

METHODS

Use of a *ROSA26:GFP* transgenic line for long-term *Xenopus* fate-mapping studies

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Abstract

Widespread and persistent marker expression is a prerequisite for many transgenic applications, including chimeric transplantation studies. Although existing transgenic tools for the clawed frog, *Xenopus laevis*, offer a number of promoters that drive widespread expression during embryonic stages, obtaining transgene expression through metamorphosis and into differentiated adult tissues has been difficult to achieve with this species. Here we report the application of the murine *ROSA26* promoter in *Xenopus*. *GFP* is expressed in every transgenic tissue and cell type examined at post-metamorphic stages. Furthermore, transgenic *ROSA26:GFP* frogs develop normally, with no apparent differences in growth or morphology relative to wild-type frogs. *ROSA26* transgenes may be used as a reliable marker for embryonic fate-mapping of adult structures in *Xenopus laevis*. Utility of this transgenic line is illustrated by its use in a chimeric grafting study that demonstrates the derivation of the adult bony jaw from embryonic cranial neural crest.

Key words bone; chimera; cranial neural crest; developmental biology; skull.

Introduction

Fate maps of embryonic tissues and other cell populations have long been important for understanding the developmental and genetic underpinnings of adult morphologies (Landacre, 1921; Hörstadius, 1950; Sadaghiani & Thiébaud, 1987; Balaban et al. 1988; Clarke & Tickle, 1999). Along with the importance of fate maps for developmental studies is their utility for comparative biology. For example, detailed comparisons of fate maps among closely related species (e.g. vertebrates) inform our understanding of how alterations in the embryonic origin of a given structure may correlate with its morphological variation among taxa (Rudel & Sommer, 2003).

To yield useful results, any methodology used to generate a fate map must satisfy several criteria (reviewed

in Kisseberth et al. 1999). For example, the method must reliably indicate a cell's lineage while minimizing the risk of false positives (i.e. ascribing an incorrect embryonic origin to a given structure) as well as false negatives (failure to detect the correct embryonic origin of a structure). A complication in metamorphic anurans, such as the well-studied model *Xenopus laevis*, is the protracted tadpole stage, which may last several months or longer (Trueb & Hanken, 1992). Any fate-mapping tool employed in this animal to assess the embryonic origin of adult-specific traits must last through this protracted life-history stage and into adulthood (Gross & Hanken, 2004).

Lineage tracers based on transgenes have the advantage of being intrinsic cellular markers; they do not dilute with time or age. However, in order to be useful for fate-mapping studies, transgenes must drive widespread and persistent expression. While promoters such as the Cytomegalovirus (CMV) promoter or the *Xenopus* elongation factor 1 α (EF1 α) gene promoter can be used to drive widespread expression during embryogenesis, the expression driven by these promoters is progressively restricted as tadpoles age and remains restricted

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Accepted for publication 7 May 2006

in adult frogs (N. Marsh-Armstrong, unpublished data). Therefore, these promoters may be used as lineage tracers in young animals (e.g. De Robertis & Kuroda, 2004), but they are inadequate for assessing the derivation of adult traits. In the absence of a stable, long-term cell marker that persists through metamorphosis, experiments that explore the adult fate of embryonic cell populations in an amphibian model system have not been possible.

The fate of embryonic neural crest cells in the adult skull has historically received the most attention in avian models (Johnston et al. 1973; Le Lièvre, 1978; Noden, 1978; Couly et al. 1993; Köntges & Lumsden, 1996), but more recently it has also been addressed in the mouse (Chai et al. 2000; Morriss-Kay, 2001; Jiang et al. 2002; Matsuoka et al. 2005). Important differences have been reported in the neural crest contribution to skull bone between avian and mammalian models, raising the question as to the nature of the neural crest contribution in more primitive (basal) vertebrates (Hanken & Gross, 2005).

We aim to evaluate the neural crest contribution to skull bone in frogs as an extant model of a basal tetrapod. To this end, we have generated *Xenopus laevis* with an intrinsic cellular marker that is based on the *ROSA26* gene. *ROSA26* was identified originally as a ubiquitous marker in a retroviral gene-trapping screen in mouse embryonic stem cells (Friedrich & Soriano, 1991). The gene trap vector was later found to have integrated into and disrupted a ubiquitously expressed gene of unknown function (Zambrowicz et al. 1997). The promoter region for the *ROSA* gene was found to drive very widespread, if not ubiquitous, expression of reporter genes in transgenic mice (Kisseberth et al. 1999), albeit at levels of expression below those seen in the original *ROSA26* mice (Zambrowicz et al. 1997). Mice carrying in their *ROSA26* locus a conditional lacZ gene that expresses the reporter gene only after recombination due to Cre activity (Soriano, 1999) have become a common tool for genetic experiments. Recently, these mice have been used to study reporter expression in bone cell lineages (Lui et al. 2004), as well as the fate of neural crest cells in the skull (Jiang et al. 2002).

We use transgenic *Xenopus laevis* that express a green fluorescent protein (GFP) reporter gene under the control of the *ROSA26* promoter, together with a chimeric grafting method, in order to obtain persistent GFP expression in explants of embryonic neural crest. The combination of embryonic grafting and a persistent and ubiquitous marker finally enables characterization

of the adult, post-metamorphic fate of neural crest cells in an amphibian model system. These data will further our understanding of the nature of the contribution of osteogenic crest cells to the skull in a more comprehensive phylogenetic context.

Materials and methods

Incorporation of the *ROSA26:GFP* transgene into the *Xenopus laevis* genome via REMI

We used the technique of restriction enzyme-mediated integration (REMI) transgenesis in *Xenopus laevis* (Kroll & Amaya, 1996) to generate a transgenic line of frogs expressing the pR26-GFP plasmid (Kisseberth et al. 1999). Enhanced GFP (EGFP) is expressed under the control of a 0.8-kb fragment containing the *ROSA26* promoter (plasmid kindly provided by Dr Eric Sandgren, University of Wisconsin, Madison). pR26-GFP, linearized with *Sall*, was purified with the GeneClean II kit (Qbiogene, Irvine, CA, USA) and used to make transgenic *Xenopus laevis* as previously described (Marsh-Armstrong et al. 1999).

Embryos and tadpoles made to express this transgene showed early and widespread GFP fluorescence (data not shown). A single female (F₀) was raised to sexual maturity and outbred to wild-type animals. The F₁ offspring of these matings were used in the subsequent chimeric grafting experiments as well as analyses of the persistence of GFP expression in adult tissues. An F₂ generation was produced from the colony of F₁ individuals via *in vitro* fertilization using wild-type male sperm. Fertilizations were carried out using previously described methods (Sive et al. 2000).

Tissue processing and histology

Transgenic *ROSA26:GFP* and wild-type animals were fixed overnight at 4 °C in 3.7% PFA, pH 7.4. Specimens were rinsed several times for ~1 h in non-sterile PBS solution. Following rinsing, several organs (heart, liver, colon, kidney) were removed from the transgenic and wild-type individuals. The tissues were rinsed again in PBS prior to immunohistological staining and processing.

Both the *ROSA26:GFP* and wild-type tissues were placed in a single 2-mL vial containing 5–10% normal goat serum (as a block) as well as a 1 : 2000 dilution of one or two of the following antibodies: Torrey Pines BioLabs rabbit polyclonal anti-GFP; AbCam rabbit polyclonal anti-GFP.

The vials were rocked on a nutator at 4 °C overnight (for at least 16 h) and then rinsed in multiple washes of PBS + 0.025% Triton X-100 at room temperature for 6–8 h. The tissues were placed back into a 2-mL vial containing a 1 : 500 dilution of goat anti-rabbit Alexa Fluor 488 (Molecular Probes, Eugene, OR, USA). Vials were again rocked at 4 °C overnight and then rinsed in multiple washes of PBS + 0.025% Triton X-100 at room temperature for 6–8 h. Once these tissues (heart, liver, colon and kidney) were analysed in whole-mount, they were cryosectioned and reanalysed in cross-section.

In addition to the tissues described above, various cranial tissues were assessed in cross-section for the presence of the GFP protein. Frontal cryosections were analysed for distribution of GFP (antibody stain, above) in a wild-type frog, a *ROSA26:GFP* transgenic frog and a mandibular stream cranial neural crest (CNC) grafted chimera (see Fig. 5). In our experiments, native GFP fluorescence did not persist strongly through the combination of protracted developmental times (i.e. after metamorphosis) and fixation employed to examine transgenic tissues. To determine the utility of antibody staining, we processed 16–20 µm-thick cryosections using an antibody staining protocol (see above) and compared them against neighbouring sections that were not processed immunohistochemically (Fig. 1). Positive label is present only on sections processed using the anti-GFP antibody staining protocol, which yielded significant amplification of the endogenous GFP signal (Fig. 1A) compared with untreated sections (Fig. 1B).

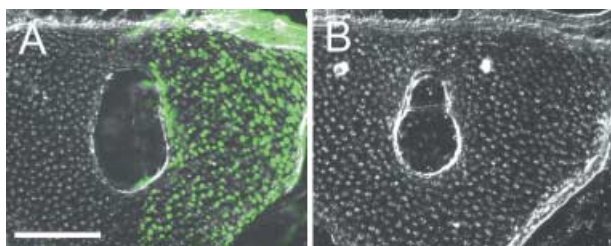


Fig. 1 Use of a polyclonal anti-GFP antibody significantly amplifies GFP signal in cryosections of chimeric *Xenopus*. (A,B) Serial cross-sections through the rostral cartilage of an adult frog (NF stage 66, +2 months). Tissue processed using anti-GFP antibody (A; green dots) demonstrates robust staining for GFP. There is no detectable GFP expression in an adjacent section from the same individual (B; separated by ~60 µm), which was processed without the antibody. In both sections, the right side received a grafted explant of cranial neural crest from a *ROSA26:GFP* transgenic donor; the left (ungrafted) side of the wild-type host provided an internal control. Scale bar, 200 µm.

Comparison of growth and morphology between transgenic *ROSA26:GFP* frogs and wild-type frogs

Transgenic *ROSA26:GFP* frogs and sibling wild-type frogs were divided among four groups: Group 1 – *ROSA26:GFP*, brightly fluorescing ($n = 12$); Group 2 – *ROSA26:GFP*, less brightly fluorescing ($n = 31$); Group 3 – wild-type set I ($n = 12$); Group 4 – wild type set II ($n = 31$). A sample of Group 1 individuals were killed and analysed for *GFP* expression in several tissues of the head and body (see below). All individual groups were reared in 4-L tanks with a separate supply of sterile, fresh 10% Holtfreter solution (Marine Biotech; Beverly, MA, USA) and fed *ad libitum* either live food (black worms) or frozen blood worms and frog brittle.

We observed no statistical difference in survival rate between the F_1 *ROSA26* and wild-type frogs (Pearson chi-square test, $P = 0.238$; Table 1). Because the frog transgenesis procedure with which the *ROSA:GFP* frog was generated is expected to increase the likelihood of developmental abnormalities, we carefully observed and characterized the normal growth and morphological development of this transgenic strain of frogs. All specimens were cleared and stained as whole-mounts according to standard procedures (Klymkowsky & Hanken, 1991).

A set of adult *ROSA26:GFP* transgenic and wild-type individuals ($n = 8$ per group) were compared on six linear and one weight measurements (Table 2). All measures were tested for significance at the $P = 0.05$ level using a Student's *t*-test. Statistical analyses were carried out using Microsoft Excel (Version X) and SPSS (version 11.5).

Southern hybridization

Genomic DNA was isolated from a wild-type and two transgenic members of the backcrossed F_2 generation

Table 1 Survivorship of *ROSA26:GFP* transgenic *Xenopus laevis* from 1 week post-fertilization (NF stage 42) to adult stage (NF 66), compared with wild-type frogs. There is no difference in survivorship between groups. Pearson chi-square test: 1 degree of freedom; $\chi^2 = 1.397$; $P = 0.238$

Individual genetic background	Early tadpoles (NF stage 42)	Adults (NF stage 66 +2 months)	Total survival rate
Wild type	44	15	34.1%
<i>ROSA26:GFP</i> (F_1 generation)	44	10	22.7%

Table 2 Comparison of external growth measurements* reveals no difference between F₁ transgenic frogs and wild-type frogs. Individual values indicate the mean measurement for each group ± standard deviation

Individual genetic background	IOD	LCD	SVL	HL1	HL2	FL	W
Wild type (<i>n</i> = 8)	4.48 ± 0.32	7.33 ± 0.46	24.36 ± 2.94	26.67 ± 2.63	32.46 ± 3.66	12.13 ± 1.01	1.74 ± 0.70
<i>ROSA26:GFP</i> (<i>n</i> = 8)	4.82 ± 0.39	7.77 ± 0.57	24.67 ± 3.73	29.22 ± 4.11	34.61 ± 5.23	12.46 ± 1.64	1.81 ± 0.95
<i>P</i> -value	0.08	0.12	0.86	0.16	0.36	0.64	0.85

*IOD, interocular distance; LCD, lateral cranial distance; SVL, snout–vent length; HL1, hind limb length, measured from the longest toe to the hip when the limb is extended 90° from the long axis of the body; HL2, hind limb length, measured from the longest toe to the vent when the limb is extended 90° from the long axis of the body; FL, forelimb length; W, weight. Linear measurements are expressed in millimeters; weight is expressed in grams.

expressing either no detectable fluorescence compared with wild-type (–), or an individual expressing the strongest levels of GFP (+++). A sample of 20 µg of each was digested overnight with *EcoRI*. Genomic DNA was precipitated and run for 14 h on a 0.7% agarose gel at 20 V. Genomic DNA was then blotted to a Hybond nylon membrane for 12 h and processed for hybridization according to the method of Sambrook et al. (1989).

Probe was prepared by cutting out a ~1-kb fragment of the GFP cassette from the R26R-GFP plasmid using *AflIII* and *BamHI*. The probe was radiolabeled with ³²P-dCTP using the Rediprime II kit (Amersham Pharmacia, Piscataway, NJ, USA). The probe was hybridized to the membrane overnight in UltraHyb buffer (Ambion, Austin, TX, USA), rinsed according to the manufacturer's instructions, and exposed for 4 days at –80 °C.

Embryonic tissue grafting

Embryo grafting was carried out at Nieuwkoop and Faber (NF) stage 14 (Nieuwkoop et al. 1994; Fig. 5B). Tissue explants of small portions of the CNC corresponding to the mandibular stream were grafted from *ROSA26:GFP* donor embryos into unlabelled wild-type hosts following standard procedures (Sadaghiani & Thiébaud, 1987; Gross & Hanken, 2005; Fig. 5B,C). All grafting was carried out on the right side; the left side of the developing head served as an internal negative control.

In each graft, a portion of the presumptive mandibular stream neural crest was first removed from an unlabelled wild-type host embryo and replaced with an equivalent-sized explant of the same region of the CNC from a labelled, donor transgenic embryo (Fig. 5B). Chimeric embryos were reared through metamorphosis, processed in cross-section, and analysed for the presence of donor-grafted GFP-positive cells in the adult bony jaw (Fig. 5F).

Image acquisition

Whole-mount specimens were placed in a Petri dish containing PBS + 0.025% Triton X-100 and photographed using a Leica MZ FLIII fluoroscope. Cryosectioned tissues were mounted with Fluoromount-G (Southern Biotech; Birmingham, AL, USA) and photographed using a Leica fluoroscope (model DMRE). High-resolution digital images were obtained with a 12-bit, black-and-white CCD camera (ORCA, Hamamatsu) using Openlab software (Improvision, UK). All tissues were stained and treated with identical antibody-staining protocols and photographic specifications.

All specimens were photographed in bright-field (cryosections were photographed using Nomarski optics) as well as fluorescence (B-filter, Leica) illumination. Black and white images were pseudo-coloured green using Openlab, merged with the light images and saved as TIFF files.

Intact adult specimens and cleared-and-stained skeletal preparations were photographed using an Automontage camera system (Syncroscopy USA, Frederick, MD, USA). All images were saved as TIFF files and compiled into figures using Adobe Photoshop 7.0.

Results

A new transgenic line of *Xenopus laevis* with widespread and persistent GFP expression

REMI transgenesis (Fig. 2A–C) was used to create embryos in which enhanced GFP (eGFP) expression was driven by a 0.8-kb proximal promoter fragment from the mouse *ROSA26* gene (Kisseberth et al. 1999). These embryos had widespread transgene expression that, on average, was lower than that typically observed for the CMV promoter (data not shown). A single F₀

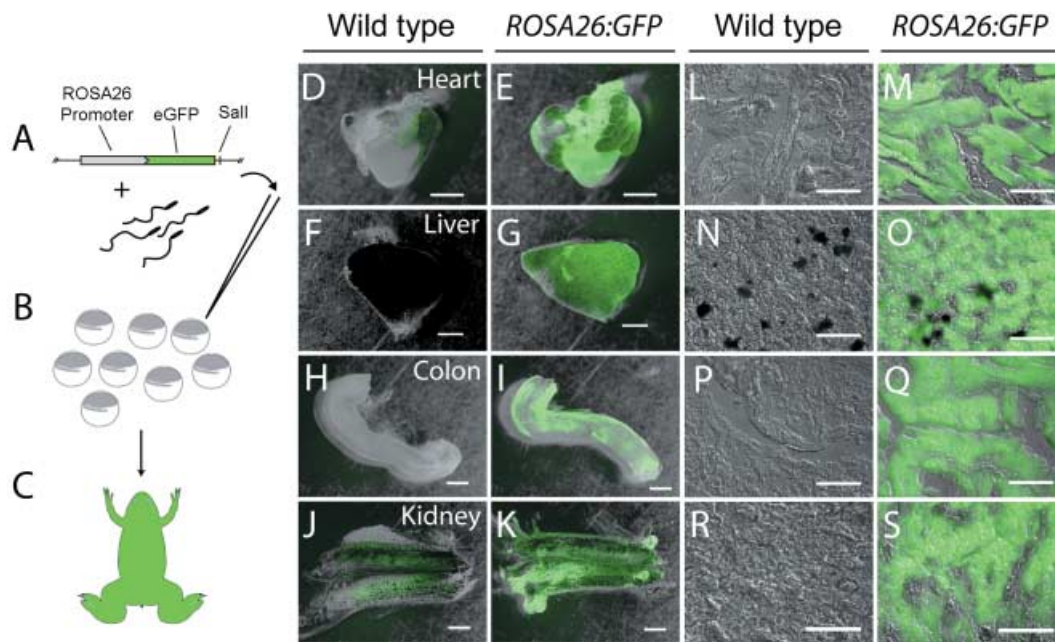


Fig. 2 Schematic depiction of REMI method of *ROSA26:GFP* transgenesis and anti-GFP staining of transgenic tissues. (A) A plasmid containing the *ROSA26:GFP* cassette is incubated with de-membranated wild-type sperm nuclei in a solution containing restriction enzyme (*Sall*). Resultant double-stranded breaks in the sperm nuclei genome allow for the introduction of the linearized plasmid. (B) Single nuclei are injected into unfertilized wild-type oocytes. (C) Animals carrying successful integration events and showing widespread GFP fluorescence are reared through metamorphosis. Offspring of a single founder female showing strong fluorescence were used as labelled, cranial neural crest (CNC) explant donors in all experiments. Several tissues were assayed for anti-GFP staining in whole-mounts and serial sections (D–S). In all tissues examined, the *ROSA26* promoter drives *GFP* expression well into adulthood. Two wild-type organs processed for whole-mount immunohistochemistry, heart (D) and kidney (J), showed minor background staining that was much lower than the signal detected in comparable tissues from animals expressing the *ROSA26:GFP* transgene (E and K, respectively). The background staining was not evident in sections (L, R) when compared with *ROSA26:GFP* tissue (M, S). Scale bars: D–K, 1 mm; L–S, 100 μ m.

ROSA26:GFP female was raised to adulthood. Upon breeding, this animal was found to carry multiple integrations of the transgene. There was variable GFP expression among the progeny: some had very weak fluorescence, whereas others had fluorescence comparable with that typically obtained with the CMV promoter (data not shown). We attribute the differences in expression to different integration sites – some allowing stronger expression than others – as well as a variable number of transgene integration sites.

The *GFP* reporter did not appear to affect either the survivability or the development of the founder female, further supporting the observation that GFP reporters are completely biosafe (Chalfie et al. 1994). We observed no developmental defects in the progeny of this founder (see Fig. 4, Table 2). As with other lines (Marsh-Armstrong et al. 1999), there appears to be no detectable change in either the pattern or the intensity of expression of the *ROSA26:GFP* transgene after passage through the germ line (data not shown). This

enabled us to use the outbred offspring (F_1) of this founder female (F_0) in subsequent grafting experiments and analyses.

GFP is detectable in a diverse array of organs and tissues throughout adulthood

To assess GFP expression in adult animals, we first analysed whole-mount, fixed tissues, including the heart, liver, colon and kidney. In each case, antibody staining revealed strong and uniform expression of the *GFP* protein (Fig. 2E,G,I,K). Minor fluorescence associated with the secondary antibody (AlexaFluor 488) was noted in the auricles of the heart (Fig. 2D) as well as restricted portions of the kidney (Fig. 2J). In both cases, this background fluorescence was negligible when compared with the intense fluorescence of *ROSA26:GFP* transgenic tissue, which had been processed with the identical staining and imaging protocols. Non-specific background staining that is occasionally present in whole-mount

preparations is readily distinguished from specific, positive staining associated with the *ROSA26:GFP* construct.

All tissues were sectioned to determine the extent to which transgenic samples may be distinguished from wild-type samples at a fine microscopic level (Fig. 2L–S). In each comparison, the transgenic tissue showed strong and specific GFP signal whereas the wild-type tissue lacked any detectable signal. The slight background staining visible in some whole-mounts of wild-type tissues was not present in stained cryosections (Fig. 2L,R).

Some regions of the transgenic colon whole-mount appear to be unstained (Fig. 2I). We cannot explain the slightly lowered expression of GFP in this transgenic tissue preparation. We do note that in the corresponding sectioned preparation of the colon the signal is not diminished (Fig. 2Q), but instead it contrasts strongly with the complete and expected absence of signal in the wild-type section (Fig. 2P). It is unclear if the transgenic section shown (Fig. 2I) is demonstrative of all transgenic individuals, or only the brightest fluorescing individuals.

Comparison of reporter gene expression between transgenic and wild-type adult cranial tissues

Expression of the *ROSA26:GFP* reporter construct in cranial tissues was of special concern because of our interest in using this construct to assess direct contributions of cranial neural crest cells to the head of adult *Xenopus laevis*. Therefore, we surveyed cryosections of the prosencephalon, pterygoideus muscle, rostral cartilage and (ossified) pars articularis of the quadrate bone (Fig. 3).

In every case, strong fluorescence was present in transgenic tissue (Fig. 3B,D,F,H) and generally absent in wild-type tissue (Fig. 3A,C,E,G). Some non-specific background staining is visible in wild-type skeletal muscle (Fig. 3C). As described above for whole-mounts, however, autofluorescence in wild-type tissue is minimal when compared with the fluorescence observed in transgenic tissue (Fig. 3D), and these two tissue types can be distinguished unequivocally according to staining intensity in chimeric grafting studies.

Transgenic, wild-type and chimeric individuals have indistinguishable cranial skeletal morphology and survival rates

Two important prerequisites for the use of *ROSA26:GFP* transgenic embryonic tissue in long-term fate-mapping

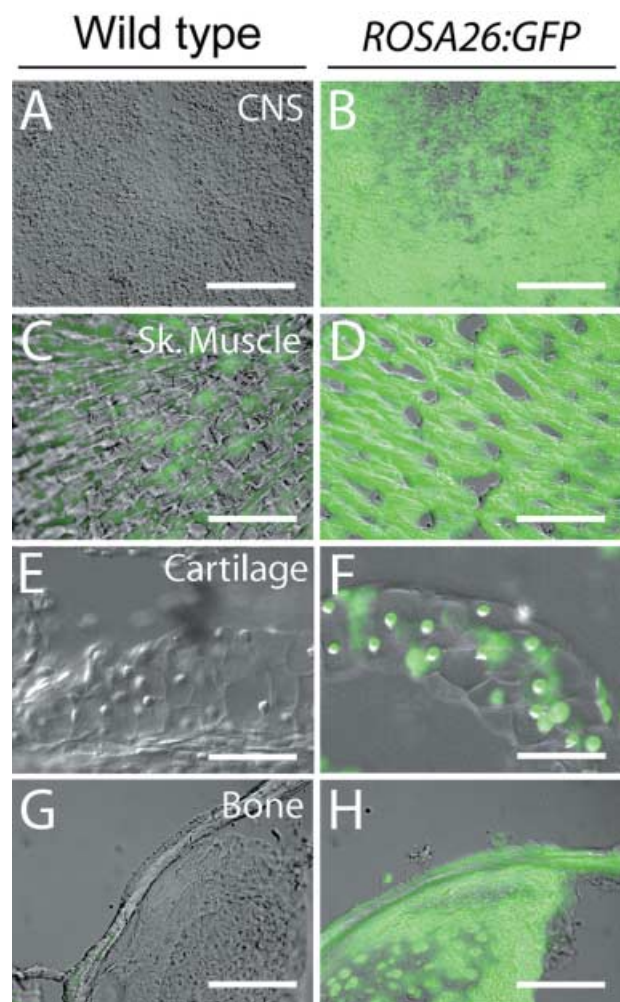


Fig. 3 Antibody staining of adult cranial tissues. Frontal sections through the prosencephalon (A,B), pterygoideus muscle (C,D), rostral cartilage (E,F) and (ossified) pars articularis of the quadrate bone were immunostained to determine if the transgenic *GFP* label persists in these tissues. In all comparisons, anti-*GFP* staining in the *ROSA26:GFP* tissues (B,D,F,H) is clearly distinguishable when compared with wild-type tissues (A,C,E,G). Wild-type pterygoideus muscle showed minor background fluorescence (C) that is clearly distinguishable from the much more intense positive *GFP* staining of *ROSA26:GFP* muscle (D). CNS, central nervous system; Sk. muscle, skeletal muscle. Scale bar, 100 μ m.

studies are that transgenic and wild-type animals develop along identical timelines, and that the adult morphologies of these two groups are indistinguishable from one another. Therefore, we reared groups of transgenic and wild-type frogs under identical conditions and assessed their survivability and cranial morphologies (Fig. 4).

The survival rate of *ROSA26:GFP* transgenic frogs was slightly lower than that of wild-type frogs, but the difference between group means was not statistically significant (Table 1). Furthermore, there was no difference

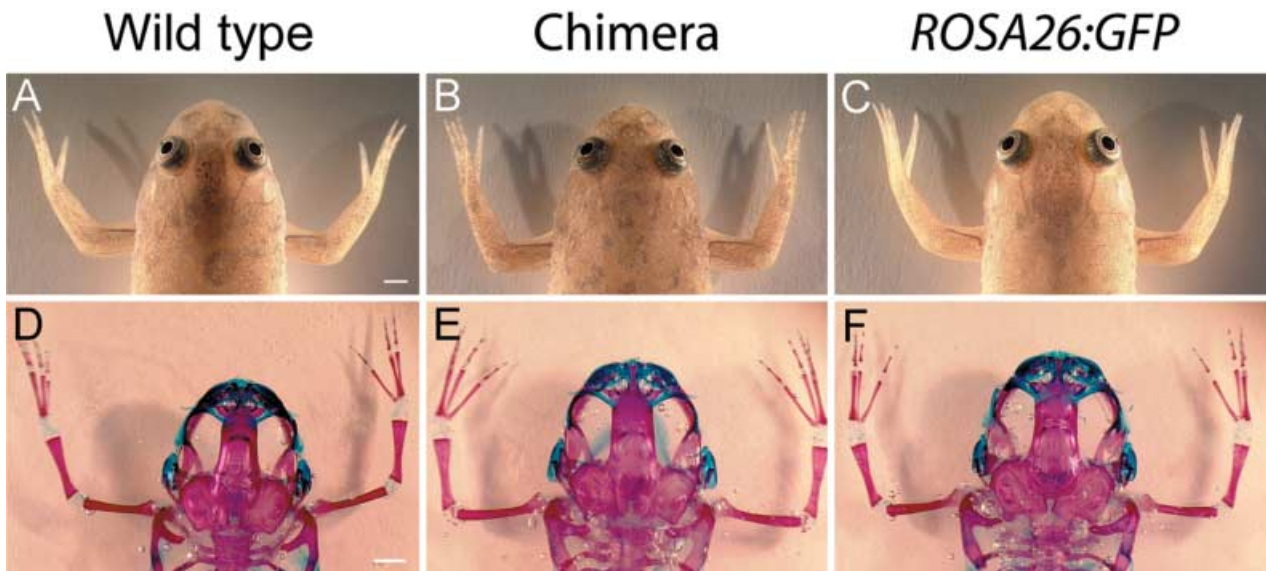


Fig. 4 External cranial morphology and skeletal development are indistinguishable among wild-type frogs (A,D), *ROSA26:GFP* transgenic frogs (C,F) and chimeric frogs (B,E). All animals were reared to at least 6 months of age; skeletal anatomy was assessed in cleared-and-stained whole-mounts (D–F). The three groups were indistinguishable from one another in terms of normal development and growth. The minor differences in cranial size among the three cleared-and-stained specimens depicted here are common even among populations of normal tadpoles reared under identical conditions but in different tanks. Scale bar, 2 mm.

in adult body size or proportions between these two groups (Table 2). Both external and internal (skeletal) cranial morphology also appear to be identical among transgenic, wild-type and chimeric individuals (Fig. 4).

Use of the *ROSA26:GFP* transgenic frog for tracing adult derivatives of embryonic neural crest

To determine if the *ROSA26:GFP* transgene offers a reliable lineage tracer for long-term mapping studies, we grafted premigratory CNC cells that give rise to the mandibular stream from a transgenic embryo into an unlabelled wild-type host (Fig. 5A–C). The neural crest has long been recognized as the principal source of the chondrocranium in larval amphibians, including mandibular cartilages that constitute the larval lower jaw (Landacre, 1921; Sadaghiani & Thiébaud, 1987). Here, however, we were interested specifically in determining whether neural crest also contributes to bones of the lower jaw, which in anurans do not differentiate until metamorphosis and are fully formed only in adults.

Beginning 2 months following the end of metamorphosis (NF stage 66), we assessed the derivation of bone and cartilage of the anterior lower jaw in chimeric frogs that received *ROSA26:GFP*-labelled neural crest grafts

(Fig. 5D). At this stage, anti-GFP staining strongly and uniformly labels all tissues in *ROSA26:GFP* frogs (Fig. 5G), whereas positive GFP label is absent in wild-type individuals (Fig. 5E). As expected, labelled chondrocytes were evident within the mandibular cartilage on the operated (right) side of the lower jaw (MC; Fig. 5F). However, label was also present within the calcified matrix of the dentary, one of two bones that form the anuran lower jaw (Fig. 5F, arrowheads). Additional label was present in the upper jaw, within the matrix of the premaxillary bone and in nearby tooth buds (Fig. 5F; arrow). GFP-positive tissues are absent on the un-operated (left) side of all chimeras (Fig. 5F).

Early survival and GFP expression among outbred F_2 progeny

As a further test of the future utility of the *ROSA26:GFP* line of transgenic *Xenopus*, we outbred F_1 females to wild-type sperm via *in vitro* fertilization and assessed early survival of the F_2 generation. The transgenic embryos suffered a significantly higher mortality compared with wild-type fertilized eggs in the first 24 h post-fertilization (hpf; Table 3). Beyond 24 hpf, however, survivorship rates were comparable, and high, in both groups. More than 95% of embryos alive at 24 hpf remained alive at

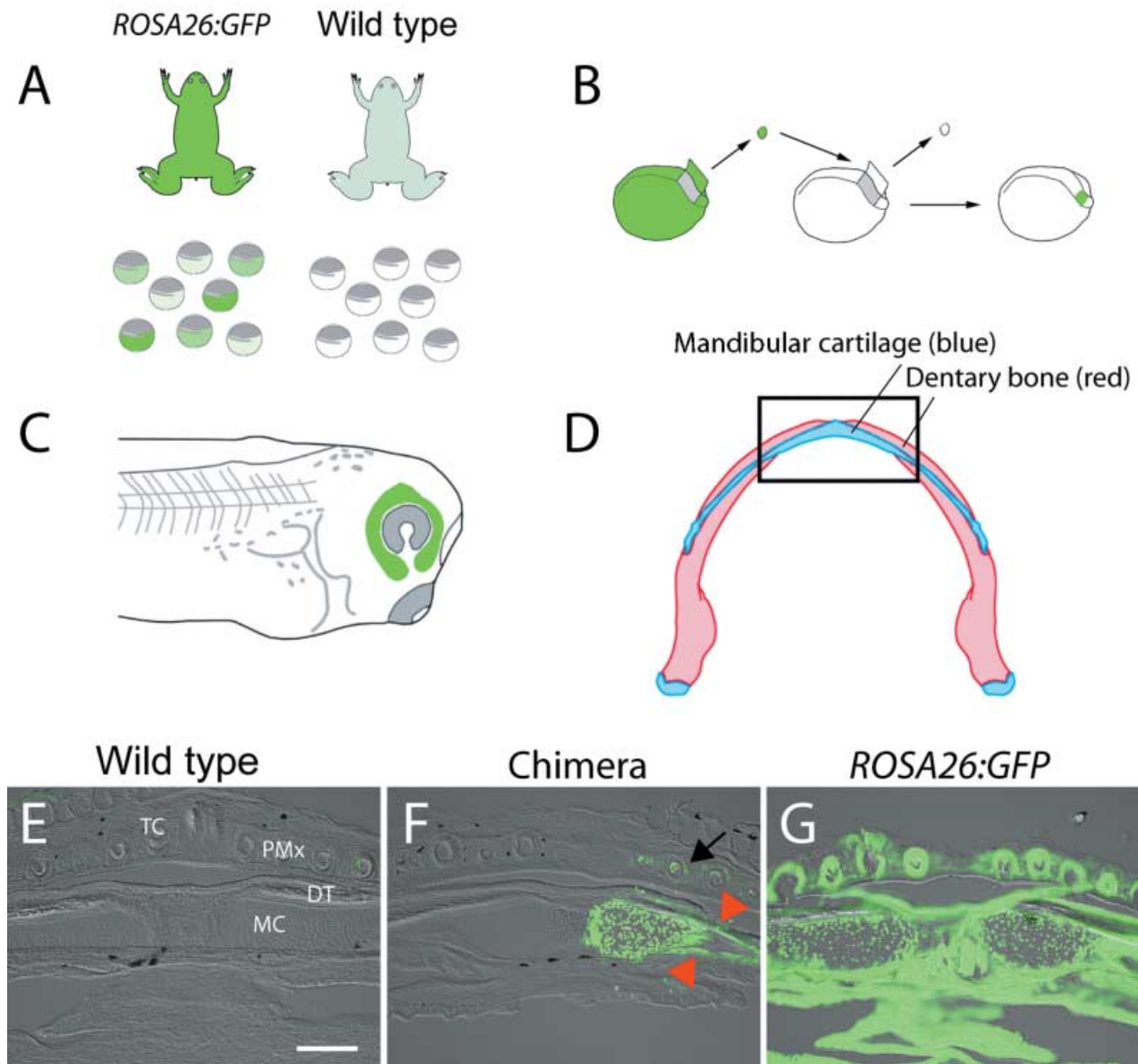


Fig. 5 Embryonic derivation of the adult lower jaw. We grafted premigratory cranial neural crest (CNC) from transgenic *ROSA26:GFP* donor embryos to wild-type host embryos at neurula stage, utilizing published fate maps (Sadaghiani & Thiébaud, 1987). In all experiments, eggs from both the *ROSA26:GFP* founder female and a wild-type female were fertilized *in vitro* with sperm from a wild-type male (A). *ROSA26:GFP* zygotes demonstrated variable intensity of *GFP* fluorescence; such variability is typical in animals with multiple transgene integration sites. Only the brightest, healthiest looking embryos were used as CNC explant donors (B). A tissue explant was removed from a transgenic *ROSA26:GFP* donor and replaced into a wild-type host. The explant was allowed to heal in place and was assayed the next morning to ensure proper development and migration of the CNC within the mandibular stream (C). Embryonic derivation of the adult mandibular cartilage and dentary bone in the lower jaw was assessed in chimeras that received a labelled graft of the mandibular stream of CNC. The jaw was sectioned in frontal plane (boxed region, D) and assessed for the presence of fluorescent label. *GFP* staining is absent in a wild-type specimen (E), whereas the entire jaw is labelled strongly in a *ROSA26:GFP* adult (G). Only the right (operated) side expresses *GFP* in a chimera that received the labelled graft (F). The left (control) side lacks the label, indicating that graft-derived cells did not cross to the contralateral side of the embryo or adult during development. Dense staining is evident within the extracellular matrix of the dentary bone (red arrowheads). Additional staining is present in the premaxillary bone of the upper jaw as well as portions of several tooth buds (black arrow). Abbreviations: DT, dentary bone; MC, mandibular cartilage; PMx, premaxillary bone; TC, tooth cusps. Scale bar (E–G), 250 μ m.

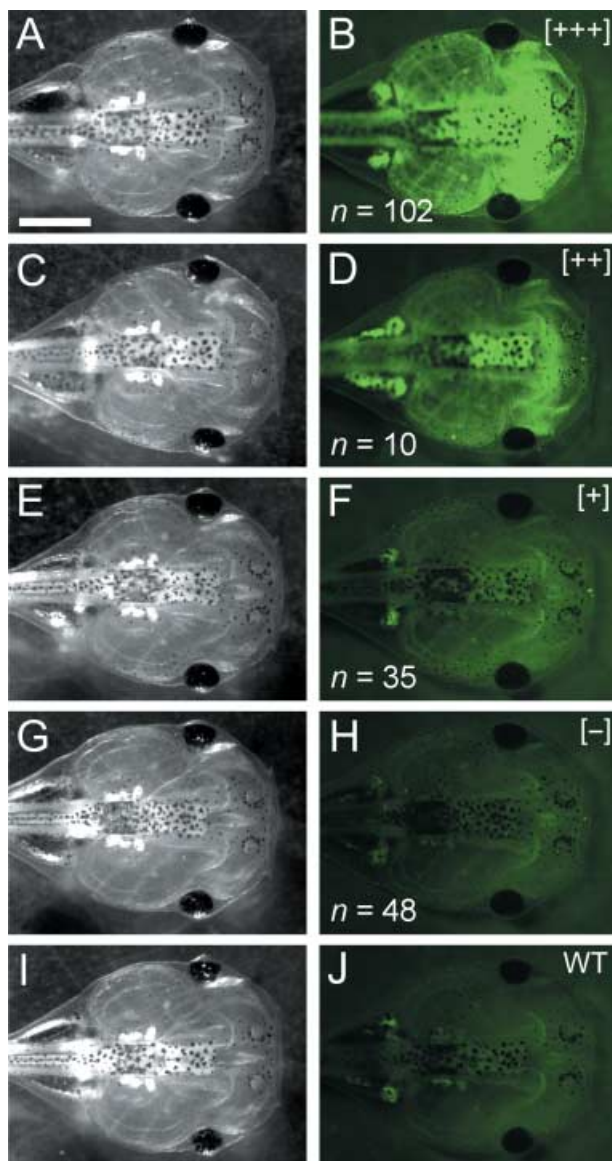
Table 3 Early survivorship of F₂ ROSA26 embryos compared with wild-type embryos

Individual genetic background	Total no. of fertilized eggs at day 0	Embryos surviving between day 0 and 24 hpf* (NF stage 15/16)	Embryos surviving between 24 and 48 hpf (NF stage 33/34)	Tadpoles surviving between 48 hpf and 6 dpf (NF stage 44)
Wild type	842	651 (77.3%)	627 (96.3%)	581 (92.7%)
F ₂	438	224 (51.1%)	216 (96.4%)	195 (90.3%)
Pearson chi-square results		$\chi^2 = 91.26$; $P < 0.001$	$\chi^2 = 0.006$; $P = 0.94$	$\chi^2 = 1.25$; $P = 0.26$

*hpf, hours post-fertilization; dpf, days post-fertilization; NF, Nieuwkoop and Faber.

48 hpf, and more than 90% of embryos alive at 48 hpf remained alive at 6 days post-fertilization (dpf).

We also assessed variability in GFP expression across the outbred F₂ generation. Endogenous fluorescence



of live tadpoles was scored qualitatively at 6 dpf (approximately NF stage 44). Individuals were exposed to the same level of UV illumination and sorted among four groups (Fig. 6): (1) brightest fluorescence (+++), (2) intermediate fluorescence (++) , (3) low-level fluorescence (+) and (4) no detectable fluorescence (-). Endogenous fluorescence in the 'brightest' and 'intermediate' individuals was very strong, especially compared with wild-type individuals (Fig. 6I,J). 'Low-level' individuals were only slightly brighter than those with no detectable fluorescence and wild-type tadpoles. Transplantation experiments that use the ROSA26:GFP transgenic line should target brightest individuals as tissue donors whenever possible.

Because members of the F₂ generation will be made available to our colleagues, we wished to determine whether the non-fluorescing (-) members of this generation carry the transgene. Towards this end, we performed a Southern hybridization of the genomic DNA from two representative members of this generation: (1) an individual not expressing fluorescence stronger than that of a wild-type individual (-), and (2)

Fig. 6 Variability of endogenous GFP fluorescence among live F₂ individuals. Eggs from a single ROSA26 transgenic female (F₁) were fertilized *in vitro* using wild-type sperm. Qualitative differences in endogenous GFP fluorescence were recorded among the resulting F₂ progeny viewed under identical conditions (constant time and intensity of exposure to UV illumination). These individuals were assigned among four groups according to qualitative measures of GFP fluorescence: brightest (B; +++), intermediate (D; ++), low (F; +) and undetectable (H; -). The last group is indistinguishable from wild-type individuals (J). There was no apparent difference in morphology among F₂ individuals, nor between transgenic and wild-type individuals. The 3 : 1 ratio of GFP-expressing (B,D,F; n = 147) to non-GFP-expressing individuals (H; n = 48) suggests that the F₁ female assayed in this experiment probably carries multiple copies of the ROSA26:GFP transgene. Scale bar, 1 mm.

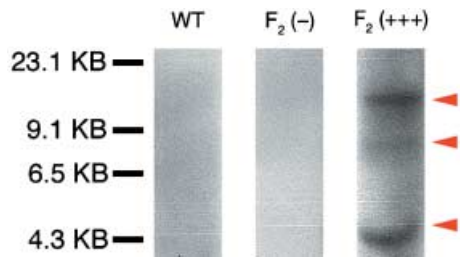


Fig. 7 Southern blot analysis of members of the F_2 generation of transgenic *ROSA26* individuals. Genomic DNA from a wild-type individual (WT), an F_2 individual expressing fluorescence at levels indistinguishable from wild-type (-), and an F_2 individual expressing the brightest level of GFP fluorescence (+++) were analysed for the presence of transgene integration using Southern blot analysis. Both the wild-type individual and the transgenic F_2 -individual are shown not to carry the transgene. The F_2 (+++) individual, however, carries multiple copies of the transgene (arrowheads). At present, it is not clear if fluorescence is correlated with the site of integration, number of copies at each integration site, or a combination of the two.

an individual displaying the strongest level of fluorescence both as an embryo and as a tadpole (+++). Our results show that the transgenic individual without apparent fluorescence [F_2 (-); Fig. 6(G,H)] does not have any integrations of the transgene (Fig. 7). Furthermore, the brightly fluorescing individual [F_2 (+++); Fig. 6(A,B)] has multiple copies of the transgene integrated into its genome (Fig. 7, arrowheads). At present we do not know the number of integration sites nor the number of copies per integration site. In light of these results, other investigators are encouraged to select the most brightly fluorescing individuals for long-term grafting projects.

Discussion

ROSA26 promoter drives widespread reporter expression in adult tissues of *Xenopus laevis*

An essential requirement of any fate-mapping technique is that 'labelled' cells be both identifiable and distinguishable from surrounding 'unlabelled' cells in a chimeric environment (Kisseberth et al. 1999). Furthermore, the ideal fate-mapping technique should yield negligible background fluorescence. In every circumstance we considered, *ROSA26:GFP* transgenic tissues were much brighter and clearly distinguishable from wild-type tissues, especially when prepared as serial sections. Certain tissues may demonstrate endogenous

autofluorescence due to a wide variety of causes unrelated to our experimental paradigm (e.g. flavoproteins and lipofuscins; Billington & Knight, 2001). In our study, this potential problem was confined principally to skeletal muscle, which displayed minor and non-specific background staining (autofluorescence) in sections. Use of *ROSA26:GFP* transgenic frogs for fate-mapping this particular tissue will require extra diligence to interpret staining patterns observed in chimeras correctly. Because there is variability in the levels of expression of different primary transgenic animals and the founder *ROSA26:GFP* female was not selected based on its expression level, it is likely that generating additional lines with the same *ROSA26:GFP* transgene will lead to lines with higher levels of expression and a higher signal-to-noise relative to endogenous autofluorescence.

We observed no statistically significant difference in survival rate to adulthood in *ROSA26:GFP* transgenic vs. wild-type frogs. The only survival difference we detected was confined to initial embryonic stages (0–24 hpf; Table 3). Large variability in survival rate is common even when breeding different wild-type animals. This early difference in survival rate might be a consequence of the sperm nuclei/REMI transgenesis procedure through which the *ROSA26:GFP* founder was created. Recently, alternative transgenesis procedures for *Xenopus* have been reported (Allen & Weeks, 2005; Pan et al. 2006), which may produce less deleterious effects than the sperm nuclei/REMI procedure. Future lines of *ROSA26:GFP Xenopus* may be generated with these novel methods as well. Finally, the presence of the transgene does not have any apparent effects on external or internal morphology (Fig. 4, Table 2).

The *ROSA26:GFP* transgene passes stably through the germ line

Numerous reporter constructs show stable passage through the germ line of *Xenopus laevis* (Marsh-Armstrong et al. 1999). We similarly find that the *ROSA26:GFP* cassette is capable of passing from F_0 to F_1 and F_2 generations with no apparent change to the expression pattern or loss of fluorescence intensity of the reporter gene. The variability we observe in different animals is attributable to the fact that the one founder line for these studies had multiple integrations of the transgene. While the optimal line for future experiments

would contain a single integration event of the transgene driving high expression of the reporter, we show here that the existing *ROSA26:GFP* line is suitable for grafting of labelled tissues, cellular progeny of which may be identified in both living and fixed specimens throughout all phases of the life history of *Xenopus laevis*.

Use of the *ROSA26:GFP* transgenic line as a novel tool for fate-mapping studies in *Xenopus laevis*

We previously reported derivation of the adult bony cranial vault of *Xenopus laevis* from embryonic neural crest using fluorescent dextran as a long-term marker (Gross & Hanken, 2005). Unfortunately, because the visible signal of fluorescent dextran diminishes over extended periods of time due to continued cell divisions and eventual autolysis of the tagged fluorophore, this method has limited applicability in studies that seek to map the embryonic derivation of adult-specific features.

Here we demonstrate that embryonic grafting, a standard technique in experimental embryology, may be combined with transgenic technology to produce chimeric embryos that express GFP in grafted tissues and their cellular progeny well into adulthood. For the first time, the embryonic origin of the bony lower jaw in adult *Xenopus laevis* is traced to the mandibular stream of cranial neural crest. Thus, along with the establishment of this line of transgenic frogs comes the ability to map the embryonic origin of a variety of adult tissues and organs, especially those that do not form until metamorphosis.

Use of the mouse *ROSA26* promoter in frog transgenic studies

Based on widespread use of the *ROSA26* locus in the mouse for genetic experiments that require ubiquitous transgene expression, we tested whether the murine *ROSA26* promoter could drive similar expression in *Xenopus*. Such widespread and persistent transgene expression, which was not found in any of the previously available promoters for use in *Xenopus*, would enable, among other things, long-term fate-mapping studies in *Xenopus* that to date have been impossible.

Available sequence databases reveal no obvious homologue to the *ROSA26* gene in *Xenopus laevis* or *Silurana tropicalis*, or any sequence in the *Silurana tropicalis* genome with high conservation to the murine *ROSA26* promoter (data not shown). Interestingly, the

murine promoter is still capable of driving widespread and persistent expression in *Xenopus* as it does in the mouse (Kisseberth et al. 1999). Thus, there appears to be functional conservation of this promoter in spite of the absence of obvious sequence conservation.

Unfortunately, little is known of the murine *ROSA26* gene. It appears to encode an RNA transcribed in the opposite strand of another ubiquitously expressed gene, *Thumpd3*. Unlike the *ROSA26* gene, *Thumpd3* does appear to be conserved outside of mammals, including *Xenopus*. Thus, it is possible that the *ROSA26* promoter works to regulate the expression of the *Thumpd3* gene in both mice and frogs, although the *Xenopus ROSA26* promoter cannot be identified with current algorithms. Alternatively, the *ROSA26* promoter may function in frogs as it does in the mouse even without the equivalent gene in frogs.

Future applications of the *ROSA26* promoter in *Xenopus laevis*

Whereas several promoters, including *ROSA26*, may drive widespread reporter expression in *Xenopus* embryos and very young tadpoles, *ROSA26* is the only promoter of which we are aware that retains widespread expression in older tadpoles, through metamorphosis, and into adulthood. Thus, *ROSA26* may be the promoter of choice for experiments that address these later ontogenetic stages and require widespread transgene expression. *ROSA26* also may be a promoter of choice to be used in *Xenopus* transgenic studies that employ binary systems such as those based on *Flp* or *Cre* recombinases (Werdien et al. 2001; Ryffel et al. 2003; Gargioli & Slack, 2004) or in conjunction with ligand-inducible transgene expression systems in *Xenopus* (Das & Brown, 2004).

Conclusions

We have established a new line of transgenic frogs that express *GFP* under the control of the murine *ROSA26* promoter. Reporter expression is present in unfertilized eggs and persists in every cell and tissue examined well into adulthood. This line of frogs offers a tractable tool for fate-mapping experiments because transgenic individuals grow along identical timelines as wild-type frogs with no apparent behavioral, developmental or morphological abnormalities. Incorporation of this transgenic line into ongoing studies of cranial development

has further allowed us to apply a novel technique to solve a previously intractable problem, the fate of embryonic cells in the adult (post-metamorphic) head. We encourage use of this line of frogs for fate-mapping studies and as a step towards the introduction into *Xenopus laevis* of other genetic techniques (e.g. crosses utilizing site-specific recombinases) that until now have been limited mainly to mammalian model systems.

Acknowledgements

We thank Sarah Whitton for help with statistical analyses and Anne Everly for help with animal rearing and skeletal preparations. We are also grateful to Rahul Kanadia, Jennifer Mansfield, Douglas Kim and Maria Samson for help with Southern blot experiments. Eric Sandgren provided the pR26-GFP plasmid. Special thanks to Paul Trainor for help with GFP antibody staining. Funding was provided by the US National Science Foundation (EF-0334846; AmphibiaTree), The William F. Milton Fund (Harvard University) and a Goelet Summer Research Fellowship (Museum of Comparative Zoology).

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