From rivers to ocean basins: the role of ocean barriers and philopatry in the genetic structuring of a cosmopolitan coastal predator

**Running title:** Global population genomics of the Bull Shark

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# Ethics approval (include appropriate approvals or waivers)

All samples presented in this study were collected for alternative purposes, such as monitoring of local fisheries or biodiversity, and samples were re-used to avoid new invasive sampling. Samples from New South Wales (Australia) were collected in accordance with NSW Department of Primary Industries (DPI) Animal Care and Ethics Committee permit 07/08 and NSW DPI Research Permit Section 37 (PO1/0059A-2.0). Sampling in Northern Territory (Australia) was approved by the Charles Darwin University Animal Ethics Committee (Approval No. A11041 and A19008) and undertaken through NT *Fisheries Act* Special Permits S17/3364 and S17/3467. Sample collection in Cape York and Victoria River (Australia) was conducted under a CSIRO animal ethics permit (A2-2016; 2017-04). Tissue samples obtained from northern Australia, as published in Tillett et al. (2012), were collected under NT *Fisheries Act* Special Permit S17/27134 and Kakadu Research Permit number RK 689, in agreement with CDU Animal Ethics Committee Approval number A07001. Samples in Papua New Guinea were collected with consent from local communities in Gulf Province, and under supervision of the Gulf Provincial Fisheries Office. All fin clips from the Gulf of Mexico were collected under two IACUC protocols: one at the University of South Alabama (IACUC protocol 974394) and one at Mississippi State University (IACUC-17-620). Samples from Japan were obtained in accordance with the Guidelines for Care and Use of Animals approved by the ethics committee of the University of Tokyo (A16-13 and P19-2). The capture of Bull Sharks is permitted by the Agriculture, Forestry and Fisheries Department of Okinawa Prefecture (26-29, 27-25, 28-18, 29-20, 30-30, 31-22, 2-7). Handling of live shark specimens from Fiji were approved under the Research Permit issued by the Fiji Ministry of Education, Heritage and Arts to Beqa Adventure Divers and performed in accordance with relevant guidelines and regulations. All samples from Brazil were collected from parts of dead bycatch specimens in local markets (2003-2005) under market sampling permit IBAMA PA 037/2002; samples corresponded to the same set as previously used in Karl et al. (2011) and no ethical agreement was required. Samples from Western North Atlantic (USA), Gulf of California, Sierra Leone, Mozambique, Indonesia, Thailand, and Queensland (Australia) were collected from local fisheries landings and did not require specific ethics approvals, as per national guidelines. All material from Sri Lanka was collected from dead fisheries bycatch specimens that are not protected by the Fauna and Flora Protection Ordinance (FFPO) or any other law in place in Sri Lanka, in accordance with the letter with reference number WL/3/2/74/17, dated 4th January 2018, issued by the Department of Wildlife Conservation, Sri Lanka. Samples from the Arabian Peninsula and adjacent waters were collected from fishery landings and no sampling permit was required. An authorization letter from the Ministry of Climate Change and Environment of the United Arab Emirates was obtained. Costa Rican samples were collected under the biodiversity permit VI-2391-2019 issued to the project VI-B7162. All samples from Réunion were collected from dead sharks caught on the Réunion Island shark control program and no ethical agreement was required. Samples from Seychelles were collected from landed artisanal fishery catch and no ethical agreement was required. All samples provided by the KZN Sharks Board from the east coast of South Africa were collected from dead sharks caught in the KwaZulu-Natal bather protection programme as constituted by Act 5 of 2008 in the province of KwaZulu-Natal and no ethical agreement was required. All shipping procedures of shark tissue samples were conducted under the relevant import and export permits issued by the Australian Government, Department of Agriculture and Water Resources. The species exported is not listed under CITES or any other international regulatory convention. Samples were imported into Australia under AQIS permit number 0001500212 and 0003253262 issued to CSIRO.

# Conflicts of interest: We note no conflict or competing interests among the authors in relation to the information provided in this manuscript.

# Abstract

Understanding the population structure of a species is important to accurately assess its conservation status and manage the risk of local extinction. The Bull Shark (*Carcharhinus leucas*) faces varying levels of exploitation around the world due to its coastal distribution. Information regarding population connectivity is crucial to evaluate its conservation status and local fishing impacts. In this study, we sampled 922 putative Bull Sharks from 19 locations in the first global assessment of population structure of this cosmopolitan species. Using a recently developed DNA-capture approach (DArTcap), samples were genotyped for 3,400 nuclear markers. Additionally, full mitochondrial genomes of 384 Indo-Pacific samples were sequenced. Reproductive isolation was found between and across ocean basins (eastern Pacific, western Atlantic, eastern Atlantic, Indo-West Pacific) with distinct island populations in Japan and Fiji. Bull Sharks appear to maintain reproductive connectivity using shallow coastal waters as dispersal corridors, whereas large oceanic distances and historical land-bridges act as barriers. Females tend to return to the same area for reproduction, making them more susceptible to local threats and an important focus for management actions. Given these behaviours, the exploitation of Bull Sharks from insular populations, such as Japan and Fiji, may instigate local decline that cannot readily be replenished by immigration, which can in turn affect ecosystem dynamics and functions. These data also supported the development of a genetic panel to ascertain the population of origin, which will be useful in monitoring the trade of fisheries products and assessing population-level impacts of this harvest.

**Keywords:** DArTseq, mitogenome, close-kin, genetic connectivity, provenance, DNA forensics

# Introduction

Understanding the population structure of a species and the barriers that disrupt dispersal is important to accurately assess the global conservation status and manage the risk of local extinction. This is especially true for species of commercial importance (Begg et al., 1999) or conservation concern (Moritz, 1994), which are impacted disproportionally by anthropogenic or environmental pressures. Dispersal can promote genetic connectivity across patches of suitable habitat (Ronce, 2007), but physical barriers and behaviours can ultimately limit gene flow, even when dispersal potential is high, and result in demographically-independent populations (Waples & Gaggiotti, 2006). When genetically isolated populations are reduced in size due to unsustainable harvest, there is a risk of inbreeding depression and the loss of genetic diversity, without the chance to be ‘rescued’ by individuals dispersing from adjacent populations, increasing the likelihood of population extinctions (Frankham et al., 2017).

In marine taxa, gene flow tends to be restricted by environmental or biogeographic barriers, movement ecologies, and habitat preferences (Bowen et al., 2016; Dudgeon et al., 2012; Hirschfeld et al., 2021; Rocha et al., 2007). In large-bodied coastal species with global distributions, such as marine turtles, cetaceans, and many elasmobranchs (sharks and rays), large-scale marine biogeographic barriers shape population genetic structure (Dutton et al., 1999; Fontaine et al., 2007). For example, the Scalloped Hammerhead (*Sphyrna lewini*)*,* shows genetic connectivity along the continental margins, yet limited gene flow across the East Pacific barrier, the Mid-Atlantic barrier, and the Isthmus of Panama (Daly-Engel et al., 2012; Green et al., 2022). The permeability of environmental barriers, such as the Indo-Australian Archipelago, changes across time and space, and consequently determines the observed distribution of genetic variation (Cowman & Bellwood, 2013). The effect of these ocean barriers on the spatial structuring of populations is essential knowledge for management, given that threats, such as overfishing and habitat modification, should be assessed and managed at biologically-relevant spatial scales. For vagile marine taxa this often requires cooperative strategies between nations (e.g., International Whaling Commission, Indian Ocean Tuna Commission, and Western and Central Pacific Fisheries Commission).

Delineation of population structure in species with high mobilities and large population sizes have recently been improved with the use of genomic data (Layton et al., 2020; Luikart et al., 2019; Oleksiak & Rajora, 2020; Ovenden et al., 2018). Complexity-reduction genome-scan methods, such as Diversity Arrays Technology sequencing (DArTseq, Jaccoud et al., 2001) and targeted DNA-capture approaches, including Rapture and DArTcap (Ali et al., 2016; Feutry et al., 2020), have been widely used to assess genetic diversity and reproductive connectivity in natural populations (e.g., Green et al., 2019; Komoroske et al., 2019). Recently, these methods have also been applied to DNA forensics or traceability studies in non-model species, with the objective to identify species, sex, provenance, and close-kin relationships (e.g., Arenas et al., 2017; Feutry et al., 2017; Nielsen et al., 2012; Stovall et al., 2018). Moreover, in taxa with slow mitochondrial DNA (mtDNA) mutation rates, such as elasmobranchs and marine turtles (Avise et al., 1992; Martin et al., 1992), the sequencing of full mitochondrial genomes (mitogenomes) instead of single genes, has improved the fine-scale resolution of matrilineal population structure (Feutry et al., 2014).

The Bull Shark (*Carcharhinus leucas*) is a cosmopolitan species that occupies tropical, subtropical, and temperate coastal waters and has an important ecological role in freshwater, estuarine, and marine environments (Matich et al., 2011; Smoothey et al., 2019; Trystram et al., 2017). This species experiences variable degrees of exploitation within its range and is assessed as Vulnerable on the IUCN Red List of Threatened Species (Rigby et al., 2021). The main threats affecting Bull Sharks are small and large-scale fisheries for meat and fins (Glaus et al., 2015; Holmes et al., 2009), and shark control programs which directly target this species (Blaison et al., 2015; Dudley & Simpfendorfer, 2006; Niella et al., 2021). Because of its global catch, the Bull Shark is also frequently found in the international shark fin and meat trade, raising questions regarding the origin and trade routes of the fished products (Cardeñosa et al., 2022; Clarke et al., 2006; Fields et al., 2018). Unsustainable exploitation may result in population declines and negative ecosystem impacts (Ferretti et al., 2010; MacNeil et al., 2020).

The Bull Shark has shown capacity for long-distance coastal movement (Brunnschweiler et al., 2010; Daly et al., 2014; Espinoza et al., 2016; Espinoza et al., 2021; Heupel et al., 2015; Lea et al., 2015) and genetic connectivity along continental shelves (Glaus et al., 2020; Pirog et al., 2019; Testerman, 2014). However, females exhibit reproductive philopatry to estuarine habitats at small spatial scales (~100 km; Karl et al., 2011; Sandoval Laurrabaquio‐Alvarado et al., 2021; Tillett et al., 2012). Genetic connectivity of Bull Sharks has been studied within specific regions, such as the western Atlantic (Karl et al., 2011; Sandoval Laurrabaquio-Alvarado et al., 2019; Sandoval Laurrabaquio‐Alvarado et al., 2021) and the Indo-West Pacific (Deng et al., 2019; Glaus et al., 2020; Kitamura et al., 1996; Pirog et al., 2019; Tillett et al., 2012). Yet, no studies have investigated genetic connectivity across its global distribution, which can provide crucial data to assess the population-level impacts of threats and identify the origin of fisheries products.

This study aims to apply genomic techniques (full mitogenomes and nuclear Single Nucleotide Polymorphisms or SNPs) to investigate the population genetic structure of Bull Sharks at global and local scales. We hypothesise that gene flow is limited by large-scale biogeographic barriers, but that connectivity occurs along continuous coastlines. We further evaluate the forensic power of genomic data to assign sample provenance and establish a diagnostic SNP panel to aid monitoring the origin and trade of fisheries products and assess population-level impacts of this trade. To achieve these goals, we analyse a global set of samples with three different genomic sequencing approaches (DArTseq, DArTcap, and mitogenomes), which allows us to assess population structure and connectivity across putative barriers for dispersal (such as open-ocean expanses, strong temperature gradients, and historical land-bridges). At a smaller spatial scale, contemporary connectivity is estimated by examining the spatial distribution of closely-related individuals, such as full siblings and cross-cohort half siblings.

# Material and Methods

## Sample collection and DNA extraction

A total of 922 putative Bull Shark samples (muscle or fin clip) were collected between 1980 and 2019 from 19 different countries or water bodies (termed ‘sampling locations’) around the globe in all major ocean basin regions: the eastern Pacific (E-PAC), western Atlantic (W-ATL), eastern Atlantic (E-ATL), and the Indo-West Pacific (IWP; Fig. 1). Samples from Brazil, eastern Indian Ocean (E-IO), Fiji, and numerous Australian samples were sourced from previously published genetic studies (n = 175; Glaus et al., 2020; Karl et al., 2011; Pirog et al., 2019; Tillett et al., 2012) while all other samples were novel. The samples from Japan included 10 individuals from the aquarium (Okinawa, locally sourced animals) with known pedigree and 38 wild-caught samples from the Urauchi River. Each sampling location had an approximately equal sex ratio, and total length (TL) ranged from 26.4 (*in utero*) to 406.0 cm (supplementary material section 2). Sharks smaller than 150 cm TL (68 % of all individuals) were considered juveniles with limited dispersal capacity (Heupel et al., 2015; Pillans & Franklin, 2004; Simpfendorfer & Milward, 1993). DNA was extracted using the Qiagen Blood and Tissue kit following the standard protocol (Qiagen Inc., Valencia, California, USA). DNA quality and quantity was assessed on a 1 % agarose gel, stained with SYBR safe (Invitrogen, USA), and with the NanoDrop ND-8000 spectrophotometer (Thermo Fisher Scientific, Waltham, Massachusetts, USA).

## SNP bait design and DArTcap genotyping

Initially, 188 samples with a minimum of 10 samples per location were genotyped following the DArTseq approach (according to Feutry et al., 2017). *De novo* SNP calling was performed with a proprietary software (DArTsoft14). The resulting markers were filtered using the ‘filter\_rad’ function, as implemented in *radiator* v1.1.5 (Gosselin et al., 2020). Filtering thresholds were chosen based on the empirical distribution of each filtering parameter (see supplementary material sections 4–6). Briefly, the data were filtered for: 1) low DArT reproducibility based on replicated libraries (technical replicates), 2) monomorphic markers, 3) high degree of missing data, high levels of inferred heterozygosity, and low sequencing coverage per individual, 4) low minor allele count, 5) low and high SNP sequencing coverage to avoid unreliable SNP calls or paralogous sequences, 6) high missing data per SNP, 7) too many SNPs per sequence due to sub-optimal sequence clustering, 8) short-distance linkage by keeping one SNP per sequence, 9) individual DNA contamination based on high proportion of heterozygous SNPs, 10) duplicated samples, and 11) departure from Hardy-Weinberg equilibrium. We further filtered out RAD-tags that were too short (< 60 bp), with low complexity (> 6 nucleotide repeats), or with high cluster size (> 10) to allow efficient capture with the RNA baits. Of the markers satisfying the filtering criteria, 3,200 population genetic markers were randomly selected for DNA-capture bait development. Secondly, the unfiltered data were run through the ‘sexy\_markers’ function from *radiator* (Devloo-Delva et al., 2022) to identify polymorphic sex-linked markers (putatively located on sex chromosomes), which were subsequently included in the DNA capture panel. One biotinylated RNA MYbait (Arbor Bioscience, USA) was synthesized per RAD-tag. DArTcap hybridization and washing followed the MYbaits standard protocol (<https://arborbiosci.com/wp-content/uploads/2020/08/myBaits_v5.0_Manual.pdf>). The DArTcap enriched libraries were sequenced on a HiSeq 2500 (Illumina, San Diego, California, USA) as described by Feutry et al. (2020).

## DArTcap SNP and individual filtering

Overall, 1,014 sample libraries, including 92 technical replicates, were genotyped with the DArTcap protocol. Additional filtering was performed to remove any unspecific enrichment or low-quality loci. This filtering was performed with the ‘filter\_rad’ function as described previously (see supplementary material sections 7–12). Sex-linked markers were removed prior to population genetic analyses.

The filtering steps were applied with different thresholds to six different subsets of the data (Table 1). A hierarchical approach was used for the clustering methods (e.g., Vähä et al., 2007). The first DArTcap data set was filtered under less stringent thresholds (e.g., allow more missing data and higher levels of heterozygosity) to permit the identification of species that were not Bull Shark (DATA1: 922 sharks). The second data set only contained confirmed Bull Sharks (DATA2: 868 sharks). Since unequal sample sizes can introduce bias in clustering algorithms (Foster et al., 2018; Puechmaille, 2016), in the third data set, all sampling locations were included but Australia was represented by a random subset of 60 individuals (DATA3: 430 sharks). The fourth and fifth data sets are comprised of sampling locations in the W-ATL and IWP, respectively (DATA4: 117 sharks; DATA5: 732 sharks). Lastly, within the IWP, samples from Japan and Fiji were removed and 60 Australian samples were again randomly selected to equalize the sample sizes and investigate unbiased signals of more subtle structure (DATA6: 221 sharks).

## Mitogenome amplification and sequencing

Samples from the E-ATL, IWP, and E-PAC were used to investigate their matrilineal evolutionary history (sample sizes indicated in Fig. 1). The full mitochondrial genomes of 384 putative Bull Sharks were amplified with two primer pairs (A and B fragments; supplementary material section 13.1). Polymerase chain reactions (PCR) were performed in 30 μL volumes, following the standard proofreading Takara LA Taq protocol (Takara, Otsu, Shiga, Japan). PCR conditions were set to 1 min at 94 °C for initial denaturation, then 40 cycles of denaturation (94 °C, 30 s), annealing (55 °C, 30 s) and extension (68 °C, 10 min); concluded with a 10 min extension at 72 °C. PCR products were cleaned following the Agencourt AMPure XP magnetic bead protocol (Beckman Coulter Inc., Indianapolis, Indiana, USA). Amplicons were quantified with a NanoDrop 8000 Spectrophotometer and the purified A and B fragments were pooled at equimolar concentrations for each individual. Subsequently, these amplicons were simultaneously fragmented and barcoded with the Nextera XT DNA Sample Preparation kits and 96 sample Nextera Index kit (Illumina). The libraries were quantified with the Qubit dsDNA BR assay kit (Life Technologies, Carlsbad, California, USA) and normalized. Libraries were then pooled and sequenced on a MiSeq desktop sequencer using the 2x250 bp paired-end reads MiSeq reagent kit v2 (Illumina).

## Mitogenome assembly and alignment

Demultiplexed fastq files were imported into Geneious prime software v2020.1 (Biomatters Ltd., Auckland, New Zealand) and the reads were paired. The Nextera adapters were removed, and the reads were quality trimmed at a phred score < 20 for a k-mer of 20 using the BBDuk tool as implemented in Geneious. Reads shorter than 60 bp after trimming were discarded from subsequent analyses. Merged reads for each individual were then mapped onto a previously published Bull Shark reference sequence (Chen et al., 2015) using “Map to Reference” tool in Geneious with the “high sensitivity” parameters and 10 iterations. The majority rule consensus (> 50 % of mapped reads for any single SNP, insertion, or deletion) for each shark was exported. All mitogenome sequences were aligned with the ‘multiple align’ tool and the MUSCLE algorithm (Edgar, 2004).

## Species identification

The mitogenomes were blasted (*megablast*) in Geneious against the GenBank nr/nt database. All sequences with a match of identity ≥ 98 % to a database entry were assigned as that species. Samples without mitogenome information (only DArTcap genotypes, i.e., DATA1) were assigned to species using a principal component analysis (PCA) as implemented in *adegenet* v2.1.1 (supplementary material section 7; Jombart & Ahmed, 2011). Here, the species identified based on mitogenomes served as a baseline to detect species clusters from PCA. To provide certainty, individuals needed to cluster on multiple PC axes with a mitochondrial-verified species. Where this was not possible, we marked the species as ‘unknown’. Only samples identified as Bull Shark were included in further analyses.

## Genetic diversity

Allelic richness (Ar), observed heterozygosity (HO), and unbiased expected heterozygosity (HE) were calculated for each sampling location with n > 1 using the R package *diveRsity* v1.9.90 (Keenan et al., 2013). Heterozygosity was calculated across the global data set (including monomorphic and polymorphic SNPs) to yield an unbiased estimate of HO (see Schmidt et al., 2021). Inbreeding coefficients (FIS) were calculated with *hierfstat* v0.04–22 (Goudet, 2005), using 1,000 bootstraps to determine a 95 % confidence interval. Mitochondrial nucleotide (πmt) and haplotype (h) diversities per location, and parsimony haplotype networks were calculated with the *pegas* v0.14 package (Paradis, 2010). All analyses were performed in R 4.0.2 (R Core Team, 2020) and provided in the supplementary material.

## Population structure

Fixation indices (mtDNA: sequence-based ΦST, SNPs: FST) were calculated between all sampling locations (n ≥ 1) and between/across ocean basins with the ‘popStructTest’ function in the *strataG* v2.4.905 package (Archer et al., 2017), and their significance assessed by 1,000 permutations. Next, dimensionality-reduction clustering analyses were conducted with *adegenet* (PCA and Discriminant Analysis of Principal Components, DAPC; Jombart & Ahmed, 2011; Jombart et al., 2010)*.* Individuals were grouped using the successive K-means algorithm implemented in the ‘find.clusters’ function. The goodness of fit, determined by the Bayesian information criterion (BIC), was employed to find the best number of clusters (K). To avoid overfitting, the optimal number of principal components was selected through cross-validation with a 10 % hold-out set and 1,000 replicates for all DAPC analyses, where individuals were grouped according to their sample location.

## Provenance assignment

The provenance assignment success of the DArTcap markers was tested with a*sssignPOP* v. 1.2.2 (Chen, Marschall, et al., 2018) and *rubias* v.0.3.2 (Anderson et al., 2008; Moran & Anderson, 2019). Assignment accuracy was tested with *assignPOP*, using both the Monte-Carlo and K-fold cross-validation procedures to test the assignment of a hold-out data set with 1,000 iterations. We tested power of the markers by selecting a subset of loci with the highest FST values (5 %, 10 %, 50 %, and 100 % of all loci) to train the assignment model. Similarly, the assignment accuracy of simulated mixed groups, based on a reference leave-one-out data set, was evaluated with *rubias* (Anderson et al., 2008). Known simulated proportions for each reporting unit were compared with the numbers estimated by *rubias*. Populations with a sample size of one (i.e., Sierra Leone, eastern Atlantic) were excluded from these analyses. We also examined the minimum number of informative markers needed to assign provenance by subsampling 5–500 markers based on loading contributions of each principal component from the DAPC analysis and testing the assignment accuracy with *rubias*.

## Kinship assignment

To investigate fine-scale contemporary connectivity, close-kin relationships were examined in each identified genetic cluster with > 100 individuals to allow accurate allele frequency estimation, namely W-ATL and IWP. Kinship was tested using a log-likelihood-ratio (LLR) approach developed by Bravington et al. (2016) and applied by Hillary et al. (2018) and Feutry et al. (2020). A statistical threshold was set to reduce the number of false positive detections (i.e., more distantly-related kin) due to the large number of pairwise comparisons. Replicated or recaptured individuals were already visually identified with the ‘filter\_rad’ function based on the number of loci with the same genotype for each pair of individuals (< 10 % genotypic difference).

# Results

## SNP genotyping, baiting success, and data filtering

One sample from Papua New Guinea (PNG) failed DArTseq library construction, resulting in 187 samples for DArTseq sequencing and genotyping. An average of 2,182,162 reads (of 69 bp length) per sample were obtained from the sequencing and a total of 250,945 SNPs, located on 168,810 unique RAD-tags, were called by the DArTsoft14 program. Data filtering discarded 233,916 SNPs and 33 sharks, leaving a total of 17,029 high-quality SNPs. Of those, we randomly selected 3,200 loci with at least one SNP (supplementary material sections 4–6). We also identified 469 sex-linked markers (three Y-linked and 466 X-linked markers; supplementary material section 4.2). Two Y- and 208 polymorphic X-linked sequences were included for DArTcap bait design.

Five samples failed DArTcap library construction. On average, 583,809 reads per sample were obtained from the DArTcap sequencing. After sequence clustering and SNP calling, we obtained 37,537 SNPs found on 26,335 RAD-tags. Quality-filtering was applied to six subsets of the DArTcap data (Table 1): the full data set (DATA1: 1,014 sharks including replicates and other species; 5,053 SNPs), the global data set with confirmed Bull Sharks (DATA2: 769 sharks; 3,409 SNPs), the global data set with equalized samples sizes (DATA3: 382 sharks; 1,849 SNPs), and regional Bull Shark data sets: W-ATL data (DATA4: 91 sharks; 931 SNPs), IWP data (DATA5: 635 sharks; 3,416 SNPs), and IWP data with equalized samples size (DATA6: 189 sharks; 1,785 SNPs).

## Mitogenome sequencing and assembly

The mitogenome was sequenced for 384 individuals with an average of 49,766 reads. Six individuals had low sequence coverage (< 5,000 mapped reads and mtDNA regions with no sequence coverage relative to the reference genome) and were subsequently removed from analyses (from South Africa, Sri Lanka, PNG, and Australia: Clarence River and Wenlock River). All reads that mapped to the reference genome were checked for ambiguous base calls at an 85 % threshold (where a base needed to be present in > 85 % of mapped reads to be called unambiguous) to detect DNA cross-contamination, barcode slippage, or heteroplasmy. Seven individuals that had ambiguous base calls were removed due to signs of DNA contamination (originating from South Africa, PNG, Seychelles, Réunion, and Australia). Overall, the mitogenome length was 16,707–16,708 bp (supplementary material section 13).

## Species identification

Mitogenome sequencing revealed 11 individuals from the Arabian Peninsula, Réunion, Sri Lanka, PNG, Costa Rica, and Fiji that were not Bull Shark: Pigeye Shark (*Carcharhinus amboinensis*, n = 5), Spinner Shark (*Carcharhinus brevipinna*, n = 1), Graceful Shark (*Carcharhinus amblyrhynchoides*, n = 1), Grey Reef Shark (*Carcharhinus amblyrhynchos*, n = 1), Pacific Smalltail Shark (*Carcharhinus cerdale*, n = 1), Dusky Shark (*Carcharhinus obscurus*, n = 1),and Speartooth Shark (*Glyphis glyphis*, n = 1). Furthermore, in the PCA analysis carried out on DATA1 an additional 54 individuals grouped with one of these species and another four individuals were neither these nor Bull Shark (supplementary material section 7.8). Consequently, these individuals were omitted, and 769 confirmed Bull Sharks were retained in the filtered DArTcap data (DATA2) and 361 in the filtered mitogenome data.

## Global genetic diversity

More than half the DArTseq and DArTcap (DATA2) markers were monomorphic at almost all locations (Table 2 and supplementary material section 5.6). In both genotyping protocols, the Ho for the E-PAC (HO = 0.036 in Costa Rica) and the W-ATL (HO range = 0.048−0.052) were lower than those for the E-ATL and IWP (HO range = 0.056−0.069; Table 2). Most sampling locations exhibited significantly positive inbreeding coefficients (Table 2), but this was most pronounced in the E-PAC (Costa Rica) and IWP (South Africa, Mozambique, and Fiji; FIS = 0.041−0.068).

A total of 165 polymorphic sites were identified across all 361 mitogenome sequences, with an average nucleotide diversity of 0.001 and a haplotype diversity of 0.890 (supplementary material section 13.3.1). Mitochondrial nucleotide diversities per sampling location ranged from 0.0001 in northern Australia (e.g., Roper and Towns rivers) to 0.003 in Sri Lanka (Table 2). Of the 165 polymorphic sites, most were only present within the IWP group (supplementary material section 13.3.4). Haplotype diversities were high in almost all sampling locations (h range = 0.80−0.99), with the lowest values in the Seychelles, Trinity Inlet (Australia), and Japan (h = 0.50−0.70.)

## Population structure

Based on the nuclear DNA data (FST, DAPC, and PCA), we identified four major genetic clusters and additional hierarchical structure within these clusters (Fig. 2, Table 3). The DAPC indicated that the optimal number of clusters for DATA3 (i.e., equal sample sizes) was K = 3–4 (supplementary material section 9.9). These clusters corresponded to the E-PAC, W-ATL, E-ATL, and IWP/Japan/Fiji (FST = 0.10−0.51, P < 0.001; Table 3). Further investigation within the latter group (i.e., DATA5; supplementary material section 11.9) revealed further clustering into three groups corresponding to IWP, Japan, and Fiji (Fig. 2). Here, the FST values ranged from 0.06 between Japan and Fiji to less than 0.02 among IWP locations (P < 0.001). We saw high divergence between the two Japan locations (Okinawa and Urauchi River; Fig. 2C; FST = 0.03, P < 0.001), where individuals from Okinawa clustered closer to the IWP group and individuals from the Urauchi River were split into two groups. While the DAPC grouped the E-ATL (one individuals from Sierra Leone) with the IWP cluster, FST showed this location as distinctly different (Table 3). Nuclear differentiation between locations in the W-ATL were small (FST < 0.007), although the DAPC and PCA analyses showed signs of differentiation between the southern (Brazil) and the northern locations (Gulf of Mexico and Western North Atlantic; supplementary material section 10). Overall, these results indicated significant genetic differentiation of the E-PAC, southern W-ATL, northern W-ATL, E-ATL, IWP, Japan, and Fiji.

The mitogenome distance-based ΦST values showed high differentiation between and across ocean basins (ΦST = 0.67−0.99, P < 0.001; Table 3) and the haplotype network demonstrated four major clusters (Fig. 3): E-PAC, E-ATL, western Indian Ocean (W-IO), and the E-IO and western Pacific combined (E-IO/W-PAC). All haplotypes, except four, were unique to a single sampling location (Fig. 3). Within the Indian Ocean, differentiation was strong between the W-IO and E-IO locations (ΦST = 0.75−0.96, P < 0.001), while the northern Indian Ocean locations (N-IO; i.e., Arabian Peninsula, Sri Lanka, and Thailand) showed intermediate divergence (ΦST = 0.46−0.52, P < 0.001; supplementary material section 13.5.3). The latter is demonstrated in the haplotype network with individuals from the N-IO split between the W-IO and E-IO/W-PAC haplogroups. Between the W-IO and E-IO/W-PAC groups, the haplotypes of several individuals indicated matrilineal movement at an evolutionary-recent time scale: two haplotypes, sampled from the Fitzroy River (Australia) and Fiji, clustered with the W-IO group and one haplotype from South Africa grouped with the E-IO/W-PAC group (indicated in Fig. 3). Haplotypes sampled from Japan (n = 4) and Fiji (n = 8) grouped with the E-IO/W-PAC cluster, where the Japanese haplotypes did not correspond to the three Japanese clusters identified with the nuclear SNPs. Within Australia, the mitochondrial ΦST showed structure between western (Fitzroy River), northern (Victoria River, Daly River, Adelaide River, Alligator Rivers — encompassing the South Alligator River and East Alligator River — Blue Mud Bay, Roper River, Towns River, Wenlock River), and eastern Australian sites (Trinity Inlet, Clarence River, and Sydney Harbour; Table 3). The haplotype network of all Australian individuals indicated three major haplotype clusters with no obvious geographic pattern, yet most haplotypes were unique to a sampling site (supplementary material section 13.4.2.3).

## Provenance assignment

After population structure was identified with the nuclear DArTcap markers, we tested the accuracy to assign provenance to a hold-out data set (Fig. 4). Using the marker contributions of the DAPC analysis, we found that at least 250–500 highly-differentiating markers were required for 100 % provenance assignment accuracy (supplementary material section 8.10.9). Individuals from Japan and Fiji were unlikely to be assigned to the inferred genetic cluster with less than 500 markers. A minimum of 50–100 markers were needed to obtain a reliable assignment (> 80 %) to each inferred population and only 5–50 markers were required to differentiate individuals from the E-PAC, W-PAC, and E-ATL, but sites within the IWP required up to 100 markers (Fig. 4).

## Individual and kinship assignment

In total, 102 duplicate individuals were identified in the data set. Most represented the technical replicates that were purposely included, but we also detected six recaptured sharks from Sydney Harbour, Japan, and Fiji (time-at-liberty = 38–741 days). These recaptures were always assigned to the same age-cohort where length data were available, thus providing confidence in the length-at-age function from Tillett et al. (2011). Within the W-ATL, we performed 4,095 pairwise comparisons and found two full-sibling pairs (FSPs) in Brazil. The IWP revealed more kin pairs due to the large number of pairwise comparisons (201,295). Overall, we found 18 full-sibling pairs (FSPs), all within the same sampling locations (Brazil, Réunion, Indonesia, Fiji, and within Australia: Daly, South Alligator, East Alligator, Towns, and Clarence rivers), and 40 half-sibling pairs (HSPs). Of the 40 HSPs, 32 were found within the same locations (Seychelles, Réunion, PNG, and within Australia: Sydney Harbour, South Alligator, East Alligator, Wenlock, and Clarence rivers), where 10 and eight HSPs had the same and different haplotypes, respectively; 14 pairs had missing haplotype information. Eight HSPs were distributed between rivers within northern Australia (Daly River/South Alligator River, n = 4; Adelaide River/South Alligator River, n = 1; South Alligator River/East Alligator River, n = 2; and Towns River/Wenlock River, n = 1). Seven of those eight ‘cross-river’ HSPs were juveniles (< 150 cm TL) from different age cohorts, and six pairs had different mitochondrial haplotypes.

# Discussion

This study provides the first global assessment of genetic population structure of a cosmopolitan coastal predator using thousands of SNP markers and full mitogenome data. We identify distinct genetic divergence driven by significant biogeographic barriers and philopatric behaviour. These results further facilitate the development of a SNP panel for species, sex, provenance, and kinship identification.

## Global population structure across ocean barriers

Population structure in large coastal sharks tends to be driven by environmental barriers, movement ecology, and habitat preference (Dudgeon et al., 2012; Hirschfeld et al., 2021). The DArTcap results revealed four global genetic clusters (E-PAC, W-ATL, E-ATL, and IWP), with Japan and Fiji each supporting additional isolated Bull Shark populations. Our results concur with previous studies in the Western Atlantic and Indo-Pacific regions, which showed genetic differentiation between the W-ATL, IWP, and Fiji using microsatellite data (Pirog et al., 2019; Testerman, 2014). We show that the E-PAC is strongly differentiated from the W-ATL, coinciding with the closure of the Isthmus of Panama (~3 Myr; Knowlton et al., 1993; O’Dea et al., 2016). Previously, Bull Shark connectivity has been suggested between the E-PAC and W-ATL, based on microsatellite data, through the Panama Canal (Pirog et al., 2019; Testerman, 2014), but our results indicate these populations are demographically isolated and that connectivity through the Panama Canal is negligible, if existent at all. Rather, we believe the differences between our results are likely due to higher power delivered by thousands of SNPs compared to fewer microsatellites (e.g., Green et al., 2019; Layton et al., 2020). In addition, the latter can suffer from homoplasy caused by high mutation rates (i.e., identical by state, not identical by descent; Estoup et al., 2002).

The greater divergence between the W-ATL and E-ATL, compared to the E-ATL and IWP, demonstrates that the Mid Atlantic Barrier (an open ocean distance barrier) forms a stronger and more enduring barrier than the Benguela Upwelling System (a thermal barrier). The permeability of the latter barrier may fluctuate over time due to oscillations in climate (Hirschfeld et al., 2021). Fiji and New Caledonia have previously been identified as differentiated island populations, although New Caledonia showed less differentiation than Fiji from other sites (Glaus et al., 2020; Pirog et al., 2019). This was also attributed to the long distances across deep water trenches and is consistent with our results for Fiji. Similarly, individuals from Japan could be isolated by deep water trenches (> 500 m deep, e.g., Okinawa Trough), since the Bull Shark generally prefers shallow waters (< 160 m; Rigby et al., 2021). Yet, sea levels between Japan and continental Asia were low until 17 Kyr ago (Voris, 2000), although the high latitudinal location of Japan may have formed a thermal barrier at that time. Additionally, historical land-bridges such as the Taiwan Strait, Tokara Strait, or Tsushima/Korea Strait could have restricted dispersal between Japan and the IWP (Chen, Wang, et al., 2018; Yin et al., 2009).

Historical land-bridges also play a potential role for the genetic structuring of Bull Sharks across northern Australia. Despite a lack of structure inferred from the nuclear markers, the mitogenome ΦST differentiation showed three separate clusters (western, northern, and eastern Australia). Here, the Torres Strait land-bridge between Australia and Papua New Guinea has been implicated for genetic structuring in a variety of marine taxa across this region (Mirams et al., 2011). This signal from the Bull Shark haplotype network is less structured, and may indicate either recent separation or historical isolation followed by connectivity (e.g., Puckridge et al., 2013).

Our study shows strong influences of land-bridges, long ocean distances, cold upwelling or high latitudes, and deep-water trenches on the genetic population structure of the Bull Shark. We hypothesize that these barriers will equally affect elasmobranch species with similar dispersal capacity, distribution, and habitat use. To date, the global population structure of only a few large-bodied shark species has been investigated (e.g., Galapagos Shark, *C. galapagensis*; Sandbar Shark, *C. plumbeus*; and Scalloped Hammerhead, *S. lewini*; Daly-Engel et al., 2012; Pazmiño et al., 2018; Portnoy et al., 2010). Known biogeographic barriers have similarly affected the population structure of these species (reviewed in Hirschfeld et al., 2021). For example, the Galapagos Shark, which occupies insular habitats, shows limited connectivity across oceanic barriers, such as the East Pacific Barrier (Pazmiño et al., 2018). Similarly, both the Scalloped Hammerhead and Sandbar Shark exhibited increased connectivity along continuous coastlines within ocean basins but showed significant divergence due to the East Pacific Barrier and closure of the Isthmus of Panama (Daly-Engel et al., 2012; Portnoy et al., 2010). Consequently, our results may provide a foundation to construct hypotheses for other exploited and threatened species with similar vagility, ecology, and life-history characteristics (such as the Spinner Shark, the Graceful Shark, or the Dusky Shark).

## Fine-scale structure and sex-biased dispersal

Minor nuclear differentiation was detected between the southern and northern W-ATL locations. This concurs with other Bull Shark studies showing strong mitochondrial and weak nuclear population structure in the W-ATL (Karl et al., 2011; Sandoval Laurrabaquio‐Alvarado et al., 2021) and that long-distance movement is rare in the Gulf of Mexico (Carlson et al., 2010). This would suggest that a weak or recent barrier to dispersal may be present in this region. Such a barrier can be attributed to oceanographic features (e.g., Caribbean Current) or reproductive asynchrony caused by seasonal differences between the hemispheres (Carrillo et al., 2015; Carrillo et al., 2017; Castro, 2011; Jensen, 1976). Similarly, genetic differentiation may be present in the IWP (as suggested by Pirog et al., 2019), but our ability to detect significant nuclear differences is constrained by high gene flow and/or recent separation (Bailleul et al., 2018; Waples, 1998).

Within the IWP, the mitogenome results demonstrated maternal structure at a finer spatial scale than the SNP results, supporting the growing evidence of long-term female philopatry and male-biased dispersal in Bull Sharks (Karl et al., 2011; Sandoval Laurrabaquio-Alvarado et al., 2019; Tillett et al., 2012) and other elasmobranchs (reviewed in Chapman et al., 2015; Phillips et al., 2021). Specifically, the mtDNA shows a separation between the W-IO and E-IO/W-PAC, not evident from the nuclear DNA, and greater differentiation of the E-ATL. The presence of haplotypes from multiple distinct mitochondrial lineages in South Africa, Sri Lanka, Thailand, Fiji, and the Fitzroy River (Australia) represents signals of secondary female contact after historical isolation or low ongoing female gene flow. However, we see that almost every haplotype is unique to a sampling location (except for locations within Australia). This could indicate that each location has experienced sufficient female isolation for the mitogenome to mutate (i.e., complete lineage sorting), and/or is a consequence of high haplotype diversity as a result of sequencing complete mitogenomes. In addition, demographic events affect the non-recombining haploid mtDNA and recombining diploid nuclear DNA differently (Heist, 2012; Lawson Handley & Perrin, 2007), and, therefore, the mito-nuclear discordance may not necessarily be driven by sex-biased dispersal (Toews & Brelsford, 2012).

The spatial distribution of close-kin, and the cross-cohort HSPs in particular, is valuable to identify contemporary dispersal and sex bias therein (e.g., Feutry et al., 2017; Feutry et al., 2020). In the IWP, we found that all FSPs and most HSPs were sampled from the same sampling location. The 32 same-river, cross-cohort HSPs had an equal likelihood of being maternally (same haplotype) or paternally (different haplotype) related, indicating contemporary biparental philopatry to a finer scale than detected by conventional population genetic methods (Tillett et al., 2012). However, the seven cross-river, cross-cohort HSPs in northern Australia were most likely paternally related, suggesting a bias towards male dispersal between rivers. This unconfounded result supports the mito-nuclear findings and suggests that male-biased dispersal may be occurring at a broader scale.

Within Japan, the SNP data grouped the individuals into three separate clusters, with individuals from Okinawa clustering more closely with the IWP and those from the Urauchi River forming two separate groups. The sharks from the Churaumi Aquarium, Okinawa were originally sourced from Japanese waters near Okinawa Island and several individuals were known to be related. The close similarity of this group to the IWP cluster may represent ongoing genetic connectivity or recent colonisation, where the strong nuclear differentiation may be caused by an accumulation of mutations at the edge of a range expansion due to the more pronounced effect of drift on a small and recently established population (Peischl et al., 2013). This divergence may be reinforced through high relatedness within a small population and seasonal differences in parturition, which would explain why we observed multiple genetic clusters within the Urauchi River. The sample size within Japan was too small to accurately estimate kinship; thus, family structure due to extremely small population size cannot be excluded (e.g., Devloo‐Delva et al., 2019; Feutry et al., 2017). Similarly, the Fiji population contains many related individuals (see Glaus et al., 2020), yet these were not removed from our analyses as they could signify an artefact of a small population size (Waples & Anderson, 2017). Mitochondrial haplotypes from individuals in Japan and Fiji are part of the E-IO/W-PAC haplogroup, supporting that these are recently established populations.

## Species, sex, provenance, and kinship identification

Increasingly, there is need to monitor wild populations and the global trade of wildlife products through DNA forensics (Cardeñosa et al., 2022). Our DArTcap panel of 3,400 SNPs can identify species, sex, provenance, and kinship, which allows future monitoring of multiple demographic aspects at low cost (~AU$15 per sample; see Feutry et al., 2020). We were able to identify most of the individuals to species-level using the DArTseq and DArTcap data (i.e., Bull Shark, Pigeye Shark, Spinner Shark, Graceful Shark, Grey Reef Shark, Smalltail Shark, Dusky Shark,and Speartooth Shark), validated alongside mitochondrial species verification and a complete mtDNA reference database. Given the global scale of shark product trade (Clarke et al., 2006) and the morphological similarity between juveniles of many carcharhinid species, a selection of these SNP markers could be developed into a rapid tool for species identification of various shark products (Johri et al., 2019; Liu et al., 2017). While the species composition of shark fin trade has been studied using mtDNA (Cardeñosa et al., 2022; Fields et al., 2018), some species, like the Galapagos Shark and Dusky Shark, show mitochondrial introgression (Corrigan et al., 2017; Naylor et al., 2012). Here, the use of nuclear SNPs can resolve the ability to separate species with such introgression or incomplete lineage sorting (e.g., Kyne & Feutry, 2017; Liu et al., 2017). Moreover, the presence of sex-linked markers, specifically Y-linked markers, in the DArTseq data allowed us to identify the genetic sex of our samples when the visual sex information was missing. Unfortunately, the RNA baits failed to capture these markers appropriately with DArTcap sequencing, although future studies could assess sex ratios from fisheries or trade samples by redesigning the RNA baits for these Y-linked markers (e.g., Stovall et al., 2018).

The DArTcap panel also shows great promise and power to assign individuals to their respective populations (ocean-basin scale), and we estimated that a minimum of 100 markers are needed to achieve an accuracy of > 80 %. Few studies have specifically investigated the power of genetic markers to assign samples from traded shark products to their population of origin, and, to date, most forensic studies of shark trade have employed mtDNA to identify broad geographic origins (e.g., Cardeñosa et al., 2021; Chapman et al., 2009; Fields et al., 2020). Here, by genetically characterizing the source populations with 769 sharks and 3409 SNPs, the regional contributions of Bull Shark products to the global trade market can be estimated in future studies and highlight key regions of harvest to inform management actions. Furthermore, our results suggest a diagnostic SNP panel of 100–500 markers could be designed to identify species, sex, and provenance from tissue samples. New genotyping technologies, such as DNA microarrays, may provide a more cost-efficient method for monitoring of Bull Shark exploitation (e.g., Arenas et al., 2017; Wenne et al., 2016). However, this panel needs further testing prior to use as an enforcement tool. This will involve obtaining additional samples from under-sampled locations (e.g., E-PAC and E-ATL) and testing the assignment accuracy on these newly acquired samples.

Finally, we were able to identify close-kin relationships with the selected DArTcap markers. These relationships are important to estimate contemporary dispersal patterns (Feutry et al., 2017; Feutry et al., 2020) or total adult abundance of a population (Bravington et al., 2016). This application requires a large sample size (relative to the total population size) and sufficient biological information (such as sex, age-at-length, and age-at-maturity), which can pose a challenge in studies of rare or threatened species. Whilst the number of kin was limited in this data set, the identified kin pairs provided insights into local connectivity in Australia.

## Conservation and management considerations

Coastal shark species, such as the Bull Shark, are particularly susceptible to anthropogenic pressures such as fishing, shark control programs, and habitat modification, due to their close proximity to human populations (Knip et al., 2010). These species fulfil an important role in maintaining the ecosystem dynamics and functions (Ferretti et al., 2010), where a reduction in the abundance of top predators may lead to cascading effects in the food web and ecosystem health (e.g., Dudley & Simpfendorfer, 2006). Understanding the patterns of reproductive isolation of coastal sharks is essential to assess the impact of these threats at the population level (Frankham et al., 2017). Given our results of large-scale connectivity, we advocate for international collaboration to monitor the impacts on dispersing individuals (predominantly males) as they maintain the gene flow and genetic diversity between populations, and likely experience threats across many different countries. Similarly, coastal development near nursery areas may affect philopatric females disproportionally, with potential consequences for a wider region if other countries depend on recruitment from those nurseries. In addition, genetically isolated and small island populations, such as Japan and Fiji, currently face a number of threats, such as targets or bycatch in fisheries (Glaus et al., 2015) and require close monitoring as even low levels of catch may cause a population reduction. Any decline in abundance is unlikely to be replenished by migration from neighbouring populations. Overall, this study suggests that coastal species with comparable characteristics exhibit similar dispersal patterns, and consequently, risk and vulnerability assessments and management actions should be considered at the smallest spatial scales, in accordance with the philopatric behaviours of these species, in order to prevent local depletion.

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# Data Accessibility and Benefit-Sharing

**Genetic data**

Mitochondrial genomes are uploaded to GenBank (accession numbers: OP006762-OP007122)

Mitochondrial DNA alignment and SNP genotypes are available on DataDryad (TBA).

**Sample metadata**

Sample metadata and Rmarkdown document, which included all performed analyses discussed in this study, are available on DataDryad (TBA).

**Benefit-Sharing Statement**

Benefits Generated: A research collaboration was developed with scientists from the countries providing genetic samples, all collaborators are included as co-authors, the results of research have been shared with the provider communities and the broader scientific community (see above), and the research addresses a priority concern, in this case the conservation of the Bull Shark. In Australia and Papua New Guinea, we consulted with the Indigenous communities providing the biodiversity resources and worked with members of the communities to collect samples based on local knowledge. More broadly, our group is committed to international scientific partnerships, as well as institutional capacity building.

# Author Contributions

The study was designed by FDD, CPB, PMK, and PF. Samples were collected and contributed by PMK, JMB, DDC, PC, XC, GC, RD, JMD, ME, DF, LGB, KG, BIGG, MIG, SIH, SH, RWJ, SJ, GJ, JTK, HM, FHM, GJPN, JEGN, NMP, RDP, AFS, KT, BJT, JAVV, PF, and all people acknowledged in the paper. DNA was extracted by FDD. Mitogenome library preparation and sequencing was performed by FDD, RMG, and JRM. FDD performed the genetic analyses with input from CPB, SM, BDP, and PF. FDD drafted the manuscript and all authors (FDD, CPB, PMK, JMB, DDC, PC, XC, GC, RD, JMD, ME, DF, LGB, KG, BIGG, MIG, RMG, SIH, SH, RWJ, SJ, GJ, JTK, HM, JRM, FHM, SM, GJPN, JEGN, NMP, RDP, BDP, AFS, KT, BJT, JAVV, PF) revised and improved the manuscript.

# Tables

Table1: Description of the different *Carcharhinus leucas* data sets used in this study. The data generated by DArTseq and DArTcap genotyping was subdivided to account for unequal samples sizes and hierarchical clustering. The number of sharks and SNPs before and after quality filtering are presented.

Table 2: Description of the genetic variation for the full *Carcharhinus leucas* data set (DATA2: 769 sharks; 3,409 SNPs) and the full mitochondrial genomes (361 sharks; 16,708 bp). Diversity indices include the number of samples for the nuclear data (Nnu), the allelic richness (Ar), the observed heterozygosity (HO), the expected heterozygosity corrected for low sample size (uHE), the inbreeding coefficient (FIS) and its 2.5 % and 97.5 % confidence intervals (FIS2.5, FIS97.5), the number of monomorphic markers (MO), the number of samples for the mitochondrial data (Nmt), the nucleotide diversity for the mitogenomes (πmt), the haplotype richness (H), and the haplotype diversity (h). ‘NA’ values indicate that no or too few samples were available to estimate genetic variation. Diversity indices are divided by geographic regions in dark grey (E-PAC = eastern Pacific; W-ATL = western Atlantic; E-ATL = eastern Atlantic; IWP = Indo-West Pacific; JAP = Japan; and FIJ = Fiji). These regions are further separated by country or oceanographic location (GOC = Gulf of California; COR = Costa Rica; BRZ = Brazil; CAR = Caribbean Sea; GOM = Gulf of Mexico; WNA = Western North Atlantic; SIL = Sierra Leone; SAF = South Africa; MOZ = Mozambique; RUN = Réunion Island; SEY = Seychelles; ARP = Arabian Peninsula; SRL = Sri Lanka; TAI = Thailand; IND = Indonesia; PNG = Papua New Guinea; and AUS = Australia). Finally, Australian sampling locations were: FZR = Fitzroy River; VIR = Victoria River; DAR = Daly River; ADR = Adelaide River; DWC = Darwin Coastal; SAR = South Alligator River; EAR = East Alligator River; BMB = Blue Mud Bay; ROR = Roper River; TOR = Towns River; WER = Wenlock River; TRI = Trinity Inlet; CLR = Clarence River; SYH = Sydney Harbour; and UNK = Australian fisheries samples from unknown origin. Japan samples were sourced from: OKI = Churaumi Aquarium, Okinawa; and URR = Urauchi River.

Table 3: Pairwise comparison of fixation indices for *Carcharhinus leucas*. FST values are presented in the bottom diagonal and sequence-based ΦST are in the top diagonal. ‘NA’ values indicate that no or too few samples were available to estimate genetic differentiation. Asterisks (\*) indicate statistical significance of p < 0.05. No pairwise comparisons were significant after Bonferroni correction. Fixation indices are coloured from high (red) to low (blue) values. GOC = Gulf of California; COR = Costa Rica; BRZ = Brazil; CAR = Caribbean Sea; GOM = Gulf of Mexico; WNA = Western North Atlantic; SIL = Sierra Leone; SAF = South Africa; MOZ = Mozambique; RUN = Réunion Island; SEY = Seychelles; ARP = Arabian Peninsula; SRL = Sri Lanka; TAI = Thailand; IND = Indonesia; PNG = Papua New Guinea; FZR = Fitzroy River; VIR = Victoria River; DAR = Daly River; ADR = Adelaide River; DWC = Darwin Coastal; SAR = South Alligator River; EAR = East Alligator River; BMB = Blue Mud Bay; ROR = Roper River; TOR = Towns River; WER = Wenlock River; TRI = Trinity Inlet; CLR = Clarence River; SYH = Sydney Harbour; and UNK = Australian fisheries samples from unknown origin; OKI = Churaumi Aquarium, Okinawa; URR = Urauchi River; and FIJ = Fiji.

# Figures

Figure 1: Map indicating the *Carcharhinus leucas* sampling locations with red circles and the known species range distribution in yellow. The sample sizes for the SNP data are underlined with the number of samples before/after data filtering. The sample sizes for the mitogenome data are in *italics* with the number of samples before/after data filtering. Putative barriers for connectivity are indicated by dashed lines. GOC = Gulf of California; COR = Costa Rica; BRZ = Brazil; CAR = Caribbean Sea; GOM = Gulf of Mexico; WNA = Western North Atlantic; SIL = Sierra Leone; SAF = South Africa; MOZ = Mozambique; RUN = Réunion Island; SEY = Seychelles; ARP = Arabian Peninsula; SRL = Sri Lanka; TAI = Thailand; IND = Indonesia; PNG = Papua New Guinea; AUS = Australia; JAP = Japan; and FIJ = Fiji. Australian sampling locations were presented as an additional inset: FZR = Fitzroy River; VIR = Victoria River; DAR = Daly River; ADR = Adelaide River; DWC = Darwin Coastal; SAR = South Alligator River; EAR = East Alligator River; BMB = Blue Mud Bay; ROR = Roper River; TOR = Towns River; WER = Wenlock River; TRI = Trinity Inlet; CLR = Clarence River; SYH = Sydney Harbour; and UNK = Australian fisheries samples from unknown origin.

Figure 2: Population clustering analysis for the global *Carcharhinus leucas* data set with a subsample of Australian sharks (DATA3: 382 sharks; 1,849 SNPs). Panel A shows the Discriminant Analysis of Principal Components (DAPC) assignment barplot for K = 6 and 51 principal components (PC). Each bar represents an individual and is coloured according to the posterior membership probabilities. Panels B–D represent the principal component analysis (PCA) scatterplot, where each point represents an individual shark, triangles indicate the mean PCA score per sampling location, and colours represent the sampling country or oceanographic location. B) PCA scatterplot with PC1 on the x-axis and PC2 on the y-axis. C) PCA scatterplot with PC3 on the x-axis and PC4 on the y-axis. D) PCA scatterplot with PC3 on the x-axis and PC5 on the y-axis.

Figure 3: *Carcharhinus leucas* mtDNAhaplotype network, based on the full mitogenome (16,707–16,708 bp). Panel A presents the network for all 361 sharks from the eastern Pacific, eastern Atlantic, Indo-West Pacific, Japan, and Fiji. The distance between haplotypes reflects the number of mutations between them. Panel B shows the ‘eastern Indian Ocean/western Pacific/Japan/Fiji’ cluster in detail (285 sharks). Here, the number of mutations between haplotypes are represented by small black dots. In panels A and B, the size of the circles is equivalent to the square root of the number of individuals that share this haplotype. The colour and shape of each haplotype corresponds to the sampling location where they were found. The three black arrows indicate haplotypes that represent recent maternal movement between haplotype clusters.

Figure 4: Reassignment success of *Carcharhinus leucas* respective to the number of informative markers to assign simulated mixtures to their population of origin (E-PAC = eastern Pacific; W-ATL = western Atlantic; E-ATL = eastern Atlantic; IWP = Indo-West Pacific; Japan = Urauchi River, Japan; and Fiji). The simulated mixtures are based on a leave-one-out resampling method in *rubias*. The assignment success was tested based on 5–500 informative markers that were selected according to their high DAPC loadings contribution from the *adegenet* package.

Table 1

|  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- |
| **Data set** | **Genotyping method** | **Data description** | **Usage** | **Before filtering** | **After filtering** |
| DArTseq1 | DArTseq | Full global data set including replicates and other species | For species identification (allow more missing data and higher levels of heterozygosity) | 218 sharks; 250,945 SNPs | 218 sharks; 28,418 SNPs |
| DArTseq2 | DArTseq | Full global data set with only Bull Shark samples | For DArTcap marker selection | 182 sharks; 250,945 SNPs | 149 sharks; 17,029 SNPs |
| DATA1 | DArTcap | Full global data set including replicates and other species | For species identification (allow more missing data and higher levels of heterozygosity) | 1014 sharks; 37,537 SNPs | 1014 sharks; 5,053 SNPs |
| DATA2 | DArTcap | Global data set with confirmed Bull Sharks | For genetic diversity, FST, and provenance assignment | 868 sharks; 37,537 SNPs | 769 sharks; 3,409 SNPs |
| DATA3 | DArTcap | Global Bull Shark data set with a subset of 60 samples from Australia | Equalised sample size to allow unbiased clustering analyses | 430 sharks; 37,537 SNPs | 382 sharks; 1,849 SNPs |
| DATA4 | DArTcap | All western Atlantic Bull Shark samples | For genetic diversity estimates without bias from population structure | 117 sharks; 37,537 SNPs | 91 sharks; 931 SNPs |
| DATA5 | DArTcap | All Indo-West Pacific Bull Shark samples | For genetic diversity estimates without bias from population structure | 732 sharks; 37,537 SNPs | 635 sharks; 3,416 SNPs |
| DATA6 | DArTcap | IWP data with a subset of 60 samples from Australia | Equalised sample size to allow unbiased clustering analyses | 221 sharks; 37,537 SNPs | 189 sharks; 1,785 SNPs |

Table 2

|  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- |
|  | **E\_PAC** | **GOC** | **COR** | **W-ATL** | **BRZ** | **CAR** | **GOM** | **WNA** | **E-ATL/SIL** | **IWP** | **SAF** | **MOZ** | **RUN** | **SEY** | **ARP** | **SRL** | **TAI** | **IND** | **PNG** |  |
| **Nnu** | 16 | 0 | 16 | 97 | 50 | 2 | 37 | 8 | 1 | 586 | 22 | 11 | 28 | 36 | 15 | 12 | 5 | 4 | 9 |  |
| **Ar** | 1.1540 | NA | 1.1540 | 1.2580 | 1.0950 | 1.0650 | 1.0870 | 1.0840 | NA | 1.3450 | 1.1120 | 1.0860 | 1.1080 | 1.1120 | 1.1140 | 1.1100 | 1.1090 | 1.0970 | 1.1050 |  |
| **HO** | 0.0360 | NA | 0.0360 | 0.0500 | 0.0520 | 0.0490 | 0.0480 | 0.0500 | NA | 0.0610 | 0.0600 | 0.0560 | 0.0590 | 0.0600 | 0.0620 | 0.0610 | 0.0660 | 0.0610 | 0.0590 |  |
| **uHE** | 0.0410 | NA | 0.0410 | 0.0550 | 0.0570 | 0.0510 | 0.0530 | 0.0540 | NA | 0.0640 | 0.0630 | 0.0640 | 0.0630 | 0.0630 | 0.0650 | 0.0640 | 0.0660 | 0.0610 | 0.0620 |  |
| **FIS** | 0.0750 | NA | 0.0750 | 0.0536 | 0.0576 | -0.0481 | 0.0638 | 0.0369 | NA | 0.0472 | 0.0465 | 0.0959 | 0.0572 | 0.0398 | 0.0370 | 0.0356 | -0.0069 | -0.0098 | 0.0285 |  |
| **FIS2.5** | 0.0781 | NA | 0.0781 | 0.0748 | 0.0694 | -0.0306 | 0.0724 | 0.0327 | NA | 0.0421 | 0.0467 | 0.0985 | 0.0528 | 0.0312 | 0.0273 | 0.0126 | -0.0320 | -0.0293 | 0.0192 |  |
| **FIS97.5** | 0.1401 | NA | 0.1401 | 0.1082 | 0.1060 | 0.1257 | 0.1165 | 0.0984 | NA | 0.0579 | 0.0811 | 0.1568 | 0.0893 | 0.0606 | 0.0668 | 0.0614 | 0.0378 | 0.0551 | 0.0703 |  |
| **Mo** | 1370 | NA | 2562 | 3340 | 1771 | 2811 | 2075 | 2512 | 3340 | 301 | 2081 | 2443 | 2010 | 1914 | 2195 | 2353 | 2699 | 2835 | 2512 |  |
| **Nmt** | 16 | 1 | 15 | 0 | 0 | 0 | 0 | 0 | 1 | 302 | 7 | 0 | 16 | 18 | 19 | 12 | 6 | 6 | 15 |  |
| π**mt** | 0.0003 | NA | 0.0003 | NA | NA | NA | NA | NA | NA | 0.0021 | 0.0019 | NA | 0.0003 | 0.0003 | 0.0018 | 0.0030 | 0.0029 | 0.0004 | 0.0006 |  |
| **H** | 10 | 1 | 9 | NA | NA | NA | NA | NA | 1 | 142 | 6 | NA | 6 | 3 | 13 | 11 | 4 | 6 | 13 |  |
| **h** | 0.9250 | NA | 0.9143 | NA | NA | NA | NA | NA | NA | 0.9873 | 0.9524 | NA | 0.8083 | 0.6601 | 0.9415 | 0.9848 | 0.8000 | 1.0000 | 0.9810 |  |
|  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
|  | **AUS** | **FZR** | **VIR** | **DAR** | **ADR** | **DWC** | **SAR** | **EAR** | **BMB** | **ROR** | **TOR** | **WER** | **TRI** | **CLR** | **SYH** | **UNK** | **JAP** | **OKI** | **URR** | **FIJ** |
| **Nnu** | 444 | 3 | 0 | 49 | 18 | 15 | 88 | 47 | 18 | 9 | 10 | 24 | 5 | 83 | 65 | 10 | 44 | 9 | 35 | 25 |
| **Ar** | 1.1160 | 1.0830 | NA | 1.1140 | 1.1120 | 1.1160 | 1.1130 | 1.1110 | 1.1160 | 1.1000 | 1.1060 | 1.1120 | 1.1110 | 1.1200 | 1.1130 | 1.1080 | 1.2800 | 1.1020 | 1.1090 | 1.2820 |
| **HO** | 0.0610 | 0.0600 | NA | 0.0600 | 0.0600 | 0.0640 | 0.0600 | 0.0590 | 0.0650 | 0.0590 | 0.0590 | 0.0600 | 0.0690 | 0.0650 | 0.0600 | 0.0610 | 0.0610 | 0.0590 | 0.0620 | 0.0570 |
| **uHE** | 0.0640 | 0.0620 | NA | 0.0630 | 0.0630 | 0.0670 | 0.0630 | 0.0620 | 0.0660 | 0.0650 | 0.0630 | 0.0640 | 0.0680 | 0.0660 | 0.0630 | 0.0640 | 0.0610 | 0.0600 | 0.0600 | 0.0640 |
| **FIS** | 0.0440 | -0.0006 | NA | 0.0470 | 0.0392 | 0.0275 | 0.0409 | 0.0383 | 0.0212 | 0.0648 | 0.0434 | 0.0456 | -0.0145 | 0.0301 | 0.0448 | 0.0342 | 0.0155 | -0.0026 | 0.0006 | 0.0809 |
| **FIS2.5** | 0.0387 | -0.0016 | NA | 0.0448 | 0.0338 | 0.0190 | 0.0370 | 0.0333 | 0.0034 | 0.0614 | 0.0449 | 0.0441 | -0.0430 | 0.0062 | 0.0431 | 0.0270 | -0.0151 | -0.0273 | -0.0440 | 0.0955 |
| **FIS97.5** | 0.0544 | 0.0940 | NA | 0.0704 | 0.0695 | 0.0588 | 0.0596 | 0.0591 | 0.0410 | 0.1174 | 0.0948 | 0.0744 | 0.0249 | 0.0296 | 0.0676 | 0.0768 | 0.0170 | 0.0315 | -0.0057 | 0.1358 |
| **Mo** | 395 | 2914 | NA | 1727 | 2178 | 2204 | 1360 | 1779 | 2120 | 2517 | 2480 | 2033 | 2681 | 1320 | 1550 | 2452 | 1969 | 2653 | 2133 | 2114 |
| **Nmt** | 203 | 5 | 21 | 25 | 18 | 0 | 38 | 17 | 13 | 4 | 5 | 19 | 4 | 21 | 13 | 0 | 31 | 10 | 21 | 9 |
| π**mt** | 0.0006 | 0.0031 | 0.0004 | 0.0006 | 0.0007 | NA | 0.0002 | 0.0005 | 0.0001 | 0.0001 | 0.0001 | 0.0006 | 0.0000 | 0.0003 | 0.0006 | NA | 0.0004 | 0.0006 | 0.0001 | 0.0014 |
| **H** | 85 | 4 | 14 | 16 | 12 | NA | 18 | 12 | 7 | 3 | 3 | 14 | 2 | 12 | 11 | NA | 4 | 3 | 3 | 8 |
| **h** | 0.9777 | 0.9000 | 0.9333 | 0.9367 | 0.9216 | NA | 0.9346 | 0.9412 | 0.8718 | 0.8333 | 0.7000 | 0.9649 | 0.5000 | 0.9333 | 0.9744 | NA | 0.6903 | 0.7111 | 0.5286 | 0.9722 |

Table 3

|  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- |
|  | E-PAC | | W-ATL | | | | E-ATL | IWP | | | | | | | | | | | | | | | | | | | | | | | | Japan | | Fiji |
|  |  | |  | | | |  |  |  |  |  |  |  |  |  |  | Australia | | | | | | | | | | | | | | |  |  |  |
|  | **GOC** | **COR** | **BRZ** | **CAR** | **GOM** | **WNA** | **SIL** | **SAF** | **MOZ** | **RUN** | **SEY** | **ARP** | **SRL** | **TAI** | **IND** | **PNG** | **FZR** | **VIR** | **DAR** | **ADR** | **DWC** | **SAR** | **EAR** | **BMB** | **ROR** | **TOR** | **WER** | **TRI** | **CLR** | **SYH** | **UNK** | **OKI** | **URR** | **FIJ** |
| **GOC** |  | 0.124 | NA | NA | NA | NA | NA | 0.788 | NA | 0.968 | \*0.967 | 0.804 | 0.664 | 0.677 | 0.951 | 0.931 | 0.649 | \*0.951 | \*0.934 | 0.923 | NA | \*0.977 | 0.944 | \*0.988 | 0.988 | 0.985 | \*0.932 | 1.000 | \*0.966 | 0.936 | NA | 0.937 | 0.989 | 0.848 |
| **COR** | NA |  | NA | NA | NA | NA | 0.958 | \*0.913 | NA | \*0.969 | \*0.968 | \*0.877 | \*0.832 | \*0.887 | \*0.965 | \*0.951 | \*0.891 | \*0.959 | \*0.947 | \*0.944 | NA | \*0.975 | \*0.956 | \*0.979 | \*0.973 | \*0.974 | \*0.949 | \*0.975 | \*0.968 | \*0.954 | NA | \*0.957 | 0.981 | \*0.925 |
| **BRZ** | \*0.220 | \*0.237 |  | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA |
| **CAR** | \*0.229 | \*0.248 | \*0.007 |  | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA |
| **GOM** | \*0.225 | \*0.243 | \*0.004 | \*0.006 |  | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA |
| **WNA** | \*0.229 | \*0.251 | \*0.005 | \*0.003 | \*0.004 |  | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA |
| **SIL** | NA | NA | 0.467 | 0.473 | 0.473 | 0.476 |  | 0.788 | NA | 0.968 | 0.967 | \*0.802 | 0.662 | 0.676 | 0.950 | 0.930 | 0.646 | \*0.950 | \*0.933 | 0.922 | NA | \*0.977 | 0.943 | 0.988 | 0.988 | 0.985 | 0.931 | 1.000 | 0.966 | 0.935 | NA | 0.937 | \*0.989 | 0.846 |
| **SAF** | \*0.478 | \*0.497 | \*0.472 | \*0.473 | \*0.476 | \*0.479 | 0.098 |  | NA | 0.211 | \*0.159 | \*0.590 | 0.113 | 0.048 | 0.747 | \*0.782 | 0.276 | \*0.828 | \*0.813 | \*0.784 | NA | \*0.897 | \*0.808 | \*0.846 | \*0.736 | \*0.757 | \*0.800 | \*0.741 | \*0.849 | \*0.781 | NA | \*0.777 | \*0.884 | \*0.643 |
| **MOZ** | \*0.489 | \*0.508 | \*0.481 | \*0.482 | \*0.485 | \*0.488 | 0.107 | \*0.004 |  | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA |
| **RUN** | \*0.485 | \*0.502 | \*0.476 | \*0.478 | \*0.481 | \*0.483 | 0.104 | \*0.002 | \*0.006 |  | \*0.606 | \*0.776 | \*0.456 | \*0.520 | \*0.941 | \*0.919 | \*0.713 | \*0.933 | \*0.915 | \*0.911 | NA | \*0.958 | \*0.929 | \*0.963 | \*0.953 | \*0.954 | \*0.918 | \*0.956 | \*0.946 | \*0.925 | NA | \*0.931 | \*0.967 | \*0.864 |
| **SEY** | \*0.486 | \*0.502 | \*0.476 | \*0.478 | \*0.481 | \*0.483 | 0.107 | \*0.002 | \*0.006 | \*0.003 |  | \*0.776 | \*0.395 | \*0.399 | \*0.937 | \*0.916 | \*0.678 | \*0.929 | \*0.912 | \*0.907 | NA | \*0.955 | \*0.925 | \*0.959 | \*0.949 | \*0.950 | \*0.915 | \*0.952 | \*0.943 | \*0.922 | NA | \*0.928 | \*0.964 | \*0.862 |
| **ARP** | \*0.479 | \*0.496 | \*0.470 | \*0.472 | \*0.475 | \*0.477 | 0.102 | \*0.002 | \*0.005 | \*0.002 | \*0.002 |  | \*0.292 | \*0.387 | 0.135 | \*0.286 | \*0.203 | \*0.322 | \*0.252 | \*0.224 | NA | \*0.483 | \*0.256 | \*0.392 | 0.256 | \*0.287 | \*0.227 | 0.228 | \*0.362 | \*0.272 | NA | \*0.256 | \*0.326 | \*0.219 |
| **SRL** | \*0.492 | \*0.507 | \*0.480 | \*0.481 | \*0.484 | \*0.487 | 0.107 | \*0.003 | \*0.006 | \*0.003 | \*0.005 | 0.001 |  | 0.000 | \*0.391 | \*0.504 | 0.000 | \*0.563 | \*0.552 | \*0.500 | NA | \*0.692 | \*0.524 | \*0.561 | \*0.404 | \*0.434 | \*0.520 | \*0.393 | \*0.585 | \*0.490 | NA | \*0.476 | \*0.610 | \*0.334 |
| **TAI** | 0.492 | \*0.509 | \*0.480 | \*0.482 | \*0.484 | \*0.487 | 0.122 | \*0.011 | \*0.013 | \*0.010 | \*0.012 | \*0.009 | \*0.011 |  | \*0.550 | \*0.648 | 0.030 | \*0.713 | \*0.691 | \*0.646 | NA | \*0.825 | \*0.677 | \*0.729 | \*0.544 | \*0.580 | \*0.666 | \*0.539 | \*0.741 | \*0.638 | NA | \*0.621 | \*0.780 | \*0.451 |
| **IND** | 0.478 | \*0.497 | \*0.468 | \*0.470 | \*0.473 | \*0.475 | 0.102 | \*0.009 | \*0.013 | \*0.012 | \*0.014 | \*0.012 | \*0.014 | \*0.020 |  | \*0.424 | \*0.345 | \*0.463 | \*0.320 | \*0.255 | NA | \*0.705 | \*0.389 | \*0.762 | \*0.651 | \*0.673 | \*0.308 | \*0.629 | \*0.571 | \*0.409 | NA | \*0.425 | \*0.660 | \*0.321 |
| **PNG** | \*0.483 | \*0.497 | \*0.471 | \*0.472 | \*0.475 | \*0.478 | 0.108 | \*0.004 | \*0.012 | \*0.005 | \*0.006 | \*0.006 | \*0.006 | \*0.013 | \*0.015 |  | \*0.405 | \*0.290 | \*0.245 | \*0.153 | NA | \*0.486 | \*0.277 | \*0.475 | \*0.343 | \*0.380 | \*0.257 | \*0.429 | \*0.427 | \*0.116 | NA | \*0.411 | \*0.639 | \*0.339 |
| **FZR** | NA | \*0.512 | \*0.483 | \*0.483 | \*0.487 | \*0.489 | NA | 0.002 | 0.006 | 0.003 | \*0.007 | 0.005 | \*0.009 | 0.016 | 0.017 | 0.005 |  | \*0.439 | \*0.433 | \*0.375 | NA | \*0.611 | \*0.404 | \*0.469 | 0.197 | \*0.256 | 0.405 | 0.288 | \*0.543 | \*0.399 | NA | \*0.431 | \*0.666 | \*0.210 |
| **VIR** | \*0.483 | \*0.497 | \*0.472 | \*0.473 | \*0.476 | \*0.479 | 0.105 | 0.001 | \*0.005 | \*0.004 | \*0.004 | \*0.003 | \*0.004 | \*0.011 | \*0.014 | \*0.006 | 0.004 |  | \*0.063 | 0.039 | NA | \*0.042 | 0.030 | \*0.133 | 0.009 | 0.073 | \*0.069 | \*0.502 | \*0.440 | \*0.317 | NA | \*0.501 | \*0.667 | \*0.410 |
| **DAR** | \*0.485 | \*0.500 | \*0.476 | \*0.477 | \*0.480 | \*0.483 | 0.103 | \*0.001 | \*0.006 | \*0.002 | \*0.002 | \*0.002 | \*0.003 | \*0.011 | \*0.012 | \*0.003 | \*0.006 | \*0.002 |  | 0.025 | NA | \*0.206 | 0.005 | \*0.268 | 0.145 | \*0.194 | \*0.029 | \*0.377 | \*0.366 | \*0.259 | NA | \*0.386 | \*0.514 | \*0.358 |
| **ADR** | \*0.485 | \*0.501 | \*0.475 | \*0.477 | \*0.479 | \*0.482 | 0.108 | \*0.002 | \*0.005 | \*0.002 | \*0.002 | 0.001 | \*0.003 | \*0.011 | \*0.009 | \*0.004 | 0.001 | \*0.003 | \*0.001 |  | NA | \*0.223 | 0.028 | \*0.234 | 0.110 | \*0.156 | \*0.018 | \*0.335 | \*0.338 | \*0.184 | NA | \*0.356 | \*0.514 | \*0.312 |
| **DWC** | \*0.475 | \*0.493 | \*0.468 | \*0.469 | \*0.472 | \*0.475 | 0.099 | 0.001 | \*0.006 | \*0.004 | \*0.003 | \*0.003 | \*0.005 | \*0.012 | \*0.011 | \*0.005 | 0.003 | 0.001 | \*0.002 | \*0.002 |  | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA | NA |
| **SAR** | \*0.485 | \*0.500 | \*0.475 | \*0.476 | \*0.479 | \*0.482 | 0.105 | 0.001 | \*0.005 | \*0.003 | \*0.002 | \*0.002 | \*0.003 | \*0.009 | \*0.012 | \*0.004 | \*0.004 | \*0.002 | \*0.001 | \*0.001 | \*0.002 |  | \*0.148 | \*0.147 | 0.062 | \*0.148 | \*0.233 | \*0.720 | \*0.603 | \*0.527 | NA | \*0.687 | \*0.802 | \*0.594 |
| **EAR** | \*0.483 | \*0.499 | \*0.475 | \*0.476 | \*0.479 | \*0.482 | 0.104 | \*0.001 | \*0.006 | \*0.003 | \*0.002 | \*0.002 | \*0.002 | \*0.011 | \*0.013 | \*0.004 | \*0.004 | \*0.002 | \*0.001 | 0.001 | \*0.002 | 0.000 |  | \*0.245 | 0.103 | 0.163 | \*0.044 | \*0.449 | \*0.411 | \*0.294 | NA | \*0.435 | \*0.615 | \*0.363 |
| **BMB** | \*0.486 | \*0.502 | \*0.476 | \*0.477 | \*0.480 | \*0.483 | 0.109 | \*0.005 | \*0.008 | \*0.005 | \*0.006 | \*0.006 | \*0.007 | \*0.013 | \*0.016 | \*0.008 | \*0.009 | \*0.006 | \*0.005 | \*0.004 | \*0.006 | \*0.005 | \*0.005 |  | \*0.296 | \*0.331 | \*0.236 | \*0.873 | \*0.646 | \*0.528 | NA | \*0.691 | \*0.883 | \*0.512 |
| **ROR** | \*0.481 | \*0.502 | \*0.475 | \*0.476 | \*0.479 | \*0.482 | 0.098 | \*0.002 | 0.006 | \*0.004 | \*0.004 | \*0.003 | 0.003 | \*0.010 | \*0.013 | \*0.006 | \*0.006 | \*0.004 | \*0.002 | 0.003 | 0.003 | \*0.002 | 0.002 | \*0.006 |  | 0.000 | \*0.132 | \*0.911 | \*0.578 | \*0.394 | NA | \*0.574 | \*0.880 | \*0.326 |
| **TOR** | \*0.495 | \*0.508 | \*0.481 | \*0.483 | \*0.485 | \*0.489 | 0.114 | \*0.006 | \*0.011 | \*0.007 | \*0.006 | \*0.004 | \*0.008 | \*0.015 | \*0.019 | \*0.010 | 0.006 | \*0.007 | \*0.006 | \*0.005 | \*0.005 | \*0.005 | \*0.005 | \*0.008 | 0.005 |  | \*0.175 | \*0.885 | \*0.597 | \*0.428 | NA | \*0.598 | \*0.879 | \*0.367 |
| **WER** | \*0.488 | \*0.503 | \*0.477 | \*0.478 | \*0.481 | \*0.484 | 0.111 | \*0.003 | \*0.008 | \*0.002 | \*0.003 | \*0.002 | \*0.003 | \*0.012 | \*0.014 | \*0.005 | 0.004 | \*0.004 | \*0.002 | \*0.002 | 0.002 | \*0.001 | \*0.001 | \*0.006 | \*0.004 | \*0.006 |  | \*0.357 | \*0.348 | \*0.255 | NA | \*0.378 | \*0.529 | \*0.328 |
| **TRI** | 0.478 | \*0.498 | \*0.470 | \*0.472 | \*0.475 | \*0.477 | 0.103 | \*0.004 | 0.010 | \*0.007 | \*0.006 | \*0.005 | \*0.011 | 0.013 | 0.013 | 0.008 | 0.002 | 0.006 | 0.002 | \*0.006 | 0.003 | \*0.004 | 0.002 | 0.007 | 0.002 | 0.010 | 0.004 |  | 0.000 | 0.186 | NA | \*0.551 | \*0.882 | 0.000 |
| **CLR** | \*0.473 | \*0.488 | \*0.465 | \*0.466 | \*0.469 | \*0.472 | 0.103 | \*0.002 | \*0.005 | \*0.003 | \*0.003 | \*0.002 | \*0.004 | \*0.010 | \*0.012 | \*0.005 | \*0.004 | \*0.003 | \*0.002 | \*0.001 | \*0.002 | \*0.001 | \*0.001 | \*0.005 | \*0.003 | \*0.005 | \*0.002 | \*0.004 |  | \*0.188 | NA | \*0.570 | \*0.750 | \*0.178 |
| **SYH** | \*0.485 | \*0.500 | \*0.475 | \*0.476 | \*0.479 | \*0.482 | 0.104 | \*0.002 | \*0.005 | \*0.003 | \*0.003 | \*0.002 | \*0.003 | \*0.011 | \*0.013 | \*0.004 | 0.003 | \*0.002 | \*0.001 | 0.000 | 0.002 | \*0.001 | \*0.001 | \*0.005 | \*0.003 | \*0.005 | \*0.002 | \*0.004 | \*0.001 |  | NA | \*0.446 | \*0.666 | \*0.178 |
| **UNK** | \*0.484 | \*0.499 | \*0.473 | \*0.474 | \*0.477 | \*0.480 | 0.104 | \*0.003 | \*0.008 | \*0.005 | \*0.004 | \*0.005 | \*0.006 | \*0.015 | \*0.013 | \*0.009 | 0.005 | \*0.005 | \*0.003 | 0.003 | 0.003 | \*0.002 | \*0.002 | \*0.007 | \*0.004 | 0.005 | \*0.004 | 0.003 | \*0.002 | \*0.002 |  | NA | NA | NA |
| **OKI** | \*0.507 | \*0.516 | \*0.486 | \*0.488 | \*0.490 | \*0.494 | 0.125 | \*0.017 | \*0.024 | \*0.017 | \*0.018 | \*0.018 | \*0.023 | \*0.028 | \*0.023 | \*0.022 | \*0.018 | \*0.019 | \*0.016 | \*0.017 | \*0.018 | \*0.017 | \*0.017 | \*0.021 | \*0.019 | \*0.023 | \*0.018 | \*0.019 | \*0.016 | \*0.017 | \*0.016 |  | \*0.555 | \*0.611 |
| **URR** | \*0.509 | \*0.519 | \*0.492 | \*0.493 | \*0.496 | \*0.499 | 0.142 | \*0.021 | \*0.026 | \*0.023 | \*0.020 | \*0.021 | \*0.025 | \*0.033 | \*0.034 | \*0.024 | \*0.023 | \*0.024 | \*0.022 | \*0.021 | \*0.021 | \*0.021 | \*0.021 | \*0.027 | \*0.024 | \*0.028 | \*0.021 | \*0.024 | \*0.021 | \*0.022 | \*0.021 | \*0.032 |  | \*0.389 |
| **FIJ** | \*0.499 | \*0.514 | \*0.486 | \*0.487 | \*0.491 | \*0.493 | 0.116 | \*0.018 | \*0.023 | \*0.017 | \*0.017 | \*0.016 | \*0.019 | \*0.027 | \*0.026 | \*0.019 | \*0.017 | \*0.018 | \*0.018 | \*0.017 | \*0.019 | \*0.017 | \*0.016 | \*0.020 | \*0.019 | \*0.021 | \*0.018 | \*0.022 | \*0.016 | \*0.017 | \*0.018 | \*0.032 | \*0.037 |  |

Fig. 1

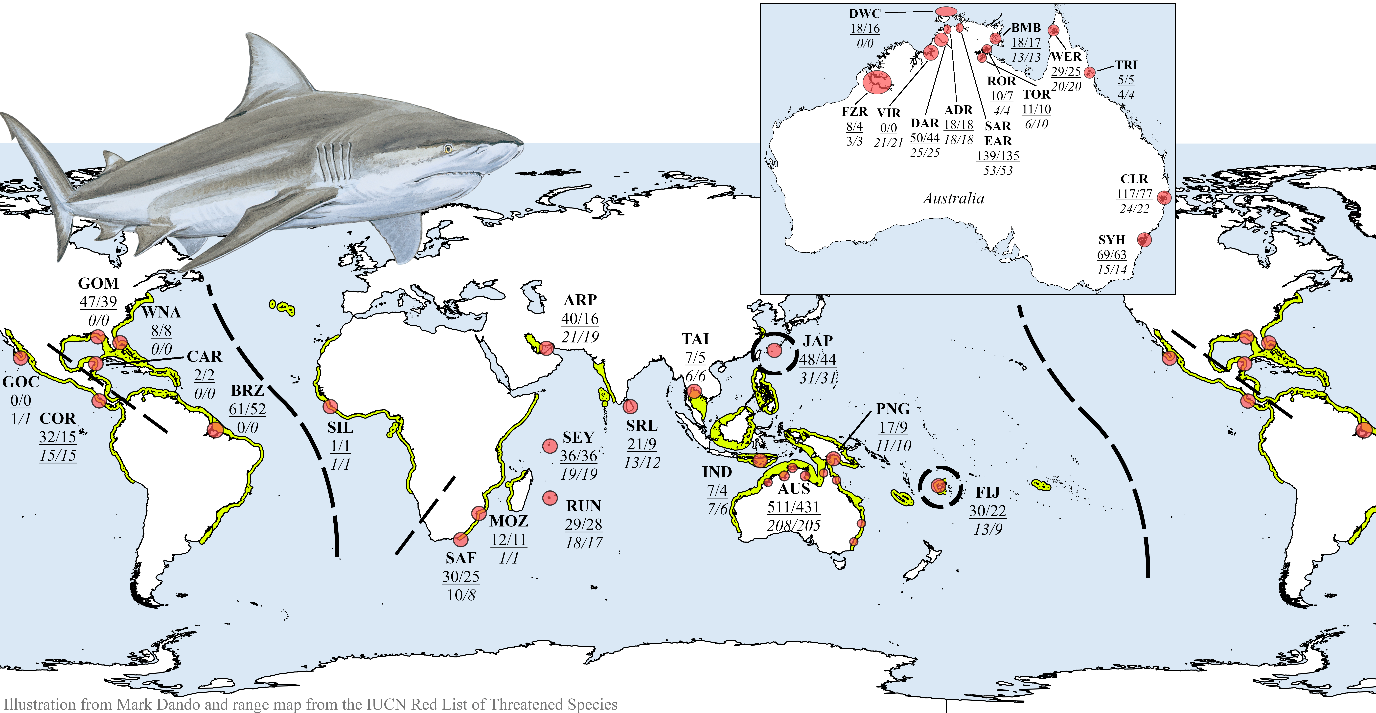


Fig. 2

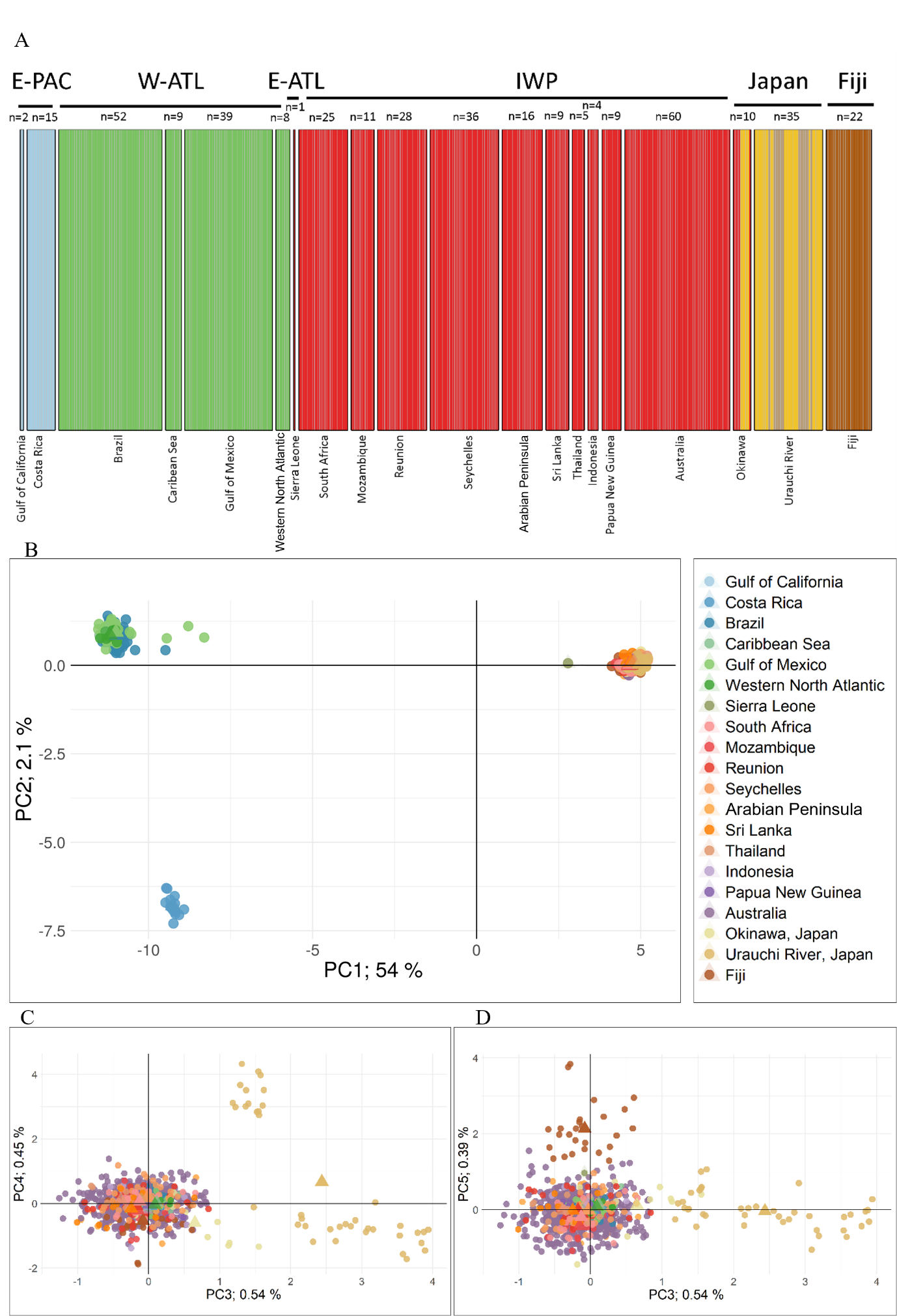


Fig. 3

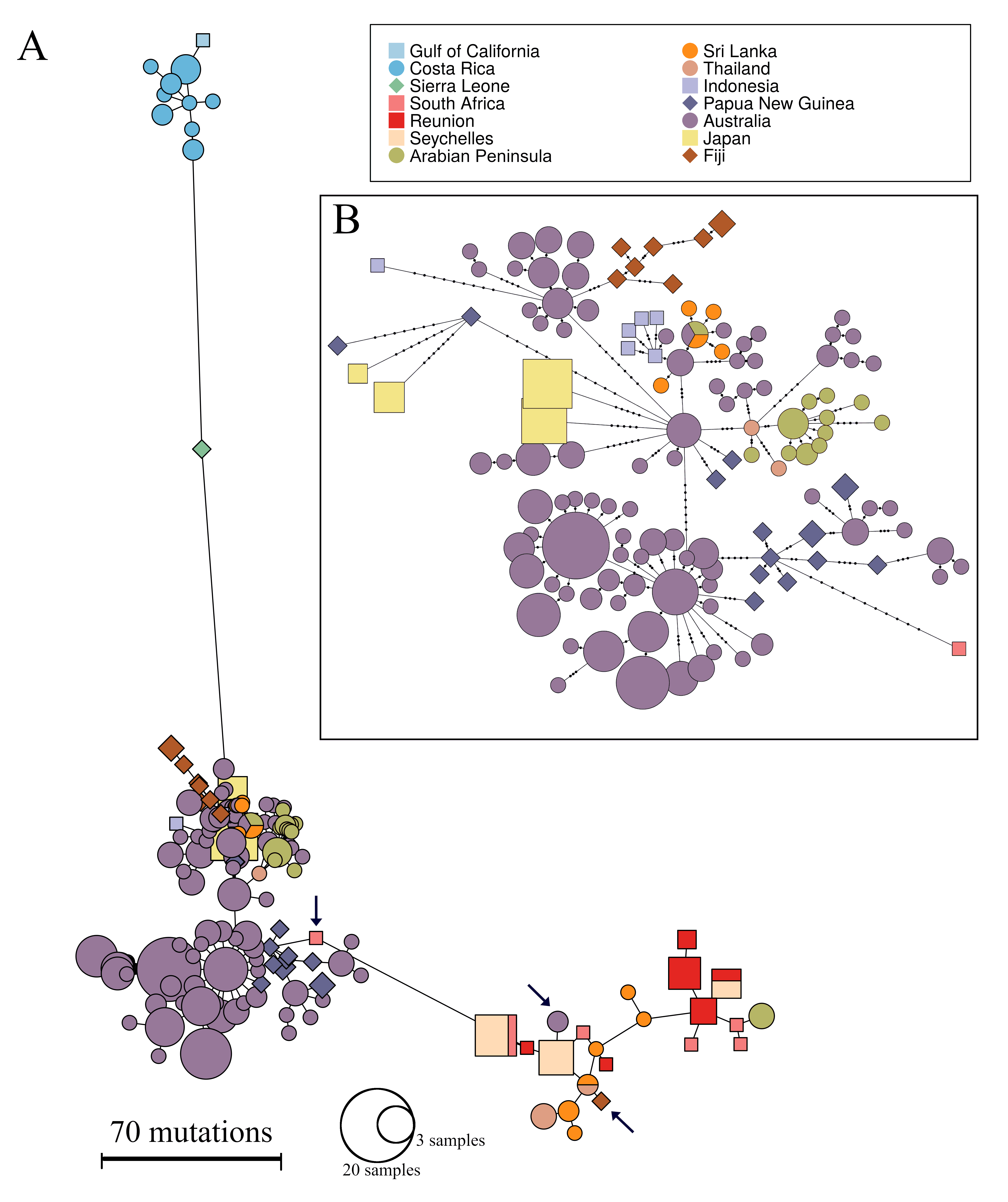


Fig. 4

