Mandibular sawing in a snail-eating snake

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The jaws of vertebrates display a striking diversity in form and function, but they typically open and close like a trapdoor rather than sliding like a saw. Here, we report unique feeding behaviour in the blunt-headed snail-eating snake, Aplopeltura boa (family Pareidae), where the snake cuts off and circumvents the indigestible part (the operculum) of its prey in the mouth using long sliding excursions of one side of the mandible, while the upper jaws and the mandible on the other side maintain a stable grasp on the prey. This behaviour, which we call ‘mandibular sawing’, is made possible by extraordinarily independent movements of the jaw elements and is a surprising departure from usual feeding behaviour in vertebrates.

Talos, the Greek mythological inventor, invented a saw inspired by a serpent jawbone or a fish skeleton1, but nonetheless, the jaws of snakes or other vertebrates usually do not act like a saw due to anatomical constraint on the jaw movements (but see2). Because the diversity in vertebrate jaws represents modifications of the homologous apparatus, their form and function are strongly constrained by the phylogeny3. Advanced snakes differ from most other vertebrates in their ability to move the left and right jaws virtually independently and swallow their prey using alternate movements of the two sides of the jaws4. Snail-eating snakes of Southeast Asia (Pareidae) or the Neotropics (Dipsadinae) retain the unilateral mobility of the jaws, but their feeding apparatus is further modified; these snakes have lost the articulation between the upper and lower jaws in contrast to most other snakes, and this makes the lower jaws of the snail-eating snakes extensively mobile and allows the mandibles to perform independent sliding excursions4–9. Several species use asynchronous retractions of the mandibles to extract snails from their shells or ingest slugs10–16. Feeding behaviors of most species of those peculiar snakes, however, have never been described perhaps due to limited access to these tropical, nocturnal, and secretive animals.

The blunt-headed snail-eating snake, Aplopeltura boa, is a pareid species that feeds on snails, including operculate species17. Gastropodan opercula are indigestible for snakes18, like their shells, and therefore are potentially harmful when consumed (e.g., they may cause intestinal obstruction). Indeed, feeding experiments showed that a snail-specialist pareid did avoid eating all of three species of sympatric operculate snails, whereas it consumed most of sympatric non-operculate snail species19. We collected A. boa and a syntopic, abundant operculate snail, Leptopoma sp. (Cyclophoridae) in a rainforest in Borneo and observed the snake feeds on the tough prey. Also, we investigated the relative abundance of operculate and non-operculate snails in the habitat of A. boa.

Results and discussion
We found that operculate snails were more abundant than were non-operculate snails: 34 and 22 individuals were encountered, respectively. Among snails of moderate size (shell width, 10–20 mm), which were often consumed by A. boa, operculate snails were more than 10 times as abundant as were non-operculate snails (25 and 2 individuals were encountered, respectively).

Individuals of A. boa (n = 8) readily preyed upon Leptopoma sp. (n = 30) in our feeding trials. Upon capture, the snakes immediately inserted their mandibles into the aperture of the shell and then extracted the operculate soft body using the mandibles. After extraction, the snakes regurgitated the extracted snail and precisely repositioned it so that the operculum protruded out of the mouth and the junction of the operculum and the soft body came to lie along the mandible on the right (n = 24, 80%) or the left (n = 6, 20%) side. From this position, the snakes moved the side of the relevant mandible backward and forward, while the snail was held in the stable

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position by the upper jaws and the mandible on the other side. These mandibular movements were especially vigorous when the mandible was being retracted. Six to 51 (median, 14) strokes of these sliding movements resulted in the removal of the operculum (Fig. 1, Supplementary Information, Videos S1, S2). Snakes consumed solely the soft part of the snails, while discarding the shell and the operculum (n = 30, 100%). Discarded opercula retained little soft tissue. The duration of extraction, repositioning, and sawing processes ranged 36–274 (median, 96), 9–459 (median, 66), and 15–428 (median, 30) seconds, respectively (Supplementary Information, Table S1). Thus, the snakes spent considerable time for handling of the extracted snail (reposition and sawing). The sequence of feeding was highly consistent among all cases, and the routine operculum-removing behaviour presumably allows *A. boa* to regularly consume the otherwise hazardous prey, which is abundant in its habitat.

The feeding apparatus of *A. boa* is illustrated in Fig. 2. *Aplopeltura boa* exhibits a set of derived morphological features known in other pareids and dipsadines, including short snout, short pterygoids, reduced supratemporals, long mandibles, and comb-like mandibular teeth. The skull of *A. boa* is short and tall, in which the snout is very short, and the orbits are exceptionally large. The pterygoids are greatly shortened, and their posterior ends are completely detached from the quadrate-mandibular joint. The quadrates are remarkably long and stout, extending ventrally rather than ventrolaterally. The mandibles are long and carry dense teeth, which are more robust than the maxillary teeth. The size and interval of mandibular teeth gradually change along the mandible; the anterior teeth are larger and sparser. The lower jaw unit (the mandible and the quadrate) is L-shaped, and the mandible travels anteroposteriorly when the quadrate swings backward and forward, as anticipated or observed in other pareids or dipsadines. This mechanism enables independent, substantial anteroposterior excursions of the mandible (Fig. 3), which is used for the extraction and the sawing processes during feeding on the operculate snails. The elongated quadrates are likely to contribute to long mandibular excursion.

Most of > 3,700 species of snakes swallow their prey whole, and prey-breaking behaviours are known only from a few species that feed either on crabs20–23 or termites24–26, which may be relatively easily broken into segments. These snakes grasp their arthropod prey with their jaws and break it apart usually using the movements of the head or the trunk. In contrast, *A. boa* cuts its mollusk prey using independent movements of the lower jaw. The mandibular sawing is, therefore, a surprising evolutionary solution for the limbless animals to utilize new food. This dexterous behaviour is especially surprising given that few other vertebrates, if any, are able to sever food in the mouth using unilateral sliding movements of a jaw element like *A. boa*. The evolutionary invention of sawing was evidently made possible by the unique feeding mechanism in the snail-eating snakes. Extensive mobility of the mandibles is a convergent trait in the two distinct lineages of snakes (pareids and dipsadines) that feed on slugs or snails, suggesting it is an adaptation to feeding on their slippery prey4–9. It is likely that acquisition of the free mandibular apparatus promoted the subsequent evolution of the novel behaviour and has resulted in functional versatility of the free-moving jaw elements.

Most pareids and some dipsadines have a larger number of teeth on the right mandible than on the left as feeding specialization to extract dextral (clockwise-coiled) snails13,15. There is a cline in the degree of the denticional asymmetry in correlation with diets, where snail-specialist species have highly asymmetrical mandibles, whereas slug-specialist species have symmetrical mandibles13,15. However, *A. boa* is an exception of this pattern...
because individuals exhibit only weak mandibular asymmetry despite its snail diet13. In the phylogeny, *A. boa* is
nested within the derived clade with mandibular asymmetry27, suggesting the presence of additional selective
forces toward the mandibular symmetry in this species. By showing the additional role of its mandibles (cutting
the prey), our results suggest functional trade-offs in *A. boa* (typical comb-like teeth in snail-eating snakes are
expected to facilitate extraction by providing a firm grip on the prey but probably are not optimal to cut the prey
tissue), highlighting the importance of behavioural studies to understand selective forces on functional units.

**Methods**
This study was conducted in the Gunung Mulu National Park, Sarawak, Malaysia (4.0440°N, 114.8144°E). We
collected *A. boa* and *Leptopoma* sp. by visual search and brought them to the Research Centre in the national
park for the feeding trials. All but three snakes that were used for other studies were released at the site of capture
after trials.

Feeding trials were conducted between 2100 and 0400 h. We fed *A. boa* with *Leptopoma* sp. and video-
recorded feeding events with infrared video cameras (HC-VX985M, Panasonic; FDR-AX30, Sony).

We conducted census survey on snails by walking the habitats of *A. boa* carefully looking for snails and
counting all snails larger than 5 mm in shell width. This survey was conducted on two nights in August and
December, respectively.

We conducted micro CT-scanning on six specimens of *A. boa* held at Kyoto University or Sarawak Forest
Department (specimen nos: KUHE27025, KUHE56016, KUHE57425, KUHE59285, SRC01008, and SRC01009).

All procedures followed the Animal Experiment Guideline of Kyoto University and were approved by the
ethical review committee of the Graduate School of Human and Environmental Studies of Kyoto University and

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**Figure 2.** Feeding apparatus of the blunt-headed snail-eating snake, *Aplopeltura boa*. CT images of the skull
and the jaws from left lateral (a), posterior (b), ventral (c), and dorsal (d) views and the quadrate and the
mandibles on the left (e) and right (f) sides. m, mandible; p, pterygoid; q, quadrate. These images are from the
specimen KUHE59285.
the Research, Development and Innovation Division of the Sarawak Forest Department (approval no. 30-A-7). All fieldwork was permitted by the State Government of Sarawak, the Sarawak Forest Department, and the Gunung Mulu National Park (permission nos. (133)JHS/NCCD/600–7/2/107, WL72/2018, WL103/2018, and WL68/2019).

Received: 27 March 2020; Accepted: 6 July 2020
Published online: 29 July 2020

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**Acknowledgements**
Funding was provided by the Asahi Glass Foundation (to K.N. and Y.K.), the Japan Society for the Promotion of Science (Grant No. 18J00809 to Y.K.), the Shikata Memorial Trust for Nature Conservation (to I.F.), and JST/JICA, SATREPS (to K.N.). We thank the State Government of Sarawak and the Sarawak Forest Department for permitting us to conduct the research, the Gunung Mulu National Park for permission and cooperation in the field, R. Fukuyama for assistance with field research, N. Morimoto, M. Nakatsukasa, and S. Hara for cooperation in CT scanning, and A. M. Durso for valuable comments on the manuscript.

**Author contributions**
Y.K. and I.F. conceived/designed the study and collected data. K.N., M.Y.H., and T.K. provided research support. Y.K. analyzed data and wrote the manuscript. All authors reviewed and edited the manuscript and approved the final version.

**Competing interests**
The authors declare no competing interests.

**Additional information**
Supplementary information is available for this paper at [https://doi.org/10.1038/s41598-020-69436-7](https://doi.org/10.1038/s41598-020-69436-7).

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