

# Spontaneous neoplasia in the western clawed frog Xenopus tropicalis

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D			Number of frogs	Number of	E	
Breeding colony	Date of birth	Sex	at stock center (May 28, 2020)	frogs with neoplasia	Breeding	Neoplasia size (mm)
NH-III-4	June 22, 2004	Female	13	1	NH-III-4	$4.9 \times 3.4 \times 3.8$
NH-VIII-7	Feb. 7, 2013	Female	19	4	NH-VIII-7 case 1	6.4 imes4.9 imes5.1
IC-III-10	Mar. 15, 2013	Female	16	2	NH-VIII-7 case 2	5.2 × 3.7 × 2.7
NA-XIV-1	Feb. 18, 2014	Female	6	1	IC-III-10 case 1	$3.2 \times 2.9 \times 2.5$
NBH-VI-11	Apr. 30, 2014	Female	21	3	IC-III-10 case 2	3.5  imes 3.2  imes 1.5
NBH-VI-7	June 11, 2014	Female	19	2	NA-XIV-1	$17.4 \times 15.4 \times 12.6$
NA-XV-1	Aug. 20, 2015	Female	12	2	NBH-VI-11	11  imes 6.4  imes 5.1

**Figure 1.**: (A–C) Representative images of spontaneous neoplasia in *Xenopus tropicalis*. (A) External morphology of the neoplasia. The arrow indicates the black stone-like nodules, and the arrowhead indicates the white-red nodules. The upper right panel depicts the frog for which the neoplasia was resected. (B) The layered structure in the black stone-like nodules. Hematoxylin–eosin (HE) staining of a histological section of the black stone-like nodule. The white arrow indicates the layered structure. Scale bar indicates 50  $\mu$ m. (C) The cyst-like structures and blood vessels in the white-red nodule. HE staining of a section of the white-red nodule. The white arrows blood vessels. Scale bar indicates 50  $\mu$ m. (D) A table showing the number of frogs with neoplasia in each colony. Regarding colony names, the first 2–3 alphabets indicate the strain names, the following Greek numbers indicate the inbred generations, and the Arabic numbers (with case numbers) indicate the colony identifiers. Most of the frogs maintained at the stock center were shipped to users as animal resources, and the remaining frogs as of May 28, 2020, were subjected to this analysis. (E) A table showing the size of neoplasia; the size was determined by measuring diameters at large and small ends, and the height above the skin surface for each neoplasia.

## Description

*Xenopus tropicalis* is an excellent model organism for studies on vertebrate development and regeneration (Horb *et al.*, 2019) and is also useful for the study of tumor formation (Van Nieuwenhuysen *et al.*, 2015; Naert *et al.*, 2016). Spontaneously occurring neoplasia in amphibians have been reported, such as in *X. laevis, Rana pipiens*, and *Andria japonicus* (McKinnell *et al.*, 1968; Meyer-Rochow *et al.*, 1991; Kawasumi *et al.*, 2012). In *X. laevis*, one of 4,000 frogs (a 2.5–3-year-old female), which were maintained in artificial outdoor ponds, displayed renal adenocarcinoma without other developmental disorders, indicating that the tumor formation was quite rare in *X. laevis* (Meyer-Rochow *et al.*, 1991). In contrast to *X. laevis*, 3.66 % of R. pipiens in the wild heterogeneous populations had renal adenocarcinoma (McKinnell *et al.*, 1968), which was experimentally induced by herpesvirus (Granoff, 1973). Currently, there are no reports of spontaneous neoplasia in *X. tropicalis*, although recently developed genome editing technologies such as TALEN and CRISPR/Cas9 offer opportunities to induce neoplasia by simple disruption of tumor suppressor genes (Van Nieuwenhuysen *et al.*, 2015; Naert *et al.*, 2016), which would allow modeling of human cancer in this species. The spontaneously occurring neoplasia in *X. tropicalis* could provide excellent opportunities to understand how the genetic background of this species influences the neoplasia phenotypes in combination with disruption experiments of tumor suppressor genes.

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The National Bioresource Project (NBRP) of X. tropicalis in Japan has successfully developed the four highly inbred wild-type strains, Nigerian A (NA), Nigerian H (NH, previously named Yasuda), Nigerian BH (NBH, previously named Golden), and Ivory Coast (IC) (Igawa et al., 2015). These strains are available for international research communities upon request. We found spontaneous neoplasia formation in the frog stocks at the NBRP (Figure 1). Analysis of frogs from seven colonies with different genetic backgrounds and chronological ages (16, 7, 6, or 5 years) identified neoplasias formed in various parts of the body, including the dorsal side, ventral side, dorsal head, dorsally on limbs, and eye (Figure 1D). Almost all frogs had a single site of neoplasia; just one frog from the NBH-VI-11 colony had multiple sites of neoplasia. The neoplasias of X. tropicalis could be divided into two types: black stone-like nodules (Figure 1A, white arrow) and white-red nodules (Figure 1A, white arrowhead). Most of the frogs were harboring only the black stone-like nodules, although a few frogs were harboring both types of nodules simultaneously (Figure 1A). Previous studies have reported that the histological analysis of amphibian and reptile neoplasias revealed the renal mass in A. japonicus exhibiting trabecular pattern and the epidermal mass in the Furcifer pardalis consisting of concentric keratin material (Kawasumi et al., 2012; Meyer et al., 2019). Histological analysis of X. tropicalis neoplasia using hematoxylin-eosin (HE) staining showed that melanocytes were enriched in the black stone-like nodules and that a layered structure was present (Figure 1B, white arrow). In contrast, cyst-like structures and blood vessels were present in the white-red nodules (Figure 1C, white arrowheads and black arrows, respectively). We then measured the neoplasia size (Figure 1E). Two frogs from the NA-XIV-1 colony had a large neoplasia, over 17 mm at the large end, over 15 mm at the small end, and over 12 mm in height (Figure 1E). The average length of the neoplasia was 7.3 mm at the large end, 4.1 mm at the small ends, and 3.5 mm in height. Therefore, it will be of interest if the frequency and size of neoplasia vary among the inbred strains on disruption of tumor suppressor genes via genome editing.

## Methods

### Request a detailed protocol

Neoplasias were fixed with 4% paraformaldehyde in phosphate-buffered saline (PBS) overnight at 4°C and were successively washed with PBS containing 0.1 % Tween 20. Fixed neoplasias were frozen in Optimum Cutting Temperature Compound, and cryosectioned with a thickness of 10 µm. Sections were stained with HE by a standard method. Each slide was photographed under a Zeiss Axio Vert. A microscope using an AxioCam MRc digital camera.

### Reagents

*X. tropicalis* strains used in this experiment were maintained at the Amphibian Research Center, Hiroshima University. All treatment procedures for *X. tropicalis* were approved by Yamagata University Animal Research Committee (#310300). For HE staining, we used the following reagents: Mayer's hematoxylin solution and 1% eosin Y solution (FUJIFILM Wako Pure Chemical Corporation, Osaka, Japan), Tissue-Tek optimum cutting temperature compound (Sakura Finetek, Tokyo, Japan).

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