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# Viviparous sea snakes can be used as bioindicators for diverse marine environments

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## Abstract

Shallow tropical marine ecosystems are under great anthropogenic pressure due to habitat destruction, overfishing, shrimping, climate change, and tourism. This is an issue of global concern as such environments hold a tremendous biodiversity much of which remains to be described. The present situation urgently calls for time- and resource-efficient methods to identify and delineate the most valuable remaining areas and to set up priorities for their management and conservation. Using indicator species can be a way to accomplish this goal. In this paper we evaluate whether viviparous sea snakes can serve as bioindicators for other rare or cryptic tropical marine fauna. Based on seven generally acknowledged criteria for bioindicators, we argue that using viviparous sea snakes as bioindicators can help monitoring marine habitats to gauge the effects of climate change, habitat change and loss, decline in biodiversity and other anthropogenic changes. However, to maximize their efficacy as bioindicators, deeper knowledge about viviparous sea snakes natural history is urgently needed. Topics for expanded research programs include the taxonomy of some groups, their breeding and feeding biology, habitat selection and their geographical distribution. Despite these gaps in our understanding, we argue that viviparous sea snakes can be utilized as bioindicators of marine ecosystem health.

**Keywords:** anthropogenic changes, conservation, herpetology, marine habitat, monitoring

## Introduction

With the current rate of biodiversity loss and the fact that there are hardly any geographical areas not subjected to global change, it is crucial to use existing information on representative species to evaluate the conservation status of a variety of ecosystems (Alcala & Russ, 1990; Brischoux et al.,

2009a; Dudgeon et al., 2006; Elfes et al., 2013; Russ & Alcala, 2003; Russ et al., 2015; Sutherland et al., 2009; Vitousek et al., 1997; Zwickl, 2006). Most marine species are cryptic and difficult to sample and/or monitor, calling for the use of indicator species in a fashion that has been successfully implemented in conservation and management of forest and agricultural ecosystems. As a case in point, there is an urgent need to develop bioindicators for tropical shallow marine environments. Previous research embracing a wide range of ecosystems and taxonomic groups indicates that dependable and useful bioindicator taxa need to fulfil as many as possible of the following seven criteria (McGeoch, 1998; Niemelä, 2000; Noss, 1990; Pearson, 1994; Pearson & Cassola, 1992):

1. well-known and stable taxonomy
2. well-known ecology and natural history
3. easy to survey
4. higher taxa widely distributed geographically and over a breadth of habitats
5. diverse number of species specializes and sensitive to habitat changes
6. patterns observed in indicator taxa are also observed in related and unrelated taxa
7. present or potential economic importance

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More specifically, in order for organisms to be useful bioindicators they should be taxonomically well-resolved, widely distributed and their ecology should be sufficiently known to allow comparisons among sites. It is futile to believe that a single group of organisms or a single species can reflect the abundance and distribution of all other co-occurring taxa in a comprehensive way (Bilton et al., 2006; Carignan & Villard, 2002; Fleishman et al., 2005; Heino, 2010). Nevertheless, biodiversity monitoring in forest and riverine ecosystems amply demonstrates that some organisms are indeed efficient indicators for a wide range of co-occurring taxonomic groups (Heino, 2010). Coral reefs, river mouths, mangroves and other shallow tropical marine waters are extremely diverse ecosystems that are subject to rapidly increasing and destructive anthropogenic pressure. These habitats are in desperate need of improved and accurate biological monitoring (Alcala & Russ, 2006; Bishop & Alsaffar, 2008; Briers & Biggs, 2003; Brischoux et al., 2009a, 2011a; Cherel & Hobson, 2007; Ellis et al., 2012; Hindell et al., 2003; Ineich et al., 2007; Salas et al., 2004; Simboura & Zenetos, 2002; Weeks et al., 2010). However, tropical seas lag behind when it comes to biodiversity monitoring, largely because the ecological roles and trophic interrelationships of their organisms are less well understood. For example, many species escape observation and study because they are sheltered in the coral matrix, or in mud and sand holes or rocky bottoms. Taxa used as indicators also need to include species that show some degree of ecological specialization, sufficient to reflect relevant qualities of different trophic levels and taxonomic groups of the ecosystem under investigation (Brischoux et al., 2009a).

Sea snakes are a group of widely distributed top predators inhabiting shallow waters from the western Indian Ocean to the eastern Pacific Ocean (Rasmussen et al., 2011a). The 70 species of sea snakes share many adaptations and ecological roles. They are paraphyletic, comprising amphibious oviparous sea kraits (the '*Laticauda* Group': 8 species) and fully aquatic viviparous sea snakes (Hydrophiinae: - about 63 species) (Elfes et al., 2013; Nankivell et al., 2020). Recent research indicates that sea kraits (*Laticauda* spp.), by virtue of their diet specialised on eels, bear the promise of being useful bioindicators for surveying Anguilliform fish (eel-like fish) in coral reefs (Brischoux et al., 2009a; Ineich et al., 2007; Reed et al., 2002; Seret et al., 2008). Several species of viviparous sea snakes co-occur with sea kraits, and the former show a variety of additional morphological and behavioural adaptations for catching prey in structurally complex and taxonomically diverse tropical shallow marine environments such as coral reefs (Sanders et al., 2013a; Udyawer et al., 2016a). To explore the utility of viviparous sea snakes in these marine environments,

we here contrast the current state of our present knowledge of their natural history with the seven general criteria for bioindicators presented by Pearson (1994). To illustrate these points, we use the knowledge about the sea kraits value as bioindicator as a contrast to the information we have about viviparous sea snakes.

### **Criterion 1: well-known and stable taxonomy**

Most authors regard the eight recognized species of oviparous sea kraits as belonging to the same genus: *Laticauda* (Cogger & Heatwole, 2006; Gherghel et al., 2016; Heatwole et al., 2005; McCarthy, 1986; Slowinski, 1989), and as late as in 2005 and 2006 newly described species were added to the genus (Cogger & Heatwole, 2006; Heatwole et al., 2005). Only very few authors have suggested the existence of more than one genus of sea kraits (Kharin & Czeblukov, 2006; Kharin & Czeblukov, 2013; Wells, 2007), so in general this group's taxonomy is regarded as stable.

In contrast, the taxonomy and phylogeny of viviparous sea snakes have been debated for many years and has been developed significantly since the first comprehensive treatment by M.A. Smith (1926); (Kharin & Czeblukov, 2006, 2009; Lukoschek & Keogh, 2006; Rasmussen, 1997, 2002; Rasmussen et al., 2011a; Sanders et al., 2008; Smith, 1926; Voris, 1977). However, the last six years have seen an increasing consensus about phylogeny as well as taxonomy in this group (Lee et al., 2016; Rasmussen et al., 2014; Sanders et al., 2013b). At present, more than 85% of all known species have been included in phylogenies using either mitochondrial or genomic DNA (Lee et al., 2016; Sanders et al., 2013b). As a result, most authors now recognize 7 genera within the viviparous sea snakes: *Aipysurus* and *Emydocephalus*, collectively called the *Aipysurus* Group, which forms the sister group of all other viviparous sea snakes and is found mainly in the Australian region (Fig. 1; see Lee et al. [2016] for a recent phylogenetic study). Although not functionally treated as sea snakes here, the semi-aquatic monotypic genera *Ephalophis*, *Hydrelaps* and *Parahydropsis*, all found in Australian and New Guinea waters, are robustly placed at the base of the branch leading to the two last genera in the viviparous sea snakes group. The final group is composed of the genera *Microcephalophis*, mainly occurring in Asia, and *Hydrophis*, which occurs in both Asian and Australian regions (Sanders et al., 2013b). Based on the phylogeny in Sanders et al. (2013a) and Lee et al. (2016), a revised taxonomy of viviparous sea snakes invalidates the formerly recognized genera *Acalyptophis*, *Astrotia*, *Disteira*, *Enhydrina*, *Kerilia*, *Lapemis*, *Kolpophis* and *Thalassophina*, because these taxa are nested within the genus *Hydrophis*. Instead of erecting many new



**Figure 1.** Two of the authors handling an *Aipysurus laevis* (Broome, Northern Australia).

genera, a single genus *Hydrophis* is now recognised for all these taxa (Lee et al., 2016; Sanders et al., 2013b). In taxonomic arrangements resulting from phylogenetic analyses of viviparous sea snakes discussed above, very few species have uncertain phylogenetic placement, and this group is now regarded as more stable than ever before. Stable taxonomy allows species to be used for repeated monitoring, without the worry that the taxonomy will undergo a drastic re-arrangement making data incomparable over time. Stable taxonomy also establishes whether a particular group is geographically situated on a large diversity of ecosystems for which it can be used as bioindicator. However, there is no doubt that research in the near future will add new species to the *Hydrophis* Group, as new areas are investigated and more specimens are analysed with respect to morphological traits and genetic variation (Nankivell et al., 2020; Udyawer et al., 2018). Many widespread species-complexes include multiple cryptic species, many of which possess narrow distributions (Lukoschek, 2018a; Sanders et al., 2014; Udyawer et al., 2018; Ukuwela et al., 2017) based on preliminary molecular studies conducted by K.L. Sanders and colleagues. An additional 20 viviparous sea snake species remain to be described, increasing global marine snake diversity potentially by as much as 30% (Udyawer et al., 2018).

In the last few years, several new checklists and identification keys based on morphology have been published providing a solid regional base for sea snake identification fulfilling Criterion 1 (Alcala, 1986; Bussarawit et al., 1989; Buzas et al., 2019; Cao et al., 2017; Carpenter et al., 1997; Cogger & Heatwole, 2006; Ganesh et al., 2019; Guinea, 1994; Ineich & Laboute, 2002; Ineich & Rasmussen, 1997; Khan,

2004; Leviton et al., 2014, 2018; Mao & Chen, 1980; Mirtschin et al., 2017; Murphy et al., 1999; Rasmussen, 2001; Rasmussen et al., 2011b, 2014; Rezaie-Atagholipour et al., 2016; Somaweera & Somaweera, 2009; Weinell et al., 2019).

## **Criterion 2: well-known ecology and natural history**

### *Breeding biology*

Sea kraits forage at sea but lay their eggs on land, although detailed information about the breeding strategy of some species is still scarce (Bonnet et al., 2014). Mark-recapture studies in New Caledonia show that female *Laticauda saintgironsi* migrate more than 50 km seasonally between their home areas and coastal nurseries to lay eggs (Bonnet et al., 2014). Coastal nurseries attracting females from a larger area have not yet been described for other *Laticauda* species, but in the Philippines *Laticauda semifasciatus* and *L. laticaudata* lay eggs in crevices on the rough surface of the walls of caves (Bacolod, 1983) and in New Caledonia *L. laticaudata* and *L. saintgironsi* lay eggs in seabird burrows (Procellariiformes), in tree cavities, and in fields of large stone blocks. The above-mentioned habitats are also used as shelter by sea kraits (Brischoux & Bonnet, 2009; Ineich & Laboute, 2002). The information about seasonality of reproduction in sea kraits is scattered, but indicates that the breeding season follows the warm season in northern and southern populations, whereas it is more diffuse and spread out over time in areas closer to the equator (Bacolod, 1983; Bonnet, 2012; Bonnet et al., 2014; Brischoux & Bonnet, 2009; Brischoux et al., 2011b; Brongersma, 1956; Greer, 1997; Pernetta, 1977; Saint Girons, 1990; Shetty & Shine, 2002a; Shine et al., 2002a; Toriba & Nakamoto, 1987; Tu et al., 1990).

The viviparous sea snakes give birth in water and they rarely or never come ashore. In this group too, there are examples of ‘nursery areas’ where many females congregate to give birth (Porter et al., 1997; Udyawer et al., 2016b; Voris, 2015), but seasonal patterns of reproduction and litter size in viviparous sea snakes remain poorly understood, despite some published data (Bacolod, 1990; Bergman, 1943; de Silva et al., 2011; Fry et al., 2001; Greer, 1997; Lemen & Voris, 1981; Masunaga & Ota, 2003; Mirtschin et al., 2017; Rasmussen, 1989; Rasmussen et al., 2011a; Voris & Jayne, 1979; Wall, 1921; Ward, 2001).

Long term studies are urgently needed to obtain a more complete picture of breeding biology in viviparous sea snakes (Udyawer et al., 2018) as some species utilize nursery sites that are suitable for a range of other species sensitive to anthropogenic threats (e.g. fishing). Very likely, locating and delineating nursery areas of viviparous sea snakes can help identify areas suitable for a range of other cryptic species. It is



also important to understand the seasonality of the reproductive effort and use of nursery sites by juveniles as change to cyclic seasonal occurrence of juveniles may indicate a disruption in the ecosystem that might influence other co-occurring species.

### Feeding Biology

In several recent studies reviewed by Brischoux et al. (2009a), stomach contents of sea kraits (*Laticauda* spp.), which forage at sea, revealed more than 45 species of eels (Anguilliformes) of which 15 were new to science (Brischoux & Bonnet, 2009; Brischoux et al., 2007ab, 2009ab; Ineich et al., 2007). Tabata et al. (2017) investigated three species of *Laticauda* in Japan and confirmed that *L. colubrina* and *L. laticaudata* are specialized for feeding on eels, whereas *L. semifasciata* consumes various other families of fish as was reported from Taiwan (Mao & Chen, 1980).

Information about the diet of viviparous sea snakes is mainly based on gut content analyses (Glodek & Voris, 1982; Greer, 1997; McCosker, 1975; Rasmussen, 2001; Voris & Voris, 1983; Voris et al., 1978). Although there is comprehensive literature available about diet, such information is limited to only a few species (Fig. 2). Interestingly, from the perspective of bioindicator utility, several species appear to have a specialised diet, such as fish eggs, catfish, gobies, or eels, whereas others are more generalist feeders, consuming fish from more than 10 families (Voris & Voris, 1983).

Many viviparous sea snakes feed on secretive fish that spend most or all of their life hidden in crevices in reefs or in mud burrows (McCosker, 1975; Sanders et al., 2013b; Voris & Voris, 1983). Fishes with these kinds of secretive lifestyle are

found in several coral reef-dwelling fish orders, and many species are known from museum specimens only (Møller & Schwarzhans, 2008). As efficient predators in these very microhabitats, viviparous sea snakes can be used as bioindicator for secretive fish that are otherwise impossible to observe and monitor.

There is therefore an urgent need for more collection, for example when tissue-samples and stomach contents are taken from sea snakes before preservation, so that molecular/genetic analyses can be used to confirm and further explore the feeding biology and dietary diversity of sea snakes. A promising alternative method to laborious collection of rare and cryptic reef fish is the use of sea snakes as “sample catchers” as mentioned above; sea snake feeding studies should focus on the analysis of sea snake gut contents, so as to establish their natural dietary niche breadth (Udyawer et al., 2018).

### Sexual dimorphism and diet in sea snakes

In sea kraits, body and head sizes are larger in females than in males (Shine et al., 2002b). This sexual dimorphism is probably an adaptation to a sex-related divergence in prey size, as females eat larger eels than do males (Shetty & Shine, 2002b; Shine et al., 2002a).

Viviparous sea snakes also show sexual dimorphism in size, females growing larger than males (Burns & Heatwole, 2000; Shine et al., 2019), but further sexual dimorphism also exists, for example in scale rugosity and tail size (Avolio et al., 2006; Crowe-Riddell et al., 2016; Shine et al., 2019; Shine et al., 2003). Because different sexes target different life stages of prey, we would expect that a change in sex ratio of viviparous sea snakes might indicate a concomitant shift in prey life history.

The concept of the ecological niche is part of an influential body of general ecological theory addressing species' co-existence in resource-limited and competition-structured communities (Amarasekare, 2003; Hutchinson, 1978; Jeffries & Lawton, 1984; Schluter, 2000). Viviparous sea snakes, especially members of the genus *Hydrophis*, include many species with sympatric, partially- or completely overlapping distributions and similar habitat preferences (Sanders et al., 2013a). However, the partitioning of resources (such as diet, nest sites, resting sites and breeding sites) among such species is a topic of major interest, which is not well understood at present (Dunson & Minton, 1978; Lukoschek et al., 2007a; Udyawer et al., 2018). Substantial morphological variation in body and relative head size have been documented among some co-occurring species (Sherratt et al., 2018, 2019). Within the genera *Microcephalophis* and *Hydrophis*, extremely small-headed (microcephalic) species co-occur with large-headed



**Figure 2.** *Hydrophis elegans*, captured on the sea surface above depths of 145 m regurgitated a benthic eel indicating recent successful foraging on the sea floor (Broome, Galathea3 Expedition).



**Figure 3.** *Hydrophis brookii*, a small-headed (microcephalic) species (Songkhla Lake, Thailand).

(macrocephalic) species, a major body-plan difference apparently related to foraging mode and prey type variation (Sanders et al., 2013a) (Fig. 3). Therefore, viviparous sea snakes are well suited to test predictions about diet overlap and feeding niche segregation, likely increasing their value as bioindicators. A new model for rapid speciation and ecomorph differentiation in sea snakes was published by Sanders et al. (2013a) further suggesting the promise of viviparous sea snakes as a model study system for interface of community ecology and speciation theory.

Despite this, we still need information about ecology and natural history, but we regard Criterion 2 as fulfilled.

### Criterion 3: easy to survey

All species of sea kraits can be observed on coral reefs and land; the most terrestrial species *Laticauda colubrina* and its closest relatives are the ones most often seen a bit inland from the seashore proper (Heatwole et al., 2005; Liu et al., 2012). When on land, these snakes prefer downwind shores, sheltered harbors, rocky jetties, cavities, and crevices (Bonnet et al., 2009; Liu et al., 2012). Some species are adapted to climbing and can be found on top of cliffs and in trees (Bonnet et al., 2016).

Viviparous sea snakes, on the other hand, with few exceptions, spend all their time in water. The exceptions include three semiaquatic species (*Hydrelaps darwiniensis*, *Parahydrophis mertonii* and *Ephalophis greyi*), which are also found on mudflats during low tide. Sea snakes that inhabit coral reefs and lagoons can be surveyed by travelling slowly (at about four knots) in a small boat and visually identifying snakes

observed within 3 m of the path of the boat and to a depth of up to 3 m if the water is not turbid (A.R. Rasmussen, pers. obs.). At low tide, surveys can be conducted on foot, for example by searching reef flats (Guinea & Whiting, 2005). Also, baited remote underwater video station samplings have been used to investigate the distribution of viviparous sea snakes (Udyawer et al., 2014). The results from this technique were very promising and in congruence with previous trawl and underwater visual surveys. Monitoring viviparous sea snakes using passive acoustic monitoring is also very useful (Udyawer et al., 2016b) and can provide information about home range, spatial movement, and habitat selection (Udyawer et al., 2018).

At night many viviparous sea snakes can be observed on the water's surface, rendering them convenient to survey from small boats equipped with a strong light. This method permits easy catching of sea snakes by use of nets, with little risk of injury for snake or handler. This method can be quite successful in turbid waters such as rivers and estuaries (Porter et al., 1997) and even at sea (Bonnet et al., 2016; Buzas et al., 2019; Rezaie-Atagholipour et al., 2016). Other methods for catching sea



**Figure 4.** Diving in clear water during the day is an excellent way to catch sea snakes (Photo: Erik Frausing).

snakes involve diving or snorkelling in clear water during the day (Bonnet et al., 2016; Heatwole et al., 1978) (Fig. 4).

Finally, the use of environmental DNA (eDNA) as a monitoring tool for the presence/absence of species has gained tremendous momentum in recent years (Foote et al., 2012; Thomsen et al., 2012; Thomsen et al., 2016; Valentini et al., 2016). By extracting DNA directly from water or sediment samples, it is possible to PCR-amplify and sequence species-specific DNA markers ('barcodes') such as segments of the highly variable mitochondrial gene cytochrome B (*cytB*). Although this technology is still in a somewhat explorative phase where validation criteria and caveats must be thoroughly



established, it is clear that there is great potential for its use in monitoring — even for animals such as viviparous sea snakes which rarely occur in high densities. First and foremost, however, this method requires DNA sequence reference databases with sufficient data/taxon coverage to allow identification of eDNA sequences to the level of sea snake species. Fortunately, extensive *cytB* sampling for sea snakes is available in Genbank (Lee et al., 2016; Sanders et al., 2013b).

Thus, the above-mentioned established methods to survey and catch sea snakes are easy to implement and we regard Criterion 3 as fulfilled.

**Criterion 4: higher taxa widely distributed geographically and over a breadth of habitat**

Globally there are some 63 species of viviparous sea snakes found in the tropical and subtropical waters of the Indian Ocean and the Pacific Ocean, from the east coast of Africa to the Gulf of Panama (David & Ineich, 1999; Nankivell et al., 2020; Rasmussen et al., 2011c). Most species are found in the Indo-Malayan Archipelago, the China Sea, Indonesia and the Australian region (David & Ineich, 1999; Heatwole et al., 2005; Kharin & Czeblukov, 2009; Nankivell et al., 2020; Rasmussen, 1992; Rasmussen et al., 2011ab, 2014; Smith, 1926; see also Elfes et al., 2013 for a global species richness map of viviparous sea snakes). The historical biogeography of viviparous sea snakes has been inferred for some taxa believed to have expanded from the Australian region into Asia and then back again (Ukuwela et al., 2016; Ukuwela et al., 2017). There is only one truly pelagic sea snake (*Hydrophis platurus*), and it has the widest range of all sea snakes. *Hydrophis platurus* is found both in the Pacific and Indian Oceans, and a few specimens have been observed in the Atlantic Ocean (Bessesen & Galbreath, 2017; Branch, 1998). Other species, such as *H. caeruleus*, *H. cyanocinctus*, *H. curtus*, *H. ornatus*, *H. peronii*, *H. spiralis*, and *H. stokesii* have a very wide distribution in both Asia as well as in the Australian region ((Buzas et al., 2019; David & Ineich, 1999; Gritis & Voris, 1990; Ineich & Rasmussen, 1997; Rasmussen et al., 2014; Rezaie-Atagholipour et al., 2016). With their wide distribution some viviparous sea snake species have potential for use in existing monitoring programs across these regions. Other species, however, are known only from very restricted areas: *H. laboutei* only at Chesterfield Reef (Rasmussen & Ineich, 2000), *H. parviceps* from a very restricted area in south Vietnam (Rasmussen et al., 2012), *H. semperi* from Taal Lake in the Philippines (Garcia et al., 2014), and *H. sibauensis* from a limited area, more than 1000 km up-river from the coast in the Sibau River (an affluent Kapuas River, on Borneo Island; Rasmussen et al., 2001).

Over the last 40 years a large number of new distributional records have been added and about 11 new sea snake species have been described (Cogger & Heatwole, 2006; Dotsenko, 2011; Heatwole et al., 2005; Kharin, 1983, 1984, 1985, 2005; Nankivell et al., 2020; Rasmussen et al., 2001; Rasmussen & Ineich, 2000; Rasmussen et al., 2011c; Sanders et al., 2012a; Ukuwela et al., 2012). We can expect that more species still will be described in the future, as many areas suitable for sea snakes have not yet been explored, especially those between the two hotspot areas in the South China Sea and Northern Australia (Elfes et al., 2013). Moreover, some of the known species may harbor morphologically cryptic species in less well studied parts of their geographical range (D'Anastasi et al., 2016; Sanders et al., 2015; Udyawer et al., 2018).

The majority of observation involving viviparous sea snakes are made at depths less than 100 m (Greer, 1997); however, a few observations of specimens have been recorded as deep as 245 m (Crowe-Riddell et al., 2019). Some species are found in shallow waters in and around coral reefs, (*Emydocephalus* spp.; Alcalá et al., 2000; Lukoschek & Shine, 2012; Fig. 5). Others are restricted to mangroves in and on mudflats (*Hydrelaps darwiniensis*; Sweet, 1989). Some species seasonally occur in rivers and river mouths (e.g., *H. brookii*, *H. melanosoma*, *H. obscurus*, *H. schistosus*, and *H. torquatus*; Voris, 2015), whereas others are recorded most often in deeper water (e.g., *H. curtus*, *H. cyanocinctus*, *H. laboutei*, *H. ornatus*, *H. ocellatus*, *H. spiralis* and *H. czeblukovi*; Buzas et al., 2019; Cao et al., 2014; Smith, 1974). Some recent studies indicate that species in the *Aipysurus* Group have a more distinct geographical population structure compared with species in the *Hydrophis* Group, which have intraspecific population differentiation lacking geographically-based genetic structure



**Figure 5.** Characteristic head of *Emydocephalus annulatus*, normally found in shallow waters around and in coral reefs forage for fish eggs (Scotts Reef, Australia).

(Lukoschek et al., 2007a; Lukoschek & Shine, 2012; Nitschke et al., 2018). Most species in the *Hydrophis* Group are found in a variety of microhabitat types, such as soft sediment habitats, sea grass beds, marine gravel bottoms, rivers, and river mouths. Species of the *Aipysurus* Group also occur in a variety of coral reef and sand bottom marine habitats, but less frequently in riverine microhabitats (Cogger, 2000; Heatwole, 1999; Lukoschek et al., 2013; Porter et al., 1997; Rasmussen, 1987).

In summary, viviparous sea snakes are undoubtedly widely distributed, with substantial variation in range size among species. Collectively, they utilize a breadth of marine and estuarine habitats and, at the same time, some species are microhabitat generalists whereas others are more specialist.

Consequently, we argue that viviparous sea snakes meet Criterion 4 for use as bioindicators.

**Criterion 5: diverse number of species specialised and sensitive to habitat changes**

Some investigations in New Caledonia indicate that species of *Laticauda* are very sensitive to environmental change and pollution from dense human populations (Goiran et al., 2017; Goiran & Shine, 2013). Specific information on habitat use primarily is limited to Australia and New Caledonia (Brischoux et al., 2011b; Brischoux & Shine, 2011; Dunson, 1975; Karthikeyan & Balasubramanian, 2008; Lukoschek et al., 2007b; Padate et al., 2009; Shine, 2005), complemented by a few interesting studies from Asia (Voris, 2015). Most species occur in shallow habitats such as coral reefs, muddy or sandy areas, sea grass beds, mangroves, river mouths or rivers, lakes, or rocky areas.

A recent case study indicates that viviparous sea snakes are very sensitive to environmental changes; Ashmore Reef in the Timor Sea was long regarded as a hotspot for viviparous sea snake diversity with at least 14 recorded species (Guinea, 1995; Guinea & Whiting, 2005; Minton & Heatwole, 1975; Smith, 1926). However, by the late 1990s or early 2000s, species began to disappear from the reef and during a survey in 2010 (Lukoschek et al., 2013) only two species were recorded (*A. laevis* and *H. stokesii*) and follow-up surveys in 2012 and 2013 found no sea snakes (Guinea, 2013; Sanders et al., 2014) demonstrating a dramatic decline. The disappearance of 14 species from a single reef in such a short time (Elfes et al., 2013; Lukoschek, 2018b; Lukoschek et al., 2013) is a mystery. Bycatch, pollution, and seabed changes due to trawling are all known to affect sea snake numbers negatively. None of the above disturbances occurred in Ashmore Reef, giving no explanation for the disappearance of so many species. However, in a presentation at the World Congress of Herpetology (2020),

M. Guinea suggested that human disturbance may be to blame, further indicating that sea snakes may be sensitive to environmental change as an ‘early warning’ system for unknown processes as indicators of localized ecosystem change. In selected areas in Australia, commercial sea snake fisheries and by-catch have been studied during the past 30 years, indicating a similar negative impact of these activities as in other areas (Fry et al., 2001; Heales et al., 2008; Ward, 1996, 2000; Wassenberg et al., 2001; Wassenberg et al., 1994; Zhou et al., 2012).

Studies in the South China Sea indicate that sea snakes have declined as well; in the Gulf of Thailand, populations disappeared from rivers (*Hydrophis torquatus*, *H. klossi*; Rasmussen et al., 2011a). Also, on the island of Borneo, the number of specimens (collected previously in great numbers at river mouths) appears to have declined (A.R. Rasmussen, pers. obs. 2005). Trawl fisheries in the Andaman Sea catch fewer specimens in recent years, and fishers state that snake size has declined, indicating possible habitat degradation (Phuket Port; Rasmussen, 1997). A small 2000/2001 survey in Cambodia suggested similar results (A.R. Rasmussen, unpublished data).

In conclusion, many indications of sea snake population sensitivity to habitat change are apparent. Combined with the fact that numerous microhabitat- and diet-specialized species exist, leads us to argue that Criterion 5 is met as well.

**Criterion 6: patterns observed in indicator taxa are also observed in other related and unrelated taxa**

As mentioned before, there are large marine areas hosting viviparous sea snakes that are under great anthropogenic pressure due to habitat destruction, pollution, over-fishing, and climate change (Elfes et al., 2013). This will without doubt also negatively affect other related and unrelated taxa, such as water snakes (Homalopsidae), wart snake (Acrochordidae), burrowing eels (Anguilliformes) and corals (Anthony, 2016; Arvedlund, 2009; de Silva et al., 2010; Kannan et al., 2019; Murphy, 2007). Longitudinal datasets for viviparous sea snakes are available from multiple locations (Ashmore Reef, New Caledonia, Malacca Strait) and these can be utilized to examine for correlations with changes in population size, relative abundances, and demography of secretive fish species (Goiran & Shine, 2013; Lukoschek et al., 2013; Shine et al., 2019; Voris, 1985, 2015).

Combining diet data from viviparous sea snakes with occurrence data of threatened fish species (e.g., IUCN Red List) is also a possible way to meet Criterion 6.

**Criterion 7: present or potential economic importance**





**Figure 6.** Sea snakes collected alive by Vietnamese fishermen for export to China (Cao et al., 2014).

“The sociological justification associated with indicator species is often one that requires mollification more than major economic impact. Without minimizing the significance of this criterion, even economic potential and minimal economic impact may be sufficient to rebut objections that might have eliminated an otherwise ideal indicator taxon” (Pearson, 1994).

Since at least 1934, sea snake meat and skins have been used commercially in the Philippines (Bacolod, 1990), Australia, Japan, Taiwan, Thailand, and Vietnam (Padate et al., 2009; Rasmussen et al., 2011a; Warrell, 1994; Wassenberg et al., 2001; Fig. 6). Vietnam squid fishermen alone harvest 82 tons of viviparous sea snakes annually in the Gulf of Thailand, roughly equal to 225,500 individual sea snakes (Cao et al., 2014). To our knowledge this constitutes one of the largest harvest activities of venomous snakes and marine reptiles in the world today.

Obviously, Criterion 7 is fulfilled when it comes to sea snakes. However, most sea snake fisheries in the Indian Ocean and in the Pacific are neither monitored nor reported in the literature and are thus beyond the control of local governments and conservation agencies. We therefore recommend that further studies also focus on the economic use of sea snakes and its impact on populations.

## Conclusions

Previous investigations of sea kraits as bioindicators show that they are excellent for surveying Anguilliform fish assemblages in coral reefs (Brischoux et al., 2007ab, 2009a; Ineich et al., 2007). Sea kraits thus meet several of the

bioindicator criteria discussed above, but we argue that viviparous sea snakes have even better prospects of being good bioindicators because: 1) they have a wide distribution in tropical and subtropical waters of the Indian and Pacific Oceans (David & Ineich, 1999; Rasmussen et al., 2011a; Ukuwela et al., 2016); 2) the group includes feeding specialists and generalists (Heatwole, 1999; McCosker, 1975; Rasmussen, 2001; Voris & Voris, 1983; Voris et al., 1978), 3) there are many specialized species, but the group as a whole covers a broad range of marine feeding niches (Cogger, 1975, 1992; Heatwole & Cogger, 1994; Heatwole et al., 1974; Lukoschek et al., 2007a; Rasmussen & Ineich, 2010; Sanders et al., 2010, 2012b), and 4) we know that they are affected by anthropogenic activities.

A further advantage of using viviparous sea snakes as bioindicators is that, being active and efficient predators, they “sample” and thus assess the diversity and the abundance of many secretive fish groups occurring in a variety of marine habitats (estuaries, sandy bottoms, muddy bottoms, as well as in coral reefs and rocky substrates). Another fundamental advantage is that viviparous sea snakes are easy to catch when they come to the surface to rest and breathe, permitting monitoring at low cost and requiring only simple equipment (Bonnet et al., 2016; Brischoux & Bonnet, 2008; Buzas et al., 2019; Porter et al., 1997; Rasmussen & Hay-Schmidt, 2008). In waters with high visibility, such as coral reefs and shallow sandy bottoms, viviparous sea snakes are easy to catch and handle, even when researchers themselves are submerged (Bonnet et al., 2016; Brischoux et al., 2009a; Guinea & Whiting, 2005; Heatwole et al., 1978; Rasmussen & Elmberg, 2009; Shine et al., 2003, 2004).

We argue that viviparous sea snakes have great potential as bioindicators for monitoring marine habitats during this time of great climate change, habitat change and loss, decline in biodiversity, and other anthropogenic changes. This effort will require deeper knowledge of the taxonomy of some groups, their breeding and feeding biology, habitat selection, and distribution. However, we assert that even given our current knowledge limitations, viviparous sea snakes should be considered as suitable bioindicators of marine ecosystem health.

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