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# **Original Article**

# Tuna aggregation dynamics at Drifting Fish Aggregating Devices: a view through the eyes of commercial echosounder buoys

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This study addresses novel questions on the dynamics of tuna aggregations around floating objects, using echosounder buoys data collected throughout the drifts of newly deployed Drifting Fish Aggregating Devices (DFADs) in the Atlantic Ocean (AO) and Indian Ocean (IO). Time series of presence/absence of tunas were obtained by supervised classification of acoustic data. To avoid biases related to the variability in individual DFAD soak times, a new approach was developed to estimate the average colonization time of new DFADs by tuna aggregations. We showed that tunas colonize DFADs after an average of 16 days in the AO, and 40 days in the IO. Moreover, the analysis indicated that the time span during which tuna aggregations occupy DFADs is driven by a time-independent process with short- and long-term residence modes. On average, DFADs were continuously occupied by tuna aggregations for 6 and 9 days in the IO and AO, respectively. The time between two consecutive aggregations at the same DFAD averaged 9 days in the IO and 5 days in the AO. Throughout their soak time after being colonized, DFADs remained occupied for a larger proportion of time in the AO (63%) than in the IO (45%).

**Keywords:** absence times, associative behaviour, colonization times, Drifting Fish Aggregating Devices (DFADs), echosounder buoys, residence times, tropical tunas

### Introduction

Substantial development in the tropical tuna purse-seine fishery has occurred over recent decades and it now accounts for the majority of the world's tropical tuna catch (ISSF, 2019). The increasing trend in catches has been accompanied by regular technological developments of vessels and fishing tools (Fonteneau *et al.*, 2000; Gaertner and Pallarés, 2002; Torres-Irineo *et al.*, 2014). Since the early 1990s, these developments have included the deployment of artificial floating objects, known as Drifting Fish Aggregating Devices (DFADs) used to increase fishery efficiency. This fishing mode exploits the behavioural trait of the target tuna species, skipjack tuna (*Katsuwonus pelamis*), yellowfin tuna (*Thunnus albacares*), and bigeye tuna (*Thunnus*  *obesus*), to naturally aggregate around floating objects (see Fréon and Dagorn, 2000; Castro *et al.*, 2001). Thousands of DFADs specifically designed to attract tunas, are deployed by purse seine fleets in all of the world's oceans to enhance their catches. Initial estimates put the number of DFADs operated annually across the three major oceans in the range of 50 000 and 100 000 (Baske *et al.*, 2012; Scott and Lopez, 2014; Gershman et al., 2015). Furthermore, the scale at which this fishing gear is used has quadrupled in less than a decade, in both the Atlantic Ocean (AO) and Indian Ocean (IO) (Fonteneau *et al.*, 2015; Maufroy *et al.*, 2017). Over the past 30 years, this fishing mode has been significantly improved through the sequential introduction of new technologies (radio, GPS, and echosounder buoys), with the latter now providing skippers and fleet managers with detailed information on the location and biomass associated with DFADs (Lopez *et al.*, 2014). In recent years, more than half of the world's purse seine catch of tropical tunas has come from DFAD fishing (Dagorn *et al.*, 2013; Fonteneau *et al.*, 2013).

The mechanisms underlying the associative behaviour of tropical tunas with floating objects remain poorly understood, despite the proposal of numerous hypotheses (see Fréon and Dagorn, 2000; Castro et al., 2001). As a consequence of the massive increase in the use of DFADs in the recent years, improving knowledge on the associative behaviour of tropical tunas with floating objects has become a key research priority. Primary areas of concern include the concomitant changes in catchability of tunas at floating objects as well as the understanding of the impact of DFADs on their ecology and that of other associated fauna. To date, most research efforts have focused on describing the associative dynamics of individual fish, primarily through electronic tagging studies, at both coastal anchored fish aggregating devices (FADs) (Holland et al., 1990; Ohta and Kakuma, 2005; Dagorn et al., 2007a; Mitsunaga et al., 2012; Robert et al., 2013; Rodriguez-Tress et al., 2017), and DFADs (Schaefer and Fuller, 2013; Matsumoto et al., 2014, 2016; Tolotti et al., 2020). In contrast, behavioural patterns of FAD-associated aggregations (i.e. entire tuna schools under a DFAD) have received considerably less research attention. When investigated, studies were primarily focused on the spatial or temporal characterization of aggregations at fine timescales, using acoustic equipment on board research vessels (Josse and Bertrand, 2000; Doray et al., 2006; Moreno et al., 2007a; Trygonis et al., 2016). The instrumentation of DFADs with satellite-linked echosounder buoys began in the late 2000s. Since then, their use has become widespread in many fleets. Currently, almost all deployed DFADs are equipped with these devices, which provide remote and near real-time information on the DFAD location and aggregated biomass (Lopez et al., 2014; Moreno et al., 2019). Echosounder buoys generate a considerable stream of data that can be used to characterize DFAD aggregations (Lopez et al., 2016; Orue et al., 2019a; Baidai et al., 2020). The availability of such data, which is continuously being collected, represents an unprecedented opportunity to observe fish aggregations associated with floating objects over long time scales. To date, only one study has provided characteristics of tuna aggregations at DFADs over the scale of weeks and months, using data from fisher's echosounder buoys (Orue et al., 2019b).

This work aims to characterize how tunas occupy DFADs, to improve the understanding of their aggregation dynamics around these objects. Using data from commercial echosounder buoys attached to DFADs deployed in the Western Indian Ocean and Eastern Atlantic Ocean, we assessed several parameters related to the association of tuna aggregations with DFADs. These include the elapsed time between deployment of a new DFAD and its colonization by tunas; the average duration of association of tuna aggregations with DFADs and the length of time that DFADs remains vacant between consecutive tuna aggregations.

## Material and methods Echosounder data

The echosounder data used in this study were collected using M3I buoys (Marine Instruments, Nigrán, Spain, www.marinein struments.es) attached to DFADs from the French tropical tuna purse seiner fleet operating in the Western Indian Ocean and

Eastern Atlantic Ocean, from 2016 to 2018. The M3I buoy is equipped with a GPS positioning device and an echosounder powered by solar panels, operating at a frequency of 50 kHz, with a power of 500 W, and a beam angle of  $36^{\circ}$ . The data output of the buoy is simplified acoustic information, designed for easy visual interpretation by fishers. The M3I buoy samples the water column in 3-m layers covering a total depth of 150 m (50 layers, with the first two corresponding to the transducer near-field). An acoustic sample consists of 50 ordered categorical scores (ranging from 0 to 7), resulting from the automatic conversion of the acoustic backscatter signal recorded per 3-m layer, with an inbuilt algorithm. In the default-operating mode, 12 samples collected at  $\sim$ 2 h intervals, are transmitted daily via satellite by the buoy.

#### Data cleaning process

The raw data provided by the echosounder buoys were cleaned using the following protocol (see Baidai *et al.*, 2017 for details on the procedure):

- Duplicated rows, inconsistent positions, data recorded on land, at shallow positions (depth <150 m), or under low voltage conditions (poor reliability of acoustic data collected below 11.5V) were omitted from the database.
- (2) A rule-based algorithm, which uses buoy speed and its variations as main classifiers, was applied to discriminate acoustic data recorded when the buoy is on board a vessel from those actually recorded when the buoy is deployed at sea.

#### Classification of tuna presence/absence

The presence or absence of tuna at a DFAD was assessed from acoustic data collected by echosounder buoys, following the methodology described in Baidai *et al.* (2020). This approach involves preliminary processing of the acoustic data, followed by a classification using random forest algorithms applied separately to each ocean. The learning datasets were constructed by crossreferencing the acoustic data with the activities of fishers at DFADs (namely sets with associated tuna catches, DFAD deployments, and DFAD visits) recorded in logbooks and by on-board observers. A detailed description of the classification procedure can be found in Supplementary Material S1. The minimum catch value representing tuna presence in the learning dataset was 1 ton. Thus, in this work, the term "tuna aggregation" refers to a fish aggregation whose tuna biomass is at least equal to this value.

An additional post-processing step was applied to improve the predictions made by the classification algorithm during the course of DFAD trajectories. Prediction results with short durations (i.e. an isolated event of presence or absence lasting a single day) were considered unlikely, attributed rather to misclassification, and corrected using the previous or following day's predicted values. This step allowed for the correction of 9 and 7% of the initial predictions made by the classification model, in the AO and the IO, respectively.

#### Newly deployed DFADs

Only newly deployed DFADs (i.e. DFADs used for the very first time) equipped with echosounder buoys were considered. Natural floating objects (e.g. logs), reinforced old DFADs found at sea, relocated DFAD and buoy transfers were all excluded. Deployments of new DFADs were identified from fishing

20 10 Latitude 0 -10 50 40 45 55 -20 -30 -20 10 40 70 100 110 120 -40 -30 -10 20 30 50 60 80 90 -50Longitude Daily buoy detection: 0 Tuna absence Tuna presence

**Figure 1.** Presence/absence of tuna aggregations along the course of the trajectories of newly deployed DFADS monitored in Western Indian Ocean and Eastern Atlantic Ocean from 2016 to 2018. Orange dots indicate days when tuna aggregations were present, white dots represent days with no tuna aggregations.

logbooks and observer data collected from 2016 to 2018 in the AO and IO. The 2016–2018 period was selected to provide a relatively homogeneous study period, while maintaining sufficient data in both oceans. Observer data were collected by the IRD-Ob7 observatory under the EU Data Collection Framework and the French OCUP program (Observateur Commun Unique et Permanent). Trajectories and time series of tuna presence/absence associated with newly deployed DFADs were then identified by cross-referencing the logbook and observer databases with the echosounder buoy database, using the unique identification code of the buoy and the deployment date. To ensure that the subset correctly identified newly deployed DFADs (and not potentially misreported reinforcement activities), records for which fishing sets were reported during the week following the deployment were removed from the dataset. An additional cleaning step was applied to the dataset to omit data with inconsistent positions between the location of the DFAD deployment recorded in the logbook or observer database, and the actual position recorded by the buoy (0.3 and 0.8% of the dataset in AO and IO, respectively). The cleaned dataset of newly deployed DFADs included 9118 trajectories with 498 276 presence/absence data points for the IO and 285 trajectories with 18 102 presence/absence data points for the AO (Figure 1).

#### Soak time and colonization time

Soak time was defined as the number of days between the deployment of a DFAD equipped with a buoy and the first reported operation on it (i.e. either a fishing set or the retrieval of the buoy). Tuna colonization time refers to the number of days between the deployment of a DFAD and the first day when a tuna aggregation is detected by the echosounder buoy (Figure 2). The term "colonized DFAD" thus refers to a DFAD that has aggregated tuna at least once (for longer than 1 day).

Due to fishing and buoy retrievals, the number of buoys at sea available for the analysis declined for increasing soak times. This can induce bias in the estimate of tuna colonization times obtained from simple averages (Figure 3). Specifically, the lower the number of DFADs with long soak times, the lower the chances of observing long colonization times, which leads to an underestimate of colonization times from simple arithmetic averages. To overcome this bias, colonization times were estimated from daily colonization rates, considering the daily fraction  $r_i$  of colonized DFADs relative to the total available DFADs, for each day *i* after deployment, see [Equation (1)]:

$$r_i = \frac{N_{\text{colonized}_i}}{N_{\text{colonized}_i} + N_{\text{uncolonized}_i}},$$
(1)

where, Ncolonized<sub>i</sub> indicates the number of DFADs colonized during day *i* after deployment and Nuncolonized<sub>i</sub> denote not-yetcolonized DFADs on day *i* after deployment. The denominator of [Equation (1)] corresponds to the total number of DFADs available for colonization on day *i*, namely, the total number of DFADs in the water that at day i-1 after deployment were not yet colonized.

Mann–Whitney *U* tests were used to compare daily colonization rates between the IO and the AO. The unbiased mean colonization time  $(T_{col})$  (in days) was then estimated as the inverse value of the average of daily colonization rates (-r):

$$T_{\rm col} = 1 - r \tag{2}$$

where -r denotes the average daily colonization rate:

$$-r = \frac{1}{D} \sum_{i=1}^{D} r_i \tag{3}$$

where *D* represents the total number of days during which the daily colonization rates  $r_i$  were calculated. When numbers of available DFADs were too low [i.e. the denominator in Equation (1)], the daily colonization rate becomes less reliable. A preliminary sensitivity analysis, included in the Supplementary Figure S1.2, showed that *D* corresponds to the number of days after



**Figure 2.** Schematic representation of the timeline of tuna aggregation dynamics at a DFAD. The term "end of trajectory" denotes here the first operation carried out on FADs likely to disturb the aggregation below (e.g. either a fishing set or the retrieval of the buoy).



Figure 3. Daily colonization rates (bars) and percentage of equipped DFADs available (solid lines) over time in the AO (a) and the IO (b). Red dashed lines indicate the number of days at which 30 DFADs were still available.

deployment when at least 30 DFADs remained available for colonization.

#### Aggregation stability

The continuous residence time (CRT) is commonly used to represent the amount of time spent by acoustically tagged individual tunas around a FAD without a day scale (>24 h) absence (Ohta and Kakuma, 2005; Capello *et al.*, 2015). Alternatively, the continuous absence time (CAT) refers to the time interval between two consecutive associations for an individual tuna (Robert *et al.*, 2012; Rodriguez-Tress *et al.*, 2017). In this work, the concepts of CRT and CAT were adapted and applied to DFAD aggregations rather than to individual fish. Accordingly, the *aggregation's continuous residence time at a floating object* (FOB-aCRT) was considered as the time span within which a tuna aggregation was continuously detected at a DFAD without a day scale (>24 h) absence. Similarly, the *continuous absence time of aggregation at a floating object* (FOB-aCAT) was defined as the period between

two consecutive detections of tuna aggregations at the same DFAD. Values occurring directly before an operation on the DFAD (fishing event or retrieval of the buoy) were excluded from the analysis as they were artificially truncated. Finally, the overall proportion of time that a tuna aggregation remained at a colonized DFAD (named *DFAD occupancy rate*), expressed as the ratio of the sum of all FOB-aCRTs against its soak time after the colonization period, was assessed and compared between oceans using Mann–Whitney *U* tests.

#### Survival analyses of FOB-aCRT and FOB-aCAT

Survival analyses (Capello *et al.*, 2015) were used to characterize the distribution of FOB-aCATs and FOB-aCRTs. Survival curves were constructed using the fraction of FOB-aCATs and FOBaCRTs shorter than a given time, and compared between oceans using the logrank statistical test (Harrington and Fleming, 1982), implemented in the "*survival*" package in R (Therneau, 2015).

**Table 1.** Summary of tuna aggregation metrics measured in the AO.

	Min.	Max.	Median	Mean	SD
DFAD soak time (days)	1	305	44	63.28	65.08
Daily colonization rate (days <sup>-1</sup> )	0	0.15	0.06	0.06	0.04
Tuna colonization time (days)	-	-	-	16.10	9.66
FOB-aCAT (days)	2	86	4	5.38	6.01
FOB-aCRT (days)	2	96	4	8.96	11.52
Occupancy rate (%)	5.13	97.59	60.49	63.27	19.86

Table 2. Summary of tuna aggregations metrics measured in the IO.

	Min.	Max.	Median	Mean	SD
DFAD soak time	1	363	43	54.24	45.52
(days)					
Daily colonization rate	0	0.07	0.02	0.02	0.01
$(days^{-1})$					
Tuna colonization	-	-	-	40.46	17.31
time (days)					
FOB-aCAT (days)	2	119	5	8.84	10.93
FOB-aCRT (days)	2	109	4	6.20	6.86
Occupancy rate (%)	2.83	98.08	46.16	45.45	21.73

Survival curves were also fitted using three models: (i) single exponential, (ii) double exponential, and (iii) power law (Supplementary Table S1.3), by adapting the methodology of Robert *et al.* (2013) to the DFAD aggregation metrics. Exponential models assume association dynamics (presence or absence of an aggregation at a DFAD) to be independent of time. Double exponential models imply the existence of two distinct time-scales occurring within aggregation presence or absence at a DFAD. Power law models indicates a time-dependent probability of presence and absence of tuna aggregations, meaning the longer the time a DFAD is occupied or vacant, the smaller the probability that a change in state will occur. Models were discarded if one or more parameters were not significant (p > 0.05 based on the *t*-statistics). The best-fitting models were chosen based on the Akaike Information Criterion (AIC) and q-q plots.

#### Results

#### Daily colonization rates and colonization times

No significant difference was found between DFAD soak times from the AO and the IO (Mann–Whitney *U* tests, p = 0.76) with mean values of 63.28 days (*SD* 65.08 days), and 54.24 days (*SD* 45.52 days), respectively. Approximately 22% DFADs in the AO (62 DFADs) and 34% (3122 DFADs) in the IO did not show any sign of colonization by tunas during their soak time. The soak time of vacant DFADs (averages of 18.66 and 28.52 days for AO and IO, respectively) was significantly lower than that of colonized DFADs (averages of 75.68 and 67.63 for AO and IO, respectively), with a *p*-values (Mann–Whitney *U* tests) lower than 0.001 in both oceans (Supplementary Figure S1.4).

For colonized DFADs, the time before the echosounder buoy detected the first aggregation of tunas averaged 13.17 days (*SD* 12.37 days) in the AO and 20.22 days (*SD* 20.83 days) in the IO. Stable trends in daily colonization rates were observed in both oceans (see Figure 3). The average daily colonization rates were significantly higher (Mann–Whitney *U* test, p < 0.001) in the AO (-r = 0.062, *SD* 0.037) than in the IO (-r = 0.025, *SD* 0.011). Calculating the unbiased average colonization times following Equations (1)–(3) resulted in colonization times that were 2.5 times shorter in the AO ( $T_{col} = 16.10$  days, *SD* 9.66 days—see Table 1) than in the IO ( $T_{col} = 40.46$  days, *SD* 17.31 days—see Table 2).

# Aggregation continuous residence (FOB-aCRT) and absence times (FOB-aCAT)

A total of 15 415 FOB-aCRTs and 13 328 FOB-aCATs events were recorded during the course of the trajectories of newly deployed DFADs in the IO. In the AO, 723 FOB-aCATs and 779

FOB-aCRTs were recorded. Distributions of FOB-aCATs and FOB-aCRTs in both oceans are shown in Figure 4. The average duration of tuna aggregations was 8.96 days (SD 11.52) around DFADs in the AO and 6.20 days (SD 6.86) in the IO. It should be noted that very long CRTs of tuna aggregations under the same DFAD were also observed in both oceans (96 and 109 days, in the AO and IO, respectively). The average time that DFADs remained vacant between two consecutive tuna aggregations (FOB-aCAT), was 5.38 days (SD 6.01 days), with a maximum duration of 86 days in the AO, and at 8.84 days (SD 10.93 days), with a maximum of 119 days in the IO (Tables 1 and 2).

Inter-ocean comparisons of FOB-aCAT and FOB-aCRT survival curves indicated significant differences in the associative dynamics of tuna aggregations (logrank test, p < 0.001 for both FOB-aCAT and FOB-aCRT ocean-comparisons—Figure 5).

In the IO double exponential models were the best fitting for survival curves of both FOB-aCATs and FOB-aCRT. Short-term residences represented 94% of the FOB-aCRTs with a mean duration of 4.58 days, while long-term residences represented 6% with a mean duration of 20.18 days. Short-term absences lasted an average of 4.43 days (representing 66% of FOB-aCATs), while longterm absences had a mean duration of 15.45 days (34% of FOBaCATs).

In the AO, a double exponential model was the best fit for the survival curve of FOB-aCRTs with averages of 3.75 days (62% of FOB-aCRTs) and 15.70 days (38% of FOB-aCRTs) for short- and long-term residence times, respectively. Conversely, a single exponential model was the best fit for absence times of tuna aggregations at DFADs with a mean duration of 4.30 days (see Table 3 and Figure 6).

#### DFAD occupancy rate

Significant differences in the proportion of time that colonized DFADs were occupied by tuna were observed between the two oceans (Mann–Whitney *U* tests p < 0.001). After colonizing DFADs, in the AO tuna aggregations were detected for an average of 63.27% (*SD* 19.86%) of the soak time (Figure 7a), while in the IO, this figure was 45.45% (*SD* 21.73%) (Figure 7b).

#### Discussion

This work aimed at characterizing the dynamics of the tuna aggregation processes around DFADs, using acoustic data collected by commercial echosounder buoys on newly deployed DFADs in the IO and AO. To date, very few studies have designed scientific protocols to quantify the time that pelagic species take to colonize newly deployed DFADs. The only previous documented observations in the AO come from Bard *et al.* (1985), who reported rapid



Figure 4. Distribution of FOB-aCATs (left) and FOB-aCRTs (right) in the AO (top) and IO (bottom). FOB-aCRT and FOB-aCAT denote the aggregation's continuous residence time at a floating object and the continuous absence time of aggregation at a floating object, respectively.



Ocean: - Atlantic - Indian

**Figure 5.** Survival curves of FOB-aCRTs (a) and FOB-aCATs (b) recorded on trajectories of newly deployed DFADs in AO (black dots) and IO (white dots). The *y*-axis is on a logarithmic scale. FOB-aCRT and FOB-aCAT denote for the aggregation's continuous residence time at a floating object and the continuous absence time of aggregation at a floating object, respectively.

colonization by tunas, ranging from 1 h to 6 days, through the monitoring of a dozen newly deployed DFADs and detecting tuna presence by visual observations, on-board echosounders, or by fishing sets. Their estimates are significantly shorter than ours for the same ocean (average of 16 days); however, interpretation of these discrepancies is complicated by the large differences in methods and the time when the studies were conducted. The Bard *et al.* study was performed before the development of the FAD-fishery, when tropical tuna stocks were only moderately exploited. Furthermore, their observation protocol could not identify whether observed individuals only visited or remained associated with the DFAD. Taquet *et al.* (2007) observed that dolphinfish (*Coryphaena hippurus*) could arrive a few hours after the deployment of a new floating object, but did not necessarily

associate with it. In the IO, using Local Ecological Knowledge, Moreno *et al.* (2007b) suggested that it typically takes 1 month before tunas aggregate under a newly deployed DFAD. Although aggregation dynamics at anchored FADs may differ from those at DFADs, it is worth noting that Macusi *et al.* (2017) reported that fishers typically wait ~22 days for tuna aggregations to form at anchored FADs in the Philippines, based on interview data. In a recent study using echosounder buoys produced by a different manufacturer, Orue *et al.* (2019b) examined acoustic data from over 900 newly deployed DFADs in the Western Indian Ocean and suggested that tunas begin to aggregate ~13 days after deployment. At three times longer, the findings of this study (an average of ~40 days in the IO) appear to be more aligned with the knowledge of purse seine skippers (Moreno *et al.*, 2007b). The

Ocean	Metric	Fitting law	Parameter	Estimate	SE	t-Value	Pr(>  <i>t</i>	)	AIC
AO	FOB-aCRT	Single exponential	a	0.14	3.61E-03	38.35	6.92E-42	***	-217.80
		Double exponential	а	0.27	8.51E-03	31.29	2.75E-36	***	-416.09
			Ь	0.06	2.13E-03	29.95	2.58E-35	***	
			р	0.62	1.89E-02	33.03	1.71E-37	***	
		Power law	а	2.29	7.19E-02	31.87	3.87E-37	***	-389.71
			b	11.73	5.11E-01	22.94	7.76E-30	***	
	FOB-aCAT	Single exponential	а	0.23	5.03E-03	46.16	7.42E-31	***	-154.98
		Double exponential	а	0.24	1.24E-02	19.59	1.20E-18	***	-155.22
			b	0.03	6.18E-02	0.43	6.68E-01		
			р	0.98	3.24E-02	30.26	4.86E-24	***	
		Power law	а	22.04	2.51E+01	0.88	3.87E-01		-153.74
			b	91.78	1.08E+02	0.85	4.03E-01		
Ю	FOB-aCRT	Single exponential	а	0.20	2.84E-03	69.54	3.63E-69	***	-404.29
		Double exponential	а	0.22	8.71E-03	25.04	2.85E-37	***	-426.38
			b	0.05	1.94E-02	2.56	1.27E-02	*	
			р	0.94	3.37E-02	27.90	2.42E-40	***	
		Power law	а	8.49	2.17E+00	3.92	2.00E-04	***	-415.36
			b	39.48	1.10E+01	3.58	6.24E-04	*	
	FOB-aCAT	Single exponential	a	0.14	2.12E-03	64.29	1.01E-80	***	-471.48
		Double exponential	а	0.23	3.10E-03	72.73	1.98E-84	***	-875.29
			b	0.06	1.11E-03	58.30	1.34E-75	***	
			р	0.66	1.02E-02	64.71	9.41E-80	***	
		Power law	а	3.06	5.51E-02	55.51	3.31E-74	***	-806.27
			b	17.36	4.04E-01	43.01	4.41E-64	***	

**Table 3.** Summary of the model fits of the survival curves of aggregation continuous residence and absence times (FOB-aCRTs and FOB-aCATs) obtained in AO and IO.

AIC values of the best-fitted models are highlighted in bold.

Significance codes: \*\*\*0.001; \*0.01;

AIC, Akaike Information Criterion; Pr(>|t|), *p*-value at *t*-tests; *t*-value, value of *t*-statistic.



**Figure 6.** Survival curves of FOB-aCAT (left) and FOB-aCRT (right) the observed in AO (top) and IO (bottom) fitted with single exponential, double exponential, and power law models. The red line indicates the best fit. FOB-aCRT and FOB-aCAT denote the aggregation's continuous residence time at a floating object and the continuous absence time of aggregation at a floating object, respectively.

discrepancy between these acoustic studies may be related to (i) differences in methodological approaches applied in the conversion of acoustic data into indicators of presence or absence of tuna, (ii) the method used in this study to estimate colonization

times, which avoids possible underestimation biases linked to the large variability in DFAD soak times, or (iii) from the differences in the specificities of the buoy models used in each study. Since their introduction into the fishery, echosounder buoys have



Figure 7. Distribution of the percentage of the DFAD soak time during which tuna aggregations were detected in the AO (a) and the IO (b).

evolved rapidly, through continuous technological innovations in both hardware and software (Lopez *et al.*, 2014). Thus, the intrinsic performance of buoys for detecting tuna aggregations may differ by model and/or manufacturer. Hardware and software differences in the design of buoys may lead to variable thresholds for the detection of aggregations, which could ultimately result in biases in the detection of small aggregations for some models. Such disparities highlight the critical need for a detailed assessment of the reliability of outputs from the different models of buoys and the accuracy of the data processing methods they use to estimate fish abundance. This is especially important when considering the growing use of echosounder buoy data for scientific purposes (Moreno *et al.*, 2016).

Until now, most scientific knowledge on the behaviour of tunas around floating objects stems from observations of individuals, using electronic tags, with the majority of studies focused on anchored FADs (Holland et al., 1990; Ohta and Kakuma, 2005; Dagorn et al., 2007a; Mitsunaga et al., 2012; Robert et al., 2013; Rodriguez-Tress et al., 2017). By exploiting the potentially massive data source that echosounder buoys on DFADs represent, this work introduces two novel metrics (FOB-aCAT and FOBaCRT), providing descriptive elements of DFAD use by entire tuna aggregations. It is particularly interesting to note that, in both oceans, the time taken for tunas to colonize new DFADs was significantly longer than durations between consecutive tuna aggregations (average FOB-aCATs: 9 and 6 days in AO and IO, respectively). This result is consistent with previous assertions regarding the role of non-tuna species in the tuna colonization process. Several authors have suggested that the colonization of FADs is a sequential process starting with the arrival of non-tuna species, which may play a key role in the attraction or retention processes of tunas (Deudero et al., 1999; Castro et al., 2001; Nelson, 2003; Moreno et al., 2007b; Taquet et al., 2007; Macusi et al., 2017). The duration of the settlement stage of these pioneer communities could be one of the major factors driving the colonization time of tunas at a new DFAD. As such, colonization time may be viewed as a unique type FOB-aCAT with an extended duration due to the requisite maturation phase of the DFAD. Further studies on interspecific relationships would be of major benefit for improving our understanding of the role played by non-tuna species in the aggregative processes of tunas with DFADs.

A review of the main findings from electronic tagging studies on the associative behaviour of tunas under DFADs reveals that the CRT of individual tuna (CRT) is subject to a degree of variability related to the species or the oceanic region under consideration (see Table 4). Off the coast of Guinea in the AO, Tolotti et al. (2020) estimated average CRT values of 9 and 19 days, for skipjack and yellowfin tuna, respectively. For bigeye tuna the reported CRTs were up to 25 days, which is longer than observations from other oceans. Shorter CRTs ( $\sim$ 1 day on average) were observed for the same three species by Dagorn et al. (2007b) in the Western Indian Ocean. However, these results are likely to be underestimated due to artificial truncation of the observation experiments. Govinden et al. (2010) reported residence times ranging from 4 to 10 days (median values) depending on the tuna species, at DFADs monitored in the Mozambique Channel. In Eastern Pacific Ocean, studies carried out by Matsumoto et al. (2014) and Matsumoto et al. (2016), both indicated that individual tunas remain associated with DFADs for <7 days. Despite this variability, tuna CRTs reported by this limited number of tagging studies appear to be lower or equal to the average FOB-aCRTs obtained in this work (9 and 6 days in the AO and the IO, respectively), especially for skipjack tuna, which is the dominant species in DFAD-associated catches (Dagorn et al., 2013).

Survival analyses of the FOB-aCRTs indicated the coexistence of two distinct modes of DFAD association by tuna aggregations: a dominant mode consisting of short durations, and a longer residence mode. Nearly all of the FOB-aCRTs measured in the IO belonged to the short-term residence mode, whereas the two modes occurred in more similar proportions in the AO. There are several possible explanations for the occurrence of these different modes and their inter-ocean variation. Individually, bigeye and yellowfin tuna generally exhibit longer residence times than skipjack tuna, as indicated by the tagging studies mentioned above. Long-term residence modes may therefore reflect aggregations with a large proportion of the two former species. Furthermore, this study was conducted at a broad spatial and temporal scale. As such, it is possible that the observed differences in modes could be a result of behavioural patterns of tuna that are driven by local environmental differences (such as prey or conspecific abundance, or densities of floating objects) between seasons or oceanic regions. The long-term residence mode could also be indicative of the occurrence of turnover processes of

Table 4. Summary of main findings from previous studies on CRT of individual tunas at drifting FADs.

Study	Location	Species	FL range (cm)	CRT
Dagorn et al. (2007b)	Western Indian Ocean	SKJ	Not provided	Average at 0.91 days (maximum: 7.03 days)
		BET		Average at 1.43 days (maximum: 3.06 days)
		YFT		Average at 1.04 days (maximum: 15.22 days)
Govinden et al. (2010)	Mozambique Chanel (Western Indian Ocean)	SKJ	47–57	Median at 4.47 days (maximum: 18.33 days)
		BET	54-56	Median at 3.89 days (maximum: 6.56 days)
		YFT	29–60	Median at 9.98 days (maximum: 26.72 days)
Matsumoto et al. (2014)	Equatorial central Pacific Ocean	SKJ	36-65	Average at 2.3 days (maximum: 6.4 days)
Matsumoto et al., (2016)	Equatorial central Pacific Ocean	SKJ	34.5-65.0	Average at 1.3 days
		BET	33.5-85.5	Average at 3.8 days (maximum: $\sim$ 11 days)
		YFT	31.6-93.5	Average at 4.1 days (maximum 14.5 days)
Scutt Phillips et al. (2019)	Western Central Pacific Ocean	SKJ	46–60	Median at 1 day (maximum: 18 days)
		BET	37–90	Median at 10 days (maximum: 30 days)
		YFT	36-98	Median at 2 days (maximum: 50 days)
Tolotti <i>et al</i> . (2020)	Eastern Atlantic Ocean	SKJ	39–61	Average at 9.19 days (maximum value to 15 days)
		BET	45–61	Average at 25.31 days (maximum value to 55 days)
		YFT	34-82	Average at 19.15 days (maximum value to 55 days)

BET, Thunnus obesus; FL, fork length; SKJ, Katsuwonus pelamis; YFT, Thunnus albacares.

schools at the same DFADs as reported by Weng *et al.* (2013). Further spatially constrained analyses combined with electronic tagging studies, conducted on DFADs equipped with echosounder buoys, will be crucial to relate the individual and the collective dynamics of tuna around DFADs.

The associative behaviour of the tuna population implies that, at any given time, the overall abundance in an area is the sum of the abundance of two permanently interacting components: the associated and the free-swimming (or unassociated) populations. At present, the underlying reasons driving the association or departure of tunas from floating objects remain unclear. Nevertheless, an improved understanding of the interactions between the two population components can be achieved through the study of the relationships between the association metrics assessed at the scale of the individual (i.e. CAT and CRT) and at the scale of aggregations (FOB-aCAT and FOB-aCRT). Rodriguez-Tress et al. (2017) suggested that high FAD densities tend to reduce the time that tuna spend in an un-associated state (CAT). While these findings may need to be interpreted with caution as they stem from observations at anchored FADs, this could suggest that the underlying trend may occurs irrespective of the FAD type. Logically, higher FAD densities would increase the probability of an individual encountering and associating with a FAD, hence reducing the time individual tuna spend in a freeswimming state (CAT). Similarly, FAD vacancy (FOB-aCATs) should be related to the abundance of the un-associated tuna population. Long FOB-aCATs would result when the unassociated population is small, either due to a low overall tuna population or a large density of FADs drawing them in to aggregate. Following this reasoning, the longer FOB-aCAT and colonization time observed in the IO may thus be indicative of a smaller size of the tuna population and/or higher densities of DFADs in this ocean than in the AO. Furthermore, the double exponential curve for FOB-aCATs observed in the IO, could be a result of regions/periods where at least one of these two factors differ. Previous work by Capello et al. (2016) demonstrated that indicators of abundance for tropical tuna populations could be derived from their individual associative dynamics (CRT and CAT). Combining our current understanding of the individual associative behaviour of tunas with the metrics describing tuna aggregation dynamics provided by this work could aid in developing new methods for obtaining direct abundance estimates of tuna populations. Such methods will depend on the availability of estimates of the total number of DFADs at sea (more specifically the total number of floating objects). Currently, obtaining these statistics is a challenge in all oceans despite the recent data reporting requirements for DFAD activities by Tuna Regional Fisheries Management Organisations.

#### Conclusion

Using data from the echosounder buoys of French purse seiner fleets, this study characterized key parameters of tuna aggregations at DFADs: colonization times, aggregation lifetimes, and time span between aggregations. In both oceans, lifespan of tuna aggregations at DFAD followed a time-independent process with two modes. This suggests that the species composition and/or the local conditions (e.g. prey, conspecifics, or density of floating objects) could play key roles in aggregation dynamics. However, opposing trends also existed between the two oceans, with shorter residence time of aggregations and longer periods of DFAD vacancy in the IO than in the AO. Further spatially restricted analyses assessing these behavioural metrics at smaller spatial and temporal scales could help in understanding the dynamics of aggregations at a local scale, as well as the role played by various environmental factors. The integration of these new findings into population assessment models which account for the associative behaviour of tunas present an opportunity for the development of alternative abundance indices (independent from catch and effort data) for tropical tunas and the construction of reliable scenarios on the impacts that DFADs have on tuna populations.

### Data availability statement

The data underlying this article were provided by Ob7— "Observatoire des Ecosystèmes Pélagiques Tropicaux exploités" from IRD/MARBEC under data exchange agreement with fishing companies. Data will be shared on request to Ob7 (admdblp@ird.fr) with permission of the owners.

# Supplementary data

Supplementary material is available at the *ICESJMS* online version of the manuscript.

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