











# Heterogeneity and genomic loci of ubiquitous transgenic Cre reporter lines in zebrafish

Robert L. Lalonde<sup>1</sup>  | Cassie L. Kemmler<sup>1</sup>  | Frédérique W. Riemsdagh<sup>1</sup>  |  
 Andrew J. Aman<sup>2,3</sup>  | Jelena Kresoja-Rakic<sup>1</sup>  | Hannah R. Moran<sup>1</sup>  |  
 Susan Nieuwenhuize<sup>1</sup>  | David M. Parichy<sup>2,3</sup>  | Alexa Burger<sup>1</sup>  |  
 Christian Mosimann<sup>1</sup> 

<sup>1</sup>Department of Pediatrics, Section of Developmental Biology, University of Colorado School of Medicine, Anschutz Medical Campus, Aurora, Colorado, USA

<sup>2</sup>Department of Biology, University of Virginia, Charlottesville, Virginia, USA

<sup>3</sup>Department of Biology and Department of Cell Biology, University of Virginia, Charlottesville, Virginia, USA

## Correspondence

Alexa Burger and Christian Mosimann, Department of Pediatrics, Section of Developmental Biology, University of Colorado School of Medicine, Anschutz Medical Campus, Aurora, Colorado, USA. Email: [alexa.burger@cuanschutz.edu](mailto:alexa.burger@cuanschutz.edu) (A. B.) and Email: [christian.mosimann@cuanschutz.edu](mailto:christian.mosimann@cuanschutz.edu) (C. M.)

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## Abstract

**Background:** The most-common strategy for zebrafish Cre/*lox*-mediated lineage labeling experiments combines ubiquitously expressed, *lox*-based *Switch* reporter transgenes with tissue-specific Cre or 4-OH-Tamoxifen-inducible CreERT2 driver lines. Although numerous Cre driver lines have been produced, only a few broadly expressed *Switch* reporters exist in zebrafish and their generation by random transgene integration has been challenging due to position-effect sensitivity of the *lox*-flanked recombination cassettes. Here, we compare commonly used *Switch* reporter lines for their recombination efficiency and reporter expression pattern during zebrafish development.

**Results:** Using different experimental setups, we show that *ubi:Switch* and *hsp70l:Switch* outperform current generations of the two additional *Switch* reporters *actb2:BFP-DsRed* and *actb2:Stop-DsRed*. Our comparisons also document preferential Cre-dependent recombination of *ubi:Switch* and *hsp70l:Switch* in distinct zebrafish tissues at early developmental stages. To investigate what genomic features may influence Cre accessibility and *lox* recombination efficiency in highly functional *Switch* lines, we mapped these transgenes and charted chromatin dynamics at their integration sites.

**Conclusions:** Our data documents the heterogeneity among *lox*-based *Switch* transgenes toward informing suitable transgene selection for lineage labeling experiments. Our work further proposes that *ubi:Switch* and *hsp70l:Switch* define genomic integration sites suitable for universal transgene or switch reporter knock-in in zebrafish.

## KEYWORDS

Cre/*lox*, CreERT2, lineage tracing, site-specific recombinases, tamoxifen, transgenes, zebrafish

## 1 | INTRODUCTION

Site-specific recombinase-based techniques provide powerful versatility to transgenic models. Techniques based on Cre/*lox*, Flp/*FRT*, or phiC31 recombinase systems allow permanent excision or rearrangement of transgene cassettes to modify their activity and function.<sup>1-5</sup> As a widely used recombinase system in vertebrate models, the Cre/*lox* system combines (i) a Cre recombinase-providing transgene driven by a tissue-specific *cis*-regulatory element and (ii) a recombination-competent reporter transgene. The Cre recombinase, originally derived from the P1 bacteriophage, directionally recombines DNA at specific 13 bp palindromic repeats called *loxP* sites in a variety of species including mice and zebrafish.<sup>2,4,6,7</sup> A major application of the Cre/*lox* system in zebrafish involves the recombination of *lox*-flanked fluorophore cassettes that switch upon Cre activity to lineage-label cell populations of interest. This strategy has been successfully employed to reveal developmental lineage origins, to follow postembryonic stem cells, and to track the lineage composition of regenerating organs.<sup>1,8-16</sup> These experiments hinge upon combining a tissue- or cell type-specific Cre or 4-OHT-Tamoxifen (4-OHT)-inducible CreERT2 driver with temporal control and a broadly active or ubiquitous *lox*-based reporter transgenic.<sup>14,17-19</sup> Particularly, simple to control in developing zebrafish by 4-OHT addition to the embryo medium, CreERT2 transgenics have rapidly increased in number across the field and include Tol2-based transgenes driving CreERT2 with tissue-specific regulatory elements, gene trap collections, and first CRISPR-based knock-in lines.<sup>1,18,20-22</sup> In contrast, the generation of suitable and reproducibly well-performing *lox*-based reporter transgenes has proven challenging.

The *heat shock protein 70-like (hsp70l)*, *beta-actin2 (actb2)*, and *ubiquitin (ubi or ubb)* promoter elements have been routinely used to drive quasi-ubiquitous transgene expression in zebrafish and have been successfully applied in several *lox*-based *Switch* reporters.<sup>1,7,14,15,23-26</sup> *hsp70l*-based transgenes are a mainstay of the zebrafish's transgenic tool kit, as heat shock induction around 37°C causes rapid ubiquitous reporter expression that is however not sustained long beyond the heat shock pulse.<sup>24</sup> The *actb2* promoter provides strong, rapid transgene expression during development and in individual tissues, yet its activity drastically diminishes or silences in select cell types such as erythrocytes.<sup>25-28</sup> The *ubi* promoter drives widespread and persistent transgene expression in various independent transgenics including the Cre-sensitive *loxP* reporter *ubi:loxP-GFP-loxP\_mCherry (ubi:Switch)* that has found widespread use.<sup>14</sup> Nonetheless, *ubi*-based transgenics including *ubi:Switch* show slow reporter accumulation at early developmental stages<sup>14,28,29</sup>; this property causes considerable latency between Cre-triggered *lox* cassette recombination and reporter

detection, rendering short lineage labeling timeframes (under 24 hours) challenging to achieve.

The advantages and drawbacks of each regulatory element require careful characterization of individual transgene insertions over multiple generations, data that is rarely available to guide experimental design. Generating reproducibly functional, single-insertion *lox*-based *Switch* transgenics by randomly integrating Tol2 or ISce-I transgenesis is also notoriously screening-intensive due to position-effect sensitivity of *lox* cassette recombination.<sup>1,30,31</sup> Consequently, together with the challenges of sharing transgenic zebrafish lines internationally, the majority of labs only have access to one or few *Switch* reporter lines. Means to efficiently generate *Switch* transgenics are therefore highly desirable, requiring the identification of suitable genomic loci for transgene knock-ins.

Here, we compared previously published and validated *Switch* reporter lines in combination with ubiquitous and tissue-specific CreERT2 drivers. Our results document the heterogeneity in recombination efficiency and preferential tissue expression of individual reporters. Together with genomic integration mapping of tested transgenes, our results define *ubi:Switch* and *hsp70l:Switch* as integrations in favorable loci for *lox* cassette recombination and transgene expression.

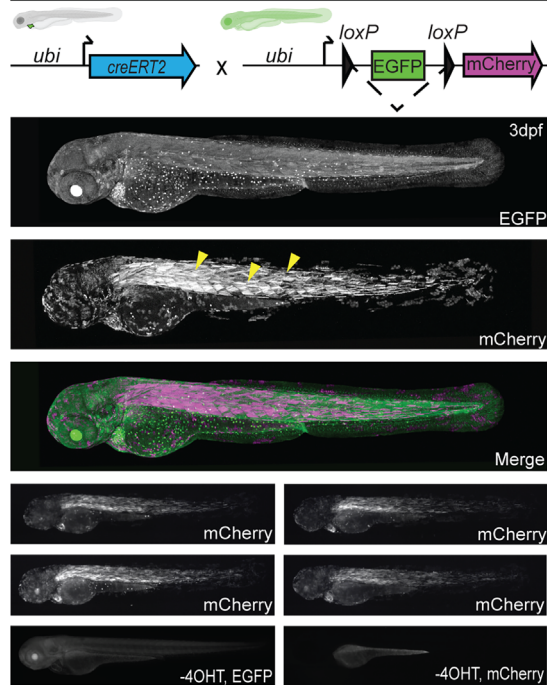
## 2 | RESULTS

### 2.1 | Ubiquitous *Switch* reporters feature variable recombination efficiency and tissue activity

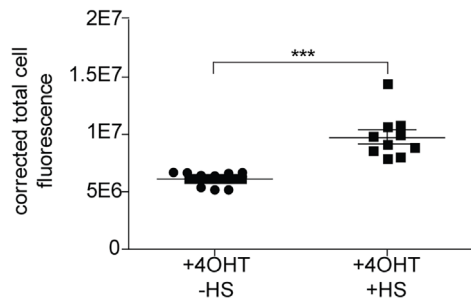
To gain insight into recombination efficiency and reporter expression across tissues by ubiquitous *Switch* reporters in development, we compared transgenic zebrafish reporters that have been previously used to follow lineage trajectories. We focused on *Switch* lines in our collection that have been documented to drive ubiquitous transgene expression: *ubi:Switch<sup>cz1701</sup>*, *hsp70l:Switch<sup>zh701</sup>*, *Tg(bactin2:loxP-Stop-LoxP-DsRed-express)<sup>sd5</sup>* (shortened to *actb2:Stop-DsRed*), and *Tg(bactin2:loxP-BFP-loxP-DsRed)<sup>sd27</sup>* (shortened to *actb2:BFP-DsRed*).<sup>14,15,23,32</sup> The *ubi:creERT2* driver has been widely used for ubiquitous CreERT2 expression in early development and provides a first assay for Cre responsiveness of individual *Switch* lines.<sup>14</sup> To establish switching efficiency under *ubi:creERT2* and homogeneity vs. heterogeneity of reporter expression, we treated *ubi:creERT2;promoter:Switch* embryos with 4-OHT at shield stage overnight, and imaged whole larvae at 3 dpf (Figure 1). In all crosses, the *ubi:creERT2*-carrying parent was male to avoid maternal CreERT2 contribution.

The *ubi:Switch* transgenic permanently recombines from EGFP to mCherry following Cre activity

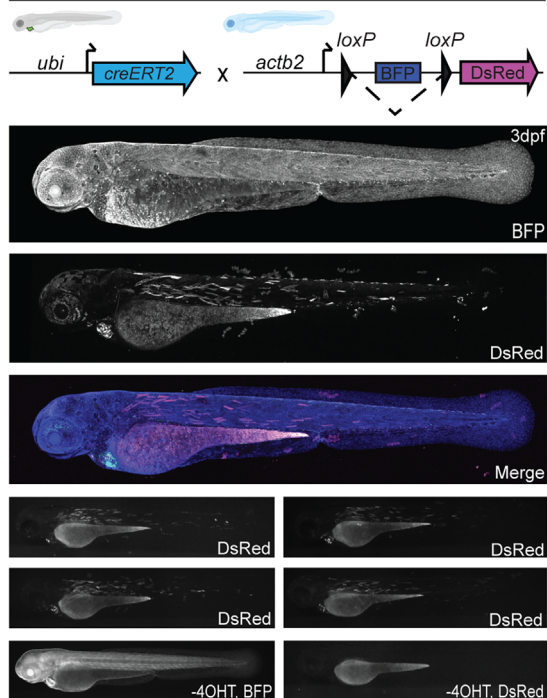
(A) *ubi:creERT2 x ubi:switch x 4OHT @ shield*



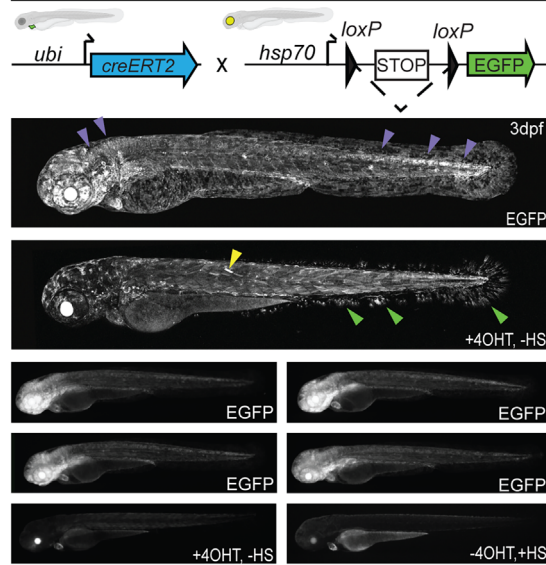
(B) *ubi:creERT2 x ubi:switch*



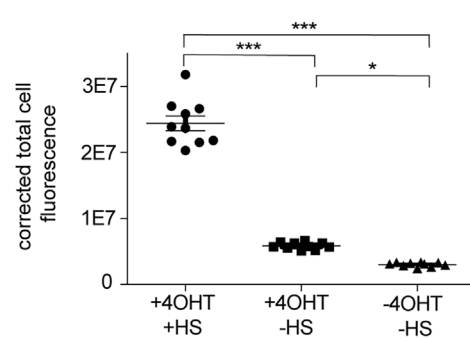
(E) *ubi:creERT2 x actb2:Switch x 4OHT @ shield*



(C) *ubi:creERT2 x hsp70:switch x 4OHT @ shield*



(D) *ubi:creERT2 x hsp70:switch*



(F) *ubi:creERT2 x actb2:stopDsRed x 4OHT @ shield*

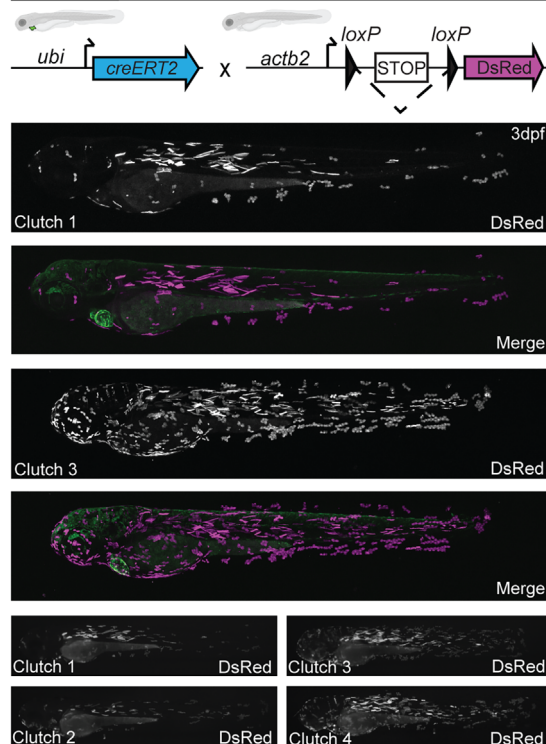


FIGURE 1 Legend on next page.

(Figure 1A), yet mCherry levels only reach detectable levels after 12–24 hours post-Cre activation.<sup>14</sup> When combined with *ubi:creERT2*, we observed strongest mCherry activity in the somitic myofibers along the entire trunk and tail (Figure 1A). Skin and fin epithelium showed more sparse switching, predominantly in the median fin and head (Figure 1A). We observed strong mCherry signal in the heart, while other tissues including the eye lens and neural tube consistently displayed comparatively lower mCherry fluorescence (Figure 1A). Promoters of *ubiquitin* genes in several species have been postulated to harbor heat shock-responsive elements, possibly to support protein degradation by increased *Ubiquitin* polypeptide production upon heat or other stress.<sup>33–38</sup> To test if *ubi:Switch* as driven by the zebrafish *ubb* gene promoter reaches higher expression levels upon heat shock, we compared the levels of mCherry fluorescence at 3 dpf with and without a 1 hour-long, 37°C heat shock prior to imaging: compared to nonheat-shocked siblings, heat-shocked *ubi:creERT2;ubi:Switch* larvae showed increased fluorescent intensity 3 hours postheat shock ( $n = 10$  for quantification, 2 clutches) (Figure 1B). These observations document that *ubi:Switch* expression can be further augmented by a brief heat shock prior to imaging.

Following Cre activity, the *hsp70l:Switch* transgenic line permanently switches from a nonfluorescent Stop cassette to EGFP.<sup>15</sup> Heat shock for 1 hour at 37°C induces prominent EGFP expression within 1 hour postheat shock treatment. To trigger reporter expression of *hsp70l:Switch*, we heat-shocked recombined embryos 2 to 3 hours prior to imaging (Figure 1C). When crossed to *ubi:creERT2* and treated with 4-OHT at shield stage, *hsp70l:Switch* showed strong EGFP signal across the entire embryo, indicating near complete switching, with the central nervous system (brain, neural tube) displaying higher EGFP signal (Figure 1C). Notably, in nonheat-shocked controls that were 4-OHT-induced, we also detected faint EGFP expression ( $n = 10$ ) in the skeletal myofibers and fin fibroblasts (Figure 1C,D); these observations indicate that tissue-specific accessibility of the

*hsp70l* element at the genomic location of *hsp70l:Switch* leads to transcriptional activity even at 28°C. Embryos that were left untreated with 4-OHT did not show any detectable EGFP expression with or without heat shock, confirming that the used Stop cassette<sup>15,39</sup> efficiently interrupts any EGFP transcription.

The *actb2:Stop-DsRed* transgene recombines from no fluorophore to DsRed and *actb2:BFP-DsRed* recombines from BFP to DsRed, enabling versatile combination with existing tissue-specific reporters based on EGFP.<sup>23,32</sup> After recombination with *ubi:creERT2* at shield stage, both *actb2 Switch* lines imaged at 3 dpf showed overall sporadic switching with the most prominent DsRed signal in the skeletal myofibers, fin epidermis, and the heart (Figure 1E,F). Of note, *actb2:Stop-DsRed* showed two distinct switching efficiencies across clutches, potentially due to multiple integrations within the current or most-recent generation (Figure 1F). Under these conditions, switching was absent from a large majority of tissues including, but not limited to, endothelial cells, skin, brain and neural tube, and fin fibroblasts (Figure 1F). Ubiquitous BFP expression is observed with *actb2:BFP-DsRed* throughout the 4-OHT induction regime (Figure 1E).

Taken together, our observations from triggering ubiquitous CreERT2 activity after onset of gastrulation reveal heterogeneous recombination efficiencies and reporter expression across four distinct, broadly expressed *lox* reporter lines. The results with *ubi:Switch* are in line with previously reported Cre-based lineage labeling experiments across various cell types and developmental stages. Nonetheless, *hsp70l:Switch* displayed distinctively more homogeneous reporter expression than *ubi:Switch*; this effect is possibly in part due to the pulsed fluorescence activation following heat shock shortly before analysis in *hsp70l:Switch* experiments, while in *ubi:Switch* the mCherry reporter dynamically accumulates and degrades in various cell types following 4-OHT treatment. In contrast, the lower switching efficiency observed with the *actb2*-based *Switch* lines indicates low Cre accessibility of the insertions selected for these transgenics over the past decade.<sup>23,32</sup>

**FIGURE 1** Ubiquitous *Switch* reporter lines show variable recombination efficiency. (A,B) *ubi:Switch*, (C,D) *hsp70l:Switch*, (E) *actb2:BFP-DsRed*, and (F) *actb2:Stop-DsRed* crossed to *ubi:creERT2*, induced with 10  $\mu$ M 4-OHT at shield stage, and imaged laterally at 3 dpf. Schematics of fluorophore cassettes for each *Switch* transgene are shown at the top of each panel and larval schematics represent secondary transgenic markers (A,B,E,F). One representative confocal image, and four representative stereo microscope images are presented per reporter. *Ubi:Switch* (A) shows preferential recombination in somitic myofiber (yellow arrowheads), *hsp70l:Switch* (C) shows preferential switching in CNS (brain, neural tube) (purple arrowheads). Nonheat-shocked (+4-OHT) controls (C, second panel) show faint EGFP expression in somitic myofiber (yellow arrowheads) and fin fibroblasts (green arrowheads). (B) Corrected total cell fluorescence (CTCF) measurements of 3 dpf *ubi:creERT2* crossed to *ubi:Switch*, induced with 10  $\mu$ M 4-OHT at shield, and heat-shocked 3 hours prior to lateral view imaging; fluorescence intensity was compared to nonheat-shocked sibling ( $n = 10$ , 2 clutches). Note increased CTCF following heat-shock at 3 dpf ( $n = 10$ , Mann-Whitney,  $P > .0001$ ). (D) CTCF *ubi:creERT2* crossed to *hsp70l:Switch*, induced with 10  $\mu$ M 4-OHT at shield. Nonheat shocked controls (center), and nonheat shocked, nontreated controls (right) are included ( $n = 10$ , 1-way ANOVA,  $P < .0001$ ). *ubi:Switch* (A) and *hsp70l:Switch* (B) show more spatially complete recombination compared to both *and actb2:BFP-DsRed* (E) and *actb2:Stop-dsRED* (F) lines that only display sparse recombination

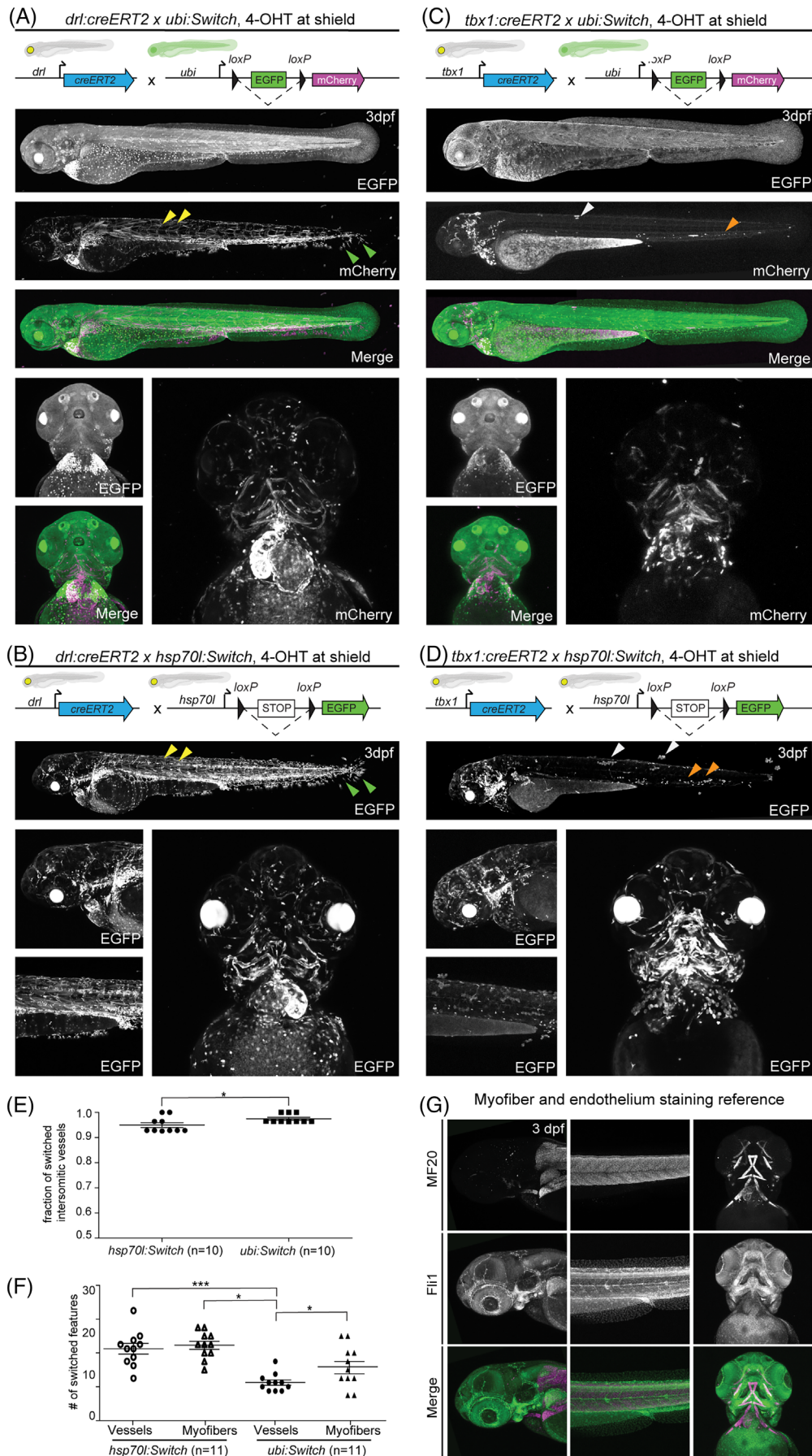


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## 2.2 | *hsp70l:Switch* shows widespread and reproducible Cre sensitivity

The widespread Cre-induced recombination and reporter expression observed with *ubi:Switch* and *hsp70l:Switch* indicate favorable genomic integration sites for *loxP* cassette recombination. To further evaluate the switching efficiencies between *ubi:Switch* and *hsp70l:Switch*, we turned to two previously established, highly active, and tissue-specific CreERT2 driver lines: (i) *Tg(drl:creERT2;cryaa:Venus)* (*drl:creERT2* for short) that expresses in lateral plate mesoderm (LPM)-primed mesendoderm during gastrulation before refining to cardiovascular lineages by mid-somitogenesis<sup>40,41</sup> (Figure 2A,B); and (ii) *Tg(tbx1:creERT2;cryaa:Venus)* (*tbx1:creERT2* for short) that starts expressing in late gastrulation and predominantly labels cardiopharyngeal progenitors together with *tbx1*-expressing ectodermal and endodermal lineages in the head<sup>15</sup> (Figure 2C,D). After crossing, we treated double-transgenic embryos with 10  $\mu$ M 4-OHT at shield stage and imaged the head region at 3 dpf laterally and ventrally to assess lineage labeling.

Combined with *drl:creERT2*, both *Switch* lines displayed switching in the same tissues and cell types, yet *hsp70l:Switch* consistently resulted in more complete recombination (lower mosaicism). This feature was most evident in the prevalence of switched fin fibroblasts, skeletal myofiber cells, macrophages, and head vasculature (Figure 2A,B). To compare recombination efficiency in a major cell type, we quantified switched vs unlabeled intersomitic vessels (ISV)<sup>29</sup>: we observed slightly more switched ISVs with *ubi:Switch* than with *hsp70l:Switch* when recombined by *drl:creERT2* ( $n = 10$ ,  $P = .0454$ ) (Figure 2E). Of note, induction of *drl:creERT2* at shield stage results in complete labeling of LPM-derived structures, but also sparse labeling of paraxial mesoderm-derived structures including somitic skeletal myofiber cells and median fin fold fibroblasts, reflecting the

starting segregation of individual mesodermal territories at shield stage<sup>40,41</sup> (Figure 2A,B).

When crossed to *tbx1:creERT2*, both *ubi:Switch* and *hsp70l:Switch* displayed broad lineage labeling in the ventricular cardiomyocytes, pharyngeal arches, cranial vasculature, head myofibers and cartilage, and hatching gland, consistent with previous work<sup>15</sup> (Figure 2C,D). Once more, *ubi:Switch* animals showed preferential switching in myofibers, vs *hsp70l:Switch* that displayed more complete switching across multiple cell types, especially head vasculature. We quantified head myofiber and endothelial switching of each *Switch* line, separately using myofiber (MF20 antibody)-stained *fli1:EGFP* embryos as reference for vessel/myofiber number and position ( $n = 11$ ) (Figure 2F,G). This analysis documented that *tbx1:creERT2* had significantly higher recombination efficiency in the head vasculature and musculature combined with *hsp70l:Switch* than when combined with *ubi:Switch* (Figure 2F). These results further underline the widespread, yet differential sensitivity to Cre activity of both *ubi:Switch* and *hsp70l:Switch*.

## 2.3 | *ubi:creERT2* shows decreased recombination activity at 48 hpf

The *ubi:creERT2*<sup>cz1702</sup> transgenic line has been widely applied to provide broad 4-OHT-inducible Cre activity, as desirable for testing new *Switch* transgenes.<sup>1,14</sup> Our data presented above further confirmed the reproducible activity of *ubi:creERT2* at shield stage paired with responsive *Switch* lines. Nevertheless, anecdotal observations have suggested that *ubi:creERT2* becomes less responsive to 4-OHT at later stages.<sup>14,29</sup> To what extent this effect depends on choice of *lox Switch* line remains uncertain.

To test the recombination capacity after gastrulation stages, we crossed *ubi:creERT2* to *ubi:Switch* and *hsp70l:Switch*, treated with 10  $\mu$ M 4-OHT at 48 hpf, and performed lateral-whole embryo imaging at 3 dpf (Figure 3).

**FIGURE 2** *hsp70l:Switch* high degrees of lineage labeling crossed to tissue-specific CreERT2 driver lines. (A,B) *drl:creERT2*, and (C,D) *tbx1:creERT2* crossed to *ubi:Switch* and *hsp70l:Switch*, respectively induced with 4-OHT at shield stage and imaged laterally and ventrally at 3 dpf. Quantifications depicted as bar diagrams with individual data points (E,F). Schematics of fluorophore cassettes for each *Switch* transgene are shown at the top of each panel and larval schematics represent secondary transgenic markers (A,B,C,D). *drl:creERT2* yields more complete recombination when crossed to *hsp70l:Switch* (B) compared to *ubi:Switch* (A). The higher degree of recombination is detectable in the somitic myofiber (yellow arrowheads), and fin fibroblasts (green arrowhead) (A,B). A higher percentage of switched ISVs is observed when combining *drl:creERT2* with *ubi:Switch* than with *hsp70l:Switch* ( $n = 10$ , unpaired *t*-test,  $P = .0345$ ) (E). *tbx1:creERT2* yields less switching mosaicism when crossed to *hsp70l:Switch* (D), compared to *ubi:Switch* (C). Higher recombination efficiency is readily observed in trunk skin (white arrowhead) and hematopoietic cells (orange arrowheads) (C,D). When crossed to *hsp70l:Switch*, *tbx1:creERT2* yields significantly more switched head vessels and myofibers compared to *ubi:Switch* ( $n = 11$ , 1-way ANOVA  $P < .0001$ ) (F). Representative lateral and ventral images (A,B,D,E). (G) Max projections of z-stack confocal images of *fli1a:EGFP* transgenic zebrafish fixed at 3 dpf and stained for myofibers using an MF20 antibody to visualize vasculature and myofiber anatomy in the zebrafish head at 3 dpf

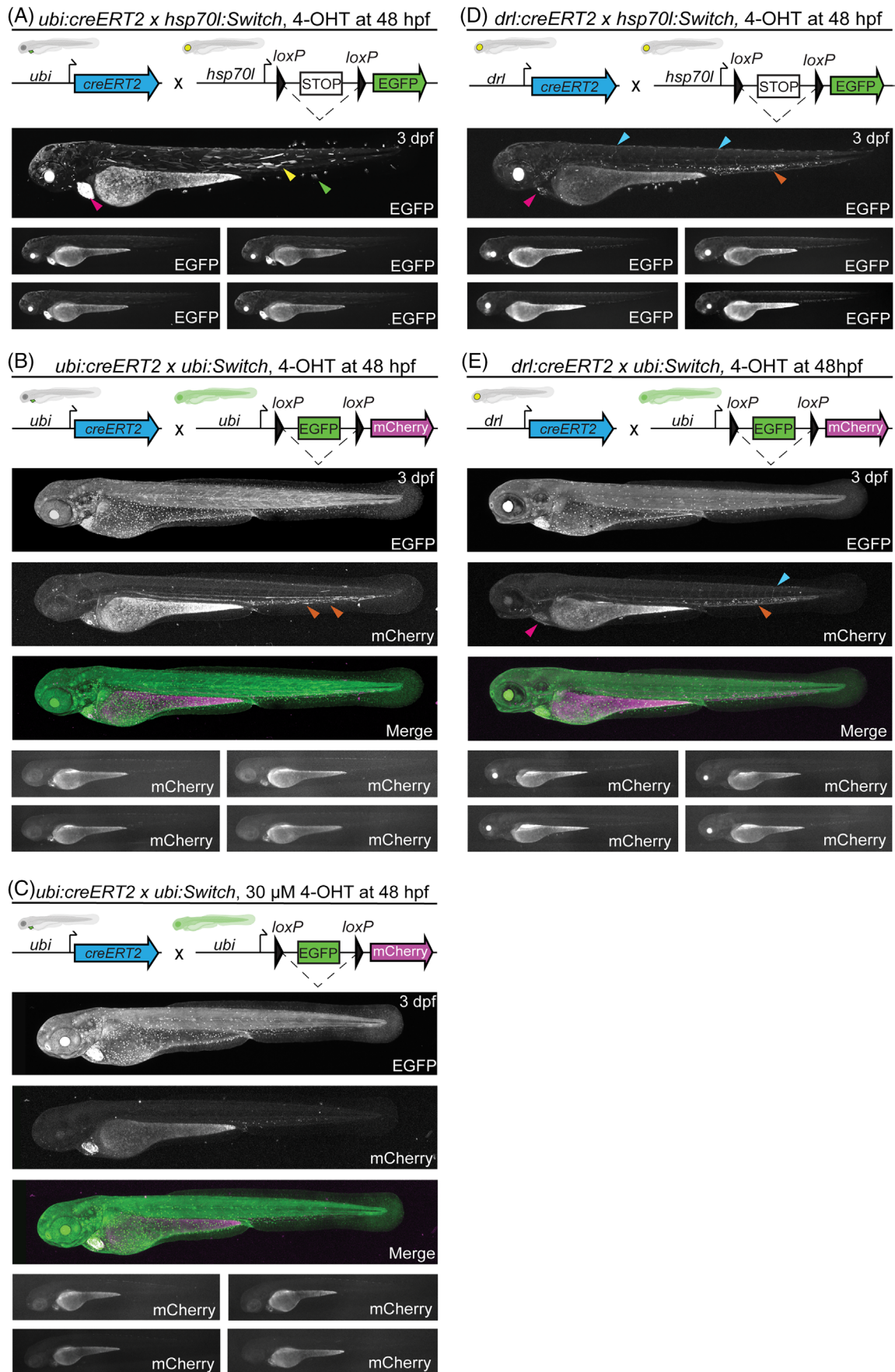


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In contrast to shield stage induction, 4-OHT induction at 48 hpf showed markedly reduced switching at 3 dpf with both *ubi:Switch* and *hsp70l:Switch* (Figure 3A,B). When crossed to *hsp70l:Switch*, 10  $\mu$ M 4-OHT induction at 48 hpf of *ubi:creERT2* resulted in switching predominantly in the skeletal myofibers and the fin fibroblasts (Figure 3A). When crossed to *ubi:Switch* and induced with this regimen, *ubi:creERT2* induced switching predominantly in the blood and heart (Figure 3B). Due to the cardiac *myl7:EGFP* reporter incorporated in the *ubi:creERT2* transgene, we could not evaluate switching efficiency in the heart using *hsp70l:Switch*. While 10  $\mu$ M results in reproducible CreERT2 activity across transgenes with negligible impact on the treated embryos, previous work has indicated that 4-OHT reaches saturation for inducing CreERT2 activity at 25  $\mu$ M.<sup>14,19,29</sup> In line with this observation, increasing 4-OHT concentration from 10  $\mu$ M to 30  $\mu$ M at 48 hpf did not notably increase switching efficiency in *ubi:creERT2;ubi:Switch* embryos (Figure 3C).

To test whether the observed decrease in switching efficiency was specifically associated with the *ubi:creERT2* line, we crossed *drl:creERT2* to *hsp70l:Switch* and *ubi:Switch*, treated with 10  $\mu$ M 4-OHT at 48 hpf, and performed lateral-whole embryo imaging at 3 dpf (Figure 3D,E). *drl*-based labeling is restricted predominantly to LPM-derived cardiovascular, hematopoietic, and fin fibroblast lineages by 48 hpf.<sup>40,41</sup> Accordingly, when crossed to *hsp70l:Switch*, 48 hpf induction of *drl:creERT2* resulted in robust switching in hematopoietic cells with sporadic labeling in the heart, endothelial cells, as well as fin fibroblasts (Figure 3D). When crossed to *ubi:Switch*, 48 hpf induction of *drl:creERT2* again resulted in robust switching in hematopoietic cells, with sporadic labeling in the heart, endothelial cells, and fin fold fibroblasts (Figure 3E). Despite maximum exposure lengths, visualizing mCherry fluorescence at 24 hours post-4OHT induction is challenging with standard stereo microscopy due to the slow reporter accumulation in *ubi:Switch* (Figure 3D,E). We did not observe any switched pectoral

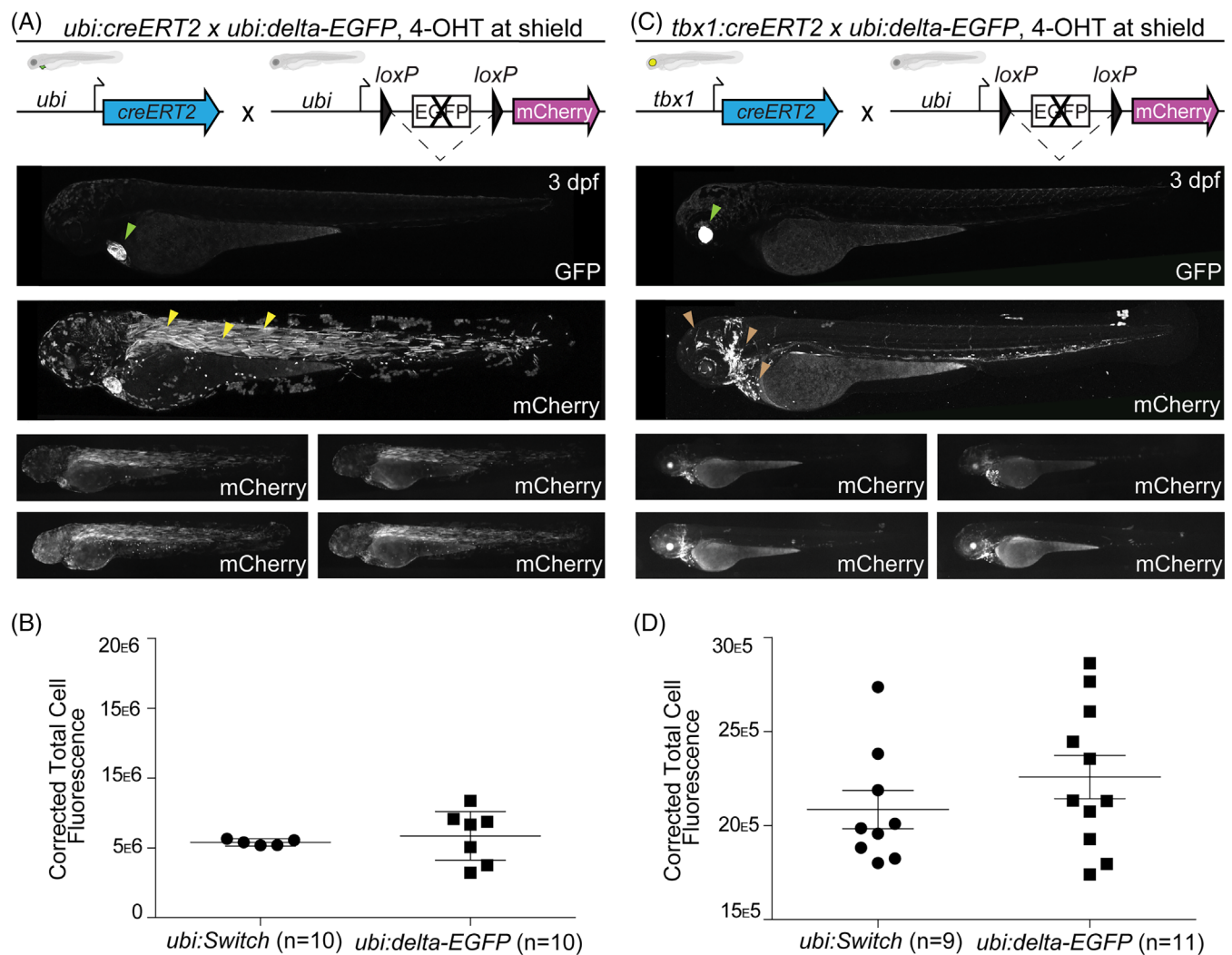
fin cells, further supporting our conclusion that *hsp70l:Switch* provides more thorough recombination reporting than *ubi:Switch*. Together, these results underscore that the original *ubi:creERT2* transgenic (*cz1702Tg*, *ZDB-ALT-110121-1*<sup>14</sup>) features diminished recombination potency at later developmental stages.

## 2.4 | Modifying *EGFP* in *ubi:Switch* does not alter recombination efficiency

CRISPR-Cas9-mediated genome editing has become a standard tool to generate zebrafish mutants. Cas9 can also be harnessed to target transgenic insertions to mutate regulatory elements and transgenic cargo such as fluorescent protein ORFs. *ubi:Switch* expresses EGFP by default, resulting in ubiquitous green fluorescence that prohibits combining the reporter with GFP-based transgenic reporters (Figure 1A). Since we predict integration site is critical to recombination efficiency, creating new stable Tol2 transgenics with modified stop cassettes lacking EGFP would likely show differential switching capacity. Instead, to disrupt EGFP translation and consequently evaluate the impact of in situ modification of a Switch locus in zebrafish, we isolated a CRISPR-induced four base-pair frameshift mutation ( $\Delta 4$ ) in the EGFP ORF, resulting in a premature stop codon (Figure 4). We derived the *ubi:Switch*-based transgenic strain *Tg(ubi:loxP-delta-EGFP-loxP\_mCherry)* (*ubi:delta-EGFP* for short) and compared this new transgene to the original *ubi:Switch* (Figure 4).

When crossed to *ubi:creERT2* and treated with 4-OHT at shield stage, *ubi:delta-EGFP* showed preferential switching in the zebrafish skeletal myofibers at 3 dpf (Figure 4A). To quantify the switching efficiencies of *ubi:Switch* and *ubi:delta-EGFP*, we performed whole-mount lateral view measurements of fluorescence intensity. These measurements did not reveal any significant differences in recombination pattern, mosaicism, or mCherry intensity at these experimental time points ( $n = 10$ ,

**FIGURE 3** *ubi:creERT2* shows decreased 4-OHT and recombination response at larval stages. (A,B,C) *ubi:creERT2* and (D,E) *drl:creERT2* crossed to *ubi:Switch*, and *hsp70l:Switch*, induced with 4-OHT at 48hpf, and imaged laterally at 3 dpf. Schematics of fluorophore cassettes for each Switch transgene and larval schematics representing secondary transgenic markers are shown at the top of each panel; one representative confocal image, and four representative stereo microscope images are presented here (A–E). *ubi:creERT2* shows sparse switching with both reporters when induced with 4-OHT at 48 hpf (A,B,C). Predominant switching occurs in somitic myofiber (yellow arrowheads), and fin fibroblasts (green arrowheads) when crossed to *hsp70l:Switch* (A), and hematopoietic cells (orange arrowhead) when crossed to *ubi:Switch*, induced with 10  $\mu$ M (B) and 30  $\mu$ M (C) 4-OHT. *drl:creERT2* with 4-OHT induction at 48 hpf results in switching in the heart (pink arrowhead), vasculature (blue arrowhead), and hematopoietic cells (orange arrowhead) when crossed to both *hsp70l:Switch* (D), and *ubi:Switch* (E). Note for *ubi:Switch* stereo microscope images that are representative of routine laboratory imaging, extended exposure (30 s) was used to capture all traces of mCherry fluorescence (B,C,E)



**FIGURE 4** Modifying *EGFP* in *ubi:Switch* in situ does not alter recombination efficiency. (A,B) *ubi:creERT2* and (C,D) *tbx1:creERT2* crossed to *ubi:delta-EGFP* that bases on *ubi:Switch* with a disrupted *EGFP* cassette, induced with 4-OHT at shield stage, and imaged laterally at 3 dpf plus corrected total cell fluorescence for quantification (B,D). Schematics of fluorophore cassettes for the *Switch* transgene are shown at the top of each panel and larval schematics represent secondary transgenic markers. One representative confocal image, and four representative stereoscopic images are presented here (A,C). *ubi:creERT2* shows high recombination efficiency with preferential switching in somitic myofiber (yellow arrowheads) (A). No significant difference is observed when *ubi:creERT2* is combined with either *ubi:Switch* or *ubi:delta-EGFP* ( $n = 10$ , unpaired  $t$ -test,  $P > .05$ ) (B). *tbx1:creERT2* switching in the ventricular cardiomyocytes, pharyngeal arches, cranial vasculature, head myofibers and cartilage, and hatching gland with *ubi:delta-EGFP* (brown arrowheads) (C). No significant difference in recombination efficiency is observed between *tbx1:creERT2* crossed to *ubi:Switch* or *ubi:delta-EGFP* ( $n = 9-11$ , unpaired  $t$ -test,  $P > .05$ ) (D). Note visibility of screening markers (*myl7:EGFP* for *ubi:creERT2*; *cryaa:Venus* for *tbx1:creERT2*) when *EGFP* ORF is removed in *ubi:delta-EGFP* (lime arrowhead) (A,C)

Figure 4B). When crossed to *tbx1:creERT2*, *ubi:delta-EGFP* showed labeling in the ventricular cardiomyocytes, pharyngeal arches, cranial vasculature, head myofibers and cartilage, and hatching gland at 3 dpf (Figure 4C), with no significant difference to *tbx1:creERT2*, *ubi:Switch* ( $n = 9-11$ , Figure 4D). From this limited experimental paradigm, we conclude that *ubi:delta-EGFP* broadly preserves the recombination efficiency of *ubi:Switch* after modification of the fluorescent ORF.

## 2.5 | Transgene mapping reveals genomic features at *Switch* reporter integrations

The rational selection of suitable locations for universal, inert, and permissive genomic integration spots for transgenes is highly desirable to minimize position effects. While the majority of zebrafish transgenesis has been performed with random integration methods including

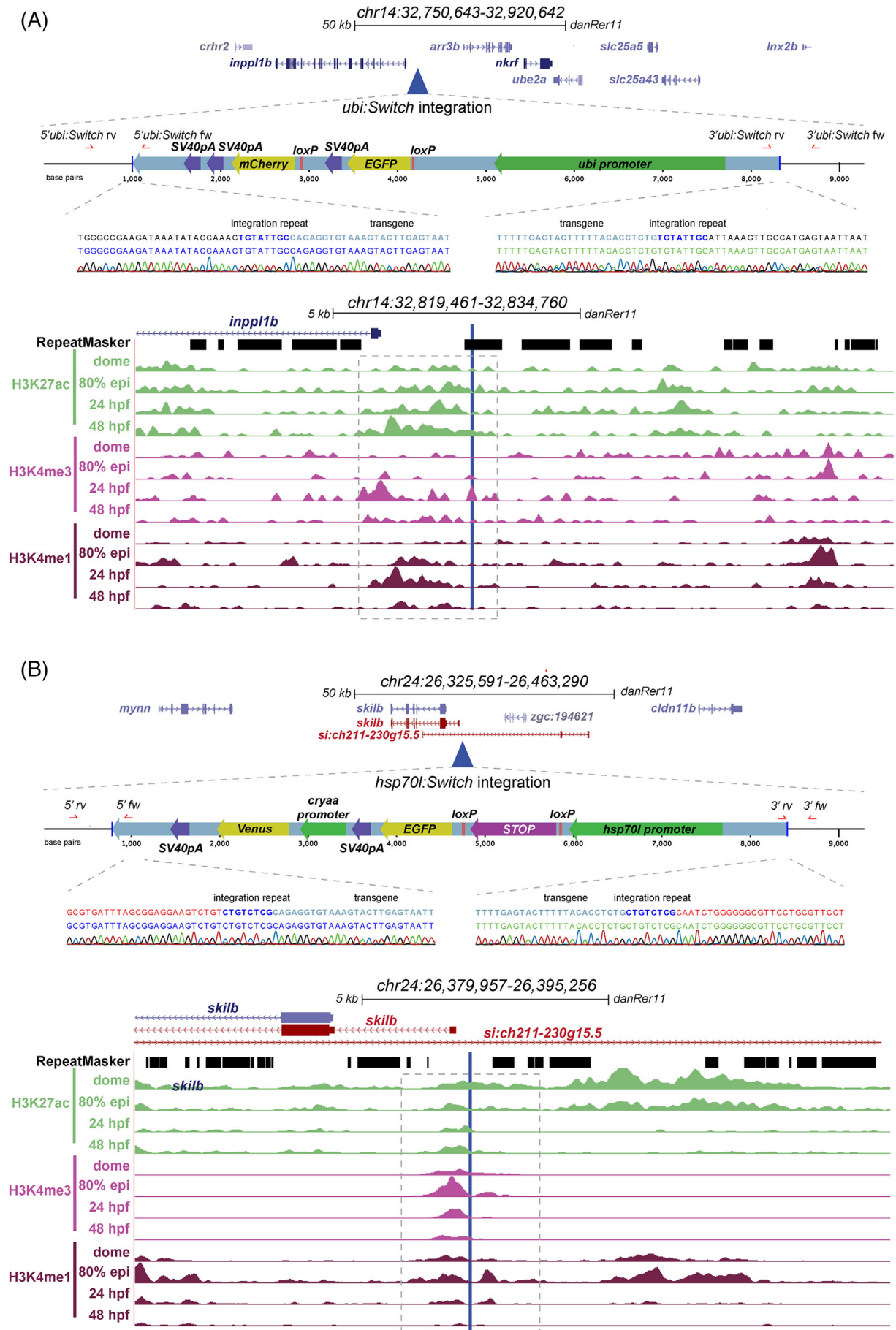


FIGURE 5 Legend on next page.

Tol2 and ISce-I, CRISPR-Cas9 now provides a first means for targeted transgene integrations. However, neither rational design principles for safe harbor sites or long term-validated loci for functional transgene integration, especially of functional *lox*-based *Switch* reporters, have been reported in zebrafish.<sup>1</sup> Our data above established that the *Tol2*-based transgenes *hsp70l:Switch* and *ubi:Switch* integrated into loci that are highly permissive to *loxP* cassette recombination and can be maintained for over a dozen generations,<sup>14,15</sup> possibly providing suitable genomic coordinates for universal transgene landing sites in the genome.

We used Thermal Asymmetric Interlaced (TAIL)-PCR to map the genomic integration loci of the *ubi:Switch* and *hsp70l:Switch* transgenes.<sup>42</sup> In contrast to ligation-mediated-PCR that involves restriction enzyme digestion, relegation, and sequencing, TAIL-PCR uses a series of nested PCRs involving transgene-specific primers and established degenerate primers. We sequenced the resulting TAIL-PCR products, charted their genomic position by BLAST, and confirmed individual transgene integrations with independent primers. The results of our transgene mapping are depicted in Figure 5A,B. The *ubi:Switch* transgene integrated at *chr14:32826253*, proximal and upstream of the gene *inpl1b* and distal and upstream of the gene *arr3b* (Figure 5A); despite this proximity to gene bodies, homozygous *ubi:Switch* zebrafish are viable with no overt morphological phenotypes. The *hsp70l:Switch* transgene integrated at *chr24:26386749*, placing it 122 bp from the transcription start of an annotated transcript of the gene *skilb*, which itself is annotated to be within the gene body of the gene *si:ch211-230g15.5* (Figure 5B). While no loss-of-function alleles for *skilb* or *si:ch211-230g15.5* have been reported to our knowledge, we found also *hsp70l:Switch* to be homozygous viable and fertile with no discernible phenotype.

In contrast to the *ubi:Switch* and *hsp70l:Switch* transgenic lines, we observed significantly less recombination with both the *actb2:Stop-DsRed* and the *actb2:BFP-DsRed* transgenic lines under the same experimental conditions (Figure 1A–D). We again used TAIL-PCR to map the genomic integration loci of the *actb2:Stop-DsRed* and *actb2:BFP-DsRed* transgenes (Figure 6A,B). The *actb2:Stop-DsRed* transgene integrated at *chr7:5941511*, within an intron of two distinct annotated genes: *dusp19b* and *si:dkey-23a13.11* (Figure 6A). The *actb2:BFP-DsRed*

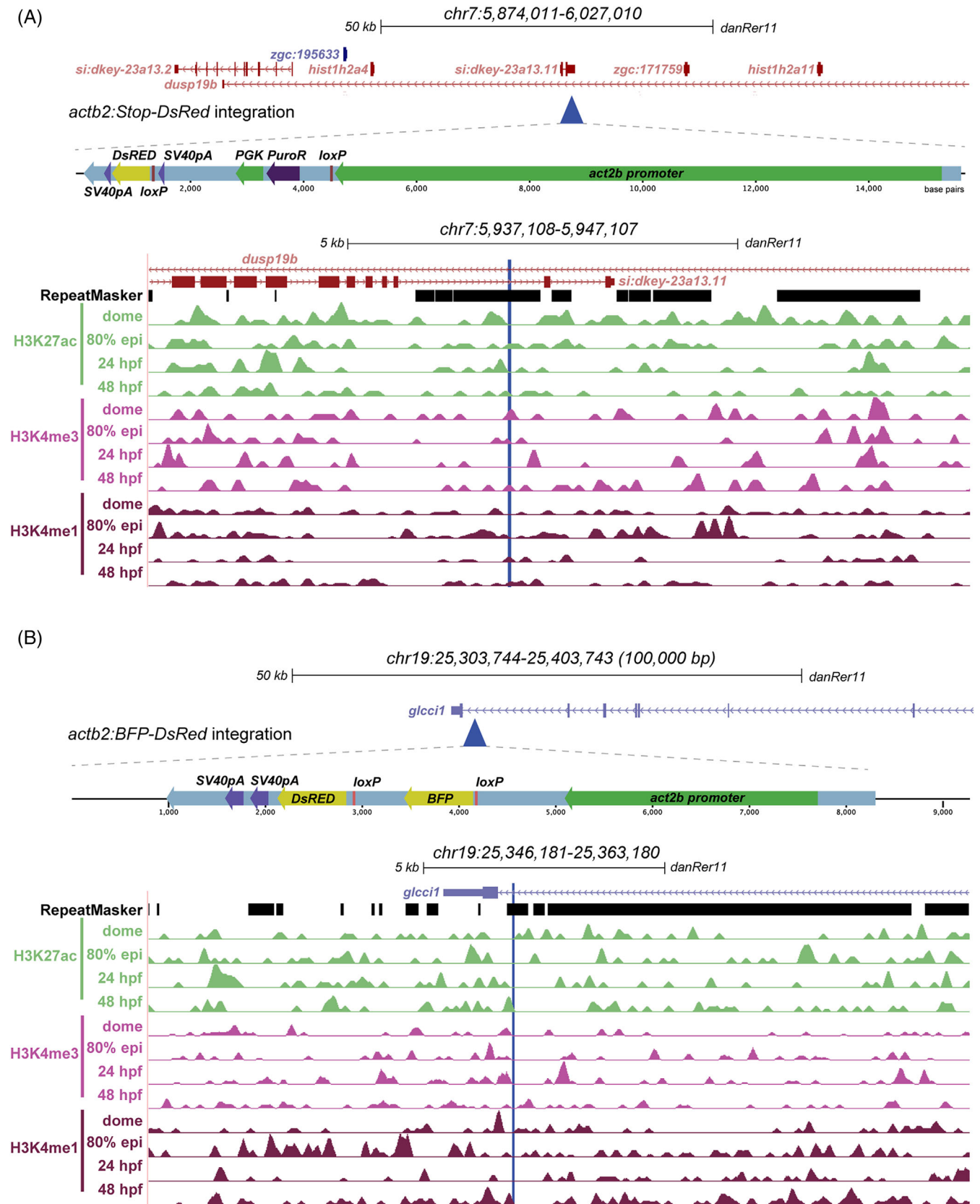
transgene integrated at *chr19:25353744*, within the intron of the gene *glcc1a* (Figure 6B). We were unable to detect any additional integrations of *actb2:Stop-DsRed* using TAIL-PCR.

To gain insight into what genomic features might influence *Switch* transgene recombination efficiency, we compared the integration sites of the mapped transgenes. All four transgenes mapped within or near protein-coding loci, with *ubi:Switch*, *actb2:Stop-DsRed*, and *actb2:BFP-DsRed* landing in highly repetitive regions (Figures 5A and Figure 6A,B). While *ubi:Switch* and *hsp70l:Switch* mapped to intergenic regions (*hsp70l:Switch* may lie within one possible gene body annotation) (Figure 5A,B), both *actb2:Switch* transgenes mapped within introns of annotated and studied genes (*dusp19b* and *glcc1a*) (Figure 6A,B). Using publicly available ChIP-seq data sets,<sup>43</sup> we explored whether distinct histone marks at the integration loci of these individual *Switch* transgenic reporters may associate with high recombination. Distinct from surrounding regions, the *ubi:Switch* integration site showed a series of peaks for H3K27ac, H3K4me3, and H3K4me1 that represent open, active chromatin between 80% epiboly and 48 hpf (Figure 5A). The immediate vicinity of the *hsp70l:Switch* integration locus featured signal for all histone marks between dome stage and 48 hpf (Figure 5B). In contrast, the integration sites of the *actb2*-based *Switch* lines did not show active histone marks at any developmental stage we plotted (Figure 6A,B). Based on a small sample size of four transgene insertions of different recombination qualities, these observations suggest that higher recombination efficiency might correspond with transgene integration in native areas containing active chromatin that may be primed for transcription. In addition, our mapping provides the genomic coordinates of the *ubi:Switch* and *hsp70l:Switch* reporters for potential future modifications and knock-ins of alternative reporters or *lox*-based *Switch* transgenes at these validated loci.

### 3 | DISCUSSION

Transgenic experiments in zebrafish depend on reliable, reproducible reagents including single-copy insertion transgenes with predictable expression patterns in space and time. *Cre/lox* experiments in particular depend on

**FIGURE 5** Transgene mapping reveals genomic features at the *ubi:Switch* and *hsp70l:Switch* loci. (A,B) Genomic integration sites of *ubi:Switch* and *hsp70l:Switch* determined by TAIL-PCR. For each transgene, the integration site is displayed in UCSC genome browser, a schematic of the transgene is depicted, the integration site is aligned to RepeatMasker track, and ChIP-seq tracks for active chromatin marks at key developmental stages. Sequencing reads for 5' and 3' transgene/genome boundaries are provided for *ubi:Switch* and *hsp70l:Switch* integration sites. *ubi:Switch* and *hsp70l:Switch* integration site loci contain peaks for all three chromatin modification signatures (H3K27ac, H3K4me3, H3K4me1, dashed boxes) (A,B)



**FIGURE 6** Transgene mapping reveals genomic features of the tested *actb2*-based *Switch* transgenes. (A,B) Genomic integration sites of *actb2:Stop-DsRed* and *actb2:BFP-DsRed* determined by TAIL-PCR. For each transgene, the integration site is displayed in UCSC genome browser, a schematic of the transgene is depicted, the integration site is aligned to RepeatMasker track, and ChIP-seq tracks for active chromatin marks at key developmental stages (A,B). Note the intronic integration sites for both transgenes and the absence of concise active chromatin marks compared to *ubi:Switch* and *hsp70l:Switch*

efficient *lox*-based *Switch* reporters, the generation of which has so far been serendipitous due to favorable Tol2-based random transgene integrations. Here, we have documented the Cre-dependent recombination or switching efficiencies of four established *lox*-based *Switch* reporters with both ubiquitous and tissue-specific CreERT2 driver lines at two different developmental time points. Further, our results suggest two long-term validated genomic loci as potential safe harbor sites suitable for *lox*-based *Switch* reporters based on their reproducible performance over generations and across labs.

While seemingly straight-forward given access to suitable transgenic lines, Cre/*lox*-based lineage tracing studies require careful consideration to ensure successful experimental outcomes.<sup>1</sup> The specific combination of fluorophores in the *lox*-based reporter may impede visualization if used in parallel with other desired transgenic reporter lines, such as when seeking to define switched cells with a tissue-specific reporter in parallel. Our conversion of *ubi:Switch* to *ubi:delta-EGFP* shows that modulating functional *Switch* transgenics in situ by small edits is feasible without drastically perturbing their properties (Figure 4). Disruption of the *EGFP* ORF in *ubi:delta-Switch* facilitates the visualization of screening markers (*myl7:EGFP*, *cryaa:Venus*, etc.) and enables the use of this line in combination with commonly used EGFP/GFP-based reporter lines.

In Cre/*lox*-based lineage labeling, the onset of fluorescent reporter expression following *loxP* recombination dictates the earliest possible analysis timepoints. As documented in previous work, *ubi*-driven transgenes feature a latency in reporter expression, restricting detectable visualization and imaging to 24 hours or more after Cre-based recombination (Figure 3C,D).<sup>1,14,28,29</sup> We note that, even with extended exposure length for imaging, visualizing mCherry fluorescence 24 hours post-4-OHT induction, remains challenging using standard stereo microscopy (Figure 3C,D). Consequently, *ubi:Switch* may not be the preferred choice of *lox* reporter for short-term trace experiments (ie, analysis within hours following first Cre activity), unless in situ hybridization for *mCherry* mRNA is used as a readout, rather than mCherry fluorescence. However, we here document that a simple heat shock prior to imaging can boost the reporter activity of *ubi:Switch* (Figure 1B); if this pragmatic way to facilitate reporter detection with this transgene is based in heat shock elements in the zebrafish *ubi* promoter or based on other effects warrants further analyses. In contrast, the *hsp70l:Switch* transgenic line bypasses long latency in fluorescent reporter expression, yielding bright EGFP expression within 2 hours postheat-shock (Figures 1B and Figure 2A,B). As a potential caveat, heat-shock efficiency has not been extensively tested in adult *hsp70l:*

*Switch* zebrafish; nonetheless the heat-shock system has been successfully used in adults with *Tg(hsp70:loxP-DsRed-Stop-loxP-EGFP)<sup>uid107</sup>*, as well as other *hsp70l*-based transgenic lines.<sup>15,44-49</sup>

*ubi:creERT2* provides a versatile tool to test new *lox*-based *Switch* lines and to establish 4-OHT regimens.<sup>1,14,29</sup> In our assays, inducing CreERT2 activity with 4-OHT during gastrulation leads to ubiquitous recombination in all tested *Switch* reporters (Figure 1). When combined with *ubi:Switch*, we noted increased mCherry signal in the heart relative to other tissues, possibly a consequence of the heart-specific *myl7:EGFP* transgenesis marker of *ubi:creERT2* that could influence creERT2 expression.<sup>50,51</sup>

Nonetheless, CreERT2 activity from *ubi:creERT2* is drastically reduced by 48 hpf despite substantial evidence that the *ubi* regulatory element remains active throughout all stages of zebrafish development.<sup>14</sup> Our data here adds to the notion that this diminished activity is due to the particular *ubi:creERT2* transgene insertion and not due to issues with 4-OHT uptake of the zebrafish embryo and larva: we have documented efficient switching with *drl:creERT2* when induced at 48 hpf, confirming 10  $\mu$ M 4-OHT penetration in superficial tissues such as fin fibroblasts and blood (Figure 3C,D). Efficient *loxP* recombination has also been documented through labeling of oligodendrocytes at 6 dpf following a 5  $\mu$ M 4-OHT induction at 5 dpf *Tg(mbpa:mCherry-T2A-CreERT2)*, and in hepatocytes in 1-2 year old zebrafish livers following 2  $\mu$ M 4-OHT induction from 5-7 dpf and 10-11 dpf (*tp1:creERT2*).<sup>47,52</sup> Labeled liver hepatocytes were observed at 10 weeks of age using *TRE:creERT2* following 1  $\mu$ M 4-OHT induction at 5 weeks for 3 consecutive days, however no description of 4-OHT delivery was described.<sup>12</sup> The maintained integration of *ubi:creERT2* is therefore best applied for basic tests and at early developmental stages to recombine *lox*-based *Switch* reporters.

Our observations here present the case that, ideally, multiple independent *Switch* lines should be used with individual Cre drivers to confirm findings and avoid biasing lineage information as not all *Switch* lines are created equal (Table 1). *lox*-based *Switch* reporters are highly sensitive to position effects, possibly arising from differential chromatin accessibility across independent transgenic insertions.<sup>1</sup> Our transgene mapping provides first details of the chromatin environment and dynamics at the integration sites of the four transgenes tested here. From this limited sample size, the well-recombining *hsp70l:Switch* stands out as having integrated adjacent to a region with native H3K4me1, K3K4me3, and H3K27Ac marks indicative of open, active chromatin from early developmental stages<sup>43</sup> (Figure 5). The *ubi:Switch* integration locus showed H3K27Ac marks across developmental timepoints, yet lacks H3K4 methylation marks. In

TABLE 1 Summary of the advantages and disadvantages of *Switch* transgenes

Line	Advantages	Disadvantages
<i>ubi:Switch</i>	Second highest recombination efficiency No need for heat-shock	Cannot be combined with other EGFP lines Latency in reporter expression
<i>ubi:delta-EGFP</i>	Can be combined with other EGFP lines Same recombination efficiency as <i>ubi:Switch</i>	No screening marker Latency in reporter expression
<i>hsp70l:Switch</i>	Highest recombination efficiency Easy screening marker cryaa:Venus Rapid transgene expression after heat-shock	Need to heat-shock
<i>actb2:BFP-DsRed</i>	Can be combined with other EGFP lines	Low recombination efficiency
<i>actb2:Stop-DsRed</i>	Can be combined with other EGFP lines	Low recombination efficiency No screening marker Multiple integrations in current generation

Note: Comparison of the relative switching efficiencies and logistical considerations for experimental design using the different *Switch* transgene lines described.

contrast, the two less Cre-responsive *actb2*-based *Switch* lines that integrated into introns show no consistent histone marks associated with open chromatin (Figure 6). Nonetheless, all observed correlation with open histone marks is based on native genome context without inserted transgene, and how the Tol2-based integrations themselves affect the chromatin context remains unknown. Our mapping further uncovered that while *hsp70l:Switch* did not integrate within a repetitive region, *ubi:Switch*, *actb2:BFP-DsRed*, and *actb2:Stop-DsRed* integrated within genomic repeat sequences of different classes (Figures 5 and 6). These first insights should encourage future mapping of Tol2 transgene insertions toward identifying features that support consistent transgene expression also beyond *Switch* lines.

So-called safe harbor sites to integrate transgenes provide a key tool for reproducible transgene deployment in any given model. Previously generated transgene integrations for repeated insertion based on the phiC31 recombinase system have shown weak to medium expression levels, have not been validated for *lox*-based recombination, or have not been maintained beyond proof-of-principle.<sup>1,53-58</sup> Consequently, versatile and validated safe harbor sites are currently missing in zebrafish, rendering Tol2- or ISce-I-based random integration and subsequent screening for functional single-copy transgenes labor-intensive. Recent work has reported the Tol2-based generation of a phiC31-targetable *attB* integration site called *SHH-SBE2* that is suitable for gene-regulatory element analysis through reproducible transgenesis into the same locus<sup>57</sup> while promising, if *SHH-SBE2* shows long-term stability and is amenable to *lox*-based *Switch* reporters warrants further investigation. Retained switching efficiency over generations is a highly desirable feature of *lox*-based *Switch* reporter transgenes.

CRISPR-Cas9-based knocking would provide means to target transgene insertion into suitable loci<sup>21,59-62</sup>; nonetheless, little predictive information is available as to what genomic loci provide safe harbor sites. In mammalian systems, the serendipitously discovered *Hipp11* (*H11*) locus is widely used for reproducible transgenesis.<sup>63-65</sup> Our data argues that, in zebrafish, functional and stable Tol2 insertions of *lox*-based *Switch* transgenes indicate potential safe harbor sites that are also suitable for recombinase-sensitive cassettes. *Ubi:Switch* and *hsp70l:Switch* have been maintained as functional lines in numerous laboratories since their isolation in 2009<sup>14</sup> and 2016,<sup>15</sup> respectively. Considering the high recombination efficiency, and retained activity over generations of both *ubi:Switch* and *hsp70l:Switch* transgenes, we propose these loci as potential safe harbor sites for future transgenic work. A possible approach involves CRISPR-Cas9-based removal of either *Switch* transgene and integration of a phiC31-targeted landing site, such as by targeting the Tol2 transposon arms 5' and 3' of the respective transgene and screening for lost reporter expression. Altogether, our observations argue that mapping of well-working Tol2 transgene insertions, especially of *lox*-based *Switch* reporters, should become more widespread in the zebrafish community to inform about suitable transgene insertion sites with validated activity.

## 4 | EXPERIMENTAL DESIGN

### 4.1 | Zebrafish husbandry and procedures

Animal care and procedures were carried out in accordance with the veterinary office of the IACUC of the

University of Colorado School of Medicine (protocol #00979), Aurora, Colorado. Females from the included Switch lines were crossed with male [regulatory-element]:creERT2 transgenic zebrafish using dividers and embryos were incubated at 28.5°C in E3 medium.

## 4.2 | Drug administration and heat shock protocol

Activity of CreERT2 was induced with 10 µM final concentration of (Z)-4-Hydroxytamoxifen (Sigma Aldrich, St. Louis, Missouri, H7904, abbreviated as 4-OHT) in E3. 4-OHT stock is stored at -20°C in the dark as 10 mM single-use aliquots dissolved in DMSO and used within 2 months of dissolving. Prior to administration, the 4-OHT aliquots were incubated at 65°C for 10 minutes and vortexed. For shield stage treatment, 4-OHT was administered overnight and then replaced with N-Phenylthiourea (Sigma Aldrich, P7629, abbreviated as PTU) at a final concentration of 200 µM in DMSO embryo medium each morning to inhibit melanogenesis. For embryos treated with 4-OHT at 2 dpf, PTU was administered at 24 hpf, 4-OHT/PTU was administered at 48hpf, replaced with PTU at 56 hpf and refreshed the next morning prior to imaging.

*Tg(hsp70l:Switch)* embryos were heat-shocked for 1 hour at 37°C in a water bath. Prior to heat shock, embryos were dechorionated and transferred to a glass vial with E3 medium. Embryos were imaged 2 hours (stereo microscope) and 3 hours (confocal microscope) later.

The *ubi:delta-EGFP* transgenic line was made using the following sgRNA to target the *EGFP* ORF in *ubi:Switch*: 5'-GAGCTGGACGGCGACGTAAA-3'. The mutant allele harbors a 4 bp deletion resulting in a frameshift mutation beginning at *Y40A*, and premature stop at *Stop42K*. *ubi:delta-EGFP* was generated by injecting in vitro-transcribed sg RNA and recombinant Cas9 protein (PNA Bio) into *ubi:Switch* zygotes.<sup>66-68</sup> Resulting F0s were screened for mosaic EGFP expression and a single, highly mosaic individual was selected to establish the line. Mutagenesis was confirmed by Sanger sequencing the PCR product amplified with Fw 5'-TTTAACATGGGAGAAGTGCAAAA-3'; Rev 5'-GTCGTCCTTGAAGAAGATGGTG-3'.

## 4.3 | Imaging

Embryos were anesthetized at 3 dpf with 0.016% Tricaine-S (MS-222, Pentair Aquatic Ecosystems, Apopka, Florida, NC0342409) in E3 embryo medium.

Basic fluorescence imaging was performed on a Leica M205FA with a DFC450 C camera. Laser scanning confocal microscopy was performed on a Zeiss LSM880 following embedding in E3 with 1% low-melting-point agarose (Sigma Aldrich, A9045) on glass bottom culture dishes (Greiner Bio-One, Kremsmunster, Austria, 627861). Images were collected with a ×10/0.8 air-objective lens with all channels captured sequentially with maximum speed in bidirectional mode, with the range of detection adjusted to avoid overlap between channels. Maximum projections of acquired Z-stacks were made using ImageJ/Fiji<sup>69</sup> and cropped and rotated using Adobe Photoshop 2021.

## 4.4 | Quantifications

For CTCF (corrected total cell fluorescence) calculations, lateral images were taken using the stereo microscope and then processed using ImageJ.<sup>29,69</sup> Whole embryos (*ubi:creERT2*) or anterior half (*tbx1:creERT2*) were traced using the magic wand tool and then measured. For every picture, the CTCF (corrected total cell fluorescence) was calculated by the formula “Integrated density whole- (area whole embryo × mean fluorescence background)”. This formula is loosely based on a method described for calculating cell-fluorescence.<sup>70</sup>

For ISV quantification lateral view confocal z-stacks were taken and then max projections were generated using imageJ. %ISVs were manually counted and calculated by dividing total switched ISV/total ISV.<sup>29</sup>

For head vessel and myofiber quantification ventral view confocal z-stacks were taken and then max projections were generated using imageJ. To obtain the total number of head myofibers and vessels, these structures were manually counted using *fli1a:EGFP* embryos stained with the MF20 antibody as a vessel/myofiber control reference, as well as published data that quantified these features.<sup>71,72</sup> Lineage-traced embryos were then imaged and processed in the same manner, and structures were again manually counted.

## 4.5 | Statistics

Unpaired nonparametric (Mann-Whitney) two-tailed *t*-test was done to compare the scores between two groups. For analyses with more than two groups, one-way ANOVA was performed to compare the scores between the groups. Adjusted *P*-values after multiple tests correction are reported and significance was set at *P* < .05.

## 4.6 | MF20 immunostaining

*fli1a:EGFP* zebrafish<sup>73</sup> were dechorionated and fixed at 3 dpf with 4% formaldehyde, 0.1% Triton-X100 in PEM (0.1 M PIPES, 2 mM MgSO<sub>4</sub>, and 1 mM EDTA) overnight at 4°C. The next day, embryos were washed in 1× PBS with 0.1% Triton-X100 and 0.1% BSA, permeabilized for 1 hour at RT with ProtK diluted in PBS to a final concentration of 5 µg/µl and next for 30 minutes in 1× PBS with 0.5% Triton-X100 for 30 minutes. After washing, embryos were blocked in PBS + 1% goat serum and 0.1% Triton-X100 for 2 hours at room temperature and incubated overnight at 4°C with primary antibody MF20 (DSHB, antibody ID# AB\_2147781) 1:50 diluted in blocking buffer. The third day, embryos were washed three times 10 minutes in 1× PBS with 0.1% Triton-X100 and 0.1% BSA and incubated overnight at 4°C with the secondary antibody goat-anti-mouse A568 1:500 (Abcam, antibody ID# ab175473). On the fourth day, embryos were washed 3× 30 minutes with 1× PBS with 0.1% Triton-X100 and 0.1% BSA and kept at 4°C in Vectashield with DAPI diluted 1:5 in the same wash buffer until imaging.

## 4.7 | TAIL-PCR

Protocol based on previous work<sup>42,53</sup>:

1. Extract DNA from fin clips or single embryos.
2. Prepare primer mixtures: Combine 1.5 µM of each *Tol2*-specific primer with 10 µM of each AD primer.
3. Pipet primary PCR reaction: 1 µL Genomic DNA, 4 µL Primer mix (primary + random), 2 µL 10× Polymerase Buffer, 2 µL dNTPs (2 mM), 0.2 µL Polymerase (Expand High Fidelity PCR System, Roche), 10.8 µL ddH<sub>2</sub>O.
4. Perform *primary* PCR step (see PCR conditions below).
5. Dilute 2 µL of the primary PCR product in 25 µL ddH<sub>2</sub>O and add to the secondary reaction: 2 µL Diluted PCR product, 4 µL Primer mix (secondary + random), 2 µL 10× Polymerase Buffer, 2 µL dNTPs (2 mM), 0.2 µL Polymerase, 9.8 µL ddH<sub>2</sub>O.
6. Perform *secondary* PCR reaction (see PCR conditions below).
7. Dilute 2 µL of the secondary PCR product in 25 µL ddH<sub>2</sub>O and add to tertiary reaction. Pipette reaction analogous to the secondary reaction with tertiary primer and secondary product.
8. Perform *tertiary* PCR reaction (see PCR conditions below).
9. Compare secondary and tertiary PCR product on a 1.5% (w/v) agarose gel. If the amplification of the

genomic region was successful, a slight band shift should occur between the secondary and the shorter tertiary product.

10. Excise bands and purify with QIAquick gel extraction kit (#NA1111-1KT).
11. Sequence purified PCR products.
12. Use BLAST (<http://blast.ncbi.nlm.nih.gov/>) to align to the zebrafish genome and identify 5' and 3' flanking regions.

The following gene-specific primers and random primers (AD3, 5, 6, 11) were used for TAIL-PCR:

*Tol2 5'1 forward* GGGAAAATAGAATGAAGTGATCTCC  
*Tol2 TIR 5'.*

*Tol2 5'2 forward* GACTGTAAATAAAAATTGTAAGGAG  
*Tol2 TIR 5'.*

*Tol2 5'3 forward* CCCCAAAATAATACTTAAGTACAG  
*Tol2 TIR 5'.*

*Tol2 3'1 reverse* CTCAAGTACAATTTTAATGGAGTAC  
*Tol2 TIR 3'.*

*Tol2 3'2 reverse* ACTCAAGTAAGATTCTAGCCAGA  
*Tol2 TIR 3'.*

*Tol2 3'3 reverse* CCTAAGTACTTGTACTTTCACCTTG  
*Tol2 TIR 3'.*

*AD-3 WGTGNAGNANCANAGA* Random.

*AD-5 WCAGNTGWTNGTNTCTG* Random.

*AD-6 STTGNTASTNCTNTGC* Random.

*AD-11 NCASGAWAGNCSWCAA* Random.

The primers needed for the mapping of *Tol2* integrations as described previously.<sup>74</sup> *W*: Weak base (*A* or *T*), *S*: Strong base (*C* or *G*), *N*: Any base (*A*, *C*, *G*, *T*).

The following PCR conditions were used to TAIL-PCR:

Primary PCR reaction:

1. 2 minutes 94°C
2. 0.5 minute 94°C
3. 1 minute 62°C
4. 2.5 minutes 72°C
5. Repetition of steps 2-4 (5×)
6. 0.5 minute 72°C
7. 3 minutes 25°C
8. Ramping 0.3°C/s to 72°C
9. 2.5 minutes 72°C
10. 10 seconds 94°C
11. 1 minute 61°C
12. 2.5 minutes 72°C
13. 10 seconds 94°C

Secondary PCR reaction:

1. 10 seconds 94°C
2. 1 minutes 61°C
3. 2.5 minutes 72°C

4. 10 seconds 94°C
5. 1 minutes 61°C
6. 2.5 minutes 72°C
7. 10 seconds 94°C
8. 1 minutes 44°C
9. Ramping 1.5°C/s to 72°C
10. 2.5 minutes 72°C
11. Repetition of steps 1-10 (15×)
12. 5 minutes 72°C

Tertiary PCR reaction:

1. 0.25 minutes 94°C
2. 1 minutes 44°C
3. Ramping 1.5°C/s to 72°C
4. 2.5 minutes 72°C
5. Repetition of steps 1-4 (30×)
6. 5 minutes 72°C

#### 4.8 | Transgene mapping confirmation

The following locus-specific primers were used to confirm the integration sites of *ubi:Switch* and *hsp70l:Switch*:

5' *ubi:Switch* Fw GGAGCATTCAGAGGTACC.

5' *ubi:Switch* Rev GACTGTAAATAAAATTGTA AGGAG.

3' *ubi:Switch* Fw CTCAAGTACAATTTTAATGG AGTAC.

3' *ubi:Switch* Rev GCTGTGAGACGATCAGGC.

5' *hsp70l:Switch* Fw GCATGACACGGCTAACCAAC.

5' *hsp70l:Switch* Rev GACTGTAAATAAAATTGTA AGGAG.

3' *hsp70l:Switch* Fw CCTAAGTACTTGTACTTTC ACTTG.

3' *hsp70l:Switch* Rev TATCAGCACACACCTTT ATCGC.

We were unable to fully confirm the integration sites for the *actb2* Switch transgenes with independent locus-specific primers. We confirmed the 5' genomic border of the *actb2:BFP-DsRed* transgene, however the 3' genomic border and both 5' and 3' genomic borders of *actb2:Stop-DsRed* could not be confirmed.

#### AUTHOR CONTRIBUTIONS

**Robert L. Lalonde:** Conceptualization (lead); data curation (lead); formal analysis (lead); investigation (lead); methodology (lead); validation (equal); visualization (equal); writing – original draft (lead); writing – review and editing (equal). **Cassie L. Kemmler:** Data curation (equal); formal analysis (equal); investigation (equal); methodology (equal); visualization (equal); writing – original draft (equal); writing – review and editing (equal).

**Frédérique W. Riemsdagh:** Data curation (equal); formal analysis (equal); investigation (equal); methodology (equal); writing – original draft (equal). **Andrew J. Aman:** Investigation (supporting); methodology (supporting); resources (supporting); writing – review and editing (supporting). **Jelena Kresoja-Rakic:** Data curation (supporting); formal analysis (supporting); investigation (supporting). **Hannah R. Moran:** Data curation (supporting); formal analysis (supporting); investigation (supporting); methodology (supporting); writing – review and editing (supporting). **Susan Nieuwenhuize:** Formal analysis (supporting); investigation (supporting); methodology (supporting). **David Parichy:** Conceptualization (supporting); data curation (supporting); methodology (supporting); resources (supporting); validation (supporting); writing – review and editing (supporting). **Alexa Burger:** Conceptualization (equal); funding acquisition (equal); project administration (equal); resources (equal); supervision (equal); Writing – review and editing (equal). **Christian Mosimann:** Conceptualization (equal); funding acquisition (equal); methodology (equal); project administration (equal); resources (lead); supervision (supporting); writing – review and editing (supporting).

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#### CONFLICT OF INTEREST

The authors declare no competing interests.

#### DATA AVAILABILITY STATEMENT

All data and reagents are contained within article, raw data of feature counts for quantifications available upon request.


#### ETHICS STATEMENT

Animal care and procedures were carried out in accordance with the veterinary office of the IACUC of the

University of Colorado School of Medicine (protocol #00979), Aurora, CO, USA.

## ORCID

Robert L. Lalonde  <https://orcid.org/0000-0002-1803-0983>

Cassie L. Kemmler  <https://orcid.org/0000-0002-4516-029X>

Fr derike W. Riemsdagh  <https://orcid.org/0000-0002-8530-3339>

Jelena Kresoja-Rakic  <https://orcid.org/0000-0003-3202-4659>

Hannah R. Moran  <https://orcid.org/0000-0001-8056-5762>

Alexa Burger  <https://orcid.org/0000-0001-7137-3910>

Christian Mosimann  <https://orcid.org/0000-0002-0749-2576>

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