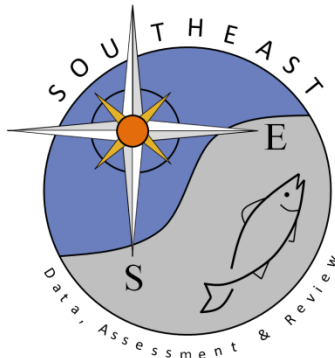


An integrated approach for assessing the survival of discarded sandbar sharks, *Carcharhinus plumbeus*, captured in scientific longlines

Taylor Grosse^{A,B,*}, Cynthia Awruch^{C,D}, Euan Harvey^B, Chris Dowling^A, Daniela Waltrick^A, Benjamin Saunders^B and Matias Braccini^A

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ABSTRACT

Context. The sandbar shark (*Carcharhinus plumbeus*) has a global distribution and is caught by commercial fishers and recreational anglers. **Aims.** To assess the stress physiology, release condition, and post-release survival of sandbar sharks caught in longline surveys conducted in Western Australia. **Methods.** Post-release survival of sandbar sharks caught in longlining surveys was assessed using an integrated approach that combined the use of hook-timers, qualitative release conditions, satellite-tagging, and blood physiology. **Key results.** Of 57 individuals examined, there was 100% post-capture survival after a maximum of 4 h on the hook. Most of these animals (88%) displayed a strong release condition, exhibiting minimal behavioural impairment. All 13 satellite-tagged individuals survived 30 days post-capture. Sharks dived up to 307 m deep and showed cyclical depth movement patterns, with some individuals moving through the water column both day and night, whereas others moved almost exclusively at night. The concentration of blood metabolites did not significantly change with time-on-hook. **Conclusion.** Post-capture and post-release survival of 100% after up to 4 h on hooks suggested that the use of longlines for surveying sandbar shark abundance had no deleterious effects on captured sharks. **Implication.** This will support future stock assessments of sharks by quantifying the survival rates in the methods used for long-term monitoring of sandbar shark populations.

Keywords: Chondrichthyes, fisheries, post-capture survival, post-release survival, PSAT tags, release condition, stress physiology, survivorship.

Introduction

Elasmobranchs are one of the most vulnerable vertebrate groups living in our oceans. An increasing number of elasmobranchs is being classified as either Endangered or Threatened by the International Union for Conservation of Nature Red List because of the life-history characteristics (slow growth rates, late age at maturity, low fecundity, and long life span) that make the group vulnerable to overfishing (Clementi *et al.* 2021; Dulvy *et al.* 2021; Pacoureaux *et al.* 2021). The economic value of fins and other elasmobranch by-products has resulted in an increased fishing pressure worldwide (Clarke *et al.* 2006, 2007; Cardeñosa *et al.* 2020). Sharks are directly targeted for the fin trade and 'flake' consumption, but are caught and discarded by fishers targeting other species (Clarke *et al.* 2007; Oliver *et al.* 2015; Dulvy *et al.* 2021).

Empirical estimates of discard mortality are often unavailable when estimating commercial or recreational fishing mortality, and the fate of discarded sharks is largely unknown (Clarke *et al.* 2006; Davidson *et al.* 2016; Braccini *et al.* 2021). This information is key for the accurate assessment of stocks and the development of effective fisheries management and conservation policies (Braccini *et al.* 2021). Different approaches have been used to assess the fate of discarded sharks on release, either by quantifying post-capture survival (PCS; survival observed at the vessel) or post-release survival (PRS; survival after discarding). For example, changes in blood chemistry (e.g. lactate, glucose,

pH, urea) can be used to estimate the likelihood of discard survival (Dapp et al. 2016), whereas methodologies such as cages to monitor survival (Rulifson 2007), tag-recapture (Hueter et al. 2006), and electronic tracking (Kneebone et al. 2013) are also used (Braccini et al. 2012; Ellis et al. 2017). However, estimates of both PCS and PRS are needed to quantify the overall mortality of the captured sharks (Braccini et al. 2012; French et al. 2015; Hutchinson et al. 2015; Hutchinson and Bigelow 2019).

Several blood-physiology indicators have been used to quantify how sharks respond to the catch-and-release. In recent years, adrenocorticotrophic hormone (ACTH) concentrations have been used to evaluate the role of ACTH, a pituitary hormone responsible for stimulating the release of glucocorticoids (GCs) associated with physical stress events. However, these studies are very limited in sharks and the roles of ACTH are not entirely understood concerning capture stress (Fuller et al. 2020). Although 1α -hydroxycorticosterone (1α -OH-B) is the primary internal corticosteroid produced by elasmobranchs (Anderson 2012) to help the body respond to physical stress (Armour et al. 1993; Anderson 2012; Ruiz-Jarabo et al. 2019; Schoen et al. 2021), corticosterone (CORT) has also been linked to initial stress-coping mechanisms in elasmobranchs (Iki et al. 2020) and has been previously used to assess the stress response in conjunction with other metabolites (Manire et al. 2007; Brinn et al. 2012). However, CORT can be separately expressed with maturity and reproductive state, which should be considered when evaluating concentrations (Rasmussen and Crow 1993; Snelson et al. 1997; Manire et al. 2007). These initial endocrine coping mechanisms to stress trigger a range of secondary stress responses such as lactate, glucose, and hydromineral balance (Karsten 2000; Heberer et al. 2010; Hammerschlag et al. 2017).

Satellite tagging, such as with pop-up satellite archival transmitting (PSAT) tags, is an effective method for assessing PRS (Mohan et al. 2020, Schaefer et al. 2021). However, the cost of satellite tags limits the number of individuals tagged (Kohler and Turner 2001; French et al. 2015). The PSAT tag transmits light, temperature, and depth data in a range of temporal frequencies that can be altered to suit the species of shark studied (Lynch et al. 2017; Sulikowski et al. 2020). Many charismatic species have been monitored using this style of tag, with varying results in PRS and spatiotemporal movements (French et al. 2015; Hutchinson et al. 2015; Hutchinson and Bigelow 2019; Sulikowski et al. 2020).

Sandbar sharks (*Carcharhinus plumbeus*) have a global distribution across temperate and tropical waters and are captured in commercial and recreational fisheries worldwide (McAuley et al. 2005). Historically, commercial shark fisheries in northern Western Australia (WA) targeted sandbar sharks, with a peak of >700 tonnes (Mg) landed in 2004–2005 (McAuley et al. 2005; Braccini et al. 2020). Annual scientific shark surveys in northern WA have been conducted for >20 years (commencing in 2001) to monitor sandbar shark

abundance (Braccini et al. 2020). Since large spatial closures were implemented in 2005 and the cessation of operations in the remaining northern shark fisheries, these scientific surveys indicate that there has been an increase in the abundance of sandbar sharks along the north-western coast (stock statuses: over-exploited in 2005 and recovering in 2021) (Braccini et al. 2020; Newman et al. 2021). Sandbar sharks are routinely tagged-and-released as part of these surveys, but there are limited reports of recaptured individuals from either recreational or commercial fisheries, and little is known about the fate of tagged sandbar sharks on release in the survey area.

In the USA, sandbar sharks caught from demersal longlines exhibited variable PRS (29–96%) owing to differences in gear specifications, shark size, soak times and fishing depth (Marshall et al. 2015; Gulak and Carlson 2021; Whitney et al. 2021). Therefore, results from previous studies may not be representative of the PRS of individuals discarded from all line fisheries (Marshall et al. 2015; Gulak and Carlson 2021). Hence, further research on the fate of sandbar sharks tagged-and-released from longlines, including experiment survey gear, is required for a more thorough understanding of its potential deleterious effects, particularly here in WA, with a recovering population. The scope of this study was to assess whether experimental surveys influence the recovering population, considering that other forms of commercial fishing in the region may have less influence (i.e. owing to shark fishing regulations and cessation of the northern shark fisheries). We took an integrated approach using qualitative release condition, PSAT tagging, and analysis of blood physiological parameters (i.e. ACTH, GC and lactate) to assess the likelihood of survival of sandbar sharks caught on demersal longline surveys.

Materials and methods

Onboard sampling

Scientific demersal longline surveys were conducted in 2020 and 2021 at 10 fixed stations along the continental shelf of northern WA (Fig. 1).

Approximately 250 hooks were deployed each day. Each deployment comprised three to five longlines of ~500 m long each. Size 12/0 J-shaped hooks baited with sea mullet (*Mugil cephalus*) (cut into two or three pieces, ~20 cm) were attached to ~2-m metal snoods to the mainline. For further details on gear configuration and deployment, refer to Braccini et al. (2020). To determine time on hook (TOH, i.e. time spent on the line after hooking), hook timers (Lindgren-Pitman Inc.) were randomly attached to 16 snoods on each deployment. The start and end latitude and longitude, depth, water temperature and time of day were recorded for each longline deployment. Thirty-nine longline

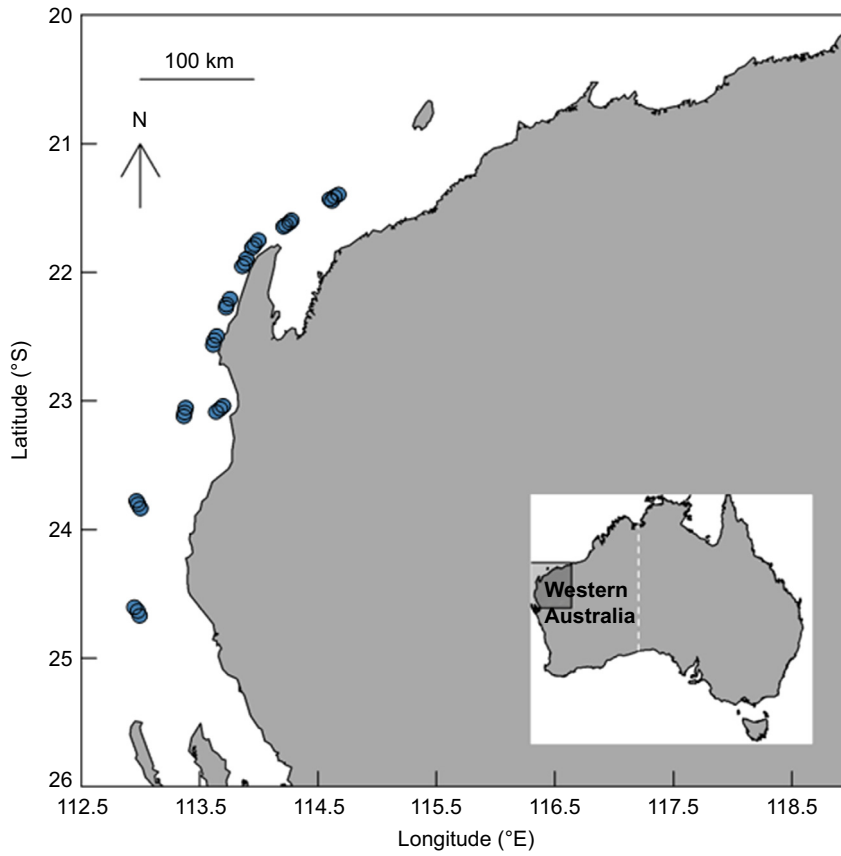


Fig. 1. Map of demersal longline (blue) survey area in northern Western Australia. Each site has five replica longline sets.

sets were conducted over 10 sites with soak times ranging from 3 to 4 h.

On capture, sandbar sharks were brought on deck and the hook was removed. The TOH was recorded, and blood samples (5 mL) were taken immediately from the caudal vein in 57 individuals randomly selected for PCS analysis by studying blood and plasma physiology, of which this process took ~1 min. Sharks were then measured (fork length, FL, cm) and sexed. Whole-blood lactate concentrations were measured *in situ* in all individuals, and the whole-blood samples from 29 of these sharks were centrifuged for 5 min at 7871g and room temperature, and plasma was separated and stored at -20°C for further analysis of ACTH and GC concentrations. Thirteen of the sharks sampled for further blood and plasma physiology were opportunistically fitted (with considerations of distributions across TOH) with survivorship pop-up satellite archival transmitting (PSAT) tags (Model sPAT, Wildlife Computers Inc., Redmond, WA, USA) to quantify PRS. Gills were not irrigated during handling to mimic commercial fishing operations. The PSAT tags were tethered to a nylon intramuscular anchor inserted at the base of the dorsal fin. All sharks captured were tagged with a plastic fin tag and released. At the time of release, the condition of each individual was classified as (1) alive and strong, (2) alive, but weak or disoriented, (3) moribund (eye

and jaw response when stimulated, otherwise exhausted and unresponsive) or (0) dead (Braccini and Waltrick 2019).

Post-capture survival (PCS) – blood and plasma physiology

Adrenocorticotrophic hormone (ACTH)

Plasma concentrations of ACTH were analysed using an ELISA kit (Product Code: CSB-E15926Fh, CUSABIO TECHNOLOGY LLC, Houston, TX, USA). Absorbance was measured with a microplate reader (Stat Fax 303 Plus Awareness Technology, Inc., Palm City, FL, USA) at a wavelength of 450 nm and converted to picograms per millilitre by using a four-parameter logistic curve. Plasma samples and assay standards (50 μL) were assayed in duplicate in a single kit. The assay kit was validated by assessment of the slope of serial dilutions of plasma samples against the assay standards. The standard curve consisted of five ACTH concentrations ranging from 75 to 1200 pg mL^{-1} and the serial dilutions consisted of two new standard curves spiked with two different 10- μL pools of plasma samples, each plasma pool was constructed by adding 5 μL of 10 individual samples (Fuller *et al.* 2020). Spiked standard curves showed good parallelism with the assay standard curve (Fig. S1 of the Supplementary material). For further information on calculating

ACTH, refer to ‘Adrenocorticotrophic hormone (ACTH)’ of the Supplementary material (Fig. S1).

Glucocorticosteroids

Total GCs were analysed using a CORT ELISA Kit (Product code: 501320, Cayman Chemical, Ann Arbor, MI, USA). This kit was previously validated to quantify 1α -OH-B, by using the cross-reactivity of the CORT antibody with 1α -OH-B and excluding other GCs by mass spectrometry (Evans et al. 2010; Lyons and Wynne-Edwards 2019; Iki et al. 2020). The cross-reactivity between CORT antibody and 1α -OH-B was reported to be 1.49% (Iki et al. 2020) and 5.2% (Evans et al. 2010). As reported by Cayman Chemical, the CORT antibody has high cross-reactivity with CORT (100%), 11-deoxycorticosterone (15.8%), 11-dehydrocorticosterone (2.9%), and cortisol (2.5%). Because we did not exclude other GCs, we assumed the CORT assay would reflect total GC concentrations of all components that immunoreacted with the CORT antibody, expressed in CORT units. For further information on calculating GCs, refer to ‘Glucocorticosteroids’ of the Supplementary material (Fig. S2).

Lactate

A hand-held Lactate Pro 2 analyser (Arkray LT-1730, ARKRAY USA, Inc., Miami, FL, USA) was used to measure whole-blood and plasma lactate concentrations *in situ*, following the manufacturer’s instructions. The Lactate Pro 2 reads in the range of 0.8–25 mmol L⁻¹, displaying ‘high’ for values higher than the upper range limit.

Post-release survival (PRS) – satellite tagging

All PSAT tags were programmed to detach from the sharks after a 30-day monitoring period. The tags are designed to record depth, temperature, and light information, which were archived at 24-h intervals for the full 30-day deployment, and 10-min intervals for the last 5 days of deployment. The PSAT tags were programmed to activate once the individual swam at a depth >10 m and to prematurely release (i.e. before the 30-day specified period) if the individual remained at a constant depth (± 2 m) for a consecutive 24 h. The tag transmits data to the ARGOS satellite system. Raw data were analysed by Wildlife Computers, which provided a report indicating the pop-off date, location, and daily values for temperature, light, and depth, as well as movement data at 10-min resolution for the final 5 days (Drymon and Wells 2017).

Data analyses

A two-way ANOVA was used to assess for differences in the blood and plasma metabolite concentrations between release condition and sex. Quantitative and semi-quantitative analytical methods such as Pearson’s correlations and means plots were used to assess how well the concentrations of blood metabolites can be used as a survival predictor. On the basis of previous work on changes in plasma/whole blood metabolites

with time (Fuller et al. 2020; Iki et al. 2020), ACTH, GCs and lactate concentrations were binned in 15-min increments for the first hour on the hook, and then bins were expanded to 30-min increments through 4 h (i.e. the maximum time on the hook).

For the 13 PSAT-tagged sharks, PRS was estimated using the transmitted depth data for the full 30-day deployment, unless the tag detached prematurely. Premature PSAT tag release occurs in one of the following three situations: sharks that sink to the seafloor and remain below a certain depth (referred to as a ‘sinker’), sharks sit at a constant depth (‘sitter’), or if a tag was floating at the surface (maximum depth was ≤ 1 m) (‘floater’) (Hutchinson et al. 2015; Lynch et al. 2017). The following two mortality scenarios accounting for the five ‘floater’ tags were considered: ‘floaters’ were (S1) considered premature tag detachments from surviving sharks, or (S2) premature detachment was the result of mortality (and scavenging) or a predation event. Kaplan–Meier (KM) survival models fit the data under these two scenarios and were used as a non-parametric approach for estimating survival. For each PSAT-tagged shark, depth profiles were plotted for the last 5 days (one record per 10 min) of the deployment periods. Changes in light, temperature, and depth values in the final 5 days of deployment were reviewed to classify the fate of ‘floater’ tags as predation events or premature detachments.

All data analysis was performed using R software (ver. 4.2.3 (2023-03-15), R Foundation for Statistical Computing, Vienna, Austria, see <https://www.R-project.org/>, accessed 12 April 2023) with packages: car (ver. 3.1-2, see <https://cran.r-project.org/package=car>; Fox and Weisberg 2019), ggplot2 (ver. 3.4.0, see <https://CRAN.R-project.org/package=ggplot2>; Wickham 2016), tidyverse (ver. 2.0.0, H. Wickham, R. François, L. Henry et al., see <https://github.com/tidyverse/dplyr>; Wickham et al. 2019), survival (ver. 3.5-3, T. Therneau, see <https://CRAN.R-project.org/package=survival>), ggpubr (ver. 0.6.0, A. Kassambara, see <https://rpkgs.datanovia.com/ggpubr/>), rstatix (ver. 0.7.2, A. Kassambara, see <https://rpkgs.datanovia.com/rstatix/>), lubridate (ver. 1.9.3, see <https://cran.r-project.org/package=lubridate>; Grolemund and Wickham 2011), ggpmisc (ver. 0.5.5, P. J. Aphalo, see <https://docs.r4photobiology.info/ggpmisc/>), maptools (ver. 1.1-8, R. Bivand and N. Lewin-Koh, see <http://maptools.r-forge.r-project.org/>), RODBC (ver. 1.3-20, B. Ripley and M. Lapsley, see <https://cran.r-project.org/package=RODBC>), chron (ver. 2.3-61, D. James and K. Hornik, see <https://cran.r-project.org/package=chron>), stringr (ver. 1.5.1, H. Wickham, see <https://github.com/tidyverse/stringr>), data.table (ver. 1.14.9, M. Dowle and A. Srinivasan, see <https://rdocumentation.org/packages/data.table/versions/1.14.8>), mgcv (ver. 1.9-0, see <https://cran.r-project.org/package=mgcv/>; Wood 2017), mgcViz (ver. 1.9-0, see <https://cran.r-project.org/package=mgcViz>; Fasiolo et al. 2019), PBSmapping (ver. 2.73.4, J. T. Schnute, N. Boers and R. Haigh, see <https://github.com/pbs-software/pbs-mapping>) and geosphere (ver. 1.5-18, R. J. Hijmans, see <https://cran.r-project.org/package=geosphere>).

Ethics

Shark capture, tagging, and the taking of blood samples were conducted under exemptions of the *Fish Resources Management Act* 1994. Exemptions were granted to the Department of Primary Industries and Regional Development (DPIRD) for research. All aspects of this study conformed to the [National Health and Medical Research Council \(2014\)](#).

Results

In total, 240 sandbar sharks (mean FL \pm s.e. 136.7 ± 0.3 cm) were captured, with TOH ranging from 46 to 250 min ($n = 28$, mean TOH \pm s.e. 105 ± 23 min). Most sharks (98.8%) were hooked cleanly (i.e. through the side or bottom jaw), with only three sharks being entangled in the line (all of which had blood samples taken). Two of these entangled sharks exhibited a physical injury to their gill slits. Fifty-seven sharks had blood samples (29 whole blood and 28 lactate only) collected (mean FL 138.2 ± 0.5 cm; range 82–162 cm), of which 13 were tagged with PSAT tags (mean FL 138.2 ± 1.0 cm; range 119–162 cm) (Fig. 2).

Post-capture survival (PCS) – blood and plasma physiology

Of the 240 sandbar sharks caught in the surveys, 74% were assigned to Release category 1. Most of the blood-sampled sandbar sharks ($n = 49$) were assigned to Release category 1, three in Category 2, and five were in Category 3. Total PCS was observed in this study, and all animals were released after sampling. An ANOVA was not performed on

release conditions because of a limited number of samples in Conditions 2 and 3 for all blood metabolites, resulting in little statistical relevance. Conditions 2 and 3 were combined to compare sharks released in good condition to those not in good conditions, although the sample size was still limited for the latter category.

No significant difference was detected between sexes for any blood-physiology metric (ACTH $t = 0.6$, d.f. = 10, $P = 0.6$; GCs $t = 0.6$, d.f. = 10, $P = 0.6$; and lactate $t = 2.2$, d.f. = 4, $P = 0.1$).

Adrenocorticotrophic hormone values ranged from 62.6 to 109.0 pg mL⁻¹, with variability being observed at each time interval. No significant ($t = 0.54$, d.f. = 23, $P = 0.6$) differences were observed in ACTH mean values through time. Glucocorticosteroid values ranged from 316.8 to 2013.7 pg mL⁻¹, and no significant ($t = 1.8$, d.f. = 23, $P = 0.1$) differences were seen through time. Neither ACTH ($\chi^2 = 0.03$, $P = 0.9$) nor GCs ($\chi^2 = 1.4$, $P = 0.2$) showed differences with sex (Fig. 3). Whole-blood lactate continued to rise with an increasing TOH ($t = 2.6$, d.f. = 24, $P = 0.02$) (Fig. 3).

Whole-blood lactate was measured for all PSAT-tagged sharks ($n = 13$, mean 16.1 ± 7.2 mmol L⁻¹), although only 11 tagged sharks had hook times available (Fig. 4). Whole-blood lactate had higher mean values after 2 h (before 2 h 13.9 ± 6.6 mmol L⁻¹; after 2 h 21.9 ± 6.1 mmol L⁻¹).

Satellite tags

Post-release survival (PRS) – satellite tagging

Post-release survival varied from 100 to 61.5% (Scenarios #1 and 2 respectively) depending on the interpretation of premature PSAT-tag releases associated with the five

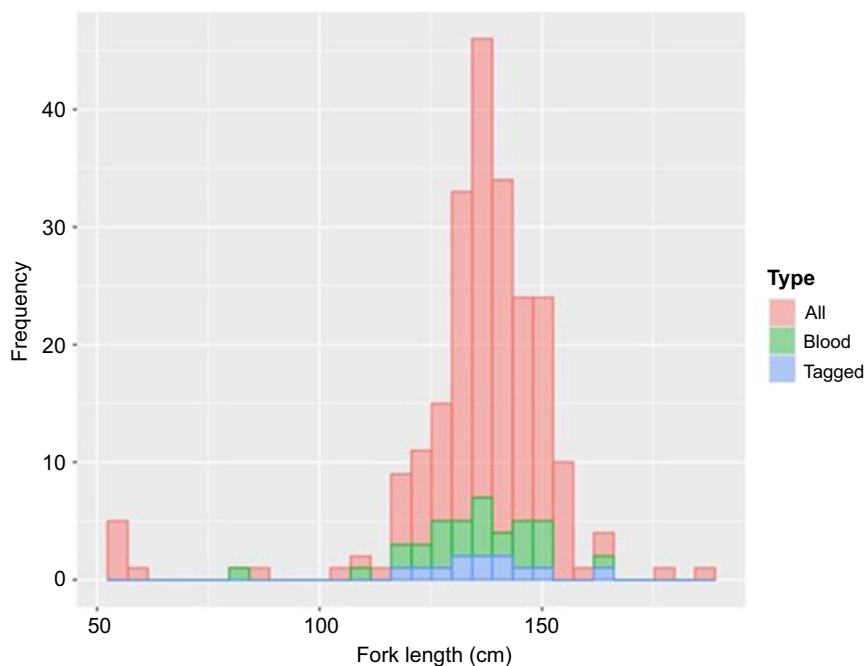


Fig. 2. Length–frequency distribution (fork length, FL, cm) of all sandbar sharks (mean FL 136.7 ± 0.3 cm, $n = 240$), with sharks that were blood sampled (mean FL 138.2 ± 0.5 cm, $n = 57$) and tagged sharks (mean FL 138.2 ± 1.0 cm, $n = 13$).

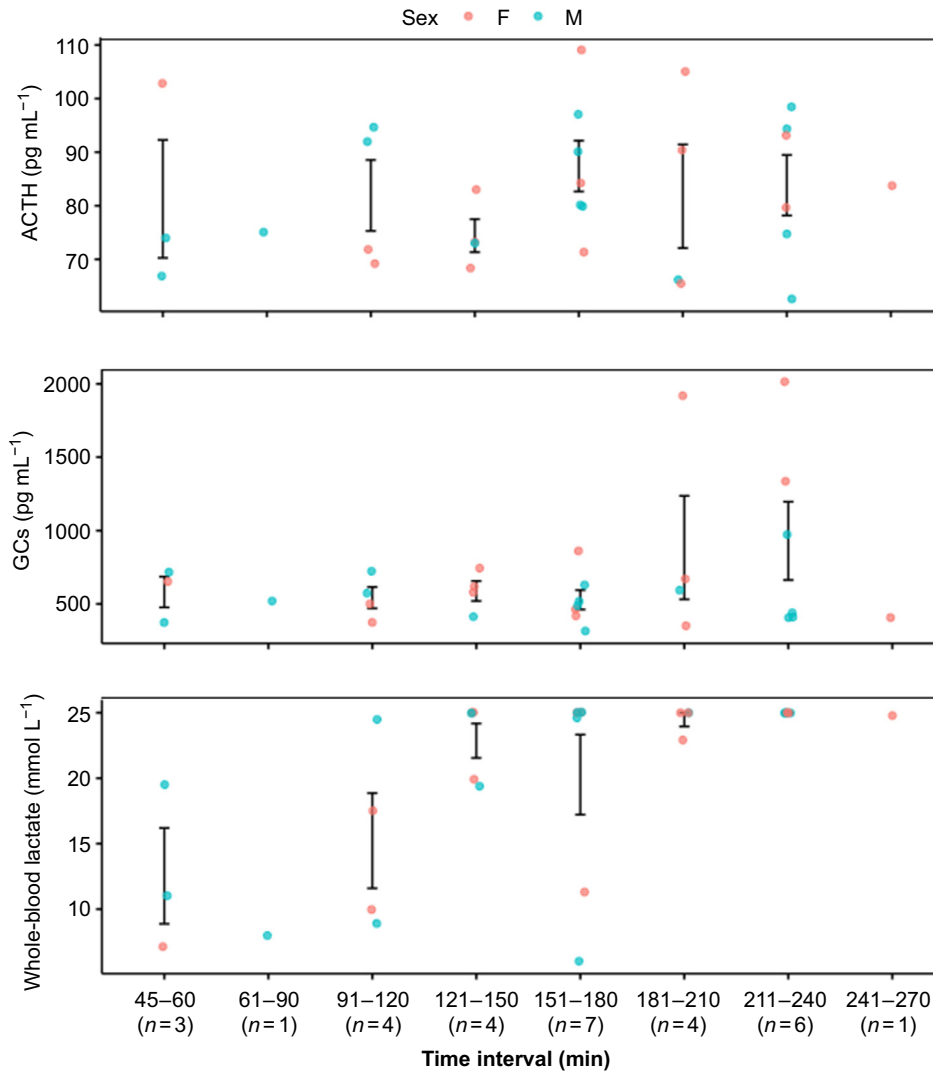


Fig. 3. Mean whole-blood and plasma metabolite concentration of 29 subsampled sharks, with all three blood and plasma metabolite values for time intervals time on hook (TOH, min) (adrenocorticotrophic hormone, ACTH; glucocorticoids, GCs). Means were calculated for sample sizes of ≥ 3 . Error bars are standard errors. Lactate field values ranged from 0 to high (≥ 25 mmol L⁻¹); high values were given a value of 25.

‘floater’ tags (Fig. S3). Under Scenario 1 (i.e. mortality equals ‘sinker’), there was no mortality after 30 days, so the KM model could not be fitted because of lack of contrast. Under Scenario 2, five of the tagged sharks had premature tag releases (i.e. mortality equals ‘floater’) and are assumed to be mortalities in KM models. The five tagged sharks that had premature tag releases were primarily released in Condition 1 ($n = 4$), with TOH ranging from 103 to 222 min (Table S1 of the Supplementary material).

Depth profiles and vertical behaviour

For the last 5 days of deployment, tagged sandbar sharks moved in the water column between the surface (0 m) and 276-m depth (Fig. S4). There was a diel movement pattern,

with some individuals moving deeper during the day (from 06:00 to 18:00 hours) and towards shallower waters at night (from 18:00 to 06:00 hours) (Fig. S4). Some sharks remained at a more constant depth during the day (e.g. SS #1) and others had larger vertical behaviours (e.g. SS #11). Shark SS #11 had the most visible diel patterns, with strong peaks in depth in the last 5 days of tag deployment (Fig. 5).

Five sharks (tag numbers SS #4, Fig. 5, and SS #2, 6, 8 and 13, Fig. S4) showed characteristics categorised as ‘floaters’, with depth being constant at 0.5 m for 24 h. All ‘floater’ tags’ temperature and light patterns remained consistent with tags at similar depth ranges that remained attached for the 30-day period. These patterns in light and temperature in the ‘floater’ tags did not conform to the values usually seen in

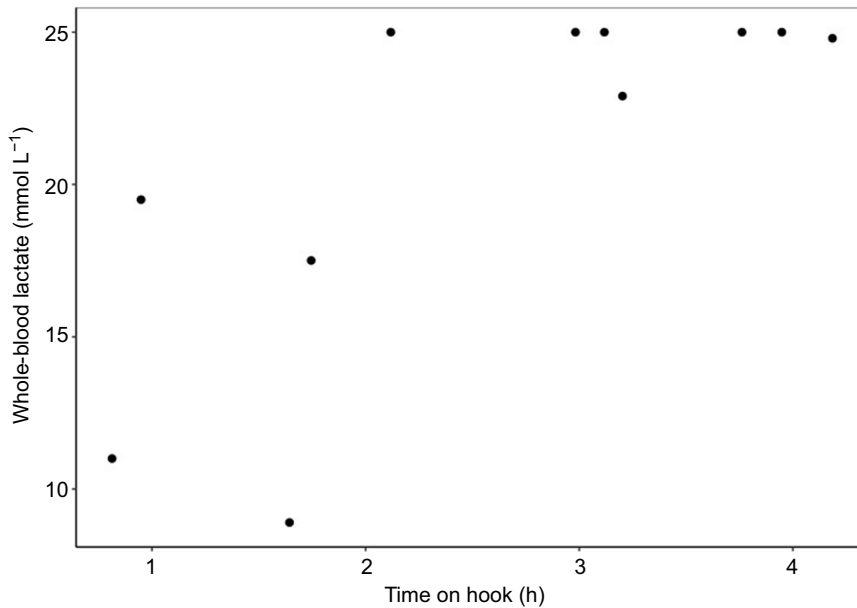


Fig. 4. Whole-blood lactate concentration of 11 tagged sharks in comparison to time on hook (TOH) (hours). Lactate field values ranged from 0 to high (≥ 25 mmol L⁻¹); high values were given a value of 25.

predation events, such as an expected in changes in vertical movement profiles, and attenuation in temperature and light level variation from ingestion. There was also no evidence of a mortality event, followed by immediate scavenging and tag detachment (i.e. the animal sinking to the seafloor for a brief period (brief constant depth) before the tag detached.) For results on the minimum and maximum depth profiles of the 13 tagged individuals over the entire 30-day period, refer to Fig. S5.

Discussion

The scope of this study was to assess whether experimental surveys influence the recovering population of sandbar sharks and, subsequently, address whether the limited recapture reports from annual scientific surveys were due to capture-related mortality. We combined the use of satellite telemetry and stress physiology to assess whether the low recapture rate of sandbar sharks was due to a low PRS in the scientific surveys (8.5%) as hypothesised by Bartes *et al.* (2021). We found that all captured animals survived a maximum 4-h hooking time.

Post-capture survival (PCS) – blood and plasma physiology

Hormones are the first indication of stress responses in vertebrates (Armour *et al.* 1993; Fuller *et al.* 2020). In a controlled environment after an acute stress event, we would have expected to observe an early peak in ACTH, followed by an increase in the GCs concentration on ACTH stimulation (Sapolsky *et al.* 2000). However, considering sharks were brought on board and sampled after at least 45 min on the

hook, we were not able to register the initial hormonal behaviour at the beginning of the stress response, as shown in other shark studies performed in controlled conditions (Fuller *et al.* 2020; Iki *et al.* 2020). Although it is impossible to get baseline concentrations, we expect low concentrations in animals that are not stressed, as has been shown in Belanger *et al.* (2001). Previous work on ACTH as a stress parameter in the Atlantic sharpnose sharks (*Rhizoprionodon terraenovae*) showed concentrations (ACTH 50–120 pg mL⁻¹) similar to those in our study (ACTH 62.64–109.04 pg mL⁻¹) (Fuller *et al.* 2020). In our study, ACTH showed variable values over time, and like in the study of Fuller *et al.* (2020), there were no differences in ACTH with sex. The time response to GCs following the production of ACTH is variable among species, but in teleost fishes, it typically take minutes (Pankhurst 2011). In this study, no clear inhibition of ACTH concentrations was observed, likely because sharks remained stressed over the entire TOH or differing levels of energy expression in individuals (i.e. routinely swimming *v.* short bursts of escape behaviours) (Gallagher *et al.* 2017; Bouyoucos *et al.* 2018). Concentrations of GCs in elasmobranch species have been reported to increase less than 1 h after an acute stress event in Japanese banded houndshark (*Triakis scyllium*; Iki *et al.* 2020) and blacktip reef sharks (*Carcharhinus melanopterus*; Schoen *et al.* 2021) and up to 24 h after the initial stress event (cururu stingrays, *Potamotrygon wallacei*; Brinn *et al.* 2012). Variable PCS rates in sandbar sharks have yet to be explored concerning hormone production. The limitation of sample size in this study and other studies has reduced the ability to observe these hormonal differences among individuals and among different species. The number of samples in this study used for hormonal blood analysis was driven by equipment shortages and difficulty with taking blood (clotting of

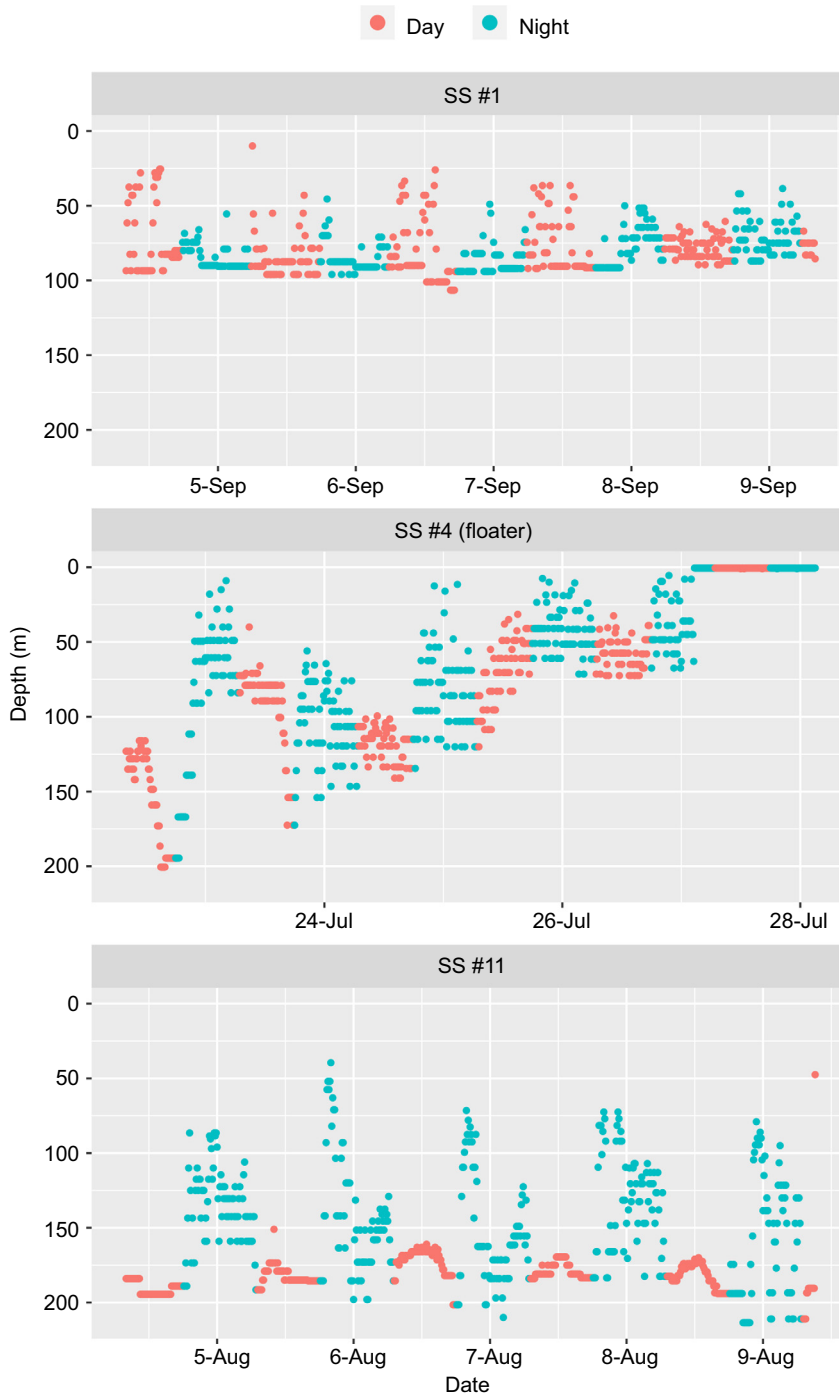


Fig. 5. Three types of temporal patterns in depth (constant, floater, and diel) for the last 5 days of deployment for a subset of satellite-tagged sharks (day 06:00 to 18:00 hours; night 18:00 to 06:00 hours).

blood) and future studies should focus on larger sample sizes ($n = 64$) to avoid this limitation ($\alpha = 0.05$, $f = 0.25$, power = 0.8).

During capture, increased energy expenditure or impaired respiration (e.g. restricted motility owing to length of gangion or net entanglement) leads to anaerobic metabolism, and its by-products, such as lactate (Brill *et al.* 2008; Skomal and Mandelman 2012). Although lactate measurements *in situ* cannot be used to forecast survival following a stress event

by capture, the whole-blood lactate values in this study ($16.1 \pm 7.2 \text{ mmol L}^{-1}$) are comparable to those of sandbar shark ($11.5 \pm 7 \text{ mmol L}^{-1}$) and to those of other Carcharhinidae species ($10.2 \pm 11 \text{ mmol L}^{-1}$) in Marshall *et al.*'s (2012) study. Marshall *et al.* (2012) compared haematological indicators in 11 species in both pelagic and demersal longlines with varying soak times of 2–12 h, although they did not use hook timers. Varying PCS was observed with 9% PCS in Atlantic sharpnose sharks and

93.2% in blue sharks (*Prionace glauca*) (Hutchinson and Bigelow 2019). With this study and the addition of hook timers, we have build on the understanding of the physiological effects of TOH and PCS of sandbar sharks (Mandelman and Skomal 2009; Gulak and Carlson 2021).

Most of the sandbar sharks swam strongly after release, with only a few being in worse-release conditions (i.e. Categories 2 or 3). Consequently, the elevated concentrations of GCs and ACTH over time reflect an adequate survival response for sandbar sharks caught in demersal longline surveys (Romero and Beattie 2022). Because of the small number of blood samples ($n = 5$, $n = 3$) and tagged sharks ($n = 1$, $n = 0$) in release conditions 2 or 3, release condition could not be assessed as an indicator of PCS or PRS. Release conditions vary among species and gear types, with dusky sharks having 52% in pelagic longline studies (Sulikowski *et al.* 2020) and Atlantic sharpnose sharks having 4% in rod and reel studies (Fuller *et al.* 2020) of catch recorded in release condition 1 (or equivalent rating), whereas sandbar sharks in the study of Marshall *et al.* (2015) had a total of 87.4%, and this study had 88% in release condition 1. Release conditions have been a useful predictor in PRS for other Carcharhinids, such as juvenile silky sharks (*Carcharhinus falciformis*), where larger sample sizes were available in each release-condition category (Hutchinson *et al.* 2015). Although qualitative metrics are difficult to compare, further research should aim to increase sample size ($n = 30$, $d = 0.37$, $\alpha = 0.05$, power = 0.8) to assess whether release condition can explain the variation in PRS.

Satellite tags

Post-release survival (PRS) – satellite tagging

Tagged individuals had 100% PRS. Sandbar sharks are known to be a robust species, with high rates of PRS (Marshall *et al.* 2015; Gulak and Carlson 2021; Whitney *et al.* 2021). Whitney *et al.* (2021) reported 96.4% PRS for sandbar sharks caught by longlines in a US fishery after TOH of 3 h. Further, Marshall *et al.* (2015) reported variable and lower PRS with a longer TOH (i.e. 100% survival with <3 h TOH; 80% survival with >4 h TOH), but this variability may also be attributed to differences in gear, handling practices, sex, and shark size, from those in other sandbar shark PRS studies (Marshall *et al.* 2015; Lynch *et al.* 2017). Our study also resulted in a higher number of adult sharks (136.7 ± 0.3 cm FL) than in the study of Marshall *et al.* (2015) (99.5 ± 21 cm FL). For some species, such as silky sharks, mature sharks are more robust, with 84.8% PRS in longline fisheries (Schaefer *et al.* 2021) compared with juveniles with 16% in purse seine fisheries (Hutchinson *et al.* 2015). However, as stated previously, further research would have to be conducted to fully understand the species-specific relationship between different factors and PRS (Knotek *et al.* 2022). Post-release survival rates of sandbar sharks are high compared with the 3 and 12% survival of

dusky sharks (*Carcharhinus obscurus*) and 25% survival of Atlantic sharpnose sharks after 3 h TOH in other longline studies (Morgan and Burgess 2007; Marshall *et al.* 2012, 2015; Sulikowski *et al.* 2020; Whitney *et al.* 2021).

Previous PSAT PRS studies have shown that constant-depth signatures reflecting a dead animal on the seafloor can be used to infer the fate of PSAT-tagged sharks (French *et al.* 2015; Hutchinson *et al.* 2015; Drymon and Wells 2017; Hutchinson and Bigelow 2019). However, in the current study, released individuals remained alive until the PSAT released from the anchor. Of these, five tags released prematurely and floated on the surface as a result of improper anchorage into the tissue or tag malfunction (Kohler and Turner 2001; Hammerschlag *et al.* 2011; Musyl *et al.* 2011). Timing of post-release mortality across studies is consistent (Ellis *et al.* 2017), with most mortalities occurring within hours of release. Although these five individuals had a premature tag detachment days after release, which can indicate mortality or predation, it is more likely that these ‘floaters’ tags are a result of tag-attachment failures. Although there was a chance for predation post-release, given predator abundance in the area, the vertical movement profiles of the five ‘floater’ tags were consistent with surviving sharks and did not display any clear evidence of a predation event or attempt (i.e. attenuated temperature and light levels or brief periods of constant seafloor depth) (Mitchell *et al.* 2018; Ryan *et al.* 2019; Braccini *et al.* 2021). The larger depth variances seen prior to the 24 h before tag detachment have previously been linked with surviving sandbar sharks (Marshall *et al.* 2015). This provides support to our assumption that 100% of the sharks tagged in this experiment survived the catch-and-release process under the longline fishing settings used as part of the survey.

Depth profiles and vertical behaviour

The behaviour exhibited by the satellite-tagged sandbar sharks mirrors that of their natural hunting behaviour (Conrath and Musick 2008). This further supports that scientific surveys are not having negative effects on sandbar sharks. The observed day–night cycle is supported by sandbar sharks’ predominant prey item of squid, which has strong diel vertical movement patterns (Stevens and McLoughlin 1991; Last and Stevens 1994; Siwabessy *et al.* 2000). The presence of squid in the mouths of tagged sandbar sharks was also noted during data collection (Chris Dowling, pers. obs).

The maximum depth recorded for the entire 30-day deployment reflects the depth profile of the continental shelf in which the sharks were caught and released (Siwabessy *et al.* 2000; McAuley *et al.* 2007). Sandbar sharks are known to inhabit depths from intertidal down to 280 m (Stevens and McLoughlin 1991), with depths of 172 m having been recorded in studies by Conrath and Musick (2008). In our study, the data transmitted from the PSAT tags recorded depths of up to 307 m, deeper than previously recorded (Last and Stevens 1994; Conrath and Musick 2008; Andrzejczek *et al.* 2018).

Conclusions

By combining the information on the immediate expression of ACTH, GCs and of whole-blood lactate concentrations continuing to rise with an increasing TOH (Hoffmayer and Parsons 2001; Fuller et al. 2020), we theorise that the sandbar sharks remained stressed throughout the capture process. However, the post-capture and post-release survival were high under current survey sampling practices, indicating that this species is resilient to capture and handling during longline surveys. The extended TOH of 4 h limited our ability to investigate the expression of ACTH and GCs to examine the relationship between the studied hormones and TOH. A shorter TOH would be required to fully explore this relationship through capturing the rapid expressions of these circulating hormones. To observe a more clearly defined relationship between lactate concentrations and TOH, lactate concentrations should be measured for a longer period of time after the initial stress event (McAuley et al. 2005; Gulak and Carlson 2021). Although a shorter TOH is recommended for testing primary stress responses for sandbar sharks, using whole-blood lactate as a secondary stress response could be an effective method of predicting mortality with longer exposure times, coupled with PRS monitoring to qualify modelling (Braccini et al. 2020; Gulak and Carlson 2021). The depth profiles of sandbar sharks in the last 5 days of tag deployment support the possibility of sharks following their prey through the water column, which could indicate the hardiness of the species after days returning to the ocean after the capture process, with some individuals exhibiting strong diel patterns of diving during the day and coming into shallower waters at night. Our results showed a clear indication that the current longline methods used in shark abundance surveys do not affect the post-capture and post-release survival of sandbar sharks. The combination of blood metabolites as stress parameters, release conditions and satellite tagging methods could be applied to other species or fisheries to develop predictive PRS modelling (Braccini et al. 2012). This could decrease the costs of survival studies as well as create an understanding of how each species reacts to fishing pressures.

Supplementary material

Supplementary material is available [online](#).

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Data availability. Data are available upon request to the authors. Please contact matias.braccini@dpird.wa.gov.au.

Conflicts of interest. The authors declare that they have no conflicts of interest.

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