

**Article** 

Not peer-reviewed version

# Spatial Ecology of the Population of Reef Manta Rays (Mobula Alfredi) In New Caledonia Using Satellite Telemetry 1 -Horizontal Behaviour

Hugo Lassauce\*, Olivier Chateau, Laurent Wantiez

Posted Date: 26 May 2023

doi: 10.20944/preprints202305.1879.v1

Keywords: Tagging; Home Range; Habitat Use; Movement Patterns



Preprints.org is a free multidiscipline platform providing preprint service that is dedicated to making early versions of research outputs permanently available and citable. Preprints posted at Preprints.org appear in Web of Science, Crossref, Google Scholar, Scilit, Europe PMC.

Copyright: This is an open access article distributed under the Creative Commons Attribution License which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Disclaimer/Publisher's Note: The statements, opinions, and data contained in all publications are solely those of the individual author(s) and contributor(s) and not of MDPI and/or the editor(s). MDPI and/or the editor(s) disclaim responsibility for any injury to people or property resulting from any ideas, methods, instructions, or products referred to in the content.

Article

# Spatial Ecology of the Population of Reef Manta Rays (Mobula Alfredi) In New Caledonia Using Satellite Telemetry 1 - Horizontal Behaviour

Hugo Lassauce 1,2,3\*, Olivier Chateau 4 and Laurent Wantiez 1,5

- 1 UMR ENTROPIE-IRD-UR-UNC-CNRS-IFREMER, Noumea, New Caledonia
- <sup>2</sup> Conservation International New-Caledonia, Noumea, New Caledonia
- <sup>3</sup> The Manta Trust, Corscombe, Dorchester, United Kingdom
- Laboratory of Marine Biology and Ecology, Aquarium des Lagons, Noumea, New Caledonia
- <sup>5</sup> University of New Caledonia, Noumea, New Caledonia

**Abstract:** The reef manta ray (*Mobula alfredi*) is a highly mobile and plankton-feeding species, classified vulnerable to extinction on the IUCN Red List for Threatened Species. Knowledge on their spatial ecology and the extent of their dispersal remain incomplete, especially within island-fragmented habitats as found in New Caledonia. Satellite telemetry was used to investigate the horizontal movement ecology of reef manta rays in New Caledonia. A total of 21 manta rays were tagged with pop-up satellite archival transmitting tags (21 Fastloc and 2 MiniPAT) that remained deployed for a duration ranging from 3 to 180 days (mean  $\pm$  SE = 76.7  $\pm$  50.3). Rays presented a strong site fidelity and an important affinity for coastal waters. Long-distance migrations (> 300 km) were also observed, mainly through coastal and shallow water paths. Horizontal movements were compared to a home range area and classified into four distinct patterns: *Fidelity, Excursion, Fidelity + Relocation* and *Relocation*. The most dominant pattern was *Fidelity* where manta rays remained within their home range for the whole duration of the tag deployment. Our findings may assist the design of more appropriate management strategies for the species in New Caledonia and other regions worldwide.

Keywords: Tagging; Home Range; Habitat Use; Movement Patterns

## 1. Introduction

Understanding the processes that influence a species distribution and its dynamics is particularly relevant for conservation [1]. This information may assist in mapping biodiversity and ecosystem services, identifying effects and potential threats of environmental changes and anthropogenic activities, enabling effective prioritization of areas for biodiversity conservation. It also provides crucial knowledge to develop tools and models used in conservation [1,2]. To obtain a comprehensive view of these processes, it is mandatory to understand the life history traits and the biology of the species as well as the environmental factors to which the species may respond. Marine species that are capable of large-scale movements have a high dispersal potential especially in continuous environment like open ocean where physical barriers are not obvious [3,4]. Yet, several studies showed high residency and site fidelity patterns in marine species where movements were influenced by environmental barriers and biological factors such as mating success, access to breeding grounds or access to consistent food resources. For instance, reef fishes show limited connectivity between reefs separated by large sand channels [5,6], killer whales display intrinsic isolation of communities due to different food resources [7], bottlenose dolphins demonstrate restricted home range linked with productive habitats [8] and oceanic whitetip sharks record high site fidelity driven by the consistent availability of prey [9]. Food resource is also a major factor that influence the movement of large-bodied filter-feeding species [10-12]. Yet, this resource depends on environmental factors resulting in different distribution patterns between regions of the world. Consequently, spatial distribution and dynamics of these highly mobile filter-feeding species may be different from one region to the next.

Reef manta rays (Mobula alfredi, (Krefft, 1868)) are filter-feeders found in tropical waters around the word. Populations are observed near coastal reefs and their movements have been documented to be driven by the availability of food resources. For instance, in the Maldives, seasonal peaks in productivity gather hundreds of individuals each year [13]. In Indonesia, reef manta rays perform long distance migration triggered by monsoon shifts and associated reduction in productivity [14]. In Australia, highly productive eddy events are likely to trigger offshore movements of individuals [15] and seasonal variations in temperature appears to initiate latitudinal migration over hundreds of kilometres along the east coast of Australia [16,17]. In contrast, when the resource is consistent throughout the year, reef manta rays seem to demonstrate strong residency patterns with only few connections between geographically close populations. In Hawaii, no connection was found over 10 years of photo-identification monitoring between two aggregation sites located only 150 km apart. Similarly, in Indonesia, acoustic telemetry suggested spatial segregation between populations that are only 150 km apart [18]. These evidences suggest that the spatial ecology of reef manta rays can be difficult to predict and that localised investigation might be necessary to obtain comprehensive understanding of their movements. This is a crucial task as the species is heavily exploited in many regions of the world for their gill plates that are used in Asian medicinal trade [19]. In addition to this fishing pressure, reef manta rays exhibit several conservative life history traits that exacerbate this vulnerability including small population size, low fecundity and fragmented distribution. Their strong affinity for coastal shallow waters increases their exposition to human activities, such as bycatch fisheries [20], uncontrolled mass tourism [21,22], habitat degradation [23-26], boat strikes [27] and fish nets entanglement [28]. Consequently, reef manta rays are classified vulnerable to extinction on the IUCN Red List of Threatened Species [29]. Providing robust information on the movement patterns and habitat use of reef manta rays would contribute to the design of effective management and conservation measures that are essential to protect the species.

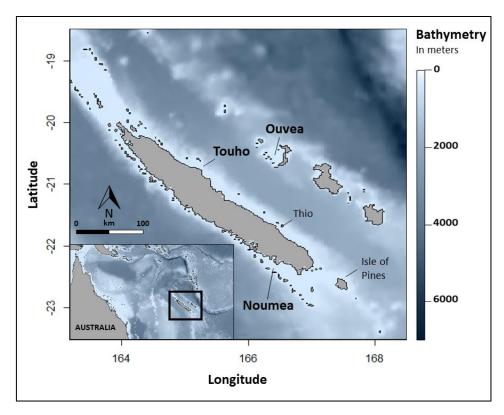
Among methods that enable spatial analysis, satellite telemetry is now a common practice with an increasing number of studies deploying tags to track movements of a wide range of species including terrestrial species, marine birds, sea turtles, marine mammals, and elasmobranchs [2,30-37]. This tool allows researchers to investigate broad-scale and fine scale movements, diving behaviour as well as preferred depth and temperature, and sometimes revealing misconceptions or unexpected findings about the ecology of a species [38-40]. Satellite tracking devices are effective tools to detect unsuspected large migrations in elasmobranchs such as the basking shark [41], the six gills shark [42] and the white shark [43], for example.

In New Caledonia, reef manta rays have not been studied until recently [46,47,48] and basic information on the population of this emblematic and threatened species are needed. Previous findings using four years of photo-identification monitoring revealed that the population of reef manta rays was distributed in all parts of the archipelago and exhibited high long-term fidelity rates at these sites. Connectivity was also limited but possible between all sites showing high dispersal potential [47]. In addition, genomic analyses found genetic differentiation between aggregation sites, which confirms the importance of site fidelity and exacerbated the lack of connectivity between geographically close habitats [48].

On this basis, we use satellite tracking to investigate the spatial ecology of *M. alfredi* in the fragmented environment of the New Caledonia archipelago. Our objectives are to assess the fine scale habitat use and potential broad scale migration patterns by addressing the following points: quantify the use of offshore versus coastal waters, show the extend and limitation of the movements from aggregation sites with high fidelity rates, define the frequency and magnitude of the connectivity between studied sites, detect potential broad scale movements and the existence of unknown key habitats.

## 2.1. Study Sites

The archipelago of New Caledonia consists in a Main Island surrounded by a barrier reef of 1,660 km that enclose shallow waters of a 16,874 km² lagoon [49]. This barrier reef marks the limits of the continental shelf where the bathymetry drops to depths greater than 2,000 m. Smaller islands with a relatively narrow continental shelf, named the Loyalty Islands, are located off the east coast of the Main Island separated by a 2,000-meter-deep channel (Fig. 1).



**Figure 1.** Locations of the deployment of pop-up satellite tags (SPLASH10 and MiniPAT) on reef manta rays ( $Mobula\ alfredi$ ) in New Caledonia, South Pacific. Study sites (in bold): Noumea (N=6), Ouvea (N=4) and Touho (N=6).

Reef Manta rays were tagged at two locations off the Main Island: Noumea in the southern part of the west coast and Touho in the northern part of the east coast. The other tagging site was off the northern island of the Loyalty Islands, Ouvea. In Noumea, tagging operations took place at two aggregation sites (24 km apart) on the barrier reef: the Boulari channel (22°29′S, 166°26′E) and the Dumbea channel (22°21′S, 166°15′E). The Boulari channel is a cleaning station on a 15-meter-deep reef flat on the north tip of the channel and the Dumbea channel is a feeding site within the 30 meters of the water column facing the reef crest of the south tip of the channel. In Touho (location undisclosed), the aggregation site is a cleaning station located at the north tip of the Great Channel of Touho on a 15-to-20- meter reef shelf less than 5 km off the coast. In Ouvea, the deployment of the tags occurred at two aggregation sites: the Northern Pleiades (20°45′S, 166°44′E) and the Southern Pleiades (20°43′S, 166°23′E). Both sites are cleaning station on the reef slope at a depth of 10 to 15 m on the continental shelf off the northern tip and southern tip of the island, respectively (Fig. 1).

## 2.2. Tagging

We deployed 21 SPLASH10-F-321A PSAT tags and two MiniPAT tags (Wildlife Computers Inc.; Redmond, Washington, USA) coated with Propspeed™ silicone coating to prevent fouling during the deployment period. Tag deployments were timed to coincide with sightings peaks for all sites

3

between December and February over 4 years from 2015 to 2020. Two additional tags were deployed opportunistically in Touho in November 2018. In Noumea and Touho tags were deployed while scuba diving, in Ouvea tagging was performed while free diving on free-swimming manta rays. The tag is tethered by a 30 cm stainless steel cable to a titanium dart-tip. A modified pole spear was used to apply dart into the dorsal musculature of the manta ray. All tags were programmed to remain attached for a maximum period of 180 days. Alternatively, detachment was programmed in case of the tag being recorded at a constant depth for more than 24 hours (in case of mortality). SPLASH10 tags collected external temperature, light level, and pressure (depth) data every 10 s. Data were transmitted via the ARGOS satellite system (www.argos-system.org) into 12-h periods. Prior to tagging, manta rays were identified using photo-identification, the sex and maturity was also determined, and the size (disc width) was visually estimated to the nearest 10 cm.

# 2.3. Data Analysis

Locations were retrieved using the Wildlife Computers location processing systems (Fastloc GPS and GPE3). Location records are assigned with a quality rank based on the number of uplinks received per satellite pass for Argos data and on the number of connected satellites for Fastloc data. Argos data quality numbers rank from 3 to 0 and then A, B and Z, with 3 being the most accurate and Z being an invalid location. Fastloc data accuracy followed a gradient indicated by quality classes corresponding to the number of satellites, starting at 4 to 7 [50]. All location data were filtered according to the following steps. First, we manually removed duplicates and on-land locations, examined duplicates, and removed the less accurate ones. Second, argos-derived locations with an accuracy class inferior to A (no accuracy estimation) and Fastloc-derived locations inferior to 5 satellites were removed. Finally, we used the sdafilter function from the R package argosfilter [51] to exclude improbable locations based on speed and distance where trips exceeding a given speed (2 m.s<sup>-1</sup>) over a given distance (5,000 m) were considered improbable.

Filtered datasets were analysed using R version 3.6.3 [52]. Maps and bathymetry data were extracted from the marmap [53] package in R.

# 2.4. Statistical Analysis

Comparison of means between grouping factors were tested using an analysis of variance (ANOVA) and Tukey's post-hoc test. Welch's F tests were used when variance where unequal. Pairwise comparisons of means were tested with Student t-tests. The assumption of homogeneity was tested using Levene's tests. Chi-squared ( $\chi^2$ ) goodness of fit tests were used to examine the number of individuals between and among movement patterns. Z-tests of proportion were used to compare proportions. Finally, Kolmogorov-Smirnov test were used to compare distributions and Pearson correlation coefficient was used to test linear relationship between variables. In this paper, mean values are followed by the standard deviation in the format: (MEAN  $\pm$  SD).

# 2.5. Ethic Statement

The tagging was conducted with authorizations from the Southern Province (permit no: 34584) and the Northern Province (permit no: 609011–33) of New Caledonia. In the Loyalty Islands Province, no permit was required by the competent authorities, though permission of the local customary representatives was granted.

# 3. Results

# 3.1. Tagging

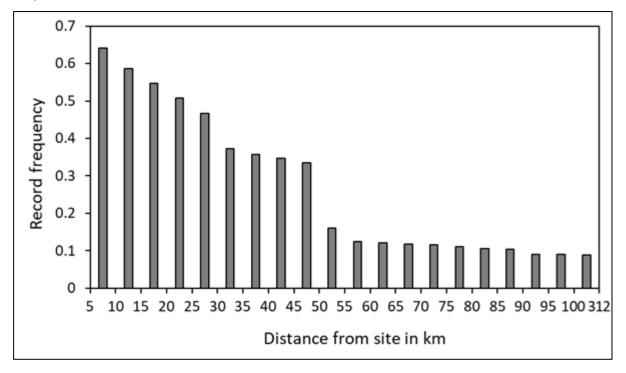
A total of 23 tags were deployed, 19 successfully transmitted data and 4 failed for unknown reasons (Table S1). Data from 16 tags were retrieved through the ARGOS server with a mean decoding rate of  $83.9 \pm 13.9$  %, and 3 tags were physically recovered allowing 100 % of the data to be decoded. After the data filtering process, 2 tags were excluded from the analysis of horizontal

movements due to inaccurate data. The mean retention period is  $76.7 \pm 50.3$  days (range from 3 to 180 days). Only the MiniPAT tag detached at the term of the programmed period (180 days), all other tag popped-up prematurely for unknown reasons. The majority of tagged individuals were females (57.9%) including a pregnant one and two were juveniles (one of each sex). Finally, all tagged individuals recorded only a few locations per day of deployment with an average of one record every 1.5 days. This number varied between sites from one record every 5.5 days and 0.9 days in Ouvea and Noumea, respectively. In Touho a location every 2.7 days of deployment was recorded.

#### 3.2. Horizontal Movements

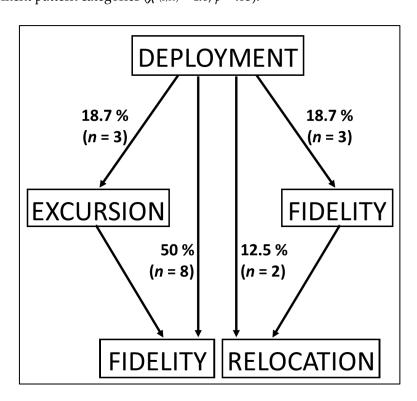
Horizontal movements varied greatly in length and direction between individuals. Manta rays travelled an average of  $4.6 \pm 3.1$  km d<sup>-1</sup> (ranging from 0.2 to 12.3 km d<sup>-1</sup>) with total track lengths varying from 23 to 688 km in 100 and 53 days, respectively. The latitudinal range extended from 7.1 to 224 km with an average of  $76.7 \pm 74.1$  km. These movements remained mostly within a relative proximity of the site of deployment with 63.5% of the recorded locations observed within 30 km from the deployment areas. The average distance from the tagging site was  $41 \pm 66$  km with a maximum at 311 km. Manta rays remained close to shallow waters, on average at a distance of  $3.5 \pm 4.4$  km from depth under 10-meter. They displayed limited offshore movements, only one individual was located offshore up to 105 km from the nearest reef. During these movements, manta rays occurred at locations with highly varying bathymetry averaging  $325 \pm 411$  m below the surface and a maximum recorded at 2,720 m deep.

Movement patterns are defined using a threshold distance from the site of origin to determinate a home range area. This value corresponds to the distance beyond which the distribution of record frequencies shows a significant drop. When combining all location data from all sites, the distribution of distance from the tagging site reveals a significant drop beyond the 50 km range (Fig. 2). This 50 km threshold from the site of origin will be used to identify movement patterns in and out of this range.



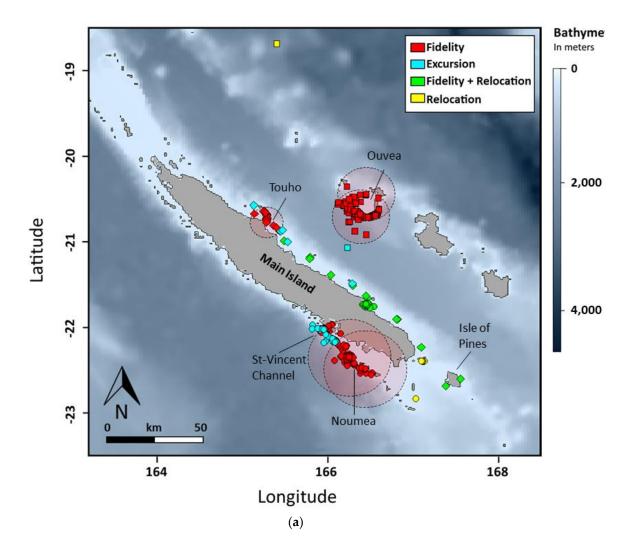
**Figure 2.** Frequency distribution of records of reef manta rays ( $Mobula\ alfredi$ ) (N = 16) at different ranges of distance from the site of tag deployment, using satellite tags (SPLASH10), in New Caledonia.

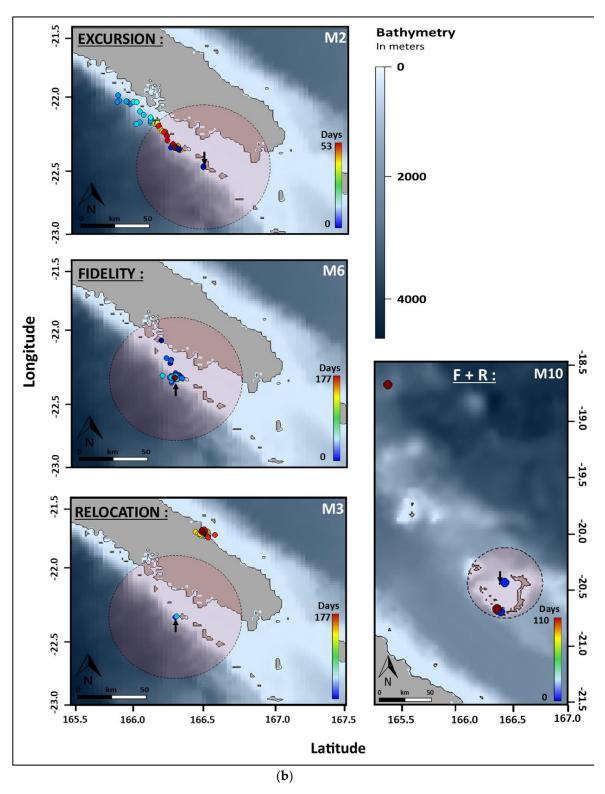
Four different movement patterns were identified from the combined records of all sites (Fig. 3, Fig. 4, Table 1). First, *fidelity* when manta rays remained within 50 km from the location of tagging. This represents half of the individuals. Second, *excursion* when manta rays extended their movement beyond 50 km off the site of origin at least once but consistently returned to this area. Three individuals followed this pattern. The number of excursions per deployment varied from 1 to 16 for a maximum duration ranging from 1 to 9 days (Fig. S1). Third, the *relocation* pattern when manta rays moved at more than 50 km off the tagging site, and remained outside this tagging site more than half of the time until release of the tag. Only two individuals (12.5 %) relocated at distance ranging from 71 to 248 km for a maximum duration of 48 and 40 days before release of the tag, respectively. Finally, the fourth pattern, *fidelity* + *relocation*, represents manta rays that had more than half of their recorded location within the 50 km range before moving further away toward the end of the deployment. This last pattern describes 18.7 % of individuals. The counts of individuals were not significantly different between movement pattern categories ( $\chi^2$ (3,16) = 4.0, p > .05).



**Figure 3.** Movement patterns recorded for reef manta rays (*Mobula alfredi*) after deployment of the tag, using satellite tag (SPLASH10), in New Caledonia. Percentage based on the total number of individuals (N = 16).







**Figure 4. a.** Movement patterns of reef manta rays ( $Mobula\ alfredi$ ) In New Caledonia using satellite telemetry (SPLASH10). Symbols shapes indicate site of origin: circle = Noumea (N = 6), square = Ouvea (N = 4) and diamond = Touho (N = 6). **b.** Examples of movement patterns of reef manta rays ( $Mobula\ alfredi$ ) In New Caledonia using satellite telemetry (SPLASH10). Circles indicate respective home range and arrows indicate deployment location. F + R = Fidelity + Relocation and Mx indicate individuals identification.

**Table 1.** Movement pattern characteristics and distribution between sites and sex of reef manta rays (*Mobula alfredi*) (*N* = 17) in New Caledonia using satellite telemetry (SPLASH10). HR: Home Range.

Movement Patterns	Pattern explanation	Noumea		Ouvea		Touho		TOTA
		Mal	Femal	Mal	Femal	Mal	Femal	L L
		e	e	e	e	e	e	
FIDELITY	100 % records in the	1	1	0	3	1	1	7
	HR							
EXCURSION	≥ 1 record outside the	0	2	1	0	1	1	5
	home range followed							
	by ≥1 record within the							
	HR							
FIDELITY +	≥ 50 % records in the							
RELOCATION	HR followed by $\geq 1$	1	0	1	0	0	0	2
	record outside at tag							
	release							
RELOCATION	≥ 50 % records outside	0	1	0	0	1	1	3
	the HR until tag release							

Comparisons between sites reveal significantly different distributions of distance frequencies (Fig. S2, Table S2). Home ranges were smaller in Touho (N = 19 records) and Ouvea (N = 105) with a limit at 20 and 35 km from the tagging site, respectively. In Noumea (N = 626) the home range was bigger with a limit at 50 km (Fig. 4, Table 1). At this site, one third of the tagged manta rays remained within the home range perimeter. Another third recorded excursions along the barrier reef up to 86.1 km north of the deployment location. The last third relocated at 96 (M4) and 250 km (M3) at the southern tip of the lagoon of the Main Island and on the east coast of the Main Island, respectively. In Ouvea, two individuals remained within the designated home range of 35 km from the site while another recorded an excursion halfway between the island of Ouvea and the Main Island over the 2000-meter-deep channel (Fig. 4, Table 1). The last manta ray relocated up to 225 km north toward the isolated reefs of Petrie and spent a maximum of 107 days outside its initial home range. In Touho, two manta rays remained within the 20 km home range (Fig. 4, Table 1). Two individuals recorded excursions, including one that moved 131 km down the coast for a maximum of 82 days outside its initial home range. Finally, the two other manta rays from Touho relocated. One travelled down to the Isle of Pines (311 km) to return later toward the tagging site where the tag detached 70 km south. The other relocated at the Isle of Pines. Differences in the number of individuals per movement patterns between site were not significant (Table S3).

Overlap in space occupation between individuals tagged at different sites was observed between manta rays from Touho and Noumea in the southern tip and the southern part of the east coast of the Main Island (Fig. 4). Among individuals tagged at the same site, overlap of movements outside the tagging sites only occurs off Noumea within an area located 50 km north of the Dumbea channel near the St-Vincent channel.

# 3.3. Sex

Females were recorded at significantly farther distance from the site of tagging than males ( $t_{(781)}$  = 5.9, p < .001). However, none of the gender was more mobile than the other with no significant difference in distances per day ( $t_{(14)}$  = 0.46, p > .05), total track lengths ( $t_{(14)}$  = 0.14, p > .05) or in the extend of the latitudinal range ( $t_{(14)}$  = 1.01, p > .05). The comparison of movement patterns among sex showed a significant difference in distributions of frequencies of recorded distance from sites (K-S:  $D_{(21)}$  = 0.67, p < .001). Females (N = 9 individuals) recorded a broader home range than males (N = 7)

with a perimeter of 50 km against 35 km, respectively. Differences in the number of individuals per movement patterns between sex were not significant (*p-values* > .05 *by Fisher's exact test*).

#### 4. Discussion

In New Caledonia, reef manta rays tend to remain close to coasts and reefs. Except for one individual, all records were within 35 km off the closest reef. Several studies that documented movements of reef manta rays reported consistent use of coastal and reef areas in Australia [15,54], in the Red Sea [55,56], in the Seychelles [57], and in the British Indian Ocean Territory MPA [58]. The use of coastal habitat by reef manta rays and other planktivorous elasmobranchs has been associated with food availability [13,24,59-63]. Coastal areas of islands surrounded by deep waters are prone to assemble the conditions to generate primary productivity hotspots combining upwellings and land inputs and generating reliable food resources which might result in strong residency pattern in reef manta rays' behaviour [57,64]. In New Caledonia, such conditions exist with little annual variations [65,66]. Limited offshore movements are likely due to the foraging ground being coastal and the probable lower productivity of the adjacent oceanic waters [67]. Other than food resources, coastal areas and shallow waters of the lagoon offer more protection from predators such as large sharks and hence, represent a suitable environment for potential nurseries. In New Caledonia, the detection of juvenile individuals remains sporadic, and neonates are even scarcer, although lagoons are places of nursery grounds for reef manta rays in Raja Ampat, Indonesia [64] and for other elasmobranchs [68-70]. Further investigations in suspected adequate nursery grounds using aerial surveys and accurate measurement methods to quantify maturation stages might lead to the discovery of nurseries in New Caledonia. In addition, coral reefs are also home to numerous cleaning stations that are essential to manta rays' well-being and health [71], as well as mating and socialization [72,73]. Sheltered waters also act as a thermal refuge where enclosed shallow waters have a significantly higher temperature than oceanic waters. In this study, manta rays were observed to favour relatively high temperatures (27 to 29 °C) but were also able to face much colder water for instance when deep diving (7.6 °C). Basking in warm shallow waters after a deep dive would allow regulation of the body temperature. Thermoregulation is an important part of the behaviour of mobulids [45,74] and other elasmobranchs [75,76] and the presence of warm-sheltered water in proximity of deep feeding grounds might be essential.

The use of offshore waters by one individual recorded in this study could be interpreted as a transiting trip toward potential alternative foraging grounds near isolated reefs. Presence of reef manta rays at isolated reefs have already been recorded during aerial surveys [77], further investigation in these areas might reveal new aggregations. Offshore movements by reef manta rays were also documented using satellite telemetry and were associated with foraging opportunities. In the east coast of Australia, [15] recorded most of the tagged rays in the offshore Capricorn Eddy Region and attributed this behaviour to be food related to the high productivity generated by the eddy. In the Red Sea, [55] observed offshore excursions to deeper water at night where manta rays performed deep dives supposedly to exploit pelagic planktonic resources.

Despite their demonstrated coastal affinity, reef manta rays have been observed over waters up to 2,720 m deep. The archipelago of New Caledonia has a relatively narrow continental shelf, especially on the east coast of the Main Island and around the Loyalty Islands, beyond which water depth drops rapidly. Even though manta rays spent most of their time within the first 50 meters, our results suggest that this species is able to use deeper water to transit between areas (e.g.,; to connect with an isolated reef) or to access to demersal food resources [78]. This supports previous findings that recorded reef manta rays commuting between islands chain separated by deep water in Indonesia [14,64] or transiting by deep water to access food resources in east Australia [15] and the Maldives [79]. However, even if there are evidence that deep water is not a strict barrier to their movements, large expanses of deep water might still be a restricting factor to connectivity. For instance, in this study, spatial occupancy overlapped only for individuals originating from sites sharing the same coastline and not for individuals separated by a 2,000-meter-deep channel. Other evidence through different spatial and temporal scale using photo-identification [47] and genomics

10

[48] showed that these connections occur but were limited, highlighting the lack of connectivity over this deep-water channel. Other studies taking place in atolls were consistent with these observations with populations recording no or only few connections between islands separated by deep waters in Hawaii [80,81], in the Maldives [82], in French Polynesia [83] or in the Seychelles [57]. On the contrary, reef manta rays showed long distance movements along continuous coastlines, up to 1150 km in east Australia [84], but also along the west coast of Australia (up to 700 km, [54]) or the coast of southern Mozambique (up to 350 km, [85]).

Horizontal movements were classified into four distinct patterns. The most dominant pattern was Fidelity where manta rays remained within their home range for the whole duration of the tag deployment. Manta rays displayed also excursions travelling, occasionally, relatively long distance as far as 131 km away from the site of origin. Moreover, total track lengths were relatively small (maximum of 688 km) in comparison to previous findings averaging 839 km in western Australia [54], 1169 km in east Australia [15] and 1074 km in the Seychelles [57]. Our results describe a behaviour that corresponds to site fidelity as defined by Chapman et al. (2015): the return of an individual to a location where it previously resided after an absence as long as or longer than the residency period [86]. Additional evidence of such behaviour was found in New Caledonia using long-term photo-identification data to find high re-sighting rates for each site [47] and genomics analysis to detect genetic structure between sites [48]. Site fidelity is often motivated by consistent foraging opportunities over time within an area [86]. This behaviour has been largely documented for reef manta rays in Hawaii [81], Mozambique [87], east Australia [60], Indonesia [64,88] or French Polynesia [83]. Although most of the tagged individuals displayed constrained movements near the aggregation site of origin, manta rays of New Caledonia demonstrate the ability to potentially connect with all parts of the archipelago, at least parts that are linked by coastlines or shallow waters. In particular, almost a third of the individuals showed Relocation and Fidelity + Relocation patterns, connecting with sites up to 311 km from the site of origin. Coupled with return trips over 200 km away from the deployment location, these observations suggest that manta rays connect to intermediate sites. For instance, while no connection was observed between studied sites during this study, overlap occurred for individuals from Touho and Noumea at in-between sites along the east coast and at the southern tip of the Main Island. Sightings of manta rays using photo-identifications over five years revealed different aggregation sites throughout the archipelago with multiple resightings from individuals originating from different sites [47]. The present results suggest the potential existence of additional aggregation sites where manta rays from different sites may encounter each other, although no field operation could be undertaken to confirm it.

The present results present a partial short-term overview of the spatial ecology of the reef manta rays of New Caledonia. The average deployment duration in this study was short (approx. 77 days) compared to what was achieved in other satellite telemetry studies on reef manta rays (e.g.; 92 days, [15]; 147 days, [55]; 116 days, [57]). This might also explain the relatively smaller track lengths recorded in this study than at other locations in the world. To complement this finding, long term monitoring of the movements and geographically extended sampling effort might help deciding whether these observations are sporadic, frequent, seasonal or to what extend long-term site fidelity is supported. They may as well reveal new connections and new potential aggregation sites. To this regard, photo-identification offers such perspectives and acoustic telemetry may narrow the gap between photo-identification and satellite telemetry by generating spatially and temporally intermediate data. In addition, genomics would provide a greater picture of the evolutionary processes that shape the population.

# 5. conclusions

This work used satellite telemetry and confirmed that reef manta rays are resident to coastal water and show strong site fidelity in New Caledonia. Yet, this species is capable of relatively long-distance migrations seemingly favouring, but not limiting them to, coastal and shallow water paths. Deep water might be a restraining factor but not a complete barrier to connectivity. These results raise concerns regarding threats associated with habitat degradation, human exploitation and

disturbance. The home range of this species may also extend over greater distances than previously thought, especially in habitat fragmented by deep waters, which raises concerns regarding potential movements outside areas under protective jurisdictions. In New Caledonia, concerns regarding the species conservation are limited since reef manta rays are not targeted by fisheries and human activity remains relatively low. Such favourable context is scarce which makes the population of reef manta rays in New Caledonia a reference to be preserved [19]. Therefore, preventive precautions should be taken at local level where coastal development is rapidly expanding and might threaten critical habitats.

**Supplementary Materials:** The following supporting information can be downloaded at the website of this paper posted on Preprints.org.

Funding: We express particular thanks to the Keidanren Nature Conservation Fund (KNCF) who funded the study through the SATO YAMA UMI Project and Conservation International. We thank the Southern Province of New Caledonia for their financial support in the form of a scholarship to HL. We also gratefully acknowledge the financial support of MAC3 Impact Philanthropies, William Brooks, Pam Rorke Levy, Audrey and Shannon Wong, Daniel Roozen and Kris Norvig in sponsoring satellite tags, and we thank OceanMax for supplying the Propspeed antifouling coating for our tags. Conservation International played a role in the study design and the data collection but had no role in analysis, decision to publish, or preparation of the manuscript. The remaining funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

Acknowledgments: We thank Conservation International, the Aquarium des Lagons, MAC3 Impact Philanthropies and The Manta Trust for their overall support for the study. We especially thank Mark Erdmann for unvaluable guidance and help on the field and continuing overall support. Special thanks to Franck Bouilleret, Mael Imirizaldu and Thomas Auger for their priceless assistance in field operations, as well as Jean-Christophe Lefeuvre and François Tron for their support. We also thank the Hô-üt Association and its members as well as all those involved in the field operations for their valuable help (Pierre Kaouma, Marino Tiaou, Jean-Baptiste Badiou, Ludovic Mazens, Abyss Plongée, Sarah Lewis, Schannel van Dijken, Minori Matsuoka). We finally gratefully acknowledge the customary representatives of Ouvéa and Touho, and the three Provinces for allowing such operations.

# References

- 1. Fletcher, R.; Fortin, M. Spatial ecology and conservation modelling. *Springer Int. Pub.* **2018**, *Chapter 1*, pp. 1–13.
- 2. Graham, R. T.; Witt, M. J.; Castellanos, D. W.; Remolina, F.; Maxwell, S.; Godley, B. J.; Hawkes, L. A. Satellite tracking of manta rays highlights challenges to their conservation. *PLoS One*. **2012**, *7*(5), *e*36834.
- 3. Palumbi, S. R. Genetic divergence, reproductive isolation, and marine speciation. *Annu. Rev. Ecol. Evol. Syst.* **2014**, *25*(1), pp. 547-572.
- 4. Palumbi, S. R. (2003). Population genetics, demographic connectivity, and the design of marine reserves. Ecol Appl.13(sp1), 146-158.
- 5. Chateau, O.; Wantiez, L. Movement patterns of four coral reef fish species in a fragmented habitat in New Caledonia: implications for the design of marine protected area networks. *ICES Mar. Sci. Symp.* **2009**, *66*(1), pp.50-55.
- 6. Meyer, C. G.; Papastamatiou, Y. P.; Clark, T. B. Differential movement patterns and site fidelity among trophic groups of reef fishes in a Hawaiian marine protected area. *Mar. Biol.* **2010**, *157*(7), pp. 1499-1511.
- 7. Foote, A. D.; Similä, T.; Víkingsson, G. A.; Stevick, P. T. Movement, site fidelity and connectivity in a top marine predator, the killer whale. *Evol. Ecol.* **2010**, 24(4), pp. 803-814.
- 8. Passadore, C.; Möller, L.; Diaz-Aguirre, F.; Parra, G. J. High site fidelity and restricted ranging patterns in southern Australian bottlenose dolphins. *Ecol. Evol.* **2018**, *8*(1), pp. 242-256.
- 9. Madigan, D. J.; Brooks, E. J.; Bond, M. E.; Gelsleichter, J.; Howey, L. A.; Abercrombie, D. L.; ...; Chapman, D. D. Diet shift and site-fidelity of oceanic whitetip sharks *Carcharhinus longimanus* along the Great Bahama Bank. *Mar. Ecol. Prog.* **2015**, *529*, pp. 185-197.
- 10. Sims, D. W.; Quayle, V. A. Selective foraging behaviour of basking sharks on zooplankton in a small-scale front. *Nature*, **1998**, *393*(*6684*), pp. 460-464.
- 11. Heyman, W. D.; Graham, R. T.; Kjerfve, B.; Johannes, R. E. Whale sharks *Rhincodon typus* aggregate to feed on fish spawn in Belize. *Mar. Ecol. Prog.* **2001,** 215, pp. 275-282.
- 12. Graham, R. T.; Roberts, C. M.; Smart, J. C. Diving behaviour of whale sharks in relation to a predictable food pulse. *J. R. Soc. Interface*, **2006**, *3*(*6*), pp. 109-116.

- 13. Anderson, R. C.; Adam, M. S.; Goes, J. I. From monsoons to mantas: seasonal distribution of *Manta alfredi* in the Maldives. *Fish. Oceanogr.* **2011**, 20(2), pp. 104-113.
- 14. Germanov, E. S.; Marshall, A. D. Running the gauntlet: regional movement patterns of *Manta alfredi* through a complex of parks and fisheries. *PLoS One*, **2014**, *9*(10), e110071.
- Jaine, F. R. A.; Rohner, C. A.; Weeks, S. J.; Couturier, L. I. E.; Bennett, M. B.; Townsend, K. A.; Richardson, A. J. Movements and habitat use of reef manta rays off eastern Australia: offshore excursions, deep diving and eddy affinity revealed by satellite telemetry. *Mar. Ecol. Prog.* 2014, 510, pp. 73-86.
- 16. Jaine, F. R.; Couturier, L. I.; Weeks, S. J.; Townsend, K. A.; Bennett, M. B.; Fiora, K.; Richardson, A. J. When giants turn up: sighting trends, environmental influences and habitat use of the manta ray *Manta alfredi* at a coral reef. *PLoS One*, **2012**, *7*(10).
- 17. Couturier, L. I.; Dudgeon, C. L.; Pollock, K. H.; Jaine, F. R. A.; Bennett, M. B.; Townsend, K. A.; ...; Richardson, A. J. Population dynamics of the reef manta ray *Manta alfredi* in eastern Australia. *Coral Reefs*, **2014**, 33(2), pp. 329-342.
- 18. Setyawan, E.; Sianipar, A. B.; Erdmann, M.; Fischer, A. M.; Haddy, J. A.; Beale, C. S.; ...; Mambrasar, R. Site fidelity and movement patterns of reef manta rays (*Mobula alfredi*: Mobulidae) using passive acoustic telemetry in northern Raja Ampat, Indonesia. *Nat. Conserv. Res.* **2018**, *3*(4), pp. 17-31
- 19. O'Malley, M. P.; Townsend, K. A.; Hilton, P.; Heinrichs, S.; Stewart, J. D. Characterization of the trade in manta and devil ray gill plates in China and South-east Asia through trader surveys. *Aquat. Conserv.: Mar. Freshw. Ecosyst.* **2017**, *27*(2), pp. 394-413.
- 20. White, W.T.; Giles, J.; Potter, I.C. Data on the bycatch fishery and reproductive biology of mobulid rays (Myliobatiformes) in Indonesia. *Fish. Res.* **2006**, *82*(*1-3*), pp.65-73.
- 21. O'Malley, M. P.; Lee-Brooks, K.; Medd, H. B. The global economic impact of manta ray watching tourism. *PLoS One*, **2013**, *8*(5), e65051.
- 22. Venables, S.; McGregor, F.; Brain, L.; van Keulen, M. Manta ray tourism management, precautionary strategies for a growing industry: a case study from the Ningaloo Marine Park, Western Australia. *Pac. Conserv. Biol.* **2016**, 22(4), pp. 295-300.
- 23. Ward-Paige, C. A.; Davis, B.; Worm, B. Global population trends and human use patterns of Manta and Mobula rays. *PLoS One*, **2013**, *8*(9), e74835.
- 24. Rohner, C. A.; Pierce, S. J.; Marshall, A. D.; Weeks, S. J.; Bennett, M. B.; Richardson, A. J. Trends in sightings and environmental influences on a coastal aggregation of manta rays and whale sharks. *Mar. Ecol. Prog.* **2013**, 482, pp. 153-168.
- 25. Croll, D. A.; Dewar, H.; Dulvy, N. K.; Fernando, D.; Francis, M. P.; Galván-Magaña, F.; ...; White, W. T. Vulnerabilities and fisheries impacts: the uncertain future of manta and devil rays. *Aquat. Conserv.: Mar. Freshw. Ecosyst.* **2016**, *26*(3), pp. 562-575.
- 26. Lawson, J. M.; Fordham, S. V.; O'Malley, M. P.; Davidson, L. N.; Walls, R. H.; Heupel, M. R.; ...; Dulvy, N. K. Sympathy for the devil: a conservation strategy for devil and manta rays. *PeerJ*, **2017**, *5*, e3027.
- 27. McGregor, F.; Richardson, A. J.; Armstrong, A. J.; Armstrong, A. O.; Dudgeon, C. L. Rapid wound healing in a reef manta ray masks the extent of vessel strike. *PLoS One*, **2019**, *14*(12), e0225681.
- 28. Stewart, J. D.; Jaine, F. R. A.; Armstrong, A. J.; Armstrong, A. O.; Bennett, M. B.; Burgess, K. B.; ...; Stevens, G. M. Research priorities to support effective manta and devil ray conservation. *Front. Mar. Sci.* **2018**, *5*, pp. 314.
- 29. Marshall, A.; Kashiwagi, T.; Bennett, M.B.; Deakos, M.; Stevens, G.; McGregor, F.; Clark, T.; Ishihara, H.; Sato, K. *Mobula alfredi* (amended version of 2011 assessment). *The IUCN Red List of Threatened Species* 2018: e.T195459A126665723. 2018 https://dx.doi.org/10.2305/IUCN.UK.2011-2.RLTS.T195459A126665723.en. Downloaded on 02 January 2021.
- 30. Baird, R. W.; Schorr, G. S.; Webster, D. L.; Mahaffy, S. D.; McSweeney, D. J.; Hanson, M. B.; Andrews, R. D. Open-ocean movements of a satellite-tagged Blainville's beaked whale (*Mesoplodon densirostris*): evidence for an offshore population in Hawaii? *Aquat. Mamm.* **2011**, *37*(4), pp. 506-511.
- 31. Block, B. A.; Costa, D. P.; Boehlert, G. W.; Kochevar, R. E. Revealing pelagic habitat use: the tagging of Pacific pelagics program. *Oceanol. Acta*, **2002**, *25*(*5*), pp.255-266.
- 32. Hart, K. M.; Hyrenbach, K. D. Satellite telemetry of marine megavertebrates: the coming of age of an experimental science. *Endanger. Species Res.* **2009**, *10*, pp. 9-20.
- 33. Mate, B. R.; Best, P. B.; Lagerquist, B. A.; Winsor, M. H. Coastal, offshore, and migratory movements of South African right whales revealed by satellite telemetry. *Mar. Mamm. Sci.* **2011**, 27(3), pp. 455-476.
- 34. Phillips, E. M.; Horne, J. K.; Adams, J.; Zamon, J. E. Selective occupancy of a persistent yet variable coastal river plume by two seabird species. *Mar. Ecol. Prog. Ser.* **2018**, *594*, pp. 245-261.

- 35. Hofman, M. P.; Hayward, M. W.; Heim, M.; Marchand, P.; Rolandsen, C. M.; Mattisson, J.; ... & Balkenhol, N. (2019). Right on track? Performance of satellite telemetry in terrestrial wildlife research. PLoS One, 14(5), e0216223.
- 36. Haywood, J. C.; Fuller, W. J.; Godley, B. J.; Margaritoulis, D.; Shutler, J. D.; Snape, R. T.; ...; Broderick, A. C. Spatial ecology of loggerhead turtles: Insights from stable isotope markers and satellite telemetry. *Divers. Distrib.* **2020**, *26*(3), pp. 368-381.
- 37. Meyers, M. M.; Francis, M. P.; Erdmann, M.; Constantine, R.; Sianipar, A. Movement patterns of whale sharks in Cenderawasih Bay, Indonesia, revealed through long-term satellite tagging. *Pac. Conserv. Biol.* **2020**, *26*(4), pp. 353-364.
- 38. Heupel, M. R.; Simpfendorfer, C. A.; Collins, A. B.; Tyminski, J. P. Residency and movement patterns of bonnethead sharks, *Sphyrna tiburo*, in a large Florida estuary. *Environ. Biol. Fishes*, **2006**, *76*(1), pp. 47-67.
- 39. Hammerschlag, N.; Gallagher, A. J.; Lazarre, D. M. A review of shark satellite tagging studies. *J. Exp. Mar. Biol. Ecol.* **2011**, 398(1-2), pp. 1-8.
- 40. Crossin, G. T.; Heupel, M. R.; Holbrook, C. M.; Hussey, N. E.; Lowerre-Barbieri, S. K.; Nguyen, V. M.; ...; Cooke, S. J. Acoustic telemetry and fisheries management. *Ecol Appl.* **2017**, 27(4), pp. 1031-1049.
- 41. Sims, D. W.; Southall, E. J.; Tarling, G. A.; Metcalfe, J. D. Habitat-specific normal and reverse diel vertical migration in the plankton-feeding basking shark. *J. Anim. Ecol.* **2005**, *74*(4), pp. 755-761.
- 42. Andrews, K. S.; Williams, G. D.; Levin, P. S. Seasonal and ontogenetic changes in movement patterns of sixgill sharks. *PLoS One*, **2010**, *5*(9), e12549.
- 43. Bonfil, R.; Meÿr, M.; Scholl, M. C.; Johnson, R.; O'Brien, S.; Oosthuizen, H.; ...; Paterson, M. Transoceanic migration, spatial dynamics, and population linkages of white sharks. *Science*, **2005**, *310*(5745), pp. 100-103.
- 44. Brunnschweiler, J. M.; Baensch, H.; Pierce, S. J.; Sims, D. W. Deep-diving behaviour of a whale shark *Rhincodon typus* during long-distance movement in the western Indian Ocean. *J. Fish Biol.* **2009**, 74(3), pp. 706-714.
- 45. Thorrold, S. R.; Afonso, P.; Fontes, J.; Braun, C. D.; Santos, R. S.; Skomal, G. B.; Berumen, M. L. Extreme diving behaviour in devil rays links surface waters and the deep ocean. *Nat. Commun.* **2014**, *5*(1), pp. 1-7.
- 46. Lassauce, H.; Chateau, O.; Erdmann, M. V.; Wantiez, L. Diving behavior of the reef manta ray (*Mobula alfredi*) in New Caledonia: More frequent and deeper night-time diving to 672 meters. *PLoS One*, **2020**, *15*(3), e0228815.
- 47. Lassauce, H.; Chateau, O.; Erdmann, M. V.; Wantiez, L. Citizen science to infer characteristics and habitat use of the population of reef manta rays (*Mobula alfredi*) in New Caledonia. University of New Caledonia, Noumea, New Caledonia, 2023, *Manuscript in preparation*.
- 48. Lassauce, H.; Dudgeon, C. L.; Armstrong, A. J.; Wantiez, L.; Carroll, E. L. Evidence of fine-scale genetic structure for reef manta rays *Mobula alfredi* in New Caledonia. *Endanger. Species Res.* **2022**, *47*, pp. 249-264.
- 49. Andréfouët, S.; Cabioch, G.; Flamand, B.; Pelletier, B. A reappraisal of the diversity of geomorphological and genetic processes of New Caledonian coral reefs: a synthesis from optical remote sensing, coring and acoustic multibeam observations. *Coral Reefs*, **2009**, *28*(3), pp. 691-707.
- 50. Van Canneyt, O.; Dorémus, G.; Laran S.; Ridoux, V.; Watremez, P. Distribution et abondance de la mégafaune marine dans le sud-ouest du Pacifique. 2015 (Scientific campaign report). Available online: https://www.observatoire-pelagis.cnrs.fr/wp-content/uploads/2021/05/9-RAPPORT\_REMMOA\_SOP\_INTER\_2015.pdf (accessed on 22 May 2020)
- 51. Freitas, C.; Lydersen, C.; Fedak, M. A.; Kovacs, K. M. A simple new algorithm to filter marine mammal Argos locations. *Mar. Mamm. Sci.* **2008**, 24(2), pp. 315-325.
- 52. R Core Team. R: A language and environment for statistical computing. R Foundation for Statistical Computing, Vienna, Austria. 2020, URL https://www.R-project.org/.
- 53. Pante, E.; Simon-Bouhet, B. marmap: a package for importing, plotting and analyzing bathymetric and topographic data in R. *PLoS One*, **2013**, *8*(9), e73051.
- 54. Armstrong, A. J.; Armstrong, A. O.; McGregor, F.; Richardson, A. J.; Bennett, M. B.; Townsend, K. A.; ...; Dudgeon, C. L. Satellite tagging and photographic identification reveal connectivity between two UNESCO World Heritage Areas for reef manta rays. *Front. Mar. Sci.* **2020**, *7*, pp. 725.
- 55. Braun, C. D.; Skomal, G. B.; Thorrold, S. R.; Berumen, M. L. Movements of the reef manta ray (*Manta alfredi*) in the Red Sea using satellite and acoustic telemetry. *Mar. Biol.* **2015**, *162*(*12*), pp. 2351-2362.
- 56. Kessel, S. T.; Elamin, N. A.; Yurkowski, D. J.; Chekchak, T.; Walter, R. P.; Klaus, R.; ...; Hussey, N. E. Conservation of reef manta rays (*Manta alfredi*) in a UNESCO World Heritage Site: Large-scale island development or sustainable tourism? *PLoS One*, **2017**, 12(10), e0185419.
- 57. Peel, L. R.; Stevens, G. M.; Daly, R.; Keating Daly, C. A.; Collin, S. P.; Nogués, J.; Meekan, M. G. Regional movements of reef manta rays (*Mobula alfredi*) in Seychelles waters. *Front. Mar. Sci.* **2020**, 7, pp. 558.

- 58. Andrzejaczek, S.; Chapple, T. K.; Curnick, D. J.; Carlisle, A. B.; Castleton, M.; Jacoby, D. M.; ...; Block, B. A. Individual variation in residency and regional movements of reef manta rays *Mobula alfredi* in a large marine protected area. *Mar. Ecol. Prog. Ser.* **2020**, *639*, pp. 137-153.
- 59. Dewar, H.; Mous, P.; Domeier, M.; Muljadi, A.; Pet, J.; Whitty, J. Movements and site fidelity of the giant manta ray, *Manta birostris*, in the Komodo Marine Park, Indonesia. *Mar. Biol.* **2008**, *155*(2), pp. 121-133.
- 60. Couturier, L. I.; Jaine, F. R.; Townsend, K. A.; Weeks, S. J.; Richardson, A. J.; Bennett, M. B. Distribution, site affinity and regional movements of the manta ray, *Manta alfredi* (Krefft, 1868), along the east coast of Australia. *Mar. Freshw. Res.* **2008**, *62*(*6*), pp. 628-637.
- 61. Couturier, L.I.E.; Newman, P.; Jaine, F.R.A.; Bennett, M.B.; Venables, W.N.; Cagua, E.F.; Townsend, K.A.; Weeks, S.J.; Richardson, A.J. Variation in occupancy and habitat use of *Mobula alfredi* at a major aggregation site. *Mar. Ecol. Prog. Ser.* **2018**, *599*, pp.125-145.
- 62. Armstrong, A. O.; Armstrong, A. J.; Jaine, F. R.; Couturier, L. I.; Fiora, K.; Uribe-Palomino, J.; ...; Richardson, A. J. Prey density threshold and tidal influence on reef manta ray foraging at an aggregation site on the Great Barrier Reef. *PLoS One*, **2016**, *11*(5), e0153393.
- 63. McCoy, E.; Burce, R.; David, D.; Aca, E. Q.; Hardy, J.; Labaja, J.; ...; Araujo, G. Long-term photo-identification reveals the population dynamics and strong site fidelity of adult whale sharks to the coastal waters of Donsol, Philippines. *Front. Mar. Sci.* 2018, 5, pp. 271.
- 64. Setyawan, E.; Erdmann, M. V.; Lewis, S. A.; Mambrasar, R.; Hasan, A. W.; Templeton, S.; ...; Cerutti-Pereyra, F. Natural history of manta rays in the Bird's Head Seascape, Indonesia, with an analysis of the demography and spatial ecology of *Mobula alfredi* (Elasmobranchii: Mobulidae). *J. Ocean Sci. Found.* 2020, 36, pp. 49-83
- 65. Torréton, J. P.; Rochelle-Newall, E.; Pringault, O.; Jacquet, S.; Faure, V.; Briand, E. Variability of primary and bacterial production in a coral reef lagoon (New Caledonia). *Mar. Pollut. Bull.* **2010**, *61*(7-12), pp. 335-348.
- 66. Le Borgne, R.; Douillet, P.; Fichez, R.; Torréton, J. P. Hydrography and plankton temporal variabilities at different time scales in the southwest lagoon of New Caledonia: A review. *Mar. Pollut. Bull.*, **2010**, *61*(7-12), pp. 297-308.
- 67. McCauley, D. J.; DeSalles, P. A.; Young, H. S.; Papastamatiou, Y. P.; Caselle, J. E.; Deakos, M. H.; ...; Micheli, F. Reliance of mobile species on sensitive habitats: a case study of manta rays (*Manta alfredi*) and lagoons. *Mar. Biol.* **2014**, *161*(9), pp. 1987-1998.
- 68. Heupel, M. R.; Carlson, J. K.; Simpfendorfer, C. A. Shark nursery areas: concepts, definition, characterization, and assumptions. *Mar. Ecol. Prog. Ser.* **2007**, *337*, pp. 287–297.
- 69. Papastamatiou, Y. P.; Lowe, C. G.; Caselle, J. E.; Friedlander, A. M. Scale-dependent effects of habitat on movements and path structure of reef sharks at a predator-dominated atoll. *Ecology* **2009**, *90*, pp. 996–1008
- 70. Dale, J. I.; Wallsgrove, N. J.; Popp, B. N.; Holland, K. N. Nursery habitat use and foraging ecology of the brown stingray *Dasyatis lata* determined from stomach contents, bulk and amino acid stable isotopes. *Mar. Ecol. Prog. Ser.* **2011**, 433, pp. 221–236.
- 71. Barr, Y.; Abelson, A. Feeding–Cleaning Trade-Off: Manta Ray "Decision-Making" as a Conservation Tool. *Front. Mar. Sci.* **2019**, *6*, pp. 88.
- 72. Stevens, G. M. W. Conservation and population ecology of manta rays in the Maldives, Doctoral dissertation, University of York, 2016.
- 73. Perryman, R. J.; Venables, S. K.; Tapilatu, R. F.; Marshall, A. D.; Brown, C.; Franks, D. W. Social preferences and network structure in a population of reef manta rays. *Behav. Ecol. Sociobiol.* **2019**, *73*(8), pp. 114.
- 74. Stewart, J. D.; Hoyos-Padilla, E. M.; Kumli, ,K. R.; Rubin, R. D. Deep-water feeding and behavioral plasticity in *Manta birostris* revealed by archival tags and submersible observations. *Zoology*, **2016**, *119*(5), pp. 406-413.
- 75. Thums, M.; Meekan, M.; Stevens, J.; Wilson, S.; Polovina, J. Evidence for behavioural thermoregulation by the world's largest fish. *J. R. Soc. Interface*, **2013**, *10*(78), 20120477.
- 76. Speed, C. W.; Meekan, M. G.; Field, I. C.; McMahon, C. R.; Harcourt, R. G.; Stevens, J. D.; ...; Bradshaw, C. J. A. Reef shark movements relative to a coastal marine protected area. *Reg. Stud. Mar. Sci.* **2016**, *3*, pp. 58-66.
- 77. Laran, S.; Doremus, G.; Mannocci, L.; Van Canneyt, O.; Watremez, P.; Cadinouche, A.; ...; Razafindrakoto, Y. Progress of the REMMOA aerial surveys conducted in the French EEZ and adjacent waters: contrasted cetacean habitats in the southwest Indian Ocean. Report, 2016.
- 78. Lassauce, H.; Chateau, O.; M. V.; Wantiez, L. Spatial ecology of the population of reef manta rays (*Mobula alfredi*) in New Caledonia using satellite telemetry: 2-Vertical behaviour. University of New Caledonia, Noumea, New Caledonia, 2023, *Manuscript in preparation*.

- 79. Harris, J. L.; McGregor, P. K.; Oates, Y.; Stevens, G. M. Gone with the wind: Seasonal distribution and habitat use by the reef manta ray (*Mobula alfredi*) in the Maldives, implications for conservation. Aquatic Conservation: *Aquat. Conserv.: Mar. Freshw. Ecosyst.* **2020**, *30*(8), pp. 1649-1664.
- 80. Clark, T. B. Abundance, home range, and movement patterns of manta rays (*Manta alfredi*, *M. birostris*) in Hawai'i. Doctoral dissertation, University of Hawaii at Manoa, Honolulu, December 2010.
- 81. Deakos, M. H.; Baker, J. D.; Bejder, L. Characteristics of a manta ray Manta alfredi population off Maui, Hawaii, and implications for management. *Mar. Ecol. Prog. Ser.* **2011**, 429, pp. 245-260.
- 82. Kitchen-Wheeler, A. M.; Ari, C.; Edwards, A. J. Population estimates of Alfred mantas (*Manta alfredi*) in central Maldives atolls: North Male, Ari and Baa. *Environ. Biol. Fishes*, **2012**, *93*(4), pp. 557-575.
- 83. Carpentier, A. S.; Berthe, C.; Ender, I.; Jaine, F. R.; Mourier, J.; Stevens, G.; ...; Clua, E. Preliminary insights into the population characteristics and distribution of reef (*Mobula alfredi*) and oceanic (*M. birostris*) manta rays in French Polynesia. *Coral Reefs*, **2019**, *38*(6), pp. 1197-1210.
- 84. Armstrong, A. O.; Armstrong, A. J.; Bennett, M. B.; Richardson, A. J.; Townsend, K. A.; & Dudgeon, C. L. Photographic identification and citizen science combine to reveal long distance movements of individual reef manta rays *Mobula alfredi* along Australia's east coast. *Mar. Biodivers. Rec.* **2019**, 12(1), pp. 14.
- 85. Venables, S. K.; van Duinkerken, D. I.; Rohner, C. A.; Marshall, A. D. Habitat use and movement patterns of reef manta rays *Mobula alfredi* in southern Mozambique. *Mar. Ecol. Prog. Ser.* **2020**, *634*, pp. 99-114.
- 86. Chapman, D. D.; Feldheim, K. A.; Papastamatiou, Y. P.; Hueter, R. E. There and back again: a review of residency and return migrations in sharks, with implications for population structure and management. *Annu. Rev. Mar. Sci.* **2015**, *7*, pp. 547-570.
- 87. Marshall, A. D.; Pierce, S. J.; Bennett, M. B. Morphological measurements of manta rays (Manta birostris) with a description of a foetus from the east coast of Southern Africa. *Zootaxa*, **2008**, *1717*(1), pp. 24-30.
- 88. Germanov, E. S.; Bejder, L.; Chabanne, D. B.; Dharmadi, D.; Hendrawan, I. G.; Marshall, A. D.; ...; Loneragan, N. R. Contrasting habitat use and population dynamics of reef manta rays within the Nusa Penida marine protected area, Indonesia. *Front. Mar. Sci.* **2019**, *6*, pp. 215.

**Disclaimer/Publisher's Note:** The statements, opinions and data contained in all publications are solely those of the individual author(s) and contributor(s) and not of MDPI and/or the editor(s). MDPI and/or the editor(s) disclaim responsibility for any injury to people or property resulting from any ideas, methods, instructions or products referred to in the content.