosario MPA	21°33.469'N	81°44.401'W	29-150
C-29	20	Cuba, S coast, Archipiélago de Los Canarreos, Cayo Largo del Sur MPA	21°34.854'N	81°33.130'W	26-157
C-31A	21	Cuba, S coast, Golfo de Cazones, Punta Brava, Sur Ciénaga de Zapata, no MPA	21°57.650'N	81°10.355'W	18-150
C-33A	22	Cuba, S coast, Bahía de Cochinos, east wall, Zapata National Park	22°07.444'N	81°07.518'W	30-150
C-38	23,24	Cuba, S coast, south of Cienfuegos, Banco Silvertown, west wall, proposed MPA	21°25.180'N	79°56.377'W	24-150
C-38A	25	Cuba, S coast, south of Cienfuegos, Banco Silvertown, south wall, proposed MPA	21°24.470'N	79°54.942'W	40-153
C-41	26	Cuba, S coast, Cayo Caballones, south slope, Jardines de La Reina MPA	20°47.495'N	78°57.323'W	30-150
C-48	27,28	Cuba, S coast, Cabo Cruz, Desembarco del Granma National Park	19°49.600'N	77°44.551'W	40-154
C-50	29,30	Cuba, S coast, Sierra Maestra, Chivirico Proposed MPA	19°56.551'N	76°23.937'W	27-155
C-52	31	Cuba, S coast, Sierra Purial, Punta Imías, no MPA	20°02.601'N	74°41.498'W	60-161
C-52A	32	Cuba, S coast, Sierra Purial, east tip of Punta Imías, 1 nmi east of ROV 31, no MPA	20°02.513'N	74°40.345'W	43-156
C-53A	33	Cuba, eastern tip of Cuba, NW Punta Maisí, Punta Silencio, no MPA	20°19.007'N	74°16.001'W	54-155
C-53B	34	Cuba, eastern tip of Cuba, NW Punta Maisí, Punta del Fraile, no MPA	20°19.482'N	74°13.940'W	50-142
C-54	35	Cuba, NE coast, Cabo Lucrecia, no MPA	21°05.508'N	75°39.821'W	45-151
C-54B	36	Cuba, NE coast, Cabo Lucrecia, proposed MPA	21°04.817'N	75°38.530'W	65-156
C-56	37,38	Cuba, NE coast, Archipiélago Sabana-Camagüey, Cayo Sabinal MPA	21°41.096'N	77°10.241'W	24-150
C-58	39	Cuba, NE coast, Archipiélago Sabana-Camagüey, Cayo Coco, proposed MPA	22°34.192'N	78°22.361'W	42-140
C-58A	40	Cuba, NE coast, Archipiélago Sabana-Camagüey, Cayo Coco, proposed MPA (just north of NW edge of MPA)	22°35.262'N	78°27.706'W	46-144
C-59A	41	Cuba, NE coast, Archipiélago Sabana-Camagüey, Cayo La Vela MPA	22°54.858'N	79°41.854'W	22-143
C-59C	42	Cuba, NE coast, Archipiélago Sabana-Camagüey, Cayo Jutía, ¼ nmi N of snorkel Station, no MPA	22°59.446'N	79°48.188'W	30-147
C-60A	43	Cuba, NE coast, Archipiélago Sabana-Camagüey, Cayo Cruz del Padre, inside and outside MPA	23°17.792'N	80°53.809'W	40-161

transit around the island covered ~2,778 km. Dives ranged from depths of 188 m to 25 m and resulted in a total dive time of 103 hours, covering approximately 27 km. A total of 21,146 digital still images documented habitat and species, and photo transect images (7,063) will be used for future analyses of percent cover and density of corals, sponges, and algae. The high-definition video will be used to document species composition and density of fish. A total of 343 specimens of benthic macro-invertebrates and macro-algae were collected with the ROV.

OCEANOGRAPHIC DATA

The surface currents along the western half of Cuba, between Havana and Cienfuegos, were likely part of the Cuban Countercurrent, which flows counterclockwise against the Caribbean and Florida Currents (Fig. 2). Along the southeastern coast from Archipelago Jardines de la Reina to the eastern end, which abuts the northern steep slope of the Cayman Trench, currents were generally eastward. Surface currents were counterclockwise around the eastern end of Cuba. Surface currents along the northeastern Cuban coast, however, did not have a prevailing direction, and were likely influenced by the competition between the Florida Current and Antilles Current and by tidal fluctuations. Historical measurements by Atkinson et al. (1995) suggested that the residual currents in the Old Bahamas Channel were predominantly westward toward the Florida Straits. Temperatures recorded by the ROV showed surface temperatures ranging from 27.09-29.52°C for all dives (Table 2). In general, temperatures were greatest along the south coast (29.52°C). The coolest temperatures were found along the northeast coast (27.09°C). At mid-depth of the dives (50 m, in the deep mesophotic zone), temperatures ranged from 25.26-28.62°C. Minimum temperatures at the deeper depths (162-182 m) were 21.35- 25.93°C. A thermocline of 25-28°C was common around 80-100 m depths. Surface salinities ranged from 35.11-36.73 PSU and at 50 m depths were 35.14-36.77 PSU. Visibility based on ROV video was excellent at most sites (estimated 30- >50 m), except sites

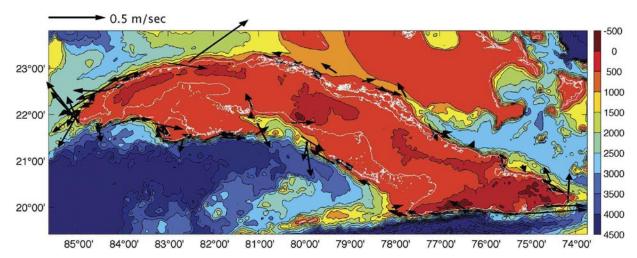


Fig. 2. Shelf-edge currents around Cuba from 6-hourly mean surface currents (arrows) measured with shipboard ADCP during the *F.G. Walton Smith* cruise, May 14 to June 12, 2017.

Table 2. Oceanographic data collected with a Seabird Fastcat 49 on the *Mohawk* ROV during the *F.G. Walton Smith* cruise, May 14 to June 12, 2017.

	Maximum Depth		Mid Depth (50 m)		Surface (1 m)		
	Depth (m) Max	Temp.(°C) Range	Salinity (PPT) Range	Temp. (°C) Range	Salinity (PPT) Range	Temp. (°C) Range	Salinity (PPT) Range
Northwest Coast	177.27	22.47-24.19	36.84-36.89	26.77-27.44	36.31-36.40	27.71-28.27	36.16-36.37
West Coast	182.13	22.53-25.93	36.65-36.90	27.50-28.25	36.29-36.37	28.17-28.58	36.28-36.31
Southwest Coast	162.29	22.69-24.15	35.59-36.89	27.68-28.46	35.14-36.41	28.53-29.52	35.11-36.41
Southeast Coast	162.06	22.53-24.41	36.80-36.93	25.99-28.62	36.11-36.67	28.42-29.48	36.04-36.35
Northeast Coast	162.01	21.35-25.56	36.72-37.41	25.26-27.88	36.39-36.77	27.09-28.54	35.64-36.73
Total	182.13	21.35-25.93	35.59-37.41	25.26-28.62	35.14-36.77	27.09-29.52	35.11-36.73

near stream runoff. Surface currents at the dives sites varied from 10 to 51 cm s⁻¹ (0.2-1 kn), but occasionally were in excess of 100 cm s⁻¹ during which the ROV could not be safely deployed and recovered.

GEOMORPHOLOGY

The ROV dives documented the deep mesophotic habitat surrounding Cuba. Many of the dive sites, especially on the eastern end of Cuba, had never been explored previously. All 43 dives confirmed the presence of MCE habitat. It appears from these dives that, just like the shallow reefs that fringe most of the Cuban coast, the deep reefs also parallel most of the shelf edge for nearly 2,800 km, and include deep fringing reefs that are close to shore (primarily along the northwest coast, southeast coast, and northeast coast), the barrier reefs along the various archipelagos (Los Colorados, Canarreos, Jardines de la Reina, and Sabana-Camagüey), and deep-water banks (Banco de San Antonio, and Banco Silvertown) (for terminology of reef types, see page 390, Zlatarski and Estalella, 1982; González Ferrer, 2004). Topographically, the most conspicuous features of most dive sites were the deep island slope (125->150 m), deep fore-reef escarpment (vertical wall; 50-125 m), and deep fore-reef (30-50 m) (Fig. 3). The geology of the deep reef features are likely carbonate limestone, and none of the sites appeared to be granitic, even at the southeastern coast off the Sierra Maestra mountain range where we expected granitic rock slopes as the coastline drops from +1,974 m (mountain peak) to -7,686 m (Cayman Trough).

The following is a general description of the deep fore-reef and slope (Figs. 3, 4).

Deep Island Slope (125- >150 m)- At many sites, this zone was a steep (70°), relatively smooth rock pavement slope intersected with low relief, vertical sediment chutes (1-m deep, 1-3 m wide), although some sites had slopes varying between 45° and 90° at 150 m. The surface rock usually showed horizontal layering and had scallop-shaped erosional features (10-30 cm diameter). This zone had the lowest diversity and density of biota. Fathometer profiles showed slopes of 10-45° extending to depths of 500 m.

Deep Fore-Reef Escarpment (the 'Wall', 50-125 m)- This zone formed a near-vertical wall (80-90° slope) at most sites. The base of the wall was commonly between 100 and 125 m depths although at many sites the wall continued to >150 m (>200 m at Site C-29 off the NE coast). At some sites, the upper wall (45-80 m) was formed of large rock "buttresses" which overhung

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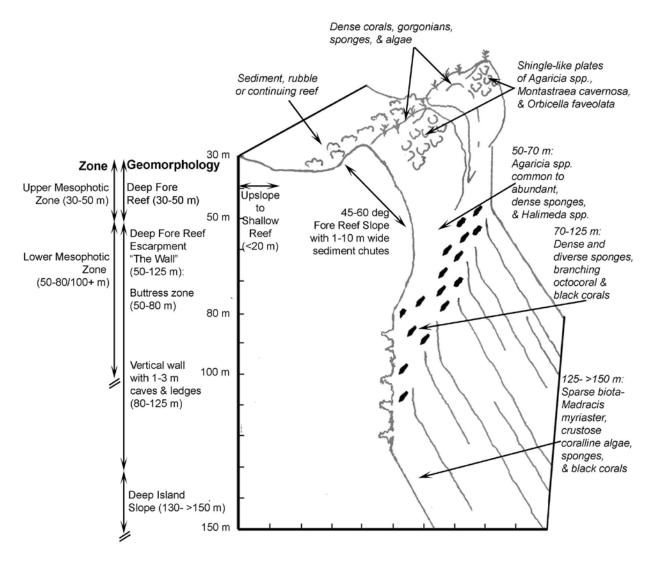


Fig. 3. Schematic of general geomorphological features and biozonation of Cuba's mesophotic reefs showing a well-developed fore reef zone and wall with buttresses.

the plane of the wall by 3-5 m. The buttresses were generally 10-20 m wide and were separated by sediment chutes which provided a conduit for the downslope transport of the shelf sediments (Hoskin et al., 1986). Fewer macrobiota occurred in the chutes. Between depths of 60 and 125 m, the wall was highly eroded at many sites, with karst-like topography forming caves (1-3 m diameter) and ledges (1-2 m wide).

Deep Fore-Reef Crest and Slope (30-50 m)-Seaward of the shallower reef zones (3-25 m), a zone of shelf-edge reefs generally paralleled the shelf-edge break. At some sites the deep fore-reef crest formed a ridge or sill-like feature (3-5 m relief). Generally, this ridge was not continuous but a series of mounds (5-10 m diameter, 3-5 m high). The fore-reef slope of this zone was typically steep (45-60°) with rocky escarpments, and spurs and grooves. The fore-reef slope

of this zone was often intersected with through the reef. At some sites these were sediment chutes (1-10 m wide) which cut actually fringing reefs close to the shore

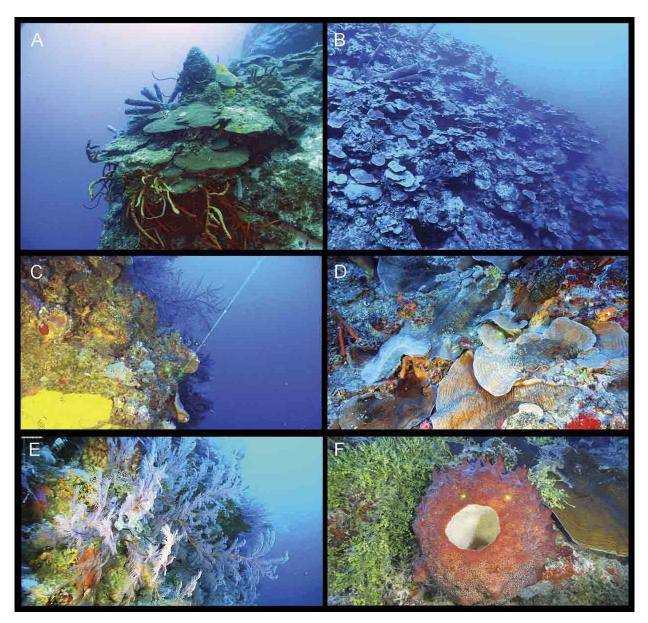


Fig. 4. Habitat and reef biota of Cuba's mesophotic reefs. Deep fore-reef slope: A) 1-m diameter plates of *Orbicella faveolata*, conical dead *Montastraea cavernosa*, tube sponges (*Aplysina archeri*), and rope sponges (*Agelas sceptrum, Aplysina cauliformis*), Punta de Pedernales MPA, Site C-21, 33 m; B) dense shingle-like corals *O. faveolata, Agaricia* spp., *Montastraea cavernosa*, Bahía de Cochinos, Site C-33A, 30 m. Deep fore-reef escarpments (the 'wall'): C) rock buttress with dense array of sponges (yellow sponge- *Siphonodictyon coralliphagum*), black corals, branching octocorals, and stylaster corals, Guanahacabibes National Park, Site C-12B, 82 m (lasers 10 cm); D) plates of *Agaricia* spp., Guanahacabibes National Park, Site C-12B, 64 m (lasers 10 cm); E) black coral bushes *Plumapathes* sp. on vertical wall, Guanahacabibes National Park, Site C-12B, 79 m; F) large barrel sponge *Xestospongia* sp., green alga *Halimeda* sp., and scleractinian *Agaricia* sp., Cayo Largo del Sur MPA, Site C-29, 61 m.

(e.g., along the SE coast off Sierra Maestra, Bahia de Cochinos, and Guanahacabibes). Elsewhere the shelf-edge reefs occurred seaward of the Archipelagos (e.g., Los Colorados, Canarreos, Jardines de la Reina, Sabana-Camagüey) and may be considered barrier reefs (Zlatarski and Estalella, 1982; González Ferrer, 2004). Also the two deep-water banks that were surveyed (San Antonio and Silvertown) had similar topography along the shelf-edge break.

BIOZONATION

Mesophotic coral ecosystems (MCEs) are characterized by light-dependent corals and associated communities typically found at depths ranging from 30-40 m and extending to >150 m in tropical and subtropical regions (Baker et al., 2016). Various researchers have divided what is now called the mesophotic zone into different subzones and depths (e.g., Goreau and Goreau, 1973; Lang, 1974; James and Ginsburg, 1979; Ginsburg et al., 1991; Lesser et al., 2009; Locker et al., 2010; Sherman et al., 2010; Clark et al., 2014; Reed, 2016; Laverick et al., 2017). For this paper, we use the depth of 30 m as the upper extent of the upper mesophotic zone and have used the depth of 50 m as the transition between the upper and lower mesophotic zones where we generally saw a fairly distinct change in geomorphological zonation and species. Therefore, the upper mesophotic zone is generally in the deep fore-reef, and the lower mesophotic zone is basically on the wall and buttresses of the deep fore-reef escarpment. The definitive maximum depth of the mesophotic zone is problematic and varies from site to site based on light, sedimentation, and substrate (Baker *et al.*, 2016).

Deep Island Slope (125- >150 m)-Although the deep island slope often had photosynthetic species such as crustose coralline red algae (CCA) and encrusting green algae, there were no zooxanthellate corals. Ahermatypic corals such as Madracis myriaster were common along with various azooxanthellate, branching octocorals (Ellisella elongata, Nicella goreaui. Swiftia exserta), black als (Antipathes sp., Tanacetipathes sp., Stichopathes sp.), and stylasterine corals. Sponges were common but not as diverse or dense as on the wall. Common sponge species include a bright yellow crustose sponge (Order Verongiida) forming abundant large patches (20-100 cm²), thick-walled vase sponge (*Xestospongia* sp. Cu-01), several species of *Oceanapia* with elongated projections, and massive sponges from various taxa (Orders Haplosclerida, Suberitida, and Tetractinellida).

Deep Fore-Reef Escarpment (the 'Wall', 50-125 m)- This is the most impressive and recurring feature of most sites along the Cuban shelf edge. This zone had the greatest diversity and density of macrobiota of the deep reefs. Starting at 100-125 m. nearly all vertical surfaces were covered with sponges (e.g., Agelas spp., Aplysina Callyspongia spp., Ceratoporella spp., nicholsoni, Clathria spp., Geodia spp., Leucetta spp., Oceanapia spp., Petrosia spp., Plakortis spp., Spirastrella spp. and Xestospongia spp.); algae (CCA, Halimeda spp., Peyssonneliaceae (several species), Lobophora spp., and Dictyota spp.), branching octocorals (*Nicella* spp., *Ellisella* spp., Swiftia exserta), and black corals (bushy Antipatharia to 1-2 m, and dense fields of Stichopathes sp.). In general, the dominant scleractinian corals on the wall were Agaricia (A. fragilis, A. grahamae, A.

lamarcki, and possibly A. undata). These typically first occurred at the maximum depths of 80-122 m, and were often quite common to abundant between 50 and 70 m and on the upper slope of the buttresses. Therefore, the bottom of the mesophotic zone generally was around 80-120 m, but this was very variable by site. Corals were less common where there was less light, e.g., under the overhanging buttresses, on north facing walls, or where the deep fore-reef did not form a sill that blocks the flow of sediments downslope. For example, one reef at Faro Roncali, Guanahacabibes National Park (Site C-12A) had no ridgelike deep fore-reef to block sediment flow from the shelf, and as a result no scleractinian corals or CCA were observed on the wall. But, Site C-12B nearby had a distinct shelf-edge reef with 5-m relief which blocked sediment flow except in the area of the chutes, and also had dense populations of plate coral on the wall and upper slope.

Deep Fore-Reef (30-50 m)- This region forms the upper mesophotic zone where higher abundance and diversity of scleractinian species were characteristically found. Agaricia spp., Montastraea cavernosa, Orbicella faveolata, Stephanocoenia intersepta and Porites astreoides were dominant. Several sites, including Banco de San Antonio (Site C-10A), Guanahacabibes (C-12B), Bahía de Cochinos (C-33A), and Cayo Sabinal (C-54B), had dense shingles of plate corals (O. faveolata or Agaricia spp.) on the fore reef slope and crest (estimated 60-80% cover at Site C-33A). Common shallow reef sponges were abundant here as well, such as Agelas spp, Aiolochoria spp., Aplysina spp., Amphimedon spp., Cinachyrella spp., Cliona spp., Ircinia spp., Niphates spp., Smenospongia spp., Spongia spp., Verongula spp., and Xestospongia spp.

BIODIVERSITY

A total of 477 taxa of benthic macrobiota and 178 fish taxa were identified from the ROV video/photo surveys and from the specimens collected (Appendices 1 and 2, respectively). These data will be quantified in the future for percent cover and for documenting densities of corals and fish. A total of 343 mesophotic benthic samples were collected during this expedition; preliminary analyses indicate that some are new species and records of depth or distribution.

Corals- A total of the 101 cnidarian taxa that were identified morphologically included 46 Scleractinia, 32 Alcyonacea (branching octocorals and Alcyoniina), and 15 Antipatharia. The deepest occurrences of identified zooxanthellate corals were Agaricia sp. (122 m), Stephanocoenia intersepta (104 m), and M. cavernosa (101 m) (Table 3). These were the most common scleractinians on the wall in the deeper mesophotic zone, whereas O. faveolata was more common in the upper mesophotic zone. The most common octocorals were in the genera Ellisella, Nicella, and Antillogorgia (syn. *Pseudopterogorgia*), and Swiftia exserta. The most common black corals included Stichopathes spp., Tanacetipathes tanacetum, Tanacetiphates sp., and Antipathidae. In addition, 123 scleractinian corals were collected, some of which will be used for genetic analysis.

Sponges- Cuban MCEs proved to be a very favorable habitat for marine sponges in both species richness and abundance. We identified 296 distinct taxa: 280 Demospongiae, 9 Homoscleromorpha, and 7 Calcarea, of which 115 (39%) were identified to species and 107 to genus (35%). The rest were only tentatively identified to families, orders, or classes, and are the subject of ongoing taxonomic research.

Table 3. Deepest observations of macrobiota during ROV dives at all Sites during the *F.G. Walton Smith* cruise, May 14 to June 12, 2017.

Taxa	Max. Depth (m)
Scleractinia	
Agaricia sp.	122
Stephanocoenia intersepta	104
Montastraea cavernosa	101
Helioseris cucullata	79
Agaricia grahamae	74
Orbicella faveolata	72
Octocorallia	
Swiftia exserta	188
Porifera	
Verongiida Cu-01 (yellow crust)	183
Xestospongia sp. Cu-01	168
Ceratoporella nicholsoni (sclerosponge)	159
Xestospongia muta	134
Leucetta sp. (Calcarea)	124
Chlorophyta	
Chlorophyta (thin green crust)	169
Halimeda sp.	127
Rhodophyta	
Crustose coralline (CCA)	169
Peyssonneliaceae	139
Ochrophyta	
Lobophora sp.	139
Dictyota sp.	100
Chordata	
Pterois volitans/miles (Lionfish)	188

The ten most frequently recorded species were the barrel sponge *Xestospongia muta*, *Xestospongia* sp. Cu-01, an encrusting unidentified bright yellow Verongiida sp. Cu-01, large tubular *Aplysina archeri*, several species of *Agelas* (*A. sceptrum*, *A. dilatata*, and *A. citrina*), and *Mycale laxissima*. Sponges colonized all depths and substrates surveyed. The deepest records were a bright yellow Verongiida sp. Cu-01 encrusting on

rock (183 m deep), Xestospongia sp. Cu-01 (168 m), Ceratoporella nicholsoni (159 m), X. muta (134 m), and Leucetta sp. (124 m) (Table 3). Peak diversity started at depths of 100-125 m in which sponge morphologies (crusts, plates, tubes, spherical, vases, and branches) and multicolor patterns abounded. From the collections, there are at least 10 sponge species that are new to science within the genera of Aplysina and Verongula (Verongiida); Amphimedon, Callyspongia, and Xestospongia (Haplosclerida); Clathria (Poecilosclerida); Cinachyrella (Tetractinellida); (Suberitida); Erylus (Tetractinellida) and *Plakortis* (Homosclerophorida).

Algae- A total of 63 taxa of macroalgae (29 Chlorophyta, 25 Rhodophyta, 9 Ochrophyta) and 1 Cyanobacteria (syn., Cyanophyta) have been identified to date. The most frequent taxa were crustose coralline algae (CCA), Halimeda copiosa, Lobophora spp., and *Dictyota* spp.. In general, the diversity and coverage of algae were very low on the deep island slope where only CCA and thin encrusting green algae occurred. Between 50 and 100 m, the algal diversity and cover increased, with H. copiosa and CCA dominating. In the upper mesophotic zone (30-50 m), the species richness and cover increased even more, with Lobophora spp. and other Dictyotales dominating. Some interesting records of maximum depth of algal occurance include an unidentified green crust (169 m), CCA (169 m), Peyssonneliaceae (139 m), Lobophora spp. (139 m), and Halimeda spp. (127 m) (Table 3). Cyanobacteria were observed at eight sites and abundant at one described below. Detrital leaves of the marine angiosperm Thalassia testudinum were also observed on the deep island slope at some sites showing the vertical connectivity between coastal-marine ecosystems.

Fish- A total of 178 species of fish were observed throughout the survey (Appendix 2). Between 60 and 150 m, fish diversity and abundances were low at all sites, whereas from 30-60 m the diversity increased, as well as the frequency and abundance of fish species. The deep island slope zone (125-150 m) was dominated by Blackcap Basslet (Gramma melacara), Cave Basslet (Liopropoma mowbrayi), Sunshinefish (Chromis insolata), Blackfin Snapper (Lutjanus buccanella), Squirrelfish (Holocentrus adscensionis) and Longspine Squirrelfish (*H. rufus*). The walls (lower mesophotic reef zone, 50-120 m) and upper deep fore-reef (30-50 m) were inhabited by species common on Cuban shallow reefs, including numerous species from the families Acanthuridae, Balistidae, Carangidae, Chaetodontidae, Haemulidae, Lutjanidae. Holocentridae. Labridae. Pomacanthidae, Pomacentridae, Serranidae. Groups of jack (*Caranx latus*, *C.* ruber and C. lugubris) were frequently observed. Ten species of grouper (Serranidae) were identified throughout the sampling area including Coney (Cephalopholis fulva), Graysby (Cephalopholis cruentata), Nassau Grouper (Epinephelus striatus) and Red Hind (*E. guttatus*). Hundreds of Ocean Triggerfish (Canthidermis sufflamen) were observed at Banco de San Antonio MPA. Courtship and nest building behavior by the triggerfish were observed leading to the conclusion that this is a spawning aggregation site. A spawning aggregation of Dog Snapper (Lutjanus jocu) was also observed at a site in Archipelago de Camagüey between 60 and 70 m depths. This site is part of the Centro y Oeste de Cayo Coco Ecological Reserve (Perera-Valderrama et al., 2018). We propose this site should be made a no-take fishery reserve or, at least,

a no-take during the spawning season(s) since other commercially important species might also spawn there. We observed Lionfish (*Pterois volitans/miles*) at most sites; the greatest depth was 188 m at Cabo Lucrecia (Site C-54B). The abundance of the invasive Lionfish was generally low, generally <10 observed during a four hour dive at most sites. An exception was at Cayo Coco (Site C-58) where they were relatively abundant with 38 counted on one dive. However, sampling by ROV video cameras could bias fish counts due to avoidance and therefore abundances of Lionfish could be greater than we observed.

REGIONAL COMPARISONS OF CUBA'S MCES

Sketches of reef profiles were drawn to illustrate various regions of Cuba (Figs. 5-8). These are not the specific coastal zones outlined in Areces (2002), but represent the reef profiles of six sites within each of the following general regions of Cuba: western, southern, eastern, and northern. Photographs taken with the ROV that represent each of these regions are provided in Appendix 3. Four general habitats were added to the sketches: rock pavement, mud/ sediment, caves/ledges, and coral shingles. The zone of eroded rock forming caves or reentrants and ledges was found on the wall of the deep fore-reef escarpment at many of the sites. These features are karst-like topography likely formed during the low sea levels of the last glacial period. The coral shingles zone consisted mostly of O. faveolata and Agaricia spp. and was found at some sites on the deep fore-reef crest and slope (generally 30-40 m). Each profile also includes the maximum depth observed at that site for the primary plate corals: M. cavernosa, Agaricia spp., O. faveolata, and S. intersepta (abbr. in Figs. 5-8).

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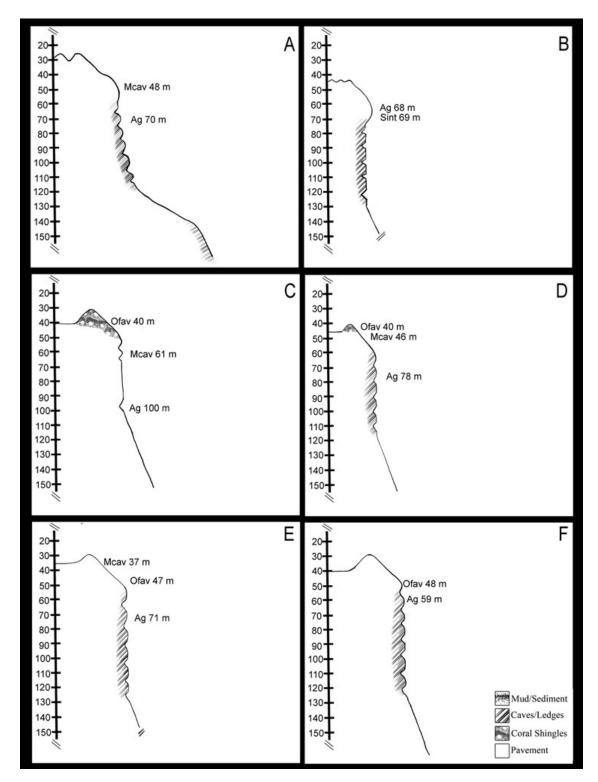


Fig. 5. Sketches of reef profiles selected to illustrate the various regions of Cuba's MCEs. (see Fig. 1). Western region (Archipelago de Los Colorados, Banco San Antonio, Guanahacabibes): A. Site C-04, B. Site C-07, C. Site C-10A, D. Site C-12B, E. Site C-15, F. Site C-16A. The specific depths and slopes were drawn to scale from observations made during the ROV dives.

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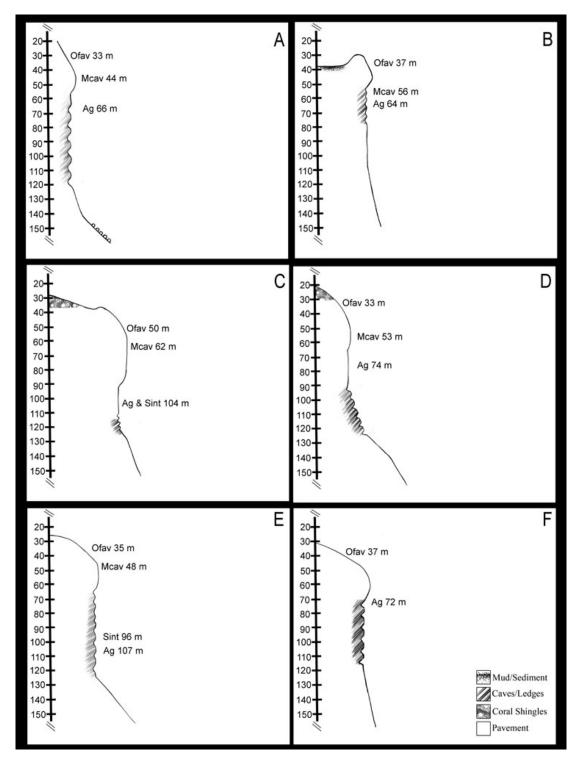


Fig. 6. Sketches of reef profiles selected to illustrate the various regions of Cuba's MCEs (see Fig. 1). Southern region (Isla de la Juventud, Archipelago de los Canarreos, Bahia de Cochinos, Banco Silvertown, Jardines de La Reina): A. Site C-21, B. Site C-23, C. Site C-28, D. Site C-33A, E. Site C-38, F. Site C-41. The specific depths and slopes were drawn to scale from observations made during the ROV dives

In the western region, the crest of the shelf-edge reefs was generally between 30 and 40 m (Fig. 5). This fore-reef zone generally consisted of a series of mounds (3-10 m relief) along the shelf-edge break that were intersected with sand chutes. Some had buttresses on the deep fore-reef slope and upper wall. Shingles of O. faveolata were found at Sites C-10A (Banco San Antonio) and C-12B (Las Tumbas, Guanahacabibes). The deep fore-reef escarpment generally was highly eroded with caves or reentrants and ledges at depths of about 60 to 110-130 m. Agaricia spp. generally was first observed around 70 m depth, but was as deep as 100 m at Banco San Antonio (Site C-10A). In addition, these shelf-edge reefs were all quite similar even though they comprise fringing reefs (Guanahacabibes), barrier reefs (Archipelago de Los Colorados), and bank reef (San Antonio).

The reefs of the southern region generally showed less pronounced mounds or ridges along the shelf-edge break (Fig. 6), and the shelf-edge break varied greatly among sites. For example, the fore-reef slope at SW Isla de Juventud (C-21) and Bahia de Cochinos (C-33A) continued up to at least 20 m depth where we had to cut off the transect for the safety of the ship. Site C-33A had the densest cover of O. faveolata coral shingles at any site, in depths of 20-30 m. Cayo Rosario (C-28) also had dense O. faveolata shingles on the deep fore-reef slope but these were mostly dead. Agaricia occurred in this region as deep as 107 m (Banco Silvertown, C-38). Also the zone of eroded topography varied greatly among the sites.

Shelf-edge reefs along the eastern region were highly variable (Fig. 7). The shelfedge break of the deep fore-reef was as deep as 50-60 m at some sites, and several sites near the eastern tip of Cuba showed extensive erosion and lacked the vertical wall (Punta Imias, C-53; Punta Maisi, C-53A; Cabo Lucrecia, C-54B). Sherman et al. (2010) also found lower topographic relief and less pronounced escarpment on the southeastern region of Puerto Rico. They attribute this due to the fact that the eastern facing slopes are more exposed to trade-wind generated swells and tropical hurricanes that typically approach from the southeast. They also found that MCEs are patchier and more widely spaced on the southeast slopes.

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The northern region of Archipelago de Sabana-Camagüey (Fig. 8) also showed variability ranging from eroded, low slope geomorphology (Sites C-58, C-58A) to the walls and buttresses (Sites C-56, C-59C, and C-60A) that were typically present at the western region. One site at Cayo Coco (C-58A) was mostly smooth rock pavement, with a relatively narrow region of highly eroded, rock ledge habitat at 70-90 m. Also the maximum depth of *Agaricia* tended to be shallower at several sites in this region perhaps due to the northern facing walls (45 m at C-58, 52 m at C-59C).

To date, only general comparisons of benthic macrobiota and fish distributions can be made for the various mesophotic regions of Cuba from these data. Quantitative analyses of the photo transects for macrobenthos cover and density as well as analyses of fish densities from the video transects are planned as part of our multinational research efforts and will allow for more detailed comparisons of any regional and depth differences in the future. Appendices 1 and 2 provide the species lists of macrobenthos (invertebrates and algae) and fish, respectively, by

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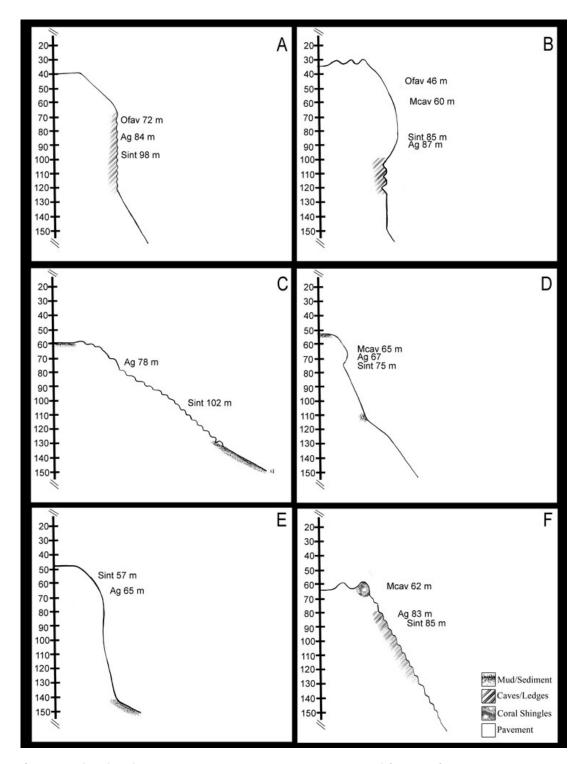


Fig. 7. Sketches of reef profiles selected to illustrate the various regions of Cuba's MCEs (see Fig. 1). Eastern region (Granma National Park, Chivirico MPA, Punta Imias, Punta Maisi, Cabo Lucrecia): A. Site C-48, B. Site C-50, C. Site C-52, D. Site C-53A, E. Site S-54, F. Site C-54B. The specific depths and slopes were drawn to scale from observations made during the ROV dives.

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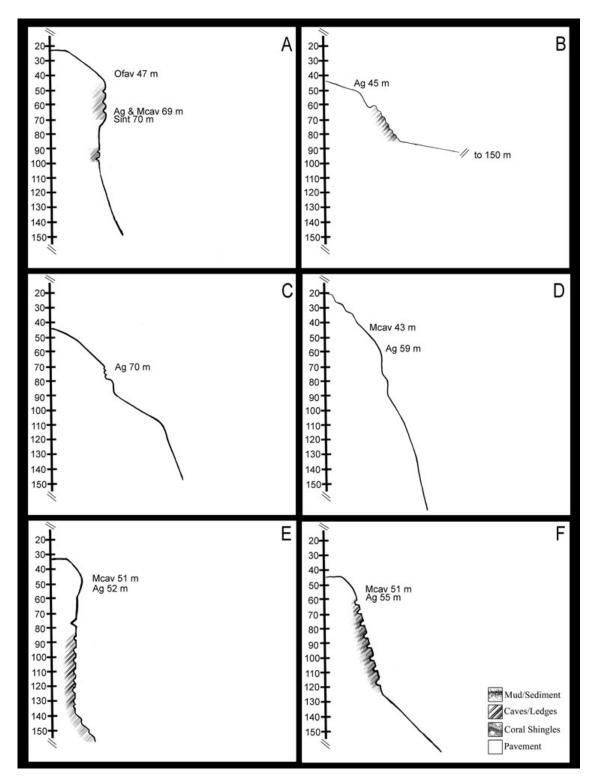


Fig. 8. Sketches of reef profiles selected to illustrate the various regions of Cuba's MCEs (see Fig. 1). Northern region (Archipelago de Sabana-Camagüey): A. Site C-56, B. Site C-58, C. Site C-58A, D. Site C-59A, E. Site C-59C, F. Site C-60A. The specific depths and slopes were drawn to scale from observations made during the ROV dives.

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site and region based on the ROV annotations and sample collections. In general, there were some regional and site specific differences, but overall the MCE communities and species distributions were relatively similar around the island. For these comparisons we have divided the dive sites into 6 regions (see Fig. 1): Northwest coast-Archipelago de Los Colorados, Sites C-04 to C-07; West coast-Banco de San Antonio, Guanahacabibes, Sites C-10 to C-16A; Southwest coast- Isla de la Juventud, Archipelago de los Canarreos, Golfo de Cazones, Bahia de Cochinos, Sites C-21 to C-31A: Southeast coast-Banco Silvertown, Jardines de La Reina, Granma National Park, Chivirico MPA, Punta Imias, Sites C-38 to C-52A; Northeast Coast-Punta Imias [north side], Cabo Lucrecia, Archipelago de Sabana-Camagüey, Sites C-53 to C-60A). Based on the presence/ absence species data of Appendix 1, both the northeastern and southeastern regions had the greatest number of macrobenthos taxa (259 taxa each). The west and southwest regions had 229 and 228 taxa, respectively, and the northwest had the least (205 taxa). However, the sites were not evenly distributed among these regions, so comparisons of species richness are only tentative. However, the number of taxa per site is more relevant, typically, each site had a similar amount of ROV time, basically one dive for the macrobenthic surveys. Sites C-48 (Granma) and C-50 (Chivirico) on the southeast coast had the most taxa (126 and 125 taxa, respectively), but C-52 (south Punta Imias), which is also on the southeast coast, had the least (57 taxa). Unlike most of the other sites in that region, Site C-52 was muddy from 160 to 130 m, and did not have the vertical rock wall and buttresses found at C-48 and C-50 (Fig. 7).

Corals- Cnidarian species richness of the mesophotic reefs based on the ROV annotations was greatest on Cuba's southeast coast, where a total of 146 taxa were observed. The sites with the greatest individual coral species richness (including Scleractinia, Octocorallia, and Antipatharia) were located on the southeast coast; 47 and 46 coral taxa having been recorded at Sites C-48 and C-50, respectively. The northwest coast had the fewest observed coral taxa (57 taxa), perhaps due in part to fewer stations within this region as compared to all other regions. The fewest coral taxa were observed at Site C-12A (11 taxa) at Guanahacabibes. This station was clearly an outlier in terms of observed coral species. Within octocorals, Ellisella sp. was observed at all sites while Nicella sp. was present in all sites except C-12A. Likewise, among the Scleractinia, Agaricia sp. were observed at all stations except C-12A. M. cavernosa was also commonly reported in the ROV annotations (32 of the 35 sites). Frequent Antipatharia taxa included Stichopathes sp. which was observed at all sites and Tanacetipathes sp. which occurred at 29 sites. Twenty taxa were only observed at single sites, the majority of which were either rare species (e.g., the scleractinians Dendrogyra cylindrus and Madracis myriaster). In terms of present/absence, few strong regional differences were observed among coral taxa, although the southeast coast showed the greatest species richness for scleractinian species and may represent a hotspot of reef-building species diversity. All other regions were generally similar in terms of observed coral taxa, with the exception of the northwest coast.

Sponges- Only two sponge species were present at all sites. The first, Xestospongia

sp. Cu-01, may be a new species and resembles X. muta, but with thicker walls and is relatively shorter and chubby in appearance. The second, as yet unknown, is a thick bright yellow verongiid crust (Verongiida sp. Cu-01) that lacks any fiber skeleton. Three species were found at 33 sites (Agelas sceptrum, Aiolochroia crassa, and X. muta). The most species-rich sites for sponges were found at Site C-07 (Archipelago Colorados) on the northwest coast with 72 taxa and, on the south coast, C-28 (Archipielago Canarreos) with 68 taxa and C-50 (Chivirico) with 66 taxa. However, overall the most species-rich region for sponges was the northeastern region (153 taxa), followed by the southeastern region (148 taxa) and western region (142 taxa). The fewest taxa were found in the northwest region (113 taxa) and southwest (131 taxa). By station the fewest sponges were found at Site C-52 (33 taxa) and C-52A (37 taxa) at Punta Imias on the southeast coast, and C-58A (36 taxa) at Archipelago de Camagüey on the northeast coast. Surprisingly, 99 sponge taxa were found exclusively at single sites (33% of the 296 morphotypes) which might reflect high level of endemism or widespread but patchy distributions. The most diversified genera were: 1) Agelas with 16 identified species (6 of which occur at least in 15 sites) and 10 unidentified morphotypes, and 2) Aplysina with 10 identified species (6 taxa occur at more than 15 stations) and 7 unidentified morphotypes. The interpretation of the regional and local patterns of sponge diversity will require further analysis that will study factors such as habitat, current patterns, and biogeographic affinities.

Algae- Macroalgal taxa that were common to all regions of the Cuban mesophotic coral reefs included the coralline red algae (Subclass Corallinophycidae) and the family

Peyssonneliaceae in the Rhodophyta; some green unidentified crustose forms, the genera Halimeda (particularly H. copiosa, H. goreaui, H. tuna), Avrainvillea, Penicillus, *Udotea* (particularly *U. cyathiformis*) in the Chlorophyta; and the genera *Dictyota* and Lobophora in the Ochrophyta. Site C-10 (Banco San Antonio, west coast) had the greatest number of taxa (24), whereas C-52 (southeast coast) which lacked the typical vertical wall had the least (4 taxa). The northeastern region had the highest number of macroalgal taxa identified so far (38 taxa), whereas the western regions had the least (24 taxa in the northwest region, and 25 in the southwest). These results could indicate better conditions for the development of macroalgae in the northeastern region of Cuba, and/or lower herbivory levels, but further work is still needed.

Fish- There were 178 fish taxa identified from the ROV annotations (Appendix 2), however only one, *Chromis insolata*, was present at all 36 sites. There were three species present at 33 sites; Gramma melacara, Halichoeres garnoti, and Stegastes partitus. An additional three species were found at 32 sites; Holacanthus tricolor, Chromis cyanea and the invasive Lionfish Pterois volitans/miles. Similar to sponges, the most species-rich site for fish was C-07 on the northwest coast, with 67 taxa observed. Site C-12B at Guanahacabibes had 63 taxa and C-48 on the southeast coast had 62 taxa. Regionally, the southeast coast was the richest with 112 fish taxa present; however, the site with the fewest taxa was on the southeast coast (Site C-52 with 17 taxa), which was also true for the sponges and algae, followed by Site C-52A (26 taxa) and C-04 on the northwest coast (27 taxa). Overall, the northwest coast had the lowest species richness with 91 taxa; however, this region had the fewest number of sites. Thirty-nine taxa were found only at single sites while 13 were found at only two sites. The number of species found at single sites will likely decrease after detailed analyses of ROV videotapes are completed and fishes overlooked during real-time dive annotations are added to the database. Fish assemblages were not significantly different from one another inside and outside the MPAs based on the non-metric multidimensional scaling ordination (MDS) plot of the presence/absence transformed fish data from the ROV observations. With the exception of one dive, fish assemblages from all dives were 40% similar to one another, regardless if they were inside or outside the MPAs. Based on the SIMPER (Similarity Percentages), the species that had the most effect on fish assemblage differences inside and outside the MPAs were higher abundances of Caranx lugubris, Haemulon plumierii, Scaridea, and Gramma loreto inside the MPAs and higher abundances of Holocentrus rufus outside the MPAs. In the western area, sites that coincided with marine protected areas (Banco de San Antonio, Guanahacabibes National Park, and Cayo Rosario) had greater abundances of commercially important species, i.e., snapper (Lutjanidae), grouper (Serranidae), jack (Carangidae) and mackerel (Scombridae). The sites outside of these marine protected areas had a lower abundance of these species, which could be an indicator of historical overfishing.

DISCUSSION

MESOPHOTIC CORAL HABITATS OF CARIBBEAN, GULF OF MEXICO AND SOUTHEASTERN U.S.

MCE habitats are quite common throughout the Western Atlantic regions of the

Caribbean, Gulf of Mexico, and southeastern U.S., and include deep island slopes, off-shore banks, and shelf-edge reefs. However, the MCE habitat and biota are quite variable within these regions. The general geomorphology and mesophotic zonation of the deep fore-reef slope and escarpment along the Cuban coast are strikingly similar to those described for the Bahamas (Porter, 1973; Reed, 1985; Ginsburg et al., 1991; Lesser and Slattery, 2011); Jamaica (Goreau and Goreau, 1973; Goreau and Land, 1974; Lang, 1974; Land and Moore, 1977; Dustan and Lang, 2018), and the Belize barrier and atoll reefs (James and Ginsburg, 1979). The 'Wall' at Belize also forms spectacular vertical cliffs from depths of 65 to 120 m, with vertical fissures, ledges and caves, and in the 1970s, the upper brow of the wall (45-65 m) had coral growth consisting of large colonies of plate corals up to 50% cover. At Discovery Bay, Jamaica, the wall forms an escarpment from 55 to 114 m, also with caves and ledges, and in the 1970s-1980s was dominated by sponges, algae, and corals (its current condition is unknown, J. Lang, pers. comm.). At that time, the deep fore-reef slope (30-65 m) at Jamaica had similar dominant coral, sponge, and algal species as do the Cuban deep reefs. The corals were predominantly Orbicella in high abundance to 45 m at some sites, Agaricia lamarcki on upper slope, A. undata, A. grahamae, Helioseris cuculata, and M. cavernosa on the lower slope; and the algae were dominated by crustose coralline algae, *Halimeda copiosa*, *H. cryptica*, plus algal turfs and fleshy macroalgae (Goreau and Goreau, 1973; Lang, 1974; Dustan and Lang, 2018). Hermatypic corals (primarily Agaricia spp.) dominated the wall around 70 m. The deep island slope (> 120 m) in Jamaica was also similar to the Cuban reefs, and had a similar heterotrophic community consisting of abundant sponges, including coralline sponges, the azooxanthellate scleractinians, alcyonaceans, fan and whip antipatharians, and stylasterine corals. In the Bahamas, Reed (1985) compared several locations including San Salvador Island that has a narrow shelf, and the windward and leeward sides of Grand and Little Bahama Banks. The wall of the deep fore-reef escarpment ranges from 45-143 m. The base generally ranges from 100-122 m. Some sites also have large buttresses occurring every 20-60 m along the top edge of the deep fore-reef slope, with sediment chutes incised between the promontories (Hoskin et al., 1986). Notches, ledges and reentrants or caves are common between 75 and 107 m, similar to Cuba. At San Salvador Island, Bahamas, these reentrants likely connect to the brackish lakes on the island which show tidal influence. These karst-like features associated with deep fore reefs of Jamaica, Belize, Bahamas and Cuba, likely formed during Late Quaternary low stands of sea level which were at least -120 m below present level during the last glacial maximum (Fairbanks, 1989). The major topographic features are apparently controlled by underlying Plio-Pleistocene features that are mantled by 1-10 m of Holocene reef accretions (James and Ginsburg, 1979; Ginsburg et al., 1991).

The Virgin Islands also support fairly extensive and diverse MCEs, predominately on steep walls around the islands and some areas of deep banks (Smith and Holstein, 2016). Extensive areas of the shelf that are potential MCE habitat (1,918 km²) at depths of 25-65 m may exceed the area of the shallow reefs. MCE

habitat around St. Croix is mostly on the steep island walls, which is typical of most Caribbean Islands and Bahamas, whereas St. Thomas and St. John have a broader shelf and deep banks. In the upper mesophotic zone (35 m), the dominant coral are *Orbicella* spp. and *Agaricia* spp. (Smith and Holstein, 2016). At deeper depths (65 m), the lettuce coral *Agaricia undata* dominates.

Off southwestern Puerto Rico, deep reefs occur at depths of 45-160 m; deep buttresses are common at depths of 45-65 m below the shallow barrier reef system (Appledoorn et al., 2016b). A terrace occurs at 80-90 m, and a steep wall extends down to 160 m (Sherman et al., 2010). This terrace feature which Sherman et al. (2010) suggests as being formed during the last deglaciation (14-15 ka) is not apparent at Cuba. Agaricia spp. and Madracis spp. corals dominate the MCE zone, and large colonies of A. undata are common. Seventeen species of scleractinian coral in the MCE zone (50-90 m).

Down current from Cuba are the mesophotic reefs of the northern Gulf of Mexico (GOM), including the Sister Sanctuaries (FGBNMS, FKNMS), and southeastern U.S. The geomorphology of these U.S. mesophotic reefs are quite different from those off Cuba. The shelf-edge reefs and banks in the northwestern GOM off Texas are built upon salt domes but also have a relatively steep mesophotic reef habitat which extends over a wide depth zone. FGBNMS is 180 km off shore Texas and is described as one of the most pristine coral ecosystems in U.S. waters (Hickerson et al., 2008). Although the top of the dome is a shallow water reef at depths of 18 m, the flanks extend into mesophotic depths down to 152 m (Clark et al. 2014). The mesophotic reefs at FGBNMS have been surveyed with ROV by some of our team (JV, JR) using similar methods as in Cuba (Voss et al., 2014). The mesophotic scleractinian coral communities on FGB and nearby banks outside the sanctuary have diverse hard coral populations (21 species) and are dominated by O. franksi, M. cavernosa, S. intersepta, and Agaricia spp. with Pseudodiploria strigosa and Colpophyllia natans also being relatively common. Also common at FGBNMS at mid-mesophotic depths (40-80 m) are algal nodules (rhodoliths) which we did not find at Cuba. The deep coral zone at FGBNMS from 85-150 m was primarily on the steep slopes of the banks, and dominated by azooxanthellate corals, antipatharian and gorgonian corals, bryozoa, and sponges. Many of these species are similar to Cuba on the lower mesophotic zone and deep island slope.

In the northeastern GOM, mesophoticdepth reefs are common all along the 70 m contour (60-90 m) from Alabama to the panhandle of Florida, and along the northwestern Florida shelf (Locker et al., 2010; Locker et al., 2016; see Reed, 2016 for map). However, for the most part, these lack zooxanthellate corals and are dominated by azooxanthellate corals such as *Madracis* sp. and *Oculina* sp., octocorals, black corals, sponges, and algae. In contrast to these reefs, Pulley Ridge, off the southwest Florida shelf, is one of the closest downstream mesophotic reefs to Cuba. It is a submerged intact barrier island that formed during the early Holocene marine transgression, and consists of the deepest photosynthetic coral reef in continental U.S. (Hine et al., 2008; Reed, 2016). The southern terminus of this geological feature is relatively flat limestone pavement and rubble at depths of 60-94 m,

and supports a MCE of photosynthetic hard corals (17 species), macroalgae (95), sponges (92), and a large variety of tropical fishes (86) (Harter et al., 2017; Reed et al., 2017a; Reed et al., 2018). A total of 12 hard coral species were identified at Pulley Ridge. Plate coral species at Pulley Ridge include Agaricia fragilis, A. lamarcki/grahamae, A. undata, H. cucullata, and M. cavernosa, all of which are also common on the Cuban reefs. Unfortunately, Reed et al. (2018) found a shocking 93.6% loss of coral cover since 2004 within the Pulley Ridge Habitat Area of Particular Concern (PR HAPC). Recent taxonomic studies of sponges collected at Pulley Ridge (Reed et al. 2017a), show a rich sponge fauna with several new species; however, there seems to be important differences between Pulley ridge and Cuban mesophotic sponge fauna which must await for comparative biodiversity studies.

Just 60 km east of Pulley Ridge and 225 km west of Key West, Florida, at the terminus of the Florida Keys island chain, are the Tortugas Ecological Reserves (TER) which are part of the 'Sister Sanctuaries' within the FKNMS that have shallow mesophotic reef habitat. TER North protects coral reef banks known as Tortugas Bank (30-50 m depths). TER South includes Riley's Hump (30-50 m) which is an isolated offshore mound that has shallow MCE habitat (Weaver et al., 2006). Scleractinians include M. cavernosa, O. annularis, and Siderastrea siderea. Individual colonies of M. cavernosa are mostly small, ranging from 10-50 cm in diameter but some were observed to 2 m in height. At deeper mesophotic depths, the benthic community is dominated by rope sponges (Aplysina cauliformis) and calcareous algae including Halimeda spp. and Penicillus spp., similar

to Cuba. The southern edge of the Hump is a sharp escarpment from 84-124 m called Miller's Ledge. The benthic assemblages of Miller's Ledge are not MCEs but are dominated by small sponges, bryozoans, small solitary corals, and the corkscrew sea whip, *Stichopathes* sp. This is more similar to what we found deeper in Cuba on the deep island slope of 120->150 m.

Further downstream are shallow shelfedge reefs which extend along most of the Florida Keys and southeastern Florida but mostly only extend to 20-25 m depths. This region of Florida has very limited MCE habitat, primarily scattered patch reefs, ledges, or rubble habitat at depths of 30-100 m. However, the mesophotic depth zone (30-150 m) has not been adequately surveyed in this region. The earliest survey of its mesophotic reefs occurred in 1979 off Key Largo in the upper Florida Keys when Jameson (1981) documented deep coral reef habitat to 39 m and coral algal habitat to 55 m. In 2015, Reed et al. found some deep reefs (125 m) with ROV off Key West, Florida, but these lacked zooxanthellate hard corals and were dominated by the octocoral genera (Diodogorgia, Thesea, Villogorgia, and Nidalia) and sponges. However, it did provide essential fish habitat for several commercially important species including snowy grouper, tilefish and snapper. Based on the presence of paleo-shorelines elsewhere, we could expect to find ledges or hard bottom habitat especially along the 70 and 125 m contours off the Florida Keys and southeastern U.S. (Locker et al., 2010). Some deep-water algal studies have also been conducted along the shelf off southeastern Florida (Hanisak and Blair, 1988; Leichter et al., 2008) which could imply the presence of mesophotic habitat. This paleo-shoreline also forms the basis for the *Oculina* reefs that extend 250 km at depths of 60-90 m off eastern Florida, and then continues as shelf-edge rock ledges from north Florida to North Carolina. The deep *Oculina* coral reefs, however, are entirely formed by azooxanthellate *O. varicosa* (Avent *et al.*, 1977; Reed, 1980; Reed *et al.*, 2007; Harter et al., 2009). The shelf edge from Florida to North Carolina is primarily hard-bottom habitat that consists of extensive ledges and escarpments at mesophotic depths, and have dense populations of sponges, algae and octocorals, but limited Scleractinia, mostly *Oculina varicosa* and *Madracis* sp. (Harter *et al.*, 2015).

DEPTH RECORDS

The maximum depth of zooxanthellate scleractinians in the world is considered to be 165 m in the Pacific, where the plate coral Leptoseris hawaiiensis occurs (Kahng and Maragos, 2006). In the Atlantic, the maximum depth is now 122 m for Agaricia sp. in Cuba. In the Atlantic Ocean and Caribbean, nine genera and at least 18 species of hermatypic scleractinia occur to depths exceeding 70 m (Reed, 1985). In the eastern GOM, the maximum depth range for 22 mesophotic scleractinian species is 31-60 m and 15 species have a maximum depth of 100 m (Jaap, 2015). Reed (1985) had found ten new depth records, including Agaricia grahamae (119 m), Helioseris (syn. Leptoseris) cucullata (108 m), Montastraea cavernosa (113 m), and Madracis sp. (probably M. decactis or M. pharensis f. luciphila, 115 m), all on the deep fore-reef escarpments off San Salvador Island, Bahamas. In comparison, in Cuba (Table 3) we found *Agaricia* sp. to a maximum depth of 122 m, which is now a new depth record in the Tropical Western Atlantic, along with the depth record for Stephanocoenia intersepta (104 m) which was previously 100 m (Reed, 1985). In the Caribbean, Kühlmann (1983) had reported S. intersepta (S. michelini, syn.) to depths of 95 m, and Agaricia spp. to 79 m. The deepest records for algae in the western Atlantic include Halimeda copiosa (130 m) and crustose coralline algae (268 m), also reported from San Salvador by Littler et al. (1985). In comparison, maximum depth of algal observations on our Cuba dives include an unidentified green crust (169 m), crustose coralline algae (169 m), Peyssonneliaceae (139 m), and Halimeda sp. (likely H. copiosa; 127 m).

CORAL DISEASE/ BLEACHING

Regarding coral health of Cuba's mesophotic reefs, of the 2,240 scleractinian colonies counted in this study, only 12 corals (0.53%, mainly Agaricia spp.) showed signs of bleaching. One Agaricia had black band disease, and one had an unidentified white syndrome disease (0.009%). In shallow water, the presence of black band, white band, and plague diseases have been recorded in Cuba (González Ferrer, 2004); white band was found in isolated colonies of Acropora cervicornis, and black band was found with relative frequency in isolated colonies (Alcolado et al., 2000). In general, the Cuban mesophotic corals appeared quite healthy compared to many shallow Caribbean reef sites. In the past decade disease in shallow corals has increased as seawater temperatures increase (Andradi-Brown, 2016) and reports now show mesophotic coral communities are not immune to disease. In the Caribbean U.S. Virgin Islands, disease in mesophotic corals has been documented down to 100 m depths (Smith et al., 2010). The type of disease in the Virgin Islands was placed in the category of 'White Syndromes' due to the characteristic white areas of recent tissue loss, similar to white plague disease (Weil and Hooten, 2008; Andradi-Brown, 2016). White syndromes disease was also reported on mesophotic reefs off southwestern Puerto Rico in Agaricia undata, A. lamarcki, A. grahamae, and A. agaricites (Appledoorn et al., 2016b). Six percent of the coral community showed signs of the disease. Dustan and Lang (2018) have compiled detailed descriptions of mesophotic reefs on the fore-reef slope at Discovery Bay, Jamaica, comparing their conditions from the 1960s with the present. Extensive changes have occurred, especially since the 1980s due to the mass mortality of the herbivorous *Diadema* (to 40 m depths), coral disease, bleaching, loss of other herbivores (from overfishing), reduced water quality, and hurricanes. Although the pinnacles still retain their three-dimensional, "shingled" morphology to ~ 45 m, partial mortality had partitioned many of the original Orbicella colonies into smaller isolates by 2013. This appears somewhat similar to the upper mesophotic zone for some of the Cuban reefs where the slopes are covered with many smaller plate corals rather than the extensive sheets that may have been common in the 1960s-70s. Zlatarski and Estalella (1982) reported colonies of Agaricia (f. unifaciata, likely A. lamarcki) to diameters of 3.5 to 5.6 m on the Cuban reefs in the 1970s.

Few MCEs worldwide have long-term records monitoring changes of temperature over time or shorter-term events such as cold-water upwelling or internal waves sufficient to cause coral impacts. Long-term records at some sites do show bleaching events and there is evidence that some mesophotic corals may have lower bleaching thresholds

than shallow colonies (Baker et al., 2008, Andradi-Brown *et al.*, 2016). In the region of the Gulf of Mexico, a mass bleaching event at the FGBNMS in 2005 impacted 45% of the coral community (Precht et al., 2006; Clark et al., 2014). But the reefs showed resiliency and the bleaching had been reduced to 4% of the population within a year. Since the millennium, coral reefs in the northeastern Caribbean have had many bleaching events including two major bleaching events, in 2005 and 2010 (Smith and Holstein, 2016). In the U.S. Virgin Islands, mesophotic coral cover in some areas dropped from >25% cover (Aronson et al., 1994) to currently less than 10% (García-Sais et al., 2014) which has been attributed to both bleaching events and several large hurricanes (Smith and Holstein, 2016). Temperature records on Puerto Rico's mesophotic reefs show a seasonal range of 26-29.8°C (Appledoorn et al., 2016a). Temperatures recorded by our ROV dives around Cuba in May-June showed a maximum surface temperature of 29.52°C along the south coast, whereas at 50 m depth, in the mesophotic zone, temperatures ranged from 25.26-28.62°C, and a thermocline of 25-28°C was common around 80-100 m depths. Although we do not have temperature records over the year, this upper temperature range would not be sufficient to cause bleaching typically for shallow corals, but the tolerance levels of mesophotic corals are less well understood. Since 1983, when the first bleaching event in Cuba was recorded, there have been 7 major events in Cuba (Jackson et al., 2014). Recent surveys of Cuba's shallow reefs show a loss of reef-building species, such as O. annularis and M. cavernosa (Aguilar Betancourt and González-Sansón, 2007). Average coral cover on the fore reefs (2003-2009) was 13.4% (maximum of 27% at Bahia de Cochinos and

Guajimico) but showed a decline of 1.75% / year which is slightly lower than estimated for the wider Caribbean (2.5-2.7% decline) (Alcolado *et al.*, 2009).

DIRECT HUMAN IMPACTS

In general, most dive sites appeared relatively pristine with little signs of direct human impact. One site at Cayo Largo del Sur MPA (Site 29) showed evidence of nutrient pollution. Landward of the shelfedge reef crest at Cayo Largo were extensive cyanobacterial mats on sediment, and the reef crest (35-40 m) had dense areas of 1-m conical mounds, which appeared to be old dead coral, probably M. cavernosa. Alcolado et al. (2001b) reported high nitrogen nutrients and dead coral at this site attributed from a tourist resort on nearby Cayo Largo where a sewage plant dumped partially treated waste water. On Cuba's shallow reefs, areas exposed to intense fishing and pollution, such as along the northwest coast and especially off Havana show reduced coral density and diversity (Duran et al., 2018). Increased nutrients have been linked to algal and cyanobacterial blooms elsewhere and on reefs in Florida and Belize (Lapointe, 2004; Lapointe et al., 2004; Littler et al., 2006). In addition, lost or discarded fishing gear were relatively uncommon; in total, fishing line or long line were found at eight sites and one lost net was observed. Most of these were found on the substrate on the upper mesophotic reefs but some were found at >100 m depths.

INVASIVE SPECIES

Several invasive species have impacted Western Atlantic MCEs; in particular, various macroalgae such as *Codium* and *Caulerpa*, the gorgonian *Carijoa riisei*,

and Lionfish *Pterois volitans* (Lapointe et al., 2005; Andradi-Brown et al., 2016). In the past decade Lionfish have become a major invasive on most MCE reefs in the Caribbean, Gulf of Mexico and southeastern U.S. (Schofield, 2010; Andradi-Brown et al. 2016). The first live Lionfish reported in northern Gulf of Mexico were documented at Pulley Ridge mesophotic reef west of the Florida Keys during submersible dives in 2010 where six fish were observed (Reed and Rogers, 2011). Since then, exponential increases in the abundance of Lionfish have been observed at Pulley Ridge mesophotic reef (Andradi-Brown et al., 2016; Harter et al., 2017). Thousands of Lionfish also have been recently documented at mesophotic depths in protected area reefs along the southeastern U.S. from Florida to North Carolina at depths of 50->100 m (Harter et al., 2015). Recent genetic analyses indicate western Atlantic Lionfish most commonly cited as *Pterois volitans* are hybrids between P. miles and P. lunulata/russelii (Wilcox et al., 2017); however, all of our identifications were based upon photographic data which precluded morphological or genetic analyses. On the Cuban mesophotic reefs, we observed Lionfish at most sites, but the abundance of Lionfish was generally low (<10 observed per site). Impacts of Lionfish on MCEs could result in declines of prey fish biomass and major shifts in the mesophotic benthic community. In studies of mesophotic reefs in the Bahamas, Lionfish predation on herbivorous fish has caused a shift in the benthic community to an increase of algae relative to corals (Lesser and Slattery, 2011). However, from ROV video surveys at Pulley Ridge, there has been no significant Lionfish effect on the fish assemblages (Harter et al., 2017). The presence of Lionfish at greater depths

makes it difficult to effectively control the species in shallow waters due to the contribution of larvae, juveniles and adults from deeper areas, where they are also difficult to catch. For this reason, there would be an intact reservoir of its populations in the mesophotic reefs which tentatively could populate shallow reefs.

CONCLUSIONS

This Joint Cuba-U.S. Expedition provides for the first time data on the extent and health of mesophotic coral reefs around the entire coast of Cuba, covering nearly 2,778 km. This research cruise is of special significance to understanding the distribution of the deep reefs and the basic oceanographic, biological and ecological processes that take place in MCEs around Cuba and at local and regional levels. Future analyses of the taxonomy and genetics of the specimens, along with quantitative analyses of the video and photo data, will allow a more precise characterization of the diversity and relative abundance of the mesophotic communities of Cuba, as well as a better understanding of the connectivity of Cuban reefs with the Sister Sanctuaries in the U.S. and elsewhere in the Caribbean. Also these data may lead to expanding or adding new Marine Protected Areas in Cuba or more fully protecting some existing MPAs. The full cruise report (Reed et al., 2017b) provides a detailed characterization of each dive site which provides benchmark data for comparisons with future studies and the effects of climate change.

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APPENDICES

May 14 to June 12, 2017. Dive Sites (see Fig. 1) compiled by regions: Northwest coast- Archipelago de Los Colorados, Sites C-04 to C-07; West coast- Banco de San Antonio, Guanahacabibes, Sites C-10 to C-16A; Southwest coast- Isla de la Juventud, Archipelago de los Canarreos, Golfo de Cazones, Bahia de Cochinos, Sites C-21 to C-31A; Southeast Archipelago de Sabana-Camagüey, Sites C-53 to C-60A. coast- Banco Silvertown, Jardines de La Reina, Granma National Park, Chivirico MPA, Punta Imias, Sites C-38 to C-52A; Northeast Coast- Punta Imias (north side), Cabo Lucrecia, Appendix 1. Species list of benthic macrobiota (invertebrates and macroalgae) identified at Cuba's mesophotic reefs from ROV dives during the F.G. Walton Smith Cuba cruise,

Microdictyon marinum (Bory) P.C.Silva, 1955	Halimeda tuna (J.Ellis & Solander) J.V.Lamouroux, 1816	Halimeda sp.	Halimeda opuntia (Linnaeus) J. V. Lamouroux, 1816	Halimeda incrassata (J.Ellis) J.V.Lamouroux, 1816	Halimeda goreaui W.R.Taylor, 1962	Halimeda discoidea Decaisne, 1842	Halimeda copiosa Goreau & E.A.Graham, 1967	Dictyosphaeria cavernosa (Forsskål) Børgesen, 1932	Cladophoropsis sp.	Cladophora sp.	Chlorophyta- unid. spp.	Chaetomorpha sp.	Caulerpa chemnitzia (Esper) J.V. Lamouroux, 1809	Avrainvillea sp.	Anadyomene stellata (Wulfen) C. Agardh, 1823	Chlorophyta	Cyanobacteria	Algae	Phylum/Class/Scientific Name		
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Demospongiae sp. Cu-25 Demospongiae unid. sp.	Demospongiae sp. Cu-24	Demospongiae sp. Cu-23	Demospongiae sp. Cu-22	Demospongiae sp. Cu-21	Demospongiae sp. Cu-20	Demospongiae sp. Cu-19	Demospongiae sp. Cu-18	Demospongiae sp. Cu-17	Demospongiae sp. Cu-16	Demospongiae sp. Cu-15	Demospongiae sp. Cu-14	Demospongiae sp. Cu-13	Demospongiae sp. Cu-12	Demospongiae sp. Cu-11	Demospongiae sp. Cu-10	Demospongiae sp. Cu-09	Demospongiae sp. Cu-08	Demospongiae sp. Cu-07	Demospongiae sp. Cu-06	Demospongiae sp. Cu-05	Demospongiae sp. Cu-04	Demospongiae sp. Cu-03	Demospongiae sp. Cu-02	Demospongiae sp. Cu-01	Cribrochalina vasculum (Lamarck, 1814)	Corallistes sp.	Cliona sp.	Cliona delitrix Pang, 1973	Cliona caribbaea Carter, 1882	Clathria venosa (Alcolado, 1984)	Clathria sp. Cu-04	Phylum/Class/Scientific Name	
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Northeast Coast	Northwest Coast Phylum/Class/Scientific Name C-04 C-04 C-06 C-07		Desmapsamma sp. X	Dictyoceratida sp. Cu-01	Dictyoceratida unid. sp.	Diplastrella megastellata Hechtel, 1965	Diplastrella sp.	Discodermia sp. Cu-01 X	Dragmacidon alvarezae Zea & Pulido, 2016	Dragmacidon cf. alvarezae Zea & X	Dragmacidon sp. Cu-01 X	Dysidea sp.	Ectyoplasia ferox (Duchassaing & Michelotti, 1864)	Erylus cf. formosus Sollas, 1886	Erylus new sp. X	Erylus sp. Cu-01	Geodia cf. cribata Rützler, Piantoni, van Soest & Díaz, 2014			Geodia sp. Cu-01	Geodía sp. Cu-01 Geodía sp. Cu-02	× ×	× ×	× ×	01 02 03 04 04 05 06 06 06 u-01 u-01)1 Nacelet & X	elet & X X
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Iotrochota birotulata (Higgin, 1877)		×	×	×		×		×										×	×	×		×	×						×			×	×	×
Ircinia campana (Lamarck, 1814)	\times			\times							×							×																×
Ircinia cf. strobilina (Lamarck, 1816)																													×					
Ircinia felix (Duchassaing & Michelotti, 1864)	\times			\times		\times				\times		\times				×				×														
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Ircinia strobilina (Lamarck, 1816)	×		×	×		×	×	×	\times	×	×	×			×			×	×		\times	×				×		\times		×	×		\times	×
Leiodermatium sp.		×		×	×						×																							
Lissodendoryx colombiensis Zea & van Soest, 1986					\times				×																									
<i>Melophlus ruber</i> Lehnert & van Soest, 1998																										\times								
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Monanchora arbuscula (Duchassaing & Michelotti, 1864)								×																					×		×			
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Mycale laevis (Carter, 1882)	×		×			×		×		×					×	×	×																	
Mycale laxissima (Duchassaing & Michelotti, 1864)	\times	\times	\times	\times	\times	\times		×	\times	\times	\times	\times	\times	\times	×	× ×			×	×		×				\times						×	×	\times
<i>Mymekioderma gyroderma</i> (Alcolado, 1984)																								\times										
Myrmekioderma sp. Cu-01																											×							
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Neofibularia nolitangere (Duchassaing & Michelotti, 1864)					\times	\times													×	×														
Neopetrosia carbonaria (Lamarck, 1814)																														×				
Neopetrosia cf. dutchi Van Soest, Meesters & Becking, 2014							\times									×																		
Neopetrosia subtriangularis (Duchassaing, 1850)																			×															
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Petrosiidae Cu-01 Petrosiidae Cu-02 Petrosiidae Cu-03 Petrosiidae Cu-04 Petrosiidae Cu-05 Petrosiidae Cu-06 Petrosiidae Cu-07 Petrosiidae Cu-08 Petrosiidae Cu-09 Petrosiidae Cu-10 Petrosiidae Cu-11 Petrosiidae Cu-12	Oceanapia sp. Cu-06 Oceanapia sp. Cu-07 Petrosia weinbergi van Soest, 1980	Oceanapia sp. Cu-03 Oceanapia sp. Cu-04 Oceanapia sp. Cu-05	Oceanapia sp. Cu-01 Oceanapia sp. Cu-02	Oceanapia peltata (Schmidt, 1870)	Niphatidae Cu-01 Oceanapia bartschi (de Laubenfels, 1934)	Niphates sp. Cu-01 Niphates sp. Cu-02	Niphates erecta Duchassaing & Michelotti, 1864	Niphates cf. erecta Duchassaing & Michelotti, 1864 Niphates digitalis (Lamarck,	Niphates arenata Rützler, Piantoni, van Soest & Díaz, 2014	Phylum/Class/Scientific Name	
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	Northwest Coast	West Coast	South	Southwest Coast	South	theast Coast		Northeast Coast	Coast	
Phylum/Class/Scientific Name	C-04 C-04A C-06 C-07 C-1	C-04 C-04A C-06 C-07 C-10 C-10A C-12A C-12B C-15 C-16A C-21 C-21A C-23 C-24 C-28 C-29 C-31A C-33A C-38 C-38A C	16A C-21 C-21A C-23 (C-24 C-28 C-29 C-31A C-33	A C-38 C-38A C-41	-41 C-48 C-50 C-52 C-52A C-53A C-53B C-54 C-54B C-56 C-58 C-58A C-59A C-59C C-60A	2A C-53A C-53B C-54	4 C-54B C-56 C-58	3 C-58A C-59A C-	59C C-60A
Petrosiidae Cu-13	×	-								
Petrosiidae Cu-14										×
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Petrosiidae unid. sp.	× ×	×	× × ×	× ×						
Phakellia folium Schmidt, 1870				×		×	×	× ×	×	×
Phorbas sp.	×									
Polymastia sp. Cu-01	× ×	×	× ×	×		×	×	×	×	×
Polymastia sp. Cu-02	×							×		
Polymastia sp. Cu-03								×		
Polymastia sp. Cu-04	×		×			×	×			
Ptilocaulis walpersi (Duchassaing & Michelotti, 1864)	× × ×	×	×	× ×	× ×	× ×	×	× ×		× ×
Scopalina ruetzleri (Wiedenmayer, 1977)	× × ×	× ×								
Siphonodictyon brevitubulatum Pang, 1973			×		× ×	×				
Siphonodictyon coralliphagum Rützler, 1971	× ×	× ×	× × ×	× × ×	× × ×	× × ×	×	×	×	×
Siphonodictyon sp. Cu-01	× ×	×		×		×				
Smenospongia aurea (Hyatt, 1875)	×		×	× ×	×					
Smenospongia conulosa Pulitzer-Finali, 1986					×	×				
Smenospongia echina (de Laubenfels, 1934)		×		× ×	×	× ×				
Smenospongia sp.			× ×							
Spheciospongia vesparium (Lamarck, 1815)				×						×
Spirastrella coccinea (Duchassaing & Michelotti, 1864)	×	× × ×	× × ×	× × × × ×	× ×	× ×	× ×	× ×	× ×	×
Spirastrella hartmani Boury- Esnault, Klautau, Bézac, Wulff & Solé-Cava, 1999	× × ×	× × ×	× × ×	× × × ×	×	× ×		×	× ×	×

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Phylum/Class/Scientific Name	Nor Nor	thwe	Northwest Coast	oast	C-10	G-104	Northwest Coast West Coast Southwest Coast Southwest Coast Southwest Coast	Coa	St	C-16A		C-914	Sou.	thwe	est C	Southwest Coast	C-31A	C-33A	C-38	Sou	othe G-41	theast Coast	Joas	-	.50 A	-53A	C-53R	C-54	Nord	heas	Northeast Coast	ast	Itheast Coast Northeast Coast	590 0	-60A	
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Spirastrella sp. Cu-02																				×																
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Spongia sp.																×																				
Spongia sp. Cu-01																												×								
Spongia sp. Cu-02																																			×	
Stylissa caribica Lehnert & van Soest, 1998																																	×			
Stylissa sp.																								×		×			×				×			
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Vansoestia caribensis Díaz, Thacker, Redmond, Pérez & Collins, 2015																		×																		
Verongiida Cu-01	×	×	×	×	×	×	\times	×	×	×	×	×	×	\times	×	×	×	×	×	×	×	×	×	×		×		×	×	×		×	\times	×	×	
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Anthozoa- non coral	Stylasteridae	<i>Millepora alcicomis</i> Linnaeus, 1758	Hydroidolina	Hydrozoa	Cnidaria	Plakortis sp. Cu-02	Plakortis sp. Cu-01	Plakortis dariae Ereskovsky, Lavrov & Willenz, 2014	Plakinidae unid. sp.	Plakinastrella onkodes Uliczka, 1929	Oscarellidae unid. sp.	Oscarella sp. Cu-02	Oscarella sp. Cu-01	Homosclerophorida unid. sp.	Homoscleromorpha	Xestospongia sp. Cu-03	Xestospongia sp. Cu-02	Xestospongia sp. Cu-01	Xestospongia purpurea Rützler, Piantoni, van Soest & Díaz, 2014	Xestospongia muta (Schmidt, 1870)	Xestospongia deweerdtae Lehnert & van Soest, 1999	Verongula sp. Cu-02	Verongula sp. Cu-01	Verongula rigida (Esper, 1794)	Verongula reiswigi Alcolado, 1984	Verongula gigantea (Hyatt, 1875)	Verongula cf. rigida (Esper, 1794)	Verongiida Cu-09	Verongiida Cu-08	Phylum/Class/Scientific Name	
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Northwest Coast
Chironephthya caribaea (Deichmann, 1936) Nidalia sp. Alcyonacea - gorgonian Acanthogorgia sp. Bebryce sp. Briareum sp. Callogorgia gracilis (Milne Edwards & Haime, 1857)
Ellisella barbadensis (Duchassaing & Michelotti, 1864) syn. elongata (Pallas, 1766) Ellisella elongata (Pallas, 1766)
Ellisella elongata (Pallas, 1766) Ellisella sp. Eunicea sp. Gorgonia mariae Bayer, 1961
Gorgoniidae Hypnogorgia sp.
Iciligorgia schrammi Duchassaing, 1870 Lignella richardi (Lamouroux, 1816) Muricea sp. Nicella goreaui Baver 1973
Nicella sp. Paramuricea sp. Paramuriceidae Placogorgia sp.

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Phylum/Class/Scientific Name	C-04	C-04 C-04A C-06 C-07	.C-06	0 ast	C-10	C-10A	C-12A	C-12B C	C-15	C-16A	NOTINVEST COAST WEST COAST SOUTHWEST COAST SOUTHWEST COAST C-34 C-04 C-04 C-06 C-07 C-10 C-10A C-12A C-12B C-15 C-16A C-21 C-21A C-23 C-24 C-28 C-29 C-31A C-33A C-38 C-38 C-38 C-38 C-38 C-38 C-38 C-38	C-21A	C-23	C-23 C-24 C-28 C-29 (ST C	29 C	-31A C	-33A	C-38 C	384 (-41 C-48 C-50 C-	02S	52 C-E	2A C-	53A C	-53B	C-54 C	Nort	Normeast Coast 3-548 C-56 C-58 C-58A	31 GO	-58A C	:-59A	THEAST COAST	C-60A
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Pseudopterogorgia (syn. Antillogorgia) bipinnata (Verrill, 1864)																						×													
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Plumapathes pennacea (Pallas, 1766)				×	×			×		×										×	×														
Plumapathes sp. Stichopathes lutkeni Brook, 1889	×												\times											×								×	×		
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Acropora cervicornis (Lamarck, 1816)									\times								\times		\times	\times	\times		×												
Agaricia agaricites (Linnaeus, 1758)			×	×	×	\times		×	×	×	×	\times						\times	×	×	×	\times	×			×	×	×		ì			į		
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Phylum/Class/Scientific Name	C-04 C-04A C-06 C-07 C-10 C-10A C-12A C-12B C-15 C-16A C-21 C-21A C-23 C-24 C-28 C-29 C-31A C-33A C-38 C-38A C-41 C-48 C-50 C-52 C-52A C-53A C-53B C-54 C-54B C-56 C-58 C-58A C-59A C-59C C-60A	1 C-06 C-0	7 C-10	C-10A C-1	C-12A C-12B (C-15 C	-16A C-	.21 C-2	1A C-2	C-23 C-24 C-28 C-29 (C-28	C-29 C	-31A C	-33A C	-38 C.	38A C-41 C-48 C-50 C-	#1 C-48	3 C-50	C-52 C	-52A C	-53A C	-53B C	-54 C-)-54B C-56 C-58 C-58A
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Agaricia lamarcki Milne Edwards & Haime, 1851	×									×						× ×		×		\times				
Agaricia sp.	×	×	\times	×	×	\times	×	×	^ ×	×	×	×	×	×	×	×	×	×	×	×	×	×	×	×
Colpophyllia natans (Houttuyn, 1772)	×		×	×	×		×	× ×	^ ×	×	×	×	×	×		×	×	\times			×			
Dendrogyra cylindrus Ehrenberg, 1834				×																				
Dichocoenia stokesii Milne Edwards & Haime, 1848	×																							
Diploria labyrinthiformis (Linnaeus, 1758)			×										\times		×									
Eusmilia fastigiata (Pallas, 1766)			×	×	×		×								×	×	×	×						×
Helioseris cucullata (Ellis & Solander, 1786)	×		×		×		×	× ×	^ ×	×	×	×	×	×				×						
Isophyllia rigida (Dana, 1848)	×			×																				
Isophyllia sinuosa (Ellis & Solander, 1786)		×	×	×	×				×															
Isophyllia sp.			×																					
Madracis asperula Milne Edwards & Haime, 1849																×	×							
Madracis auretenra Locke, Weil & Coates, 2007		×															×	×						
Madracis decactis (Lyman, 1859)				×							×							×						
Madracis formosa Wells, 1973															×	×	×	×			×	×	×	×
Madracis myriaster (Milne Edwards & Haime, 1850)																		×						
Madracis sp.																^		×	×	×	×	×		×
Madrepora sp.															×	×	\times	\times						
Meandrina meandrites (Linnaeus, 1758)	× ×	×	×	×	×		×	× ×	~ ×	×	×		×	×		× ×	×	×		×			×	×
Meandrina sp.																								×
Montastraea cavemosa (Linnaeus, 1767)	× ×	× ×	×	×	×	×	×	× ×	^ ×	×	×	×	×	×	×	× ×	×	×		×	×	×	×	× ×
Mussa angulosa (Pallas, 1766)		×				×	×							\times										
Mycetophyllia aliciae Wells, 1973																×								
Mycetophyllia lamarkiana Milne															×	×								

Other Annelida Filograna sp. Polychaeta Mollusca Entemnotrochus adansonianus (Crosse & P. Fischer, 1861) Gastropoda Strombus (syn. Lobatus) gigas Linnaeus, 1758	Solenastrea bournoni Milne Edwards & Haime, 1849 Stephanocoenia intersepta (Lamarck, 1836)	Scolymia lacera (Pallas, 1766) Scolymia sp. Siderastrea siderea (Ellis & Solander, 1768)	Scleractinia- unid colonial Scleractinia- unid cup Scolymia cubensis (Milne Edwards & Haime, 1848)	Pontes astreoides Lamarck, 1816 Porties divaricata Le Sueur, 1820 Porties furcata Lamarck, 1816 Porties porties (Pallas, 1766) Porties sp. Pseudodiploria strigosa (Dana, 1846)	Orbicella annularis (Ellis & Solander, 1786) Orbicella faveolata (Ellis & Solander, 1786) Orbicella fanksi (Gregory, 1895) Orbicella franksi (Gregory, 1895) Oxysmilia rotundifolia (Milne Edwards & Haime, 1848)	Phylum/Class/Scientific Name
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Bryozoa									
Bryozoa unid. sp.						× ×	×	×	×
Colatooecia serrulata (Smitt, 1 873)								×	
Arthropoda									
Majidae	×								
Panulirus argus (Latreille, 1804)		×		×					
Penaeidae	×								
Echinodermata									
Asteroidea							×		
Comatulida	×			\times			×		×
Crinoidea						×			
Davidaster discoideus (Carpenter, 1888)									×
Ophiuroidea	×	×				×	×	×	
Chordata - Invertebrate									
Ascidiacea	×		×			× ×	×		×
Non-Fauna									
Human debris									
Human debris- fishing line	×					×		×	×
Human debris- long line	×								
Human debris- net	×								
Disease									
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Bleaching		× × ×	×	×		×	×		
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Appendix 2. Species list of fish identified at Cuba's mesophotic reefs from ROV dives during the F.G. Walton Smith Cuba cruise, May 14 to June 12, 2017. Dive sites (see Fig. 1) compiled by regions (see Appendix 1 caption).

1758) Acanthurus chirurgus (Bloch, 1787)	Abudefduf saxatilis (Linnaeus,	Cuvier, 1029)	Ogcocephalus nasutus	_ophiiformes	Sargocentron vexillarium (Poey, 1860)	Plectrypops retrospinis (Guichenot, 1853)	Neoniphon marianus (Cuvier, 1829)	<i>Myripristis jacobus</i> Cuvier, 1829	Holocentrus sp.	Holocentrus rufus (Walbaum, 1792)	Holocentrus adscensionis (Osbeck, 1765)	Beryciformes	Synodus synodus (Linnaeus, 1758)	Synodus sp.	Aulopiformes	Gymnothorax sp.	<i>Gymnothorax funebris</i> Ranzani, 1839	Anguilliformes	Actinopterygii- Larval	Actinopterygii	Actinopterygii	hordata	Phylum/Class/Order/ Scientific Name	
Major Doctorfish	Sergeant	Datiloii	Shortnose		Dusky Squirrelfish	Cardinal Soldierfish	Longjaw Squirrelfish	Blackbar Soldierfish	Squirrelfish	Longspine Squirrelfish	Squirrelfish		Red Lizardfish	Lizardfish		Moray Eel	Green Moray Eel		Larval Unid Fish	Unid Fish			Common Name	
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Chaetodon sedentarius Poey, 1860	Chaetodon ocellatus Bloch, 1787	Chaetodon capistratus Linnaeus, 1758	Cephalopholis fulva (Linnaeus, 1758)	Cephalopholis cruentata (Lacepède, 1802)	Centropyge argi Woods & Kanazawa, 1951	Centropristis ocyurus (Jordan & Evermann, 1887)	Caranx sp.	Caranx ruber (Bloch, 1793)	Carangoides bartholomaei (Cuvier, 1833)	Carangidae	Calamus sp.	Calamus calamus (Valenciennes, 1830)	Bodianus rufus (Linnaeus, 1758)	Bodianus pulchellus (Poey, 1860)	Apsilus dentatus Guichenot, 1853	Apogon sp.	Apogon pseudomaculatus Longley, 1932	Anisotremus virginicus (Linnaeus, 1758)	(Bloch, 1791)	Acanthurus sp.	Acanthurus coeruleus Bloch & Schneider, 1801	Phylum/Class/Order/ Scientific Name	
Reef Butterflyfish	Spotfin Butterflyfish	Foureye Butterflyfish	Coney	Graysby	Cherubfish	Bank Sea Bass	Jack	Bar Jack	Yellow Jack	Jack	Porgy	Saucereye Porgy	Spanish Hogfish	Spotfin Hogfish	Black Snapper	Cardinalfish	Twospot Cardinalfish	Porkfish	Margate	Surgeonfish	Blue Tang	Common Name	
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Equetus lanceolatus (Linnaeus, 1758)	Epinephelus striatus (Bloch, 1792)	Epinephelus morio (Valenciennes, 1828)	Epinephelus guttatus (Linnaeus, 1758)	Elagatis bipinnulata (Quoy & Gaimard, 1825)	Elacatinus sp.	Elacatinus louisae (Böhlke & Robins, 1968)	Elacatinus horsti (Metzelaar, 1922)	Elacatinus genie (Böhlke & Robins, 1968)	Diplectrum formosum (Linnaeus, 1766)	Coryphopterus personatus (Jordan & Thompson, 1905)	Coryphopterus glaucofraenum Gill, 1863	Clepticus parrae (Bloch & Schneider, 1801)	Chromis scotti Emery, 1968	Chromis multilineata (Guichenot, 1853)	Chromis insolata (Cuvier, 1830)	<i>Chromis enchrysura</i> Jordan & Gilbert, 1882	Chromis cyanea (Poey, 1860)	Chloroscombrus chrysurus (Linnaeus, 1766)	Chaetodontidae	Chaetodon striatus Linnaeus, 1758	Phylum/Class/Order/ Scientific Name	
Jacknife Fish	Nassau Grouper	Red Grouper	Red Hind	Rainbow Runner	Goby	Spotlight Goby	Yellowline Goby	Cleaning Goby	Sand Perch	Masked/ Glass Goby	Bridled Goby	Creole Wrasse	Purple Reeffish	Brown Chromis	Sunshinefish	Yellowtail Reeffish	Blue Chromis	Atlantic Bumper	Butterflyfish	Banded Butterflyfish	Common Name	
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	Phylum/Class/Order/ Scientific Name	Equetus punctatus (Bloch & Schneider, 1801)	Gobiidae	Gobiosoma sp.	Gramma loreto Poey, 1868	<i>Gramma melacara</i> Böhlke & Randall, 1963	Haemulon aurolineatum Cuvier, 1830	<i>Haemulon carbonarium</i> Poey, 1860	Haemulon flavolineatum (Desmarest. 1823)	<i>Haemulon melanurum</i> (Linnaeus, 1758)	Haemulon parra (Desmarest, 1823)	Haemulon plumierii (Lacepède, 1801)	Haemulon sciurus (Shaw, 1803)	Haemulon sp.	Haemulon striatum (Linnaeus, 1758)	Haemulon vittatum (Poey, 1860)	Halichoeres bathyphilus (Beebe & Tee-Van, 1932)	Halichoeres bivittatus (Bloch, 1791)	Halichoeres garnoti (Valenciennes, 1839)	Halichoeres maculipinna (Müller & Troschel, 1848)	Halichoeres pictus (Poey, 1860)	Holacanthus bermudensis Goode, 1876
	Common Name	Spotted Drum	Goby	Goby	Fairy Basslet	Blackcap Basslet	Tomtate	Caesar Grunt	French	Cottonwick	Sailors Choice	White Grunt	Bluestriped Grunt	Grunt	Striped Grunt	Boga	Greenband Wrasse	Slippery Dick	Yellowhead Wrasse	Clown Wrasse	Rainbow Wrasse	Blue Angelfish
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1828)	•	Lutjanus buccanella (Cuvier, 1828)	Lutjanus apodus (Walbaum, 1792)	Lutjanus analis (Cuvier, 1828)	Lucayablennius zingaro (Böhlke, 1957)	Liopropoma sp.	Liopropoma rubre Poey, 1861	Liopropoma mowbrayi Woods & Kanazawa, 1951	Lachnolaimus maximus (Walbaum, 1792)	Labrisomus filamentosus Springer, 1960	Labridae	<i>Kyphosus</i> sp.	Hypoplectrus unicolor (Walbaum, 1792)	Hypoplectrus sp.	Hypoplectrus <i>puella</i> (Cuvier, 1828)	Hypoplectrus nigricans (Poey, 1852)	indigo (Poey,	Hypoplectrus guttavarius (Poey, 1852)	Holacanthus tricolor (Bloch, 1795)	Holacanthus sp.	Holacanthus ciliaris (Linnaeus, 1758)	Phylum/Class/Order/ Scientific Name	
Snapper	Red Snapper	Blackfin Snapper	School- master	Mutton Snapper	Arrow Blenny	Basslet	Peppermint Bass	Cave Basslet	Hogfish	Quillfin Blenny	Wrasse	Chub	Butter Hamlet	Hamlet	Barred Hamlet	Black Hamlet	Indigo Hamlet	Shy Hamlet	Rock Beauty	Angelfish	Queen Angelfish	Common Name	
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Dhvlum/Clace/Order/	Phylum/Class/Order/ Scientific Name	s (Linnaeus,	Lutjanus jocu (Bloch & Schneider, 1801)	ni (Cuvier,	<i>us</i> sp.	anus (Cuvier,	anthus plumieri (Bloch,	Microspathodon chrysurus	Mulloidichthys martinicus	Mycteroperca bonaci (Poey, 1860)	Mycteroperca interstitialis (Poev. 1860)	<i>Mycteroperca phenax</i> Jordan & Swain, 1884	Mycteroperca tigris (Valenciennes, 1833)	<i>Mycteroperca venenosa</i> (Linnaeus, 1758)	<i>Ocyurus chrysurus</i> (Bloch, 1791)	Pagrus sp.	Paranthias furcifer (Valenciennes, 1828)	Pomacanthus arcuatus (Linnaeus, 1758)	Pomacanthus paru (Bloch, 1787)	Pomacentrus sp.	Pristigenys alta (Gill, 1862)	Prognathodes aculeatus (Poey, 1860)
Common	Common Name	Gray Snapper	Dog Snapper	Mahogany Snapper	Snapper	Silk Snapper	Sand Tilefish	Yellowtail Damselfish	Yellow	Black Grouper	Yellowmouth Grouper	Scamp	Tiger Grouper	Yellowfin Grouper	Yellowtail Snapper	Porgy	Atlantic Creolefish	Gray Angelfish	French Angelfish	Damselfish	Short Bigeye	Longsnout Butterflyfish
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Serranus tabacarius (Cuvier, 1829)	Serranus sp.	Serranus phoebe Poey, 1851	Serranus notospilus Longley, 1935	Serranus annularis (Günther, 1880)	Serranidae	Seriola sp.	Scomberomorus sp.	Scomberomorus regalis (Bloch, 1793)	Scarus vetula Bloch & Schneider, 1801	Scarus taeniopterus Lesson, 1829	Scarus iseri (Bloch, 1789)	Scarus coelestinus Valenciennes, 1840	Scaridae	Rypticus saponaceus (Bloch & Schneider, 1801)	Rypticus maculatus Holbrook, 1855	Ptereleotris helenae (Randall, 1968)	Ptereleotris calliura (Jordan & Gilbert, 1882)	Pseudupeneus maculatus (Bloch, 1793)	Pronotogrammus martinicensis (Guichenot, 1868)	Prognathodes guyanensis (Durand, 1960)	<i>ya</i> (Jordan,	Phylum/Class/Order/ Scientific Name	
Tobaccofish	Sea Bass	Tattler	Saddle Bass	Orangeback Bass	Grouper	Amberjack	Mackeral	Cero	Queen Parrotfish	Princess Parrotfish	Striped Parrotfish	Midnight Parrotfish	Parrotfish	Greater Soapfish	Whitespotted Soapfish	Hovering Dartfish	Blue Dartfish	Spotted Goatfish	Rough- tongue Bass	French Butterflyfish	Bank Butterflyfish	Common Name	
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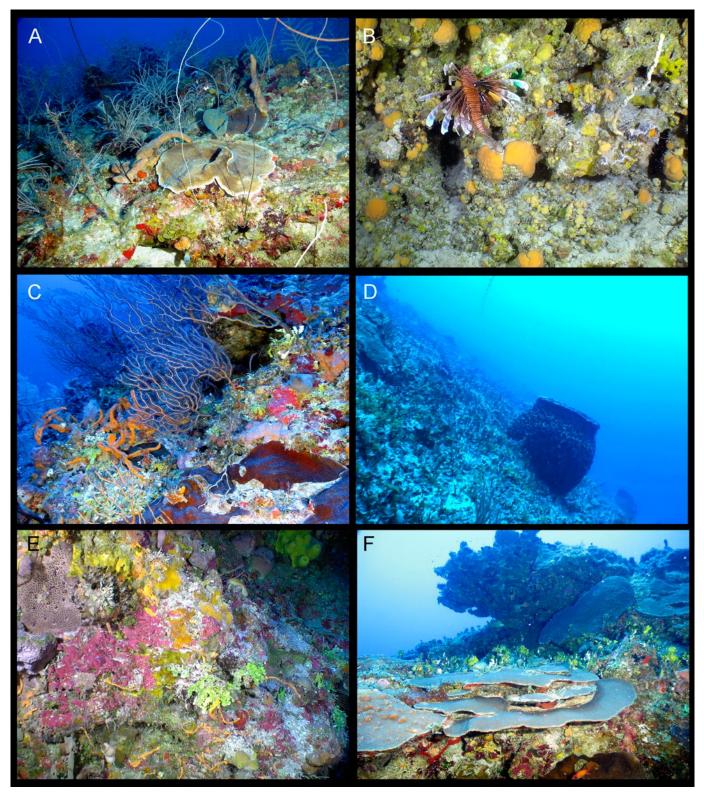
		Nor	thwe	Northwest Coast	oast		\leq	West Coast	Coas	~			S	MT.	Southwest Coast	Coa	ङ			Sot	Southeast Coast	ıst C	oast					Nor	thea	Northeast Coast	oast				
Phylum/Class/Order/ Scientific Name	Common Name	C-04	C-04 C-04A	C-06	C-07	C-10	C-10A	C-12A	C-12B	C-15 C	-16A C	-21 0-	21A C-	23 C-2	4 C-28	3 C-29	C-31A	C-33A	C-38 ()-38A (0-41 0	48 C-5	50 C-5	2 C-52	A C-53	A C-53	3 C-54	C-54B	C-56	C-58 (C-58A (C-10 C-10A C-12A C-12B C-15 C-16A C-21 C-21A C-23 C-24 C-28 C-29 C-31A C-33A C-38 C-38A C-41 C-48 C-50 C-52 C-52A C-53A C-53B C-54 C-54B C-56 C-58 C-58A C-59A C-59A C-59C-60A	C-59C	C-60A	
Seranus tigrinus (Bloch, 1790)	Harlequin Bass					×		\times											\times	\times									\times	\times	\times		×		
Serranus tortugarum Longley, 1935	Chalk Bass		\times	\times														\times				×													
Sparisoma atomarium (Poey, 1861)	Greenblotch Parrotfish																		\times		×	×													
Sparisoma aurofrenatum (Valenciennes, 1840)	Redband Parrotfish							\times	\times				~	× ×				×	\times	\times	×	× ×	×				\times	\times	\times			×			
Sparisoma rubripinne (Valenciennes, 1840)	Yellowtail Parrotfish		\times																																
Sparisoma viride (Bonnaterre, 1788)	Stoplight Parrotfish			×	\times	\times	\times		\times	\times	\times		×	×			\times	×	\times	\times	×	× ×						\times	\times			×	×		
Sphyraena barracuda (Edwards, 1771)	Great Barracuda	\times	\times	\times	\times	×	\times	\times	\times	\times	\times	×	×	× ×	×	\times		\times	\times	\times	×	× ×			\times		\times	\times	\times	×	\times	×	×		
Stegastes adustus (Troschel, 1865)	Dusky Damselfish		\times	\times															\times			×									\times			×	
Stegastes diencaeus (Jordan & Rutter, 1897)	Longfin Damselfish																		\times																
Stegastes leucostictus (Müller & Troschel, 1848)	Beaugregory											\times			\times			\times		\times	×	× ×				\times			\times						
Stegastes partitus (Poey, 1868)	Bicolor Damselfish	\times	\times	\times	\times	×	\times	\times	\times	\times	\times	×	×	× ×	\times	\times	\times	\times	\times	\times	×	× ×		×		\times	\times	\times	\times	\times	\times	×	\times	×	
Stegastes planifrons (Cuvier, 1830)	Threespot Damselfish																				\times					\times									
Thalassoma bifasciatum (Bloch, 1791)	Bluehead Wrasse			\times	\times	\times	\times	\times	\times	\times	\times	\times		×	\times	\times	\times	\times	\times	\times	×	× ×				\times			\times	\times					
Trachinotus blochii (Lacepède, 1801)	Permit											×	×																						
Trachinotus goodei Jordan & Evermann, 1896	Palometa														×																				
Scorpaeniformes																																			
Pterois volitans (Linnaeus, 1758)	Lionfish	×	\times	×	×	×	\times		\times	\times	×	×	×	×	\times	\times	\times	\times	\times		×	× ×	×	×	\times	\times	\times	\times	\times	\times	\times	×	×	×	
Syngnathiformes																																			
Aulostomus maculatus Valenciennes, 1841	Atlantic Trumpetfish																																×		
Tetraodontiformes																																			
Acanthostracion polygonius Poey, 1876	Honeycomb Cowfish		\times																																

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		Northwest Coast	ast		≶	West Coast)oas	_			တ	Southwest Coast	hwe	st C)oas	~			လ	E H	Southeast Coast	င္ပ	ast					8	₩	Northeast Coast	Coa	st			
Phylum/Class/Order/ Scientific Name	Common Name	C-04 C-04A C-06 C	C-07	C-07 C-10 C-10A C-12A C-12B C-15 C-16A C-21 C-21A C-23 C-24 C-28 C-29 C-31A C-33A)-10A ()-12A ()-12B	0-15)-16A	C-21 (-21A	C-23 I	C-24 (C-28 (0-29	C-31A	C-33/	\ C-38	C-38,	۸ C-41	C-48	C-50	C-52	C-52A	\ C-53,	\C-53	3 C-54	1 C-54	В С-5	5 C-58	8 C-58	3A C-5	C-38 C-38A C-41 C-48 C-50 C-52 C-52A C-53A C-53B C-54 C-54B C-56 C-58 C-58A C-59A C-59C C-60A	59C C-	60A
Aluterus scriptus (Osbeck, 1765)	Scrawled Filefish																		×																
Balistes vetula Linnaeus, 1758	Queen Triggerfish		\times	×		\times	\times												\times		\times									\times					
Balistidae	Triggerfish			×	×				×				×					×																	
Cantherhines macrocerus (Hollard, 1853)	Whitespot- ted Filefish			×																															
Canthidermis sufflamen (Mitchill, 1815)	Ocean Triggerfish				\times	\times		\times	\times				\times		\times			\times	\times		\times	\times			\times	\times	\times		\times			×			×
Canthigaster rostrata (Bloch, 1786)	Sharpnose Puffer	×	\times	×	\times	\times	\times		\times	\times	\times				\times	\times	\times	\times		\times		\times	\times	\times					\times	\times	×	×	~ ×		×
Lactophrys triqueter (Linnaeus, 1758)	Smooth Trunkfish			\times																	\times														
<i>Melichthys niger</i> (Bloch, 1786)	Black Durgon			\times	\times	\times		\times	\times									\times	\times			\times							\times			×			
Sphoeroides spengleri (Bloch, 1785)	Bandtail Puffer	×																	\times																
Xanthichthys ringens Linnaeus, 1758)	Sargassum Triggerfish	× ×	\times	×		\times	\times	\times	\times					\times					\times		\times		\times	\times	\times	\times	\times	\times			\times				
Elasmobranchii																																			
Carcharhinidae	Shark			×																															
Carchathinus leucas (Müller & Henle, 1839)	Bull Shark													\times																					
Myliobatiformes																																			
Dasyatis americana Hildebrand & Schroeder, 1928 Orectolobiformes	Southern Stingray	×										\times																							
Ginglymostoma cirratum (Bonnaterre, 1788) Reptilia	Nurse Shark	×	×				\times																												
Testudines																																			
Caretta caretta (Linnaeus, 1758)	Loggerhead Turtle					\times	\times																												
Eretmochelys imbricata (Linnaeus, 1766)	Hawksbill Sea Turtle						×																												

Appendix 3-1. Images from ROV dives to illustrate various regions of Cuba's MCEs.

Western region: A. Site C-04, 49 m, *Agaricia* spp., octocorals (*Antillogorgia* spp., *Ellisella* spp.), black coral- (*Stichopathes* sp.), sponges (tan finger- *Agelas sceptrum*, vase- N*iphates digitalis*); B. Site C-07, 112 m, Lionfish, sclerosponge (*Ceratoporella nicholsoni*); C. Site C-12B, 66 m, *Agaricia* sp, octocoral (*Icilogorgia schrammi*), algae (*Halimeda* sp.), sponge (branching- *Agelas sceptrum*, encrusting tan- *Spirastrella hartmani*, encrusting red- *S. coccinea*); D. Site C-10, 35 m, large *Xestospongia muta*; E. Site C-10A, 104 m, algae (crustose coralline algae, *Halimeda copiosa*), sponges (spikey- *Oceanapia bartschi*, tan with zoanthids- Petrosida, thin rope- *Agelas repens*, encrusting yellow- Calcarea, tan sphere-*Geodia cribata*, yellow tubes- *Verongula rigida*); F. Site C-10A, 38 m, *Orbicella faveolata, Montastraea cavernosa*.



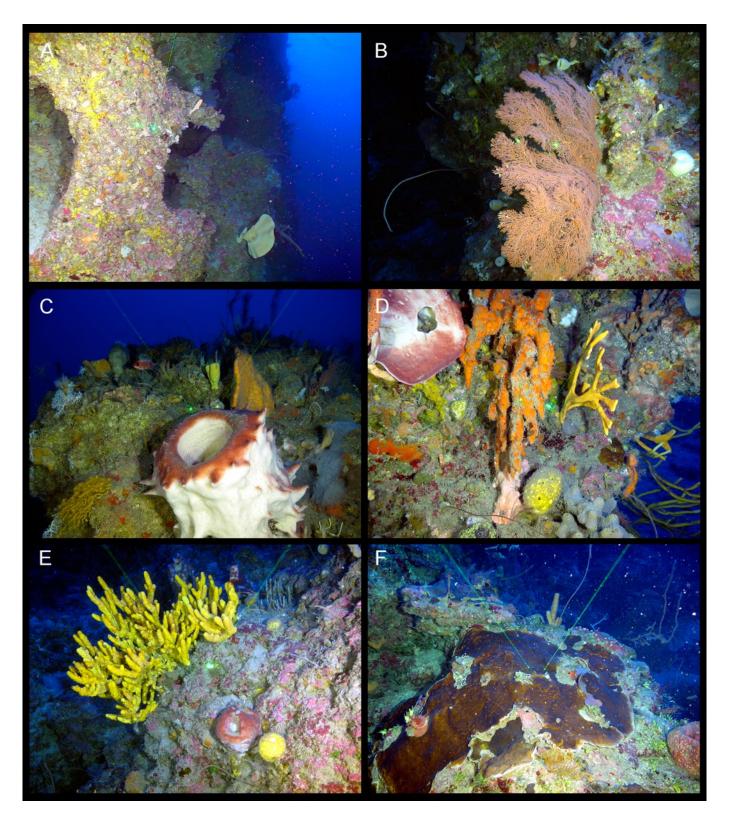
Appendix 3-2. Images from ROV dives to illustrate various regions of Cuba's MCEs.

Southern region: A. Site C-21, 32 m, *Orbicella faveolata*, sponges (orange lumpy-*Ectyoplasia ferox*, large sphere- unid. demosponge), algae (*Halimeda copiosa*), fish (Blackcap Basslet, *Gramma melacara*); B. Site C-24, 73 m, *Agaricia lamarcki*, sponges (purple tube- *Aplysina acheri*, plate- *Agelas dilatata*, sphere- *Geodia* sp.); C. Site C-29, 69 m, *Agaricia lamarcki*, sponge (tube- *Aplysina archeri*); D. Site C-38A, 108 m. sponges (elkhorn- *Agelas cervicomis*, tan fingers- *Agelas sceptrum*, black- *Asteropus* sp., purple lobate- *Aiolochroia crassa*, yellow encrusting- *Siphonodictyon coralliophagum*; E. Site C-41, 70 m, *Agaricia* sp., sponges (tan rope- *A. sceptrum*, red orange finger- *Amphimedon compressa*, orange rope- *Aplysina cauliformis*, *A. fulva*, orange branching- *Agelas cervicomis*, purple lobate- *A. crassa*); F. Site C-41, 142 m, scalloped rock, deep island slope.



Appendix 3-3. Images from ROV dives to illustrate various regions of Cuba's MCEs.

Eastern region: A. Site C-48, depth 96 m, eroded wall, crustose coralline algae, sponges (fan- *Agelas*? sp., yellow encrusting-Verongiida sp.); B. Site C-50, 78 m, octocoral (*Nicella goreaui*); C. Site C-52, 105 m, sponges (vase- *Xestospongia* sp. Cu-01, fan- *Agelas clathrodes*), fish (Longspine Squirrelfish, *Holocentrus rufus*); D. Site C-52, 81 m, sponges (large orange- unid. demosponge, fan- *Cribrochalina vasculum*, yellow- *Agelas cervicornis*, sphere- *Cinachyrella* sp., tan lobate- *Smenospongia conulosa*); E. Site C-50, 92 m, sponges (yellow- undescribed Verongiida, tan vase- *Xestospongia* sp. 1, yellow sphere- *Cinachyrella* sp.); F. Site C-50, 70 m; large *Agaricia lamarcki* (lasers 10 cm), sponge (encrusting- *Spirastrella coccinea*).



Appendix 3-4. Images from ROV dives to illustrate various regions of Cuba's MCEs.

Northern region: A. Site C-56, 67 m, school of Silversides or Anchovies at mouth of cave; B. Site C-58, 44 m, algae (*Halimeda* spp., *Udotea* sp.); C. Site C-58A, 67 m, school of Dog Snapper (*Lutjanus jocu*); D. Site C-59A, 65 m, longline wrapped on wall; E. Site C-56, 88 m, overhang of buttress on wall; F. Site C-60A, 44 m, shelf-edge reef crest with dense octocorals and sponges.

