

The amazing and anomalous axolotls as scientific models

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Abstract

Ambystoma mexicanum (axolotl) embryos and juveniles have been used as model organisms for developmental and regenerative research for many years. This neotenic aquatic species maintains the unique capability to regenerate most, if not all, of its tissues well into adulthood. With large externally developing embryos, axolotls were one of the original model species for developmental biology. However, increased access to, and use of, organisms with sequenced and annotated genomes, such as *Xenopus laevis* and *tropicalis* and *Danio rerio*, reduced the prevalence of axolotls as models in embryogenesis studies. Recent sequencing of the large axolotl genome opens up new possibilities for defining the recipes that drive the formation and regeneration of tissues like the limbs and spinal cord. However, to decode the large *A. mexicanum* genome will take a herculean effort, community resources, and the development of novel techniques. Here, we provide an updated axolotl-staging chart ranging from one-cell stage to immature adult, paired with a perspective on both historical and current axolotl research that spans from their use in early studies of development to the recent cutting-edge research, employment of transgenesis, high-resolution imaging, and study of mechanisms deployed in regeneration.

KEYWORDS

axolotl, *Ambystoma mexicanum*, embryonic development, staging, regeneration

1 | INTRODUCTION

Ambystoma mexicanum, the Mexican salamander commonly known as the axolotl, is an endangered amphibian with amazing regenerative capabilities. Although adult animals range in size, axolotls typically grow up to 20–35 cm in length from head to tail and live an aquatic lifestyle.¹ As natural predators, axolotls are capable of eating many organisms that cross their path. However, due to deterioration of their natural ecosystem, axolotls currently hold a critically endangered status.^{2,3}

Unlike most other amphibious salamanders, axolotls are neotenic animals that retain juvenile traits throughout their lifetime.⁴ Their close relatives, like the tiger

salamander (*Ambystoma tigrinum*), undergo metamorphosis as they mature, losing the fringed gills and caudal fin that make axolotls so distinctive.⁵ Their neoteny is possible because axolotls lack thyroid-stimulating hormone, a precursor to thyroxine: the necessary component to kick-start metamorphosis. In fact, an injection of iodine or thyroxine can be used to stimulate the transition, but the results are stochastic and regeneration capacity can be reduced in some cases.^{6,7} Demircan et al. identified reductions in some tissues' regenerative capacities and complete inhibition in others after thyroid hormone treatment.⁴ In contrast, Monaghan et al. found that while body size had no effect on regeneration, thyroxine-induced metamorphosis reduced regeneration rates by twofold and produced forelimb and digit abnormalities.⁷

Further research is needed to pinpoint the regenerative outcome of induced metamorphosis in axolotl and understand the mechanisms of growth regulation during regeneration.

A current gap in axolotl research is the application of new, precise tools for understanding cells at the individual and collective levels, and studies of gene and protein expression and function across development and regeneration. These experiments are key to support comparative developmental studies and identify conserved and divergent regulatory modules controlling axolotl developmental and regenerative programs. So much can be done with these animals, such as gain and loss of function,^{8–10} CRISPR-mediated transgenesis,^{10,11} single-cell characterization,^{12–16} live imaging,^{17–20} transplants and cell lineage tracing,^{20–27} and the use of *ex vivo* explants²⁸ to compare the developmental processes of this unique salamander to the robust body of avian, frog, zebrafish, and mouse developmental research. For example, axolotl gills can regenerate, yet they are one of three modes of respiration at the organism's disposal, and their full role remains unknown.²⁹

Axolotls have the potential to become more prevalent developmental models, similar to *Xenopus*, but the continuity of tools and resources is lacking. Specifically, there are few stock resource centers for ease of stocking, housing, and sharing transgenic animals. However, the NIH-funded Ambystoma Genetic Stock Center (AGSC) has created Sal-Site (<https://ambystoma.uky.edu/quick-links/sal-site>) to provide information and access to resources for investigators.^{30,31} Recent work characterizing mutant laboratory axolotls identified that all individuals from the AGSC contain small portions of the tiger salamander (*Ambystoma tigrinum*) genome, as there appears to be genetic cross-contamination.³² These findings provoke the questions: how similar are laboratory axolotls to those in the wild, and how might discoveries differ based on genomic differences? Other tools like cross-species tools for gain and loss of function, validated antibodies that recognize axolotl proteins, and interspecies comparison of transcriptional networks would greatly improve knowledge acquisition capabilities.

2 | NATURAL HISTORY OF THE AXOLOTL

Axolotls have been used as research organisms for over 150 years, and their vast potential stems from a humble beginning.³³ Native axolotls originated in Lake Texcoco, thriving along the lake's banks, and later, the canals of the Aztec city-state of Xochimilco. The Spaniards took notice of the Aztec people's fondness for eating this

aquatic species in the mid-16th century.³⁴ Then, during the Spanish conquest and colonial rule, Xochimilco expanded rapidly, to the detriment of axolotls. The 19th century saw the first scientific interest in this species, and in 1863, six axolotls were transported from Mexico to the Jardin Des Plantes in Paris. These original six individuals propagated the majority of the present-day research axolotl population. As a result, many axolotls used in research have high genetic similarity, with an inbreeding coefficient of 35%, which substantially exceeds the threshold of 12% that indicates breeding between first cousins and concerns ecologists and geneticists greatly.³⁵ This inbreeding decreases the viability of axolotls as a genetic model and potentially increases their disease susceptibility. Despite this challenge, axolotls spread as laboratory animals due to the ease of year-round breeding in captivity and naturally-occurring developmental mutations.³⁶ However, axolotls are near extinction in their native Mexico, begetting a lack of genetic diversity in wild axolotls as well. Therefore, without remediation, science may lose secrets hidden in the axolotl's genetic diversity on both fronts.³⁷ Despite these challenges, the axolotl makes up for these genetic pitfalls with many redeeming qualities.

3 | BACK TO THE FUTURE: AXOLOTLS AS MODELS FOR EMBRYONIC DEVELOPMENT

Amphibian embryos are an excellent model system for the study of cell and developmental biology due to their large sizes, resiliency, and large clutch sizes. As the major limb and spinal cord regenerative model system, axolotls are well studied in juvenile and larval stages;^{10,38} however, we lack detailed information about the molecular mechanisms that drive their early developmental processes. While axolotls retain their juvenile characteristics into adulthood, their development proceeds through distinct stages (Figure 1).^{39,40} In contrast to many other aquatic research models, axolotls utilize internal fertilization, externally laying single one-cell embryos over the course of several hours.⁴¹ Morphologically, early axolotl development appears to proceed similarly to other aquatic amphibians like *Xenopus laevis* or *Xenopus tropicalis*. For example, axolotls develop more rapidly at warmer temperatures, as noted previously.^{39,40} At their preferred cooler temperatures (17–18°C), axolotls progress through rapid cleavage stages (Figure 1, cleavage-blastula, 0–24 hours post laying, HPL), stereotypical gastrulation (Figure 1, gastrula, 24–54 hours HPL) and neurulation (Figure 1, neurula, 54–72 hours HPL). Neurulation is followed by tailbud and tadpole stages

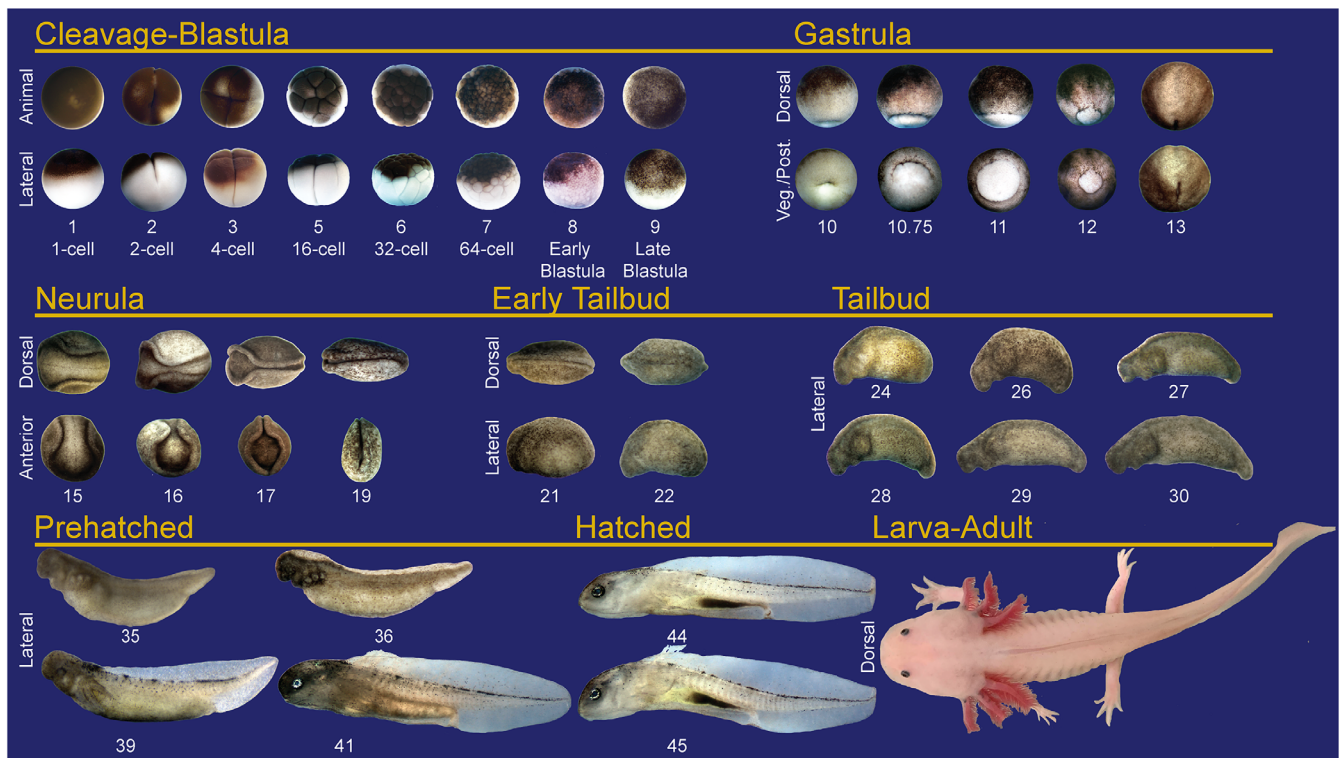


FIGURE 1 Staging series for *Ambystoma mexicanum* from fertilization through maturation. Stages were grouped into eight different categories: cleavage-blastula (0–24 hours post laying, HPL), gastrula (24–54 hours HPL), neurula (54–72 hours HPL), early tailbud (72–83 hours HPL), tailbud (83–122 hours HPL), prehatched (122–342 + HPL), hatched (342 hours to 1–3 months postlaying/fertilization), and larva-adult. Hatched larva to sexually mature adult can take up to 18 months depending on tank density. At cleavage/blastula stages, both animal and lateral views are shown. Similar to *Xenopus laevis* and *tropicalis*, *Ambystoma* embryos have pigment differentials on the animal and vegetal poles, the animal poles being dark and the vegetal poles light colored and filled with yolk even in leucistic animals as shown. At gastrula stages, we show both dorsal and posterior views. As gastrulation proceeds, the blastopore closes and the animals begin to neurulate. At neurula stages, we show dorsal and anterior views. After neural tube closure, the embryos begin axis elongation throughout the early tailbud and tailbud stages. At these stages, we show lateral views. The gill arches become visible during tailbud stages and further develop in the prehatch tadpoles (lateral view). At these stages, gills become more pronounced and eyes gain pigment and become visible. Prehatch embryos remain in their thick jelly coats in natural settings. In hatched tadpoles, the dorsal fin becomes more transparent, the gills branch and grow outward (lateral view). We show a dorsal view of a larva-adult. This animal is approximately 1 year old, but is not yet fully grown or sexually mature

(prehatched and hatched, 72–340 hours + HPL) where organogenesis and growth occur. After hatching, limb morphogenesis (characterized in detail by the literature⁴²) occurs and animals continue to grow and mature (Figure 1, larva-adult, or juvenile adult, 2–18 months post-fertilization). However, axolotl development proceeds at a slower rate during early development than *Xenopus*, and the animals are not sexually mature until they are approximately 1 year of age or older.^{40,43,44}

A current gap exists in axolotl research in the study of embryogenesis at the cell and tissue levels and the comparative understanding of the epigenetic, transcriptomic, and proteomic changes that occur in early stages and primary formation compared to regeneration. A regulatory map of the processes that control primary development during embryogenesis in axolotl would be informative in

its own right but would also inform regeneration studies in larvae, juveniles, and adult animals. It is difficult to ascertain which developmental or regenerative mechanisms are conserved with other vertebrates, including in “distantly” related fish, avians, and rodents, or even with more closely related amphibians such as *X. laevis* and *tropicalis* with the current gaps in knowledge. For the most fruitful research outcomes, it is important to know where these animals come from (primary development) in order to understand where they are going (regeneration).

Quantitative three-dimensional imaging and reconstruction of axolotl neurula embryos has provided some detail of the mechanical forces and changes that occur at the earliest developmental stages.^{45,46} Additional work focused on the role of ion transport during

neurulation.^{47,48} However, more work is needed to validate structural and molecular similarities between axolotl development and other vertebrates. Without validated antibodies and molecular stains to spatiotemporally mark specific tissue derivatives in axolotl embryos, characterization and comparative analyses of cell and tissue specification and differentiation timing will be difficult. Although there is a single study of transcriptomic changes during embryogenesis in axolotl,⁴⁹ the work was published prior to the axolotl genome sequencing and should be reassessed with this recent knowledge. Further, scientists are creating additional molecular tools for spatiotemporal profiling of gene and protein expression, but more research is needed to understand the molecular mechanisms driving development in these early stages.

Early embryonic fate specification in axolotls is somewhat varied from their closely related frog embryos. Axolotl ectoderm forms neural tissue in the absence of an organizer signal²⁸ whereas in frog embryos, ectoderm forms non-neural ectoderm or epidermis in the absence of an organizer.⁵⁰ However, neural induction and continued development requires additional instructive signaling in both organisms.^{28,51}

The study of neural development is often paired with analyses of neural crest cell formation and migration, as the two tissues arise from the same ectodermal germ layer. Axolotls were a well-used model for the first wave of neural crest research that lasted roughly from the 1920s to 1950s, in which several scientists observed the migration and differentiation of this stem-like cell population through trunk and cranial neural crest transplantation experiments. Newth's 1954 study supported the idea that neural crest cells are specified but not determined at the neurula stage.⁵² Delving further into the realm of understanding these migratory cells, Epperlein and Löfberg discovered that trunk neural crest-derived melanophores and xanthophores create the pigment pattern in axolotls.⁵³ Cell tracing studies have also been performed on axolotl trunk neural crest cells and have shown that axolotl neural crest cells have different potentials depending on where they originate in the anterior-posterior axis, similar to chicken embryos.^{22,24,54} In addition, CRISPR-mediated genome editing and morpholino knockdowns have been used successfully in axolotl embryos, suggesting that it is a tractable model for functional studies.^{10,55-58} Moury and Jacobson identified that competence to become neural crest cells is not limited to the neural folds. They also proposed that epidermis-neural plate signaling and forces generated by local interactions between the two tissues induce neural crest cells, thus supporting a hypothesis of mixed intrinsic and extrinsic factors controlling neural crest cell development in axolotl embryos.⁵⁸ These experiments established the

nature of neural crest cells and began to answer questions on the timing of neural crest specification and migration. Between the 1960s and 2000s, the shift toward *X. laevis* as a developmental model and abundance of research in the avian neural crest contributed to a shortage of publications in this field.

The brink of the 21st century saw a resurgence in axolotl developmental research, picking up from the surge of neural crest cell research in urodele amphibians from the 1920s to 1950s.⁵⁹ This second wave of neural crest cell research in axolotls resumed in the early 2000s, as new techniques gave scientists the ability to reconsider classic questions in neural crest development. This era also upheld the importance of comparative studies and centered the axolotl in the field of evolution of development ("evo-devo"). Epperlein et al. took advantage of lipophilic dye injections and the novel neural crest cell marker, AP-2, to define the order of neural crest cell migratory streams in axolotl and the potential of neural crest cells to change their migration path when displaced off course.²² Scanning electron microscopy (SEM) identified detailed information on neural crest cell migratory stream routes and stream assembly, as well as cell shape and orientation across stages.⁵⁹ Comparing axolotl neural crest cell migration to other vertebrates identified that axolotls exhibit earlier neural crest cell migration than newts, but retain the distinct migratory streams common to amphibians.⁵⁹ Epperlein et al. demonstrated the unique migratory patterns and timing of neural crest cell development in axolotls.⁵³ Explant experiments showed intrinsic neural crest cell patterning and segmental migration, but the support of extrinsic signaling from the neighboring epidermis was necessary to maintain distinct streams.⁶⁰ These two waves of axolotl neural crest cell research provide a strong basis for future axolotl studies, and it is clear this model organism still has much untapped potential.

4 | AXOLOTLS AS MODELS FOR EVOLUTION OF DEVELOPMENT (EVO-DEVO)

While lamprey and hydra are renowned model organisms for studies of evolutionarily conserved and divergent developmental and regeneration mechanisms, axolotls carve out their own niche in the field of evo-devo studies.⁵⁹ As developmental models, *X. laevis* and *D. rerio* are currently more prominent than the axolotl, but the newly sequenced axolotl genome paired with epigenetic and transcriptomic analyses provide traction to re-invigorate the use of axolotl in both comparative and mechanistic studies of conserved and divergent evolutionary

processes.⁶¹⁻⁶³ Rationale for the use of zebrafish and frog embryos over axolotls in developmental studies may include the ease of *in vitro* fertilization and jelly coat removal in frogs^{64,65} and genetic tractability in zebrafish,⁶⁶ but axolotls provide a unique evolutionary niche. Frogs and fish develop faster and can be more resilient to cleavage-stage manipulation than axolotls. As a result, many laboratory techniques have been created or adapted for *Xenopus*, including *in situ* hybridization, immunohistochemistry, transgenic methods, and expression cloning.⁶⁷⁻⁶⁹ However, the axolotl reigns supreme as a vertebrate regenerative model, and techniques pioneered in other aquatic models can, and have been, readily be adapted for use in axolotl studies.

Their large genome, neoteny, ability to regenerate, and lack of a genome-wide duplication make axolotls the perfect tetrapod models that fill a unique evolutionary niche compared to other vertebrate models.^{62,63,70-72} To date, multiple studies have used axolotls as a comparative model of development. Axolotls are unique in their lack of *Pax3*, which drives neural crest and mesoderm development in multiple vertebrate species.⁶² Its absence suggests that there are some differences in the molecular mechanisms driving complex developmental processes across species. However, multiple studies identified developmental similarities as well. Analysis of pelvic development in axolotl and lungfish identified conserved mechanisms of chondrogenesis and myogenesis between the lobe-finned fish and the tetrapod.⁷³ In addition, studies of axolotl dentition identified a conserved ecto-endodermal boundary as the potential mediator of tooth development across multiple species.²⁶ Further, loss of function studies identified that *Bapx1* is necessary for jaw joint formation in multiple vertebrates, including axolotls.⁷⁴ The current body of evo-devo research using axolotl as a bridge animal have identified similarities and differences from genomic to morphological scales. Access to newly developing tools and an annotated genome will advance these studies tremendously in the future.

5 | THE ULTIMATE REGENERATOR

Although multiple animals have limited capabilities to regenerate, the axolotl has devised unique, extensive, and elegant methods of regenerating multiple tissues (Table 1).¹¹¹⁻¹¹³ Here, we will focus on a subset of regeneration studies that span both historical and current work. Neoteny places axolotls at a unique intersection of developmental and regenerative research potential. They exhibit scarless wound healing and a lower cancer

incidence,^{6,87} and they are an experimentally accessible organism for cell plasticity studies.^{3,107} They also develop quite slowly compared to other aquatic organisms such as zebrafish and frogs, lending them to experimental embryology studies, including cell fate tracing methods.^{21,60,114-119}

Most tissues within the axolotl regenerate in some capacity, making the species highly valuable for such studies. However, the axolotl does not necessarily employ the same mode of regeneration across tissues (Table 1). For example, dentectomy studies show that tooth regeneration is a nerve-dependent process while the lower jaw can regenerate without innervation.¹⁰³ Also, axolotls can only regenerate their lens within the first 2 weeks of life, contrasting with other tissues' extended regenerative capacity.⁷⁹ Apoptotic tissue degradation is a precursor to the regenerative process following axolotl limb injuries, and an axolotl must remove injured cells and reduce immune cell counts to a specific balance to avoid unwanted damage.¹²⁰ There are multiple in-depth recent reviews on the subject of axolotl regeneration,^{112,113,121,122} but the current body of research suggests that salamanders respond to injury signals in a way that is distinct from mammals. One of the secrets to axolotl limb and tail regeneration lies in its ability to form a blastema after injury. The blastema is a region of dedifferentiated cells that forms from underlying tissues at the site of injury.^{13,21,38,119,123-125} This transient structure interacts with the wound epidermis in similar ways to mesodermal-epidermal interactions seen during embryonic development.^{55,88} However, dedifferentiation and blastemal formation are not sufficient to drive the formation of a fully functional and patterned limb or tail after amputation. Although not an exhaustive list, signaling from immune cells, nerve cells, the inflammation response, and whole organism proliferation responses also occur and are important for regeneration after injury.^{14,15,102,126,127}

While the body of research on axolotls and other salamanders has uncovered many details of their regenerative potential, the mechanistic basis of neoteny remains largely unknown. However, there may be developmental origins linking the neotenic state of axolotls with their exemplary regeneration capacity.^{128,129} Recent work has identified that many tissues maintain populations of stem-like cells, allowing for growth, wound healing, and regeneration. Embryonic stem-like cells, including neural crest cells, may be a key to a subset of axolotl regenerative capabilities.¹³⁰⁻¹³³ Understanding these mechanisms is crucial to bridge the gap in knowledge between axolotl developmental and regenerative programs.

TABLE 1 Summary of tissues and structures that regenerate in axolotls after injury

Tissue	Type of injury	Regeneration (Y/N)	References
Ectoderm derived			
Brain	Incision to left dorsal pallium	Yes, regenerate multiple original neuron populations, but connectivity anomalies do occur	75-78
Lens	Lenectomy	Yes* (lost 2 weeks after hatching)	79,80
Retina	Retinectomy	Yes	81-83
Spinal cord	Amputation and spinal cord transection	Yes	20,84-86
Skin	Full thickness excisional (FTE) wounding with 4 mm biopsy punches	Yes	87-90
Sensory hair cells	Laser ablation	Yes	91-93
Mesoderm derived			
Heart	Ventricular resection and cryoinjury	Yes	94-98
Ovary	Partial ovariectomy	Yes	99
Skeletal muscle	Limb amputation	Yes	16,100,101
Endoderm derived			
Gills	Partial amputation	Yes	29
Lung	Partial amputation	Yes	102
Derived from multiple germ layers			
Jaw/tooth	Dentectomy	Yes, teeth are nerve dependent, lower jaw is independent	103
Limb	Amputation	Yes	12,13,15,38,42,104-106
Tail	Amputation	Yes	10,27,84,91,104,107-110

6 | FUTURE OF *AMBYSTOMA* AS A RESEARCH ORGANISM

New tools broaden the horizons for comparative and functional research in axolotls even further (Table 2). Obtaining the sequenced axolotl genome in 2018 and multiple bulk and single-cell transcriptomic atlases of developing and regenerating embryos and tissues have provided a baseline for functional studies, such as the analysis of genes and proteins that aid in regeneration. Specifically, single-cell sequencing atlases have been created from lineage-specific^{12,14-16} and unbiased¹³ samples during forelimb regeneration. By performing single-cell characterization of changes in gene expression during multiple stages of axolotl limb regeneration, previous studies identified that connective tissue cells revert to embryonic profiles,¹² there are mitochondrial-specific genes supporting regeneration,¹⁶ and that this process is paired with changes in the immune response in lineage-specific and unbiased cell populations.^{13,14} In addition to transcriptomic analyses, recent characterization of the chromatin landscape using the assay for transposase-

accessible chromatin using sequencing (ATAC-seq) in regenerating axolotl limbs has added complexity to the story by identifying changes during the eight stages of regeneration.¹³⁹

A wide topic for future axolotl research is the investigation of the unique programming axolotls hold that makes their scarless wound healing, lower incidence of cancer, and regenerative capabilities currently unattainable in other vertebrates. Current research shows that regeneration is not as simple as rebooting embryonic programming; rather, it appears as though wound healing and regeneration responses are much more complex than simple dedifferentiation and redifferentiation, and depend on the type of cells, stage of regeneration, environment, and tissue-type. The process of dedifferentiation is necessary for regeneration of certain tissues, but it does not provide a complete picture of the process. Dedifferentiated cells can cross over to new cell lineages, and single-cell atlases support the dedifferentiation concept in specific tissues.^{12,138} However, the limits of this fate-switching are yet to be completely tested, and further work is needed to confirm this process across

TABLE 2 Summary of recent developments in molecular tools in axolotl research

Tool	Use	References
Sequenced genome	Key to study of sequences that regulate development, aid regeneration, and so on	61-63
Foamy virus	Gene transfer method used for regeneration studies	134
Baculo virus	Gene transfer and gene overexpression	135
Vesicular stomatitis virus	Gene transfer and neural cell labeling	136
Retroviruses	Infection in vivo and in vitro to target specific cell types in regeneration	137
Germline transgenic strains	Cell tracing and mutagenesis during development and regeneration	20,23,134,137
Click chemistry	Lung injury in axolotl salamanders induces an organ-wide proliferation response	20,102,110
Single-cell sequencing	Single-cell transcriptomic datasets from multiple germ layers and cell types in regenerating axolotl tissues	12,14-16,138
CRISPR-mediated genomic mutation	Implementation of genomic deletion via CRISPR/Cas9-mediated genome editing in axolotl	10,25,57

species.^{27,100} A related gap in knowledge is the exploration of the axolotl immune system in the regeneration of different tissues, although there is strong groundwork laid for future studies in limb, spinal cord, craniofacial, and heart tissues, among others.^{14,15,97,140} A decline in the immune system is correlated with aging and lack of regeneration in other vertebrates,¹⁴¹ and axolotls must maintain their immune systems in a careful balance to regenerate injured tissues successfully. As most regeneration studies have been performed in immature axolotls, it will be interesting to see how future studies determine whether the same reduction in immune response occurs in mature animals.

Even with the identification of novel genes and cell types during regeneration, the field still lacks many of the tools that would allow for fast analysis of functional relevance to define mechanisms, identify gene regulatory networks, and link the similarities and differences between development and regeneration in axolotls. However, with the founding of the International Society for Regenerative Biology (<https://internationalsocietyforregenerativebiology.org/>), the creation of AxoBase, a new online axolotl resource (<https://www.axobase.org/>), the NIH-funded support for Sal-Site (<https://ambystoma.uky.edu/quick-links/sal-site>) and annual Salamander Meetings bringing together researchers from across the globe, the future of research using the axolotl as a model organism is promising.

7 | POP CULTURE ICONS

Secondary to their obvious importance in discovering the secrets of vertebrate regeneration and to the potential

discoveries of new developmental pathways, these anomalous salamanders are pop culture icons. Axolotls are conspicuously adorable and hold a high status in the modern world. They are incorporated into video games, cartoons, and social media posts. We would be remiss to omit the charismatic draw of this animal in society. In Japan, the axolotl is known as the wooper looper/rooper (<https://www.caudata.org>), made popular by a commercial marketing campaign that was then followed by the creation of an axolotl named Wooper in Pokémon cartoons and video games. Further, other popular video games such as Fortnite and Minecraft introduced axolotls as passive characters in 2021,¹⁴² and Build-A-Bear created a buildable axolotl plush toy. On social media outlets like Twitter, axolotls are used for scientific communication and public engagement (e.g., #ChonkTheAxolotl), but are also popular in avatars and art. Most recently, the axolotl has been featured on the 50 peso bill from the Bank of Mexico as a representative of ancient Mexico. These animals provide the scientific community with answers while they provide the world with joy.

8 | EXPERIMENTAL PROCEDURES

8.1 | Animal Husbandry

All use of axolotl adults and embryos was performed in accordance with the UC Davis approved IACUC protocol #21448. Axolotls were bred in house and embryos were collected for fixation at multiple stages in preparation for imaging. Embryos were fixed in 4% paraformaldehyde solution in 2% phosphate buffer for 1 hour and then were

either dehydrated step-wise into 100% methanol for storage or were imaged immediately in 1× TBS (500 mM Tris-HCl pH 7.4, 1.5 M NaCl, and 10 mM CaCl₂) containing 0.1% Triton X-100 (TBST+ Ca²⁺).

8.2 | Imaging

All whole mount embryos (Figure 1) were imaged using a Zeiss Microscopy Camera AxioCam 208 color mounted to a Zeiss Stemi 305 dissecting microscope. Zen Blue software was used for processing.

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AUTHOR CONTRIBUTIONS

Carly J. Adamson: Conceptualization (equal); data curation (equal); writing – original draft (equal); writing – review and editing (equal). **Nikolas Morrison-Welch:** Conceptualization (equal); data curation (equal); writing – original draft (supporting). **Crystal D. Rogers:** Conceptualization (equal); data curation (equal); formal analysis (equal); funding acquisition (equal); methodology (equal); writing – original draft (equal); writing – review and editing (equal).

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