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2	imbricata) nesting in mangrove estuaries in the eastern Pacific Ocean
3	
4	Short title: Blood values and toxicology of EP hawksbills
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6	Aubrey M. Tauer ¹ , Michael J. Liles ^{2,3,4} , Sofía Chavarría ^{2,3,5} , Melissa Valle ^{2,3,5} , Sada Amaya ⁵ ,
7	Gabriela Quijada ⁵ , Oscar Meléndez ⁵ , Stanley Rodríguez ⁶ , Eric F. Lock ⁷ , Ana V. Henríquez ^{2,3} ,
8	Alexander R. Gaos ^{3,9,10} and Jeffrey A. Seminoff ⁸ .
9	
10	¹ Cūra Earth, Minneapolis, MN, USA;
11	² ProCosta, San Salvador, El Salvador
12	³ Eastern Pacific Hawksbill Initiative, San Diego, CA, USA;
13	⁴ Department of Biological Sciences, University of Texas at El Paso, El Paso, TX, USA
14	⁵ Departmento de Medicina Veterinaria, Universidad de El Salvador, San Salvador, El Salvador;
15	⁶ Centro de Investigación y Desarrollo en Salud, Universidad de El Salvador, San Salvador, El
16	Salvador;
17	⁷ Division of Biostatistics, School of Public Health, University of Minnesota, Minneapolis, MN,
18	USA;
19	⁸ National Oceanic and Atmospheric Administration – National Marine Fisheries Service,
20	Southwest Fisheries Science Center, La Jolla, CA, USA
21	⁹ Biology Department, San Diego State University, San Diego, CA, USA
22	¹⁰ Graduate Group in Ecology, University of California at Davis, Davis, CA, USA
23	
24	

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30 Abstract

31 Sea turtles are a keystone species and are highly sensitive to changes in their environment, 32 making them excellent environmental indicators. In light of environmental and climate changes, 33 species are increasingly threatened by pollution, changes in ocean health, habitat alteration, and 34 plastic ingestion. There may be additional health related threats and understanding these threats is 35 key in directing future management and conservation efforts, particularly for severely reduced 36 sea turtle populations. Hawksbill turtles (*Eretmochelys imbricata*) are critically endangered, with 37 those in the eastern Pacific Ocean (Mexico-Peru) considered one of the most threatened sea turtle 38 populations in the world. This study establishes baseline health parameters in hematology and 39 blood biochemistry as well as tested for heavy metals and persitent organic pollutants in eastern 40 Pacific hawksbills at a primary nesting colony located in a mangrove estuary. Whereas 41 hematology and biochemistry results are consistent with healthy populations of other species of 42 sea turtles, we identified differences in packed cell volume, heterophils and lympohcyte counts, 43 and glucose when comparing our data to other adult hawksbill analysis (1), (2), (3). Our analysis 44 of heavy metal contamination revealed a mean blood level of 0.245 ppm of arsenic, 0.045 ppm of 45 lead, and 0.008 ppm of mercury. Blood levels of persistent organic pollutants were below the 46 laboratory detection limit for all turtles. Our results suggest that differences in the feeding 47 ecology of eastern Pacific hawksbills in mangrove estuaries may make them less likely to 48 accumulate persistent organic pollutants and heavy metals in their blood. These baseline data on 49 blood values in hawksbills nesting within a mangrove estuary in the eastern Pacific offer 50 important guidance for health assessments of the species in the wild and in clinical rehabilitation 51 facilities, and underscore the importance of preventing contamination from point and non-point 52 sources in mangrove estuaries, which represent primary habitat to hawksbills and myriad other 53 marine species in the eastern Pacific Ocean.

55 Introduction

56	Disease can cause declines in wildlife populations, especially those that are already threatened or
57	vulnerable (4) (5) (6) (7). Baseline hematology and biochemistry blood parameters are useful
58	indicators for the assessment of the health status of wild nesting sea turtle populations (8) and are
59	especially helpful in clinical rehabilitation facilities (9). However, reference ranges for
60	hematology and blood biochemistry are not widely available, with many reported values derived
61	from captive animals that may not be representative of wild individuals. Additionally, data from
62	one population of a species often are used as references for other populations, despite potential
63	within-species variation (10) (11) (12) (13).
64	Hawksbill turtles (Eretmochelys imbricata) exemplify a species whose life history may
65	vary widely among populations in distinct ocean basins (14) (15). In the Atlantic and Indo-
66	Pacific, adult hawksbills primarily inhabit coral reef ecosystems (16) (17) (18) and can embark
67	on long-distance (>2,000 km), offshore migrations between nesting and foraging areas (e.g., (19)
68	(20). Hawksbills in the eastern Pacific, however, often associate with mangrove ecosystems (21)
69	(14) (22) (15) and undertake particularly short (<300 km) and neritic (<5 km) post-nesting
70	migrations (23) (24). The marked difference in life history among hawksbills in these ocean
71	basins could greatly influence general health parameters, which are largely unknown for adult
72	hawksbills (3) and which have never been analyzed for individuals inhabititing mangrove
73	estuaries. The availability of reference ranges is paramount for different populations of the same
74	species and even subspecies, as values may even vary amongst a small population depending on
75	diet and ecological variables (25).
76	Hawksbills are critically endangered globally according to the International Union for the
77	Conservation of Nature's (IUCN) Red List (26) and the population in the eastern Pacific is
78	among the most endangered Regional Management Units (27) for sea turtles worldwide (28).

79	Fewer than 700 adult female hawksbills are estimated to remain in the entire eastern Pacific
80	Ocean (29) (15), where >80% of these individuals nest on beaches in mangrove estuaries of El
81	Salvador and Nicaragua (30) (31) (15). These same mangrove ecosystems also provide important
82	developmental habitat for juvenile and sub-adult hawksbills (14) (32). Known threats to this
83	species in the region include incidental capture in coastal fisheries, human consumption of eggs,
84	and alteration of nesting habitat (29) (31). An additional, albeit understudied potential threat to
85	hawksbills inhabiting mangrove estuaries, is contamination by chemicals used in aquacultural
86	and agricultural operations, including persistent pesticide residues from shrimp ponds (33) and
87	toxic compounds dicharged by surrounding rivers (34). These contaminants have been
88	documented as negatively influencing myriad species, including estuarine fish species (35)
89	mollusks (36) and marine turtles (37). If these contaminants are present in mangrove estuaries,
90	reliance on such habitats could have direct impacts on health of hawksbills.
91	In this study, we measured blood biochemistry, hematology, and toxicological parameters
92	in wild adult female hawksbills nesting in the Bahía de Jiquilisco mangrove estuary complex in
93	El Salvador to establish baseline health data for one of the most important hawksbill nesting
94	areas in the eastern Pacific. This information will establish a baseline for these parameters and
95	aid in long-term evaluation of the health status of this severely depleted population and serve to
96	guide future management and conservation efforts, as well as to facilitate comparisons among
97	hawksbill populations in other oceanic basins.
98	

99

100 Materials and Methods

101 Study site

102 Bahía de Jiquilisco (13°13'N, 88°32'W) is located in the Department of Usulután on the south-103 central coast of El Salvador (Fig. 1), and is a National Conservation Area, RAMSAR wetland, 104 and UNESCO Biosphere Reserve. It contains the largest mangrove forest in El Salvador (635 105 km^{2}), and includes numerous islands, channels, and estuaries, with moderate development at 106 some nesting beaches (31). Bahía de Jiquilisco has 42.1 km of hawksbill nesting habitat that 107 includes eight discernable fine grained sand beaches with fragmented second growth coastal 108 forest and fruit tree plantations adjacent to the high water line (15) which host ~40% of hawksbill 109 nesting activity in the eastern Pacific (29) (31) (38).

110

Fig. 1. Locations of hawksbill nesting beaches with patrolled shoreline (black lines) at Bahía de
Jiquilisco, El Salvador, 2013–2014.

113

114 Beach Monitoring and Turtle Measurements

115 Hawksbill nesting occurs primarily between April and October, with a peak in June-July. We 116 conducted beach patrols from 1 April to 15 October 2013-2014 at Bahía de Jiquilisco, where 117 project personnel and an extensive network of >100 trained local egg collectors monitored 118 nesting habitat from 18:00 to 06:00 daily by foot and boat in search of female hawksbills. We 119 identified turtles by Inconel tags (Style 681, National Brand & Tag, Newport, KY, USA) located 120 on the second proximal scale of both front flippers and internal passive integrated transponders 121 (PIT tags; Biomark, Boise, ID, USA) in the right front flipper; Inconel and PIT tags were either 122 present from application during previous tagging seasons or were applied after egg laying was

123 completed (15). For each female hawksbill encountered, we measured curved carapace length 124 (nuchal notch to posterior-most tip of marginal scutes; CCL) and in 2013 we performed a 125 complete visual and physical examination, noting all epi-biota on the turtle and body condition.

126

127 Sample Collection and Analyses

We collected up to 12 ml of blood from the dorsal cervical sinus using a 10 ml syringe and 18 gauge 1.5 inch needle and immediately transferred the sample into a red-top glass serum separator tube and sodium heparin vacutainer tubes. They were not refrigerated prior to processing. Blood smears were made in our field base camp from sodium heparin-treated blood and were fixed with 99% methanol on glass slides and air dried.

133 We initially processed the blood in the field within 6–8 hours of blood collection. Packed 134 cell volumes were performed using a tabletop centrifuge and whole blood in sodium heparin 135 tubes was transferred to 1 ml cryotubes and frozen in liquid nitrogen for heavy metal analysis. 136 The remaining blood was spun for 10 minutes at 2000 RPMs and the serum separated and frozen 137 in cryotubes in liquid nitrogen in the field, which were subsequently stored in -20° C freezers at 138 the University of El Salvador. Samples collected in 2013 were shipped in dry ice to the United 139 States for hematology, serum biochemistry, heavy metal, and toxicology analyses, whereas in 140 2014, plasma biochemistry analyses were conducted at Centro Scan (San Salvador, El Salvador). 141 The results were pooled for determining biochemistry reference ranges after determining that 142 there was no statistical difference between the two sample sets.

For hematology, blood films were stained at the Minnesota Zoo with DipQuick stain (Jorgenson Laboratories, Loveland, CO, USA) for manual differential accounts of circulating white blood cells and for hemo-parasite identification. Total white blood cell counts were estimated. Samples for serum biochemistry were shipped on dry ice for processeing at Marshfield

147 Laboratories (Marshfield, WI, USA). The biochemical panel included alanine aminotransferase 148 (ALT), aspartate aminotransferase (AST), alkaline phosphatase, cholesterol, CO2, creatine kinase 149 (CK), glucose, lactate dehydrogenase (LDH), calcium, phosphorous, potassium, sodium, 150 chloride, bicarbonate, total protein, anion gap and uric acid (UA). Reference intervals for 151 biochemistry and hematology variables were computed using the package referenceIntervals for 152 R (39) Parametric 95% reference intervals were computed, and the one-sample Kolmogorov-153 Smirnoff test (40) was used to assess the distributional assumption. For variables with a 154 Kolmogorov-Smirnoff p-value less than 0.01, a non-parametric 95% reference interval was 155 determined instead, with endpoints given by the 0.025 and 0.975 sample quantiles of the 156 observed data.

157 Blood samples were screened at the California Animal Health and Food Safety 158 Laboratory (San Bernadino, CA, USA) for heavy metals (arsenic [detection limit = 0.010 ppm], 159 lead [0.050], and mercury [0.010]) and persistent organic pollutants (POP), including 160 organochlorine insecticides (aldrin [0.010], alpha-BHC [0.010], gamma-chlordane [0.010], 161 technical chlordane [0.050], pp-DDE [0.020], pp-DDD [0.020], pp-DDT [0.020], dicofol [0.020], 162 op-DDE [0.020], op-DDD [0.020], op-DDT [0.020], dieldrin [0.010], endosulfan I [0.010], endosulfan II [0.010], endrin [0.010], HCB [0.010], heptachlor [0.010], heptachlor epoxide 163 164 [0.010], lindane [0.010], methoxychlor [0.010]), mirex [0.010], toxaphene [0.400] and 165 polychlorinated biphenyl (Arochlor 1221, 1232, 1242, 1248, 1254, 1260, 1262 [0.200 and 166 0.400]). Mean toxicity levels were determined, with 95% confidence intervals, for arsenic, lead, 167 and mercury. If observations were missing below the limit of detection, the mean and standard 168 deviation were inferred via maximum likelihood under the assumption that the data have a log-169 normal distribution that is left-censored below the limit of detection using the package censeReg 170 for R (41). All data and analysis is publicly available as an annotated reproducible R code file at

and

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- 173
- 174 Results

175 We encountered and examined 66 nesting hawksbills at Bahía de Jiquilisco in 2013–2014, which 176 had a mean carapace length of 84.9 cm (SD 5.8, range = 71.0-96.6) and appeared in good general 177 health. Physical exam findings in 2013 included two turtles that were covered in approximately 178 5% of epi-biotic growth; nearly all other individuals were less than 1%. Six individuals exhibited 179 carapace damage, including missing scutes, although all appeared to have healed from the 180 injuries. One individual had a fairly large deformity of her distal carapace, but was mobile, in 181 good body condition, and did not have difficulty depositing eggs. Additionally, one individual 182 had a small tumor on the right rear flipper, but logistical limitations prevented biopsy collection. 183 Hematologic values are presented in Table 1. No hemo-parasites were observed for the 28 184 hawksbills evaluated in 2013. Table 2 provides the serum biochemistry reference ranges for 185 blood collected in plain serum separator tubes in 2013 and blood plasma from sodium heparin 186 tubes in 2014, including liver enzymes, total protein, electroyltes, and uric acid.

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199

200 Table 1. Hematology reference intervals for wild hawksbill turtles (Eretmochelys imbricata)

201 nesting at Bahía de Jiquilisco, El Salvador, 2013 (n = 28).

Parameter ^a	95% referen	KS p-value ^b	
	Low	High	
PCV (%)	23.0	33.6	0.7130
WBC (× $10^3/\mu$ L)	1228.2	10,871.8	0.0528
Heterophils (%)	43.4	93.6	0.3306
Heterophils ($\times 10^3/\mu L$)	0	8549.7	0.2120
Lymphocytes (%)	41.8	42.6	0.8327
Lymphocytes (× $10^{3}/\mu$ L)	87.5	2517.7	0.8282
Monocytes (%)	0.3	11.8	0.1358
Monocytes (× $10^3/\mu$ L)	0	748.3	0.4461
Basophils (%)	0	7.1	0.5905
Basophils ($\times 10^3/\mu L$)	0	398.4	0.8808
Eosinophils (%) ^c	0	5	< 0.0001
Eosinophils (× $10^3/\mu$ L) ^c	0	290	< 0.0001

^aPCV, packed cell volume; WBC, white blood cells.

^bKS p-value, Kolmogorov-Smirnoff p-value.

²⁰⁵ ^cIndicates non-parametric 95% reference interval.

229 Table 2. Serum chemistry reference intervals for wild hawksbill turtles (*Eretmochelys imbricata*)

nesting at Bahía de Jiquilisco, El Salvador, 2013-2014 (n = 66).

231

Parameter ^a	n	95% refere	nce interval	KS p-value ^b
		Low	High	
Glucose (mg/dL)	51	57.5	142.0	0.1771
AST (U/L)	51	18.3	74.3	0.8810
ALT (U/L)	33	16.3	78.1	0.8429
ALP (U/L)	51	25.7	93.6	0.4668
$CK (U/L)^{c}$	51	121	1296.2	< 0.0001
LDH (U/L) ^c	51	135.6	1645.7	0.0009
Cholesterol (mg/dL)	51	107.1	366.8	0.8260
TP (g/dL)	51	2.6	5.0	0.6850
Phosphorus (mg/dL)	18	3.4	16.1	0.6876
Calcium (mg/dL)	36	1.0	21.2	0.0379
Sodium (mmol/L)	51	140.7	169.4	0.8217
Potassium (mmol/L)	18	3.7	5.7	0.9910
Chloride (mmol/L)	51	96.9	148.5	0.1106
Bicarbonate (mmol/L)	18	9.8	33.5	0.9538
Uric Acid (mg/dL)	33	1.0	1.8	0.1846
Anion Gap (mmol/L)	18	7.6	36.8	0.2089

^aAST, aspartate aminotransferase; ALT, alanine aminotransferase; ALP, alkaline phosphatase;

233 CK, creatine kinase; LDH, lactate dehydrogenase; TP, total protein.

^bKS p-value, Kolmogorov-Smirnoff p-value.

²³⁵ ^cIndicates non-parametric 95% reference interval.

236

237 Levels of arsenic, lead, and mercury are presented in Table 3. Arsenic had the highest level, with 238 a mean of 0.245 ppm (95% confidence interval = (0.10, 0.39)). Arsenic was detectable in all of 239 the samples collected (n = 28). Lead and mercury had lower mean levels of 0.045 (95%) 240 confidence interval = (0.038, 0.056)) and 0.008 (95% confidence interval = (0.004, 0.017)) 241 respectively. Samples from all turtles tested for POPs (n = 28) were below the detectable limits. 242 243 244 245 246 247 248 249

250 Table 3. Heavy metal blood values for wild hawksbill turtles (*Eretmochelys imbricata*) nesting at

251 Bahía de Jiquilisco, El Salvador, 2013 (n = 28).

252

Parameter	Mean	95%	6 CI
		Low	High
Arsenic	0.245	0.100	0.390
Lead	0.045	0.038	0.056
Mercury	0.008	0.004	0.017

253

254

255 **Discussion**

256 Our results provide the first assessment of hematology, biochemistry, heavy metal, and persistent 257 organic pollutant levels in the blood of wild hawksbills nesting in mangrove estuaries and 258 establish baseline values for mature female hawksbills in these habitats in the eastern Pacific 259 Ocean. The population sampled in this study was rated overall as healthy, as nesting hawksbills 260 were in good body condition, had minimal epibiota, and generally had normal physical exam 261 findings. While interpreting the parameters in this study, it is important to note that it is common 262 that highly contaminated reptiles show no acute signs of health distress, thus our results should 263 not be misinterpreted as confirming the species is healthy in the region.

264

265 Hematology and Biochemistry

The hematological and biochemistry results are generally comparable to those of other species of sea turtles sampled with healthy populations (42) (43) (44) (45) including hawksbill nesting females at open-coast beaches in Brazil (1) and for hawksbill foraging aggregations at coral reefs in the eastern Pacific (Table 4). Some differences are notable in comparing values between studies, for example glucose in the (3) study is significantly higher than that of all other studies and the Packed Cell Volume is lower in our study than in (1). Notably only eight individuals were sampled in the (3) study and the sea turtles were caught in the open water and brought on to

the beach instead of testing nesting females. The authors speculate that handling stress induced a stress hyperglycemia. This study also is the only other adult wild hawksbill study to include a white blood cell differential count, which varies from ours in numbers of heterophils and lymphocytes. The other two studies, (2) and (1) have similar values for biochemistries to our study.

278 One female in our study had white blood cell count and heterophil count twice as high as 279 the lowest WBC and heterophil count sampled, so occult illness in one or more individuals of our 280 studied population may be possible (46). Biochemistry reference ranges were established (2) for 281 juvenile hawksbills occupying a coral reef ecosystem off the Pacific coast of Colombia, with 282 calcium, total protein, phosphorus, glucose values similar to our data, but with much wider 283 ranges of LDH, AST, and cholesterol. Some differences were noted between several 284 hematological and biochemistry values when compared to published data from juvenile 285 hawksbills undergoing rehabilitation in the United Arab Emirates (47). For example, juvenile 286 hawksbills had lower mean PCVs, lower total white blood cell counts, and higher AST, CK, and 287 uric acid levels. Additionally, mean calcium, phosphorus, and total protein levels were lower in 288 the rehabilitated animals when compared to our study sample. These differences may be due, at 289 least in part, to the impaired health of animals in rehabilitation, as well as possible geographic 290 variation in environmental variables or in the life-history characteristics of hawksbills in distinct 291 oceanic regions. Variation in biochemistry ranges may reflect differences in physiological 292 requirements between life stages (i.e. juvenile vs. adults) and/or behavior/habitats (nesting in 293 mangrove estuaries vs foraging at coral reef ecosystems) of each studied hawksbill population 294 (15).

	Wrobel	Goldberg	et al. 2013	Tobón-Ló	pez & Amoroc	ho Llanos 2014	Muñoz-l	Pérez et al	. 2017	Tauer et	al. this stu	dy
Parameter ^a	Mean	SD	n ^b	Mean	SD	n ^c	Mean	SD	\mathbf{n}^{d}	Mean	SD	n ^e
Hematology												
PCV (%)	39.4	2.9	41	_	_	_	-	_	-	28.34	7.71	28
RBC (× 10^{12} /L)	_	_	_	_	_	_	0.35	0.09	8	_	-	_
WBC ($\times 10^{9}/L$)	_	_	_	_	_	_	5.31	3.86	8	_	-	_
Heterophils (%)	_	_	_	_	_	_	32.3	6.9	8	65.5	12.81	28
Lymphocytes (%)	_	_	_	_	_	_	45.9	6.1	8	22.21	10.4	28
Monocytes (%)	_	_	_	_	_	_	3.6	1.9	8	6.04	2.92	28
Basophils (%)	_	_	_	_	_	_	0.1	0.2	8	2.82	2.20	28
Eosinophils (%)	-	_	-	-	-	-	18.5	4.5	8	0.36	0.99	28
Biochemistry												
Glucose (mg/dL)	98.6	14.6	41	103.5	16.6	11	1567.6	180.2	7	99.71	21.56	51
AST (U/L)	55.4	7.1	41	132.6	111.2	11	196	54	8	46.33	14.29	51
ALT (U/L)	6.6	2.4	41	_	_	_	38	15	8	47.18	15.75	33
ALP (U/L)	15.9	3.7	41	_	_	_	53	26	8	56.63	17.33	51
LDH (U/L)	_	_	-	136.5	78.9	11	-	_	-	394.41	288.12	51
Cholesterol (mg/dL)	287	42	41	84.5	30.9	11	_	-	_	236.92	66.24	51
TP (g/dL)	5.45	0.63	41	2.5	0.7	11	4.8	0.7	8	3.79	0.62	51
Phosphorus (mg/dL)	11.3	1.4	41	6.7	2	11	_	-	_	9.76	3.23	18
Calcium (mg/dL)	11.6	1.5	41	7.8	1.8	11	-	_	_	11.09	5.14	36
Sodium (mmol/L)	139.6	3.5	41	_	_	_	157	2	7	155.04	7.33	51
Potassium (mmol/L)	5.09	0.76	41	_	_	_	4.2	0.4	7	4.68	0.51	18
Chloride (mmol/L)	_	_	-	_	_	_	-	_	-	122.71	13.14	51
Biocarbonate (mmol/L)	-	-	_	_	_	_	-	_	_	21.61	6.05	18
Uric Acid (mg/dL)	0.95	0.17	41	3.7	2.7	11	_	_	_	1.39	0.20	33
Anion Gap (mmol/L)	_	_	_	_	_	_	_	_	_	22.22	7.45	18

296 Table 4. Available blood values for hawksbill turtles.

²⁹⁷ ^aPCV, packed cell volume; RBC, red blood cells; WBC, white blood cells; AST, aspartate aminotransferase; ALT, alanine aminotransferase; ALP, alkaline

phosphatase; CK, creatine kinase; LDH, lactate dehydrogenase; TP, total protein.

²⁹⁹ ^bHawksbills nesting on open-coast beaches in Brazil, Atlantic Ocean.

300 ^cHawksbills foraging at coral reefs in Colombia, Pacific Ocean.

301 ^dHawksbills foraging at coral reefs in Ecuador, Pacific Ocean.

302 ^eHawksbills nesting on beaches within mangrove estuaries in El Salvador, Pacific Ocean.

303

304 Heavy Metals

305 Heavy metal values appear variable amongst species and subpopultations in loggerhead (*Caretta* 306 caretta), kemp's ridley (Lepidochelys kempii), and green (Chelonia mydas) turtles (48) (49) (50), 307 and are likely related to environmental effects, diet, age, and geography. While hawksbills are 308 omnivorous, their diet worldwide is primarily composed of sponges (51), which are of low 309 trophic level and may explain lower levels of contaminants than sea turtle species that eat items 310 higher up on the food chain, such as olive ridley (Lepidochelys olivacea) and kemp's ridley 311 turtles (48). Adult hawksbills have been documented having relatively low concentrations of the 312 heavy metals in their blood, although maternal transfer of heavy metals from adult hawksbills to 313 their eggs is known to occur (43).

314 Higher levels of arsenic were found in adult hawksbill tissues in Japan, particularly in 315 muscle, than compared to adult green turtles (52). Additionally, arsenic levels of marine sponges 316 were found to range from 0.8–157 mgm/gram of dry weight, suggesting that sponges may be a 317 significant source of arsenic in adult hawksbills. It is unclear the role that sponges may play in 318 accumulation of other heavy metals or persistent organic pollutants, such as the low levels of lead 319 and mercury found in our study population. Importantly, hawksbills in our study area utilize 320 mangrove estuaries and are believed to feed predominantly on mangrove seeds and roots (M. 321 Liles, pers. obs.), indicating that they may feed at an even lower trophic level than populations of 322 hawksbills in other regions. The tendency to feed at low trophic levels may enable eastern Pacific 323 hawksbills to avoid higher levels of blood pollutants seen in conspecifics in other habitats, as 324 well as other sea turtle species.

326 Persistent Organic Pollutants

Organic and inorganic pollutants have been more frequently studied in loggerheads than other sea turtle species (53) (54) (55). Studies on loggerheads have found detectable POP and PCB results in which several of the individual contaminants had correlations with changes in clinical parameters such as packed cell volume (56). Further studies are needed on all sea turtle species to determine the individual and population level effects on health and reproductive outcomes in animals exposed to inorganic and organic pollutants.

333 The trophic level of food items consumed by sea turtle species at different life stages may 334 impact levels of POP and PCBs. For instance, green turtles consume marine invertebrates as 335 juveniles before transitioning to primarily algae and sea grass as adults, whereas adult 336 leatherback and hawksbills forage on jellyfish and primarily marine sponges, respectively (57). 337 For hawksbills and leatherbacks, this may mean they tend to accumulate more polluntants. More 338 recently, however, leatherback turtles (Dermochelys coriacea) in Gabon with evaluated levels of 339 POP and PCB in the blood of nesting and all turtles had levels below the detectable limit (42), a 340 recent study (58) noff the west coast of Senegal in the Cape Verde Islands comparing POP and 341 PCB levels in juvenile green and hawksbill turtles found detectable levels in both species, 342 although green turtles had both higher levels and a greater prevalence of contamination. Trophic 343 levels might not reflect higher levels of POP in adult green and hawksbill turtles and viable turtle 344 eggs (59).

345

346 Conclusions

347 Our study provides baseline health data for hawksbills nesting at a primary rookery located in a 348 mangrove estuary in the eastern Pacific Ocean, which can provide a starting point for long-term 349 monitoring of health status of hawksbills in the region and offer diagnostic indications for 350 treatment of individuals in clinical rehabilitation. Additional studies between healthy juvenile and 351 adult hawksbills in both mangrove estuaries and other habitats should be conducted to delineate 352 size or age related differences in biochemistry and hematologic values in this species, as apparent 353 health status may not reflect contaminant loads. We suggest that future research determine 354 contaminant loads of marine sponges and mangrove vegetation in the Bahía de Jiquilisco and the 355 potential role they play in accumulation of toxins in the environment. It is possible ecosystem 356 processes are occurring that prevent uptake of toxins in the environment to the sea turtles 357 themselves, or through their diet, which, contrary to most hawksbill populations, includes 358 substantially more vegetation (60). Further studies at Bahia de Jiquilisco utilizing skin, muscle, 359 carapace, fat and liver may provide different results than those obtained in this study.

360

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369 Fisheries Science Center (NMFS-NOAA) in United States (sample import: CITES

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