Specific and spatial labeling of P0-Cre versus Wnt1-Cre in cranial neural crest in early mouse embryos

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Abstract

P0-Cre and Wnt1-Cre mouse lines have been widely used in combination with loxP-flanked mice to label and genetically modify neural crest (NC) cells and their derivatives. Wnt1-Cre has been regarded as the gold standard and there have been concerns about the specificity of P0-Cre because it is not clear about the timing and spatial distribution of the P0-Cre transgene in labeling NC cells at early embryonic stages. We re-visited P0-Cre and Wnt1-Cre models in the labeling of NC cells in early mouse embryos with a focus on cranial NC. We found that R26-lacZ Cre reporter responded to Cre activity more reliably than CAAG-lacZ Cre reporter during early embryogenesis. Cre immunosignals in P0-Cre and reporter (lacZ and RFP) activity in P0-Cre/R26-lacZ and P0-Cre/R26-RFP embryos was detected in the cranial NC and notochord regions in E8.0–9.5 (4–19 somites) embryos. P0-Cre transgene expression was observed in migrating NC cells and was more extensive in the forebrain and hindbrain but not apparent in the midbrain. Differences in the Cre distribution patterns of P0-Cre and Wnt1-Cre were profound in the midbrain and hindbrain regions, that is, extensive in the midbrain of Wnt1-Cre and in the hindbrain of P0-Cre embryos. The difference between P0-Cre and Wnt1-Cre in labeling cranial NC may provide a better explanation of the differential distributions of their NC derivatives and of the phenotypes caused by Cre-driven genetic modifications.

KEYWORDS
derivation, lineage tracing, mouse, neural crest, Wnt1-Cre, P0-Cre

1 INTRODUCTION

Neural crest (NC) is a multipotent cell population derived from the lateral ridges of the neural plate in early vertebrate embryos (Leikola, 1976; Trainor, 2015). After delamination from the neural folds, NC cells migrate ventrally and extensively, giving rise to a wide variety of differentiated cell types, for example, neurons, glia, bone, cartilage, and connective tissue of the head (Bronner-Fraser, 2004; Crane and Trainor, 2006; Meulemans and Bronner-Fraser, 2004; Munoz and Trainor, 2015; Trainor, 2005b). Labeling and tracking NC have been essential for understanding the NC contribution to different tissue and cell types, which is fundamental for organogenesis.
Recently, cell lineage tracing using NC-specific promoter driven Cre transgenic mouse lines in combination with Cre reporter mouse lines has facilitated genetic marking of NC cells and their derivatives. Multiple Cre transgenic mouse lines have been generated using a NC marker gene promoter, for example, Wnt1-Cre (Echelard, Vassileva, & McMahon, 1994; Jiang, Iseki, Maxson, Sucov, & Morris-Kay, 2002; McMahon, Joyner, Bradley, & McMahon, 1992), P0-Cre (Wang, Komatsu, & Mishina, 2011; Yamauchi et al., 1999; Zhang, Marsh, Ratner, & Brackenbury, 1995), Dhh-Cre (Gershon et al., 2009; Wang, Kumar, Mitsios, Slevin, & Kumar, 2007), Pax3-Cre (Jarad and Miner, 2009), HtPA-Cre (Lee et al., 2013), Sox10-Cre (Simon, Likert, Gotz, & Dimou, 2012), Mef2c-F10N-Cre (Aoto et al., 2015). Use of these models has yielded new data pertaining to NC cell specificities in mice, for example, distinct genetics of skin-derived precursors in craniofacial and dorsal skin from NC and mesoderm, respectively (Jinno et al., 2010). NC and placodal derivation of the otic vesicle (Freyer, Aggarwal, & Morrow, 2011), a dual origin of sensory organs that include olfactory epithelium (Katoh et al., 2011) and taste bud cells (Bogs et al., 2016; Liu, Komatsu, Mishina, & Mistretta, 2012), and a dual origin of tooth bud epithelium (Wang et al., 2011).

However, variations/discrepancies in the different models that label NC cell lineage have been reported, such as in tooth (Wang et al., 2011); olfactory epithelium (Suzuki, Yoshizaki, Kobayashi, & Osumi, 2013); heart (Cavanaugh, Huang, & Chen, 2015; Millgram-Hoffman et al., 2014; Nakamura, Colbert, & Robbins, 2006; Tomita et al., 2005), and taste organs (Boggs et al., 2016; Liu et al., 2012). In comparative studies on NC contributions to specific lineages, the Wnt1-Cre and P0-Cre lines have been widely used (Katoh et al., 2011; Liu et al., 2012; Morikawa et al., 2009; Nagoshi et al., 2008, Nagoshi et al., 2011; Nakamura et al., 2006; Yoshida et al., 2006; Yoshida et al., 2008). Wnt1-Cre has been regarded as the gold standard for NC lineage, however, Wnt1-Cre also labels cells in the neural tube that makes researchers question the specificity of this line and other models when a difference is observed compared to Wnt1-Cre (Trainor, 2005a). In fact, a difference between Wnt1-Cre and P0-Cre in labeling NC has been found in multiple organs (Freen et al., 2010; Kawakami et al., 2011; Lewis et al., 2013; Liu et al., 2012; Wang et al., 2011). In contrast to Wnt1-Cre, which has been reported to trace NC cells as early as E8.5 using a ROSA26 (hereafter, R26-lacZ) Cre reporter in mice (Chai et al., 2000; Ikeya et al., 1997; Jiang et al., 2002; McMahon et al., 1992), P0-Cre has been reported to label NC derivatives at stages later than E9.0 using CAAG-CAT-lacZ (hereafter, CAAG-lacZ) as a reporter system (Yamauchi et al., 1999). Although in situ hybridization for P0-Cre transgene expression was performed (Yamauchi et al., 1999), there is no detailed information available regarding the labeling of P0-Cre in NC cells at early stages.

In the present study, the expression of two commonly used reporter transgenes, CAAG-lacZ and R26-lacZ, were compared in early embryos, and we found that DNA recombination was not consistent in the CAAG-lacZ reporter when driven by Mec2-Cre in early embryos. To further evaluate the timing, specificity and spatial distribution of P0-Cre and Wnt1-Cre activity, we used the R26-lacZ and R26-tdTomato (R26-RFP) reporters and carried out immunoreactions for Cre. Subsequently, we thoroughly compared the presence of Cre recombinase in pre- and post-migratory NC cells in P0-Cre and Wnt1-Cre embryos. Our data indicate that the P0-Cre transgene expression is specifically distributed in migrating cranial NC in addition to the notochord, and Cre immunosignals were especially extensive in the forebrain and hindbrain regions. In contrast, Wnt1-Cre transgene expression was obvious in forebrain and midbrain NC cells but was sparse in migrating NC in the hindbrain regions.

2 MATERIALS AND METHODS

2.1 Animals

Animals were maintained and used in compliance with institutional animal care protocols of the University of Georgia, the University of Michigan and the National Institute of Environmental Health, and in accordance with the National Institutes of Health Guidelines for Care and Use of Animals in research.

Mice carrying Cre recombinase driven by the protein zero (P0) promoter (P0-Cre mouse, C57BL/6J-Tg(P0-Cre)94Imeg (ID 148)), were provided by CARD, Kumamoto University, Japan (Yamauchi et al., 1999) and were mated with C57BL/6 wild type (WT) or Cre-reporter mice R26-lacZ (Soriano, 1999) or ZEG (LacZ/EGFP) (Novak et al., 2000) or R26-tdTomato (RFP) (B6.Cg-Gt ROSA26Sor tm14(CAG-tdTomato)1Hze/J, Jackson Lab, Stock #007914) to generate P0-Cre/WT, P0-Cre/R26-lacZ, P0-Cre/ZEG, and P0-Cre/RFP mice. Mec2-Cre mice, which express Cre in the epiblast from E5.5 (Tallquist and Soriano, 2000), were also bred with R26-lacZ or Cre-reporter CAAG-lacZ mouse line (Sakai and Miyazaki, 1997). Wnt1-Cre mice (B6C3-TgWnt1-cre1Rth Tg(Wnt1-1Gal4)11Rth/J, Jackson Lab, Stock #009107) were bred with C57BL/6 WT or Cre-reporter mice R26-lacZ (Soriano, 1999) or R26-RFP to generate Wnt1-Cre/WT or Wnt1-Cre/R26-lacZ or Wnt1-Cre/RFP. R26-lacZ reporter mice were maintained on a mixed background of C57BL/6 (majority) and 129S6.

2.2 Tissue collection

Timed pregnant mice were euthanized using CO2 followed by cervical dislocation to ensure death. The uterus with embryos was removed and embryos were retrieved under a microscope. The tissues were further dissected and processed for analysis, as described below.

Embryos from P0-Cre mice were harvested at embryonic day (E) 8.0, E8.5, E9.5, and E10.5. Newborn P0-Cre/ZEG mice were harvested immediately after birth. Mec2-Cre mouse embryos carrying a Cre-reporter were obtained from E6.5 to E11.5. Wnt1-Cre mouse embryos were collected at E8.0–E8.5. Noon of the day on which the dam was positive for vaginal plug was designated as E0.5. The stages of these embryos were confirmed by comparing the sizes and morphologies of multiple organs. The number of pairs of somites was counted under a microscope at the time of dissection.
2.3 | Genotyping

Specific primers were used to genotype the mice. Primers A (5′-CTG GTG ATC TGC AAC TCC AGTC-3′) and B (5′-GAG ACT AGT GAG ACG TGC TACT-3′) were used to amplify fragments from the un-recombined R26-lacZ reporter allele (~550 bp). Primers A and C (5′-TGT GAG CGA GTA GAC ACA ACC-3′) were used to detect the Cre-recombined allele (~680 bp). Primers E (5′-CAG TCA GTT GCT CAA TGT ACC-3′) and F (5′-ACT GGT GAA ACT CAC CCA-3′) were used to amplify fragments from the un-recombined CAAG-lacZ allele (~450 bp), whereas primers D (5′-GTG CTG GTT ATT GTG CTG TCTC-3′) and C were used to detect Cre-dependent deletion of the floxed region (~680 bp). To detect a Cre transgene, Cre1 (5′-GGA CAT GTT CGG GAT CGC CAG GCG-3′) and Cre2 (5′-GCA TAA CCA CGA TAA CAG CAT TGC TG-3′) primers were used.

2.4 | X-gal staining

After dissection, the embryos were fixed with 4% paraformaldehyde for 5 min at 4°C, and washed three times with 0.1 M sodium phosphate buffer (PBS, pH 7.5), then stained with 1 mg mL⁻¹ of 5-bromo-4-chloro-3-indolyl-beta-D-galactoside (X-gal, Sigma, Cat# B-4252) in 0.1 M PBS, 0.1% sodium deoxycholate, 0.2% Nonidet P-40, 2 mM magnesium chloride, 5 mM potassium ferricyanide, and 5 mM potassium ferrocyanide at 37°C.

2.5 | Immunohistochemistry

Embryos were fixed with 4% paraformaldehyde in 0.1 M PBS (pH 7.3) for 2 hr at 4°C, rinsed in PBS and then infiltrated with 20-30% sucrose prior to embedding in OCT/Tissue Tek (Sakura Finetek). Serial sections were cut at 10-μm thickness. Sections were air dried, rehydrated and blocked with 10% normal donkey serum for 1 hr at room temperature. Primary antibodies against the following proteins were applied: Cre recombinase (1:500, MAB3120, EMD Millipore, Billerica, MA), FOXA2 (1:500, 07-633, EMD Millipore, Billerica, MA), GFP (1:100, ab13970, abcam, Cambridge, MA), p75 (1:200, sc-6188, Santa Cruz Biotechnology, Dallas, TX), SOX9 (1:1000, sc-20095, Santa Cruz Biotechnology, Dallas, TX; 1:1000, ab185966, abcam, Cambridge, MA). Primary antibodies were diluted with 0.1 M PBS that contained 0.3% Triton X-100 and 1% normal donkey serum. After incubation with primary antibody overnight, sections were washed three times with 0.1 M PBS, and then incubated with secondary antibodies that were conjugated with Alexa Fluor 488, Alexa Fluor 546 or 647 (1:500, Invitrogen) for 1 hr at room temperature in 0.1 M PBS containing 0.3% Triton X-100 and 1% normal donkey serum. Sections were counterstained with DAPI solution for 5-10 min to visualize nuclei and were then rinsed in PBS. After air drying, the slides were mounted with ProLong® Gold antifade medium (Invitrogen, Eugene, OR), and the sections were examined thoroughly using a light microscope (EVOS FL, Life Technologies). Colocalization of different immunosignals was confirmed and photographed using a laser-scanning confocal microscope (Zeiss LSM 710).

To confirm the specificity of immunosignals, slides without the addition of primary antibodies were used as negative staining controls for all immunostainings in the present study. For determining specificity of Cre immunolabeling, Cre negative littermate embryos were sectioned and immunoreacted with the same concentration of primary and secondary antibodies. Additionally, Western blotting was conducted using E9.5 P0-Cre embryos (Cre⁺ embryos and Cre⁻ littermate control) to further investigate the specificity of antibodies against Cre recombinase (1:5000, MAB3120, EMD Millipore, Billerica, MA). A ubiquitously expressed protein, IκB kinase complex associated protein (IKAP) (1:5000, sc-8336, Santa Cruz Biotechnology) was used as a loading control. Equal amounts of protein were used for electrophoresis in sodium dodecyl sulfate-polyacrylamide gel and transferred to a nitrocellulose membrane. Procedures for blocking and antibody probing were performed as described (Li et al., 2011). A single band at the expected molecular weight was detected in a genotype-dependent manner with the Cre antibody (Supporting Information Figure 1).

3 | RESULTS

3.1 | Meox2-Cre driven R26-lacZ reporter expression emerges earlier and is more consistent than CAAG-lacZ

ROSA26 Cre reporter (hereafter, R26-lacZ) and CAAG-CAT-lacZ Cre reporter (hereafter, CAAG-lacZ) mice have been widely used for cell fate mapping. First, we compared the reporter sensitivity of these two lines at early embryo stages (E6.5–E11.5, Figure 1) using the Meox2-Cre transgenic line in which Cre activity emerges as early as E5.5 in the epiblast (Tallquist and Soriano, 2000). In Meox2-Cre/R26-lacZ mice, X-gal staining of whole embryos showed that signals appeared in the earliest stage examined (E6.5) and that labeled cells were extensively distributed in the whole embryos throughout all the stages examined (Figure 1a) similarly to the previous report (Tallquist and Soriano, 2000). In contrast, β-galactosidase (β-gal) activity in Meox2-Cre/CAAG-lacZ embryos was not detected until E7.5 (n = 5 or more for each stage). Weak signals were detected in the cardiac region at E8.5 (2/5) and in the somites and heart region at E9.5 (2/5) (Figure 1b), but were not apparent in head and other tissues in the trunk regions. At E10.5 and E11.5 (6/6 for each stage), the distribution of CAAG-lacZ expression in Meox2-Cre/CAAG-lacZ embryos was similar to that of Meox2-Cre/R26-lacZ throughout the entire embryo (Figure 1b).

To examine whether the discrepancy between Meox2-Cre driven R26-lacZ and CAAG-lacZ reporter activity was caused by DNA recombination efficiency, promoter silencing, or both, specific primers were designed to detect unrecombined and recombined reporter alleles (Figure 1c,d). A fragment of ~550 bp was produced using primers A and B to detect the un-recombined R26-lacZ alleles. A Cre-recombined R26-lacZ allele was detected using primers A and C, and the product was approximately 680 bp (Figure 1c). For the CAAG-lacZ reporter system, primers E and F were used to generate an ~450 bp fragment from unrecombined allele whereas primers D and C produced an ~680 bp fragment from the Cre-mediated allele (Figure 1d). Both Cre transgenes
were detected using primers Cre1 and Cre2 to generate an 270 bp fragment (Figure 1c,d).

In E8.5 Meox2-Cre/R26-lacZ embryos, the recombined allele was reliably detected when both Cre and R26-lacZ transgenes were present (Figure 1e, lane 1–3) compared to the absence of bands without Cre transgene (Figure 1e, lane 4–7). In contrast, in Meox2-Cre/CAAG-lacZ mice only half of the double transgenic embryos showed detectable recombined allele at E8.5 (Figure 1f, E8.5, lane 1–2 vs. 4–5). The
A recombined allele was reliably detected at E9.5 (Figure 1f, E9.5, lane 5–7). These data suggest that Cre-dependent DNA recombination is inconsistent in CAAG-lacZ early stage (≤E8.5) embryos.

3.2 | P0-Cre/R26-lacZ labels NC cells and derivatives in early embryos

P0-Cre/CAAG-lacZ has been reported to label NC cells and derivatives after E9.0 (Yamauchi et al., 1999). Because the R26-lacZ reporter is more accessible to Cre-mediated DNA recombination than the CAAG-lacZ described above, we used the R26-lacZ reporter to re-examine P0-Cre transgene activity in early embryos at different somite stages using X-gal staining in whole embryo tissue.

At the 4-somite stage (E8.5) (Figure 2b,b'), β-gal positive signals were found in the forebrain, hindbrain (arrows, Figure 2b'), first branchial arch and midline of the trunk neural groove region (Figure 2b,b'). The staining pattern in the 9-somite embryo was similar to that of the 6-somite stage but extended more caudally and β-gal signals were obvious in the midline and otic placodes (arrows, Figure 2c,c'). At the 12-somite (Figure 2d,d'), intense staining was found in the hindbrain region, branchial arches, otic placodes (arrows, Figure 2d'), and optic eminences. In the trunk region, signals were apparent in the neural groove area and in lateral tissues (Figure 2d,d'). At the 17-somite stage, positive signals were extensively distributed in the embryo, adding more labeled tissue regions that included the heart, frontonasal, and eye regions (Figure 2e). At the 22–26 somite stages, β-gal positive signals were extensively distributed in NC derivatives throughout the body, for example, frontonasal and eye regions, branchial arches, heart, and dorsal root ganglia (Figure 2f,g).
Notably, signals in the midbrain region were not apparent throughout the examined stages (Figure 2a–g). In littermate control embryos, no positive signals were observed within the embryos at the stages examined (Figure 2a–g).

### 3.3 | P0-Cre transgene expression is restricted to migrating NC cells and notochord in early embryos

To analyze P0-Cre expression in cranial NC, we used a monoclonal antibody against Cre to examine Cre immunoreactivity at different early somite stages of embryos. Specificity of Cre immunosignals were validated by three different methods: (1) Western blotting analysis to demonstrate the presence of a single band of expected size in a genotypedependent manner (Supporting Information Figure 1), (2) immunohistochemistry using Cre embryos labeled with primary antibody to show no signals (Supporting Information Figure 2), and (3) immunohistochemistry using Cre embryos without the primary antibody to determine background levels (Figures 3 and 4).

β-gal signals in the P0-Cre/R26-lacZ embryo emerged in the neural groove region during the 4-somite stage (Figure 2a). We further identified the location and types of cells that express P0-Cre transgene through investigating immunoreactivity for Cre recombinase and colocalization with a NC cell marker SOX9 (Figure 3a,b) and a notochord marker FOXA2 (Figure 3c) in serial sections of 4-somite embryos (n = 7). Cre+ signals were seen in a cluster of cells ventrally adjacent to the neural groove at the midbrain (Figure 3a, white arrowheads) and hindbrain levels (Figure 3b,c, white arrowheads), which is consistent with the P0-Cre/R26-lacZ whole mount staining for the reporter activity (Figure 2a). Cre immunosignals in clustered cells in the midline and ventrally adjacent to floor plate colocalized with SOX9 (arrowheads, Figure 3a,b) and FOXA2 (arrowheads, Figure 3c). However, Cre immunosignals were not detected in SOX9+ cells in the neural plate border or mesoderm layer (arrows, Figure 3a,b) nor in the FOXA2+ cells seen in the floor plate (arrows, Figure 3c). Nonspecific immunosignals from the secondary antibody were seen in the foregut diverticulum (dashed outlines, Figure 3b,c), which were confirmed by staining while omitting primary antibody (dashed outlines, Figure 3d).

At the 7–8 somite stage (n = 3), another NC marker p75+ or and SOX9+ cells were seen at all brain levels examined, that is, forebrain, midbrain (Figure 4a), and hindbrain (Figure 4b). No Cre immunosignals were seen in the midbrain (Figure 4a). In the hindbrain region Cre immunosignals were detected in a small population of p75+SOX9+ cells in the mesodermal layer (arrows, Figure 4b), and in the menenchyme of prospective branchial arch 1 (white arrowheads, Figure 4b). Cre immunoreactivity was also apparent in FOXA2+ cells in the notochord region, that is, clustered cells ventrally adjacent to the floor plate (white arrowheads, Figure 4d) but not in the FOXA2+ cells seen in the floor plate (arrows, Figure 4d). Again, nonspecific signals were seen in the foregut diverticulum when using secondary antibody for Cre immunoreactions, which was confirmed by staining while omitting primary antibody (white dashed outlines, Figure 4b,d,e).

In 14-somite embryos (n = 3), Cre+ cells were abundantly distributed in the forebrain region and the immunosignals colocalized with SOX9 and p75 (Figure 5a, arrowheads). Additionally, Cre signals were seen in some of the SOX9+ p75+ cells in the anterior hindbrain region (Figure 5a, arrows). At the posterior hindbrain level (Figure 5b), an increasing intensity gradient of Cre immunoreactivity was observed from the dorsal to ventral region, that is, no immunosignals in the dorsal most part, faint immunosignals in the trigeminal NC region (Figure 5b, tr), obvious immunosignals in the branchial arch 1 region (Figure 5b, ba1), and intense immunosignals in the optic eminence (Figure 5b, oe). Similarly, the proportions of Cre+ SOX9+ p75+ cells relative to total SOX9+ or/and p75+ cells in different regions were different. In the dorsal most part, some SOX9+ cells in the neural fold region, presumably pre-migratory NC cells, were observed and negative for Cre immunoreactivity (open arrowheads, Figure 5c). In the trigeminal NC region, a subpopulation of SOX9+ p75+ cells was Cre+ (white arrows, Figure 5c). In the primordium of branchial arch 1, Cre immunosignals were apparent in a significant population of SOX9+ p75+ cells (arrows, Figure 5d). In the optic eminence most, if not all, SOX9+ p75+ cells were brightly labeled with Cre (white arrows, Figure 5e). Importantly, Cre+ cells were not seen in the Sox9 and p75+ cells in any tissue regions. In the negative control slides (Cre+ embryo sections with primary antibody staining and Cre+ embryo sections without primary antibody), no immunosignals were seen in the 14-somite embryo tissue (Supporting Information Figure 2).

Compared to the 14-somite stage, 19–somite embryos (n = 4) had a similar distribution pattern of Cre immunosignals, that is, an increasing intensity gradient from the dorsal to ventral regions of the tissue (Figure 6a,b). Additionally, increasing proportions of Cre+ SOX9+ cells relative to the total number of SOX9+ cells were observed from dorsal to ventral regions (Figure 6c–e). However, the proportions of Cre+ SOX9+ cells relative to total SOX9+ cells in all regions were lower than those at the 14-somite stage. At the anterior hindbrain level, Cre+ immunosignals were seen in some of the SOX9+ cells in the hindbrain and intense signals were distributed in the majority of SOX9+ cells in the forebrain (Figure 6a, arrows). At the posterior hindbrain level, Cre+ cells were seen in the trigeminal NC (Figure 6b,c, tr), branchial arch 1 (Figure 6b,d, ba1) and optic eminence (Figure 6b,e, oe). The intensity of Cre immunosignals in the optic eminence was diverse in the SOX9+ cells, some of which were negative for Cre (arrowheads, Figure 6e), which is in contrast to almost all of SOX9+ cells that showed presence of Cre immunosignals in the 14-somite embryos (Figure 5e).

### 3.4 | P0-Cre labeled cells are distributed in cranial NC and derivatives in early embryos and newborn mice

To confirm the specific labeling of P0-Cre/R26-lacZ and Cre immunosignals, we used another Cre reporter line (R26-RFP) and found that the distribution pattern of P0-Cre/R26-RFP signals (11s, Figure 7) was the same as P0-Cre/R26-lacZ (Figure 2d, d′), that is, signals were found in the hindbrain region, branchial arches, in the neural groove area and lateral tissues in the trunk region (Figure 7a). Similar to lacZ expression in P0-Cre/R26-lacZ embryos, RFP signals were not apparent in the midbrain region (mb). On tissue sections at the posterior hindbrain level, Cre immunosignals and RFP signals largely colocalized (Figure 7b). Again, an increasing gradient of Cre and RFP signal intensity and
proportions of labeled cells was observed from the dorsal to ventral region, that is, no signals in the dorsal-most part, faintly labeled cells were sparsely found in the trigeminal NC region (Figure 7b, tn), obviously and frequent labeled cells (Cre\textsuperscript{+}RFP\textsuperscript{+}) in the branchial arch 1 region (Figure 7b, ba1), and intensely and mostly labeled in the optic eminence (Figure 7b, oe). Auto-fluorescent blood cells and noncellular

**FIGURE 3** Single-plane laser scanning confocal photomicrographs of transverse sections of a 4-somite P0-Cre embryo. Sections were immunostained using an antibody against Cre (green) and were double labeled with SOX9 (purple, a,b) or FOXA2 (purple, c). Arrowheads point to Cre immunoreactive cells co-labeled with SOX9 (a,b) or FOXA2 (c) in the notochord region. Arrows point to single labeled SOX9\textsuperscript{+} cells in the NC cell region (a,b) and FOXA2\textsuperscript{+} cells in the floor plate of neural tube (c). White dashed lines outline the foregut diverticulum (b–d). In the negative control slide (d), primary antibodies were omitted. Scale bar: 50 μm for all images.
**FIGURE 4** Immunoreactivity of Cre (green, a–d), SOX9 (purple, a–c) or FOXA2 (purple, d), and p75 (red, a–d) in transverse sections of a 7-somite P0-Cre embryo at the midbrain (a) and hindbrain-forebrain (b–e) levels. Arrows in (c) point to some of the triple labeled Cre<sup>+</sup>SOX9<sup>+</sup>p75<sup>+</sup> cells in the NC cell region. Arrowheads in (d) point to Cre<sup>+</sup> cells colabeled with FOXA2 in the notochord region and arrows to FOXA2<sup>+</sup> cells in the floor plate of neural tube (d). White dashed lines outline the foregut diverticulum (b, d, and e). In the negative control slide (e), primary antibodies were omitted and only secondary antibodies were applied. Scale bar: 50 μm for all images (single-plane laser scanning confocal)
FIGURE 5  Photomicrographs of transverse sections of a P0-Cre embryo at the 14-somite stage. Sections were immunostained for Cre (green), SOX9 (purple), and p75 (red) at the anterior hindbrain-forebrain level (a) and posterior hindbrain-forebrain level (b). (c–d) are higher magnification images of trigeminal NC (c, tn), branchial arch 1 (d, ba1), and optic eminence (d, oe). Arrows (a) point to Cre⁺ cells that were colabeled with SOX9 and p75 immunosignals in the anterior hindbrain regions, and arrowheads (a) point to triple labeled Cre⁺SOX9⁺p75⁺ cells in the forebrain regions. Open arrowheads (c) point to SOX9⁺ cells in the neural fold region, presumably premigratory NC cells. Arrows (c–e) point to Cre⁺ cells colabeled with SOX9⁺ and p75⁺, presumably migratory NC, in trigeminal NC regions (tn, c), first branchial arch 1 (ba1, d) and optic eminence (oe, e). Scale bars: 50 μm for all images (single-plane laser scanning confocal)
FIGURE 6  Single-plane laser scanning confocal photomicrographs in transverse sections of a 19-somite P0-Cre embryo. Sections were immunostained using an antibody against Cre (green) and were double labeled with SOX9 (purple, a–e). (c–d) are higher magnification images of trigeminal NC (c, tn), branchial arch 1 (d, ba1), and optic eminence (d, oe). Arrows (a) point to Cre$^+$ cells that were co-labeled with SOX9 in the forebrain. Arrows (c–e) point to Cre$^+$ cells co-labeled with SOX9 in trigeminal NC regions (tn, c), branchial arch 1 (ba1, d), and optic eminence (oe, e). Arrowheads in e point to SOX9$^+$ cells without Cre immunosignals. Scale bars: 50 μm, also applies to other images in the same panel.
FIGURE 7  Consistent distribution of Cre and RFP signals in P0-Cre/R26-RFP embryos with that in P0-Cre and P0-Cre/R26-lacZ embryos.
(a): Bright-field (top panel) and fluorescent (bottom panel) images of whole P0-Cre/R26-RFP embryos at 10–11 somite stages. The midbrain (mb) region was devoid of RFP signals. Arrows point to the hindbrain region labeled with RFP. (b): Single-plane laser scanning confocal photomicrographs of a transverse section of an 11-somite P0-Cre(1)/R26-RFP(1) embryo at the hindbrain level. An increasing intensity gradient of Cre and RFP signals from the dorsal to ventral region, that is, faint and sparse in the trigeminal NC (tn), clear and frequent in the branchial arch 1 (ba1), bright and almost all in the optic eminence (oe). Arrowheads point to the autofluorescent blood cells. Arrows point to some noncellular RFP\(^+\) fragments. (c): Confocal images of a transverse section of a 10-somite Cre\(^-\) embryo to illustrate the autofluorescent blood cells and non-cellular RFP\(^+\) fragments (arrows). Scale bars: 200 \(\mu\)m in (a); 50 \(\mu\)m in (b and c)
RFP+ fragments were also seen (Figure 7b, arrowheads) and confirmed in the Cre− tissue sections (Figure 7c, arrowheads).

To verify the specificity of P0-Cre in labeling NC cells and derivatives, we examined whether P0-Cre faithfully labels the nasal and frontal bones in the skull that are well-characterized cranial NC derivatives. Consistent with a previous report using Wnt1-Cre mice (Jiang et al., 2002), P0-Cre successfully labeled both nasal and frontal bones along with dura mater (Figure 8a–c). Importantly, P0-Cre did not label the parietal bones that are known as paraxial mesoderm derivatives (Figure 8c,d) (Chai and Maxson, 2006; Noden and Trainor, 2005). The fluorescent signals in the parietal bone region of whole mount tissue (Figure 8a) were confirmed on sections to be from the underlying meninges (arrows, Figure 8d), which are NC derivatives (Jiang et al., 2002).

3.5 | Different distribution pattern of Wnt1-Cre labeling from P0-Cre in cranial regions

In P0-Cre/R26-lacZ embryos (Figure 2), reporter labeling was not apparent in the midbrain region. To further examine our observation, R26-lacZ was crossed with Wnt1-Cre, which is well known for labeling midbrain.

At the 6-somite stage (Figure 9a), the signals in Wnt1-Cre/R26-lacZ mouse embryos were intense in the midbrain and along the midline of the neural groove (Figure 9a). In the 8-somite embryo (Figure 9b), β-gal positive signals were extensively distributed in the midbrain (Figure 8b). However, signals in the hindbrain were absent at the 6-somite stage (Figure 9b, arrows) and were sparse at the 8-somite stage (Figure 9b, arrows). At E10.5, the signals extended to the forebrain (fb), midbrain (mb), hindbrain (hb) and trunk regions of NC and NC derivatives in Wnt1-Cre/R26-lacZ embryos (Figure 9c), for example, frontonasal and eye regions, branchial arches, heart, and dorsal root ganglia. In contrast to the extensive labeling of Wnt1-Cre in the midbrain, P0-Cre labeling in midbrain was not obvious (Figure 9c), which is consistent with our earlier observations (Figure 2).

Next, we further analyzed the expression patterns of Wnt1-Cre transgene in cranial NC using the antibody for Cre recombinase. In the 4-somite embryos (Figure 10a), Cre immunosignals were found in SOX9+ pre- and post-migratory NC cells and in the neural epithelium at the midbrain level (Figure 10a). In contrast, SOX9+ migrating NC cells were negative for Cre in the hindbrain region (arrows, Figure 10b). At the 7-somite stage (Figure 10c), Cre immunosignals were extensively distributed in migrating NC cells in the midbrain and forebrain regions, in addition to labeling of part of the neuroepithelium. Again, in hindbrain regions, Cre immunosignals were observed only in the neural fold and were not apparent in SOX9+ migrating NC cells (arrows, Figure 10d). This is an interesting contrast to the abundant Cre immunosignals found in the hindbrain region of P0-Cre embryos at the 7-somite stage (arrows, Figure 4b,c). At the presumptive notochord regions, no Cre immunosignals were seen in Wnt1-Cre embryos (arrowheads, Figure 10b,d). In the 16-somite embryos, Cre immunosignals were found in the midbrain and forebrain but were not apparent in the hindbrain (Figure 10e), which is in contrast to the distribution of Cre immunosignals in the 16–19 somite P0-Cre embryos (Figures 5 and 6).
In the trigeminal NC regions (tn), first branchial arch 1 (ba1) and optic eminence region (oe), Cre immunosignals were sparsely seen in migratory NC cells (Figure 10f).

Also, R26-RFP Cre reporter was used to confirm the specific labeling of Wnt1-Cre driven R26-lacZ and Cre immunoreactivity. The distribution pattern of signals in Wnt1-Cre/R26-RFP embryos (Figure 11a, 5–7s) was identical to that of Wnt1-Cre/R26-lacZ embryos (Figure 9a,b, 6–8s). The signals in Wnt1-Cre/R26-RFP mouse embryos were intense in the midbrain (Figure 11a, arrowheads), and signals in the hindbrain were not obvious at the 6- and 7-somite stages (Figure 11a, arrows).

On tissue sections of a 6-somite Wnt1-Cre/RFP embryo, the distribution of RFP+ cells was coincident with that of Cre+ cells (Figure 11b,c), such as abundance within migrating NC cells in the midbrain and in the forebrain neuroepithelium (Figure 11b). The labeling was sparse among migrating NC cells in the hindbrain region (arrows, Figure 11c).

Taken together (Table 1 and Supporting Information Figure 3), both P0-Cre and Wnt1-Cre label cranial NC cells, peaked at 7-somite stage in Wnt1-Cre and 14-somite in P0-Cre. An overlap (in forebrain) but distinct (Wnt1-Cre in midbrain and P0-Cre in hindbrain) distribution patterns of the Cre activities were observed. In addition to NC, both P0-Cre and Wnt1-Cre labeled other cells, for example, notochord was labeled by P0-Cre and neuroepithelium by Wnt1-Cre.

FIGURE 9 Whole mount X-Gal visualization of lacZ gene product β-galactosidase in Wnt1-Cre/R26-lacZ mouse embryos at different stages (E8-10.5). β-gal positive signals (blue) were extensive in the midbrain (mb) at the 6s (a) and 8s (b) stages, but were sparse in the hindbrain. At later stages, signals extended to the forebrain (fb), midbrain (mb), hindbrain (hb) and trunk regions of NC and NC derivatives in Wnt1-Cre/R26-lacZ tissue at E10.5 (c). In E9 P0-Cre/R26-lacZ mouse embryos (c), β-gal positive signals (blue) were found in the forebrain (fb), hindbrain (hb) and trunk regions of NC and NC derivatives, but were not apparent in the midbrain (mb). Scar bars: 200 μm in (a and b); 1 mm in (c).

4 | DISCUSSION

The P0-Cre transgenic mouse line has been widely used for NC cell fate mapping (Feltri et al., 1999b; Ono et al., 2015; Sommer and Suter, 1998) and genetic modifications of NC-derived cells (Hu, Strobl-Mazzulla, & Bronner, 2014; Liu and Xiao, 2011). However, questions remain about how early, how specifically, and to what extent P0-Cre transgene labels NC cells in early embryos. Moreover, concerns about the specificity of P0-Cre and other Cre models in labeling NC lineage have been raised (Trainor, 2005a,b). Therefore, a detailed analysis of P0-Cre transgene activity is essential for the use and data interpretation of this model. In the present study, we found that the CAAG-CAT-Z (CAAG-lacZ) Cre reporter mice used in the original report (Yamauchi et al., 1999) did not appropriately reflect Cre activity in early stage embryos. We then investigated Cre activity using a R26-lacZ reporter and Cre immunoreactivity to find P0-
FIGURE 10  Photomicrographs from sections of E8.5 Wnt1-Cre embryos at different somite stages immunostained for the Cre and NC cell marker SOX9. At the 4–7 somite stages (a–d), Cre immunosignals (green) were bright in the midbrain (a, c) and derived NC cells and colocalized with SOX9 (purple). In contrast, in the hindbrain-forebrain levels (b, d) Cre immunosignals were seen in SOX9− premigratory NC cells but were absent in migrating NC cells. In 16-somite embryos (e, f), Cre immunosignals were seen in migrating NC cells in the midbrain and forebrain regions (e). In the trigeminal NC regions (tn), first branchial arch 1 (ba1), and optic eminence region (oe), Cre immunosignals were sparse in migratory NC cells. Scale bars: 50 μm (single-plane laser scanning confocal)
Cre expression as early as the 4-somite stage in notochord labeled with FOXA2 and in the migrating NC cells at the 7-somite stage using the commonly used NC cell markers SOX9 (Mori-Akiyama, Akiyama, Rowitch, & de Crombrugghe, 2003; Sahar, Longaker, & Quarto, 2005; Spokony et al., 2002) and p75 (Young, 2000; Young, Ciampoli, Hsuan, & Canty, 1999). Cre immunosignals were especially extensive in the forebrain and hindbrain but not apparent in the midbrain. Importantly, Cre immunosignals were not seen outside of the NC cell population and notochord. Given the fact that notochord derivatives are well-known to be restricted to the intervertebral disc (McCann, Tamplin, Rossant, & Seguin, 2012; Yamauchi et al., 1999), our data demonstrated that P0-Cre can serve as a valuable tool for studies on NC cell lineage, especially for forebrain and hindbrain NC derivatives. In contrast, Wnt1-Cre extensively labeled pre- and post-migratory NC cells at the midbrain level, but only sparsely in the trigeminal NC region at the hindbrain level. Additionally, Wnt1-Cre labeled a large population of non-NC cells in the neural plate, which is consistent with previous reports (Echelard et al., 1994; Rowitch et al., 1998).

4.1 P0-Cre specifically labels a significant population of, if not all, migrating NC cells and notochord

The specificity of Cre expression in the NC cell population was examined with two commonly used NC cell markers, p75 (Menendez, Yatskievych, Antin, & Dalton, 2011) to label migrating NC and SOX9 (Nakanishi, Chan, & Ito, 2007; Pomp, Brokhman, Ben-Dor, Reubinoff, & Goldstein, 2005) to label both pre- and post-migratory NC cells. Our data support the idea that P0-Cre specifically labels migrating NC cells and notochord based on the following observations: (1) Cre immunosignals were only detected in p75-“SOX9” double labeled cells, and were more intense in the NC cells that were close to their destinations; (2) importantly, Cre immunosignals were not seen outside of the p75-“SOX9” double labeled cells; (3) Cre immunosignals were not seen.
in SOX9\(^+\) cells in the neural fold region, which presumably premigratory NC cells, and Cre immunosignals were rarely detectable in the immediately delaminated NC cell region; and (4) Cre immunosignals only colocalized with FOXA2 in the notochord region, and were absent in FOXA2\(^+\) cells in the floor plate. These data demonstrate that P0-Cre is specific in labeling migrating NC cells and notochord. An increasing gradient of intensity of Cre immunosignals and RFP signals was observed from the dorsal to ventral regions suggesting that the expression of P0-Cre transgene in the NC cells is obtained during migration. Our data are valuable in demonstrating the specificity of P0-Cre in labeling NC cells in early embryos. The specificity of P0-Cre in labeling cranial NC cells was further confirmed by the distribution of P0-Cre labeled cells in the well-known NC derivatives in the skull, for example, frontal bones and sutures, in contrast to the absence of labeled cells in the parietal bone that is known to be non-NC derived.

The notochord is a rod-shaped structure in the midline that is ventrally adjacent to the neural tube (Jurand, 1974). Studies using a Cre mouse line driven by the notochord-specific homeobox gene Noto that traced the cell fate of notochord demonstrated that notochord serves as a source of embryonic precursors of cells within the nucleus pulposus of the mature intervertebral disc (McCann et al., 2012). In our study, P0-Cre transgene expression was detected in the notochord region in the 4–7 somite stage embryos, which is consistent with a previous report demonstrating that P0-Cre transgene labels notochord in addition to NC cells and derivatives (Yamauchi et al., 1999). With exception to the labeling of notochord, P0-Cre labels migrating NC cells specifically. Indeed, our data using P0-Cre to mark cranial bones, that is, nasal and frontal but not parietal bone, further demonstrated the specificity of P0-Cre in labeling cranial NC derivatives and that P0-Cre mice can be considered as a useful tool to trace NC lineages, at least cranial NC-derived bones (Komatsu et al., 2013), during mouse embryonic development.

However, P0-Cre did not label all of the NC cells in all of the tissue regions. The proportions of Cre\(^+\) cells relative to the total number of p75\(^-\)SOX9\(^-\) cells were different in different tissue regions. NC cells migrate ventrally to their target tissue/organs and follow certain paths after delamination. Thus, the increasing intensity gradient of Cre immunosignals and RFP signals and increasing cell proportions of Cre\(^+\) and RFP\(^+\) cells from dorsal to ventral regions suggest that P0-Cre transgene expression emerged during NC migration. Although Cre immunosignals were not detected in all the p75\(^-\)SOX9\(^-\) cells, we cannot exclude the possibility that P0-Cre does indeed label all NC cells. It would be difficult to evaluate the exact proportion of P0-Cre transgene in labeling migrating NC cells because the expression is transient and because P0-Cre can be expressed in a NC cell for a short time frame at any time point. Our observation that detection of Cre immunosignals in all the p75\(^-\)SOX9\(^-\) cells in the optic eminence and the majority of mesenchymal cells immediately under the epithelium of prospective branchial arch 1 at the 14 somite stage supports the idea that P0-Cre transgene is expressed in a significant population of, if not all, migrating NC cells in the forebrain and hindbrain regions.

Several Cre reporter lines have been generated and are widely used for cell fate mapping (Ikeya et al., 1997; Kawamoto et al., 2000; McMahon et al., 1992; Novak et al., 2000; Weissman and Pan, 2015; Weissman et al., 2011). Among these reporter lines, ROSA26 and CAAG-lacZ are two commonly used promoters for driving reporter gene expression (Araki, Araki, Miyazaki, & Vassalli, 1995; Kawamoto et al., 2000). In the present study, the efficiency and reliability of two Cre reporters, R26-lacZ and CAAG-lacZ, were compared using Mef2c-Cre, in which Cre is expressed as early as E5.5 in epiblast (Tallquist and Soriano, 2000). We found that R26-lacZ reporter expression emerges earlier than CAAG-lacZ reporter expression and that DNA recombination in R26-lacZ was reliable and consistent with Cre expression. One interesting observation is that although all E9.5 Mef2c-Cre/CAAG-lacZ embryos showed evidence of DNA recombination, the lacZ signals were weak, and were restricted to the heart and somites. It is possible that the promoter activity to drive lacZ expression in the CAAG-lacZ cassette was silenced or weaker than that of R26-lacZ in early stage embryos. Together, the data suggest that less efficient and inconsistent DNA recombination, and variable CAAG promoter activity might be the major reasons for this discrepancy.

### 4.2 P0-Cre labels cranial NC cells in a cranial region-specific manner

It has been reported that NC cells are generated by the 7-somite stage at all levels of the forebrain, midbrain, and hindbrain (Chan and Tam, 1988; Nichols, 1981; Serbedzija, Bronner-Fraser, & Fraser, 1992). By detecting both R26-lacZ reporter expression and Cre immunoreactivity we found that P0-Cre transgene expression emerges at the 7-somite stage in hindbrain and forebrain migratory NC cells. In the 14-somite embryos, we found strong Cre immunosignals in a considerable subpopulation of migrating NC cells labeled with SOX9 and p75 in the forebrain and hindbrain NC regions.

However, P0-Cre expression was rarely detected in the midbrain region, which is in contrast to extensive distribution of Cre immunosignals and RFP signals in the midbrain NC of Wnt1-Cre (Huang, Liu, Huang, Zhao, & Cheng, 2010; Lewis et al., 2013). At E8.5, when cranial NC cells emerge, Wnt1-Cre labels NC cells extensively in midbrain but not in other NC derivatives. Although Cre immunosignals were detected in the midbrain NC region, the intensity of Cre immunosignals was rarely detected compared with other regions, which is consistent with previous reports that have demonstrated that midbrain NC cells are late-born NC cells in comparison to cranial NC cells. However, it is still possible that Cre immunosignals were not detected in the midbrain NC region due to the use of Cre reporter lines such as R26-lacZ and CAAG-lacZ, which are not specific to the midbrain NC region. Therefore, to detect Cre immunosignals in the midbrain NC region, other Cre reporter lines that are specific to the midbrain NC region should be used.

### Table 1

Comparison between P0-Cre and Wnt1-Cre in labeling cranial neural crest (NC), neural epithelium (NE), and notochord to various extent

<table>
<thead>
<tr>
<th>Cranial regions</th>
<th>P0-Cre</th>
<th>Wnt1-Cre</th>
</tr>
</thead>
<tbody>
<tr>
<td>Forebrain</td>
<td>NC</td>
<td>++</td>
</tr>
<tr>
<td></td>
<td>NE</td>
<td>+</td>
</tr>
<tr>
<td>Midbrain</td>
<td>NC</td>
<td>+</td>
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<tr>
<td></td>
<td>NE</td>
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<tr>
<td>Hindbrain</td>
<td>NC</td>
<td>++</td>
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<tr>
<td></td>
<td>NE</td>
<td>+</td>
</tr>
<tr>
<td>Notochord</td>
<td>+</td>
<td>+</td>
</tr>
</tbody>
</table>

++ extensively; + clearly; ± rarely; − negatively.
many fewer cells were seen in hindbrain (Danielian, Muccino, Rowitch, Michael, & McMahon, 1998; Echelard et al., 1994). Wnt1 and Wnt1-Cre expression is not restricted to the NC domain but also labels dorsal neural stem cells that contribute to both the central nervous system and to neural progenitors (Barriga, Trainor, Bronner, & Mayor, 2015; McMahon et al., 1992). In the present study, we crossed the same R26-lacZ reporter mouse with P0-Cre and Wnt1-Cre and confirmed the labeling pattern (Figure 9). The difference between P0-Cre and Wnt1-Cre in labeling NC cells was profound in the midbrain and hindbrain, which may be why P0-Cre and Wnt1-Cre label different populations of cells in the orofacial organs, for example, tooth buds (Wang et al., 2011) and taste buds (Boggs et al., 2016; Liu et al., 2012).

It has been reported that forebrain crest cells appear by the 8–14 somite stages and migrate dorsally over the presumptive eye where they meet the ventrally migrating midbrain crest cells. Although we did not see much Cre reporter activity or Cre immunoreactivity in the midbrain, Cre immunoreactivity was detected in all the SOX9”p75” cells in the optic eminence, which suggests that forebrain is the source of NC cells that contribute to optic eminence development.

5 | CONCLUSIONS

The P0-Cre transgenic mouse model was first generated to label NC derivatives (Yamuchi et al., 1999) and has been widely used for NC lineage tracing (Boggs et al., 2016; Kawakami et al., 2011; Liu et al., 2012; Ogawa et al., 2015; Wang et al., 2011) and genetic modification (Feltri et al., 1999a; Komatsu et al., 2013; Nomura-Kitabayashi et al., 2009; Ogawa et al., 2015). Differences have been found between P0-Cre and Wnt1-Cre in labeling NC lineage. Here, we re-visited the P0-Cre model at early embryonic stages and demonstrated that P0-Cre transgene specifically labels migrating cranial NC cells in the forebrain and hindbrain as early as the 7-somite stage and reaches its peak expression at the 14-somite stage. Additionally, P0-Cre labels notochord at the 4–7 somite stages in early embryos. Moreover, the distribution pattern of P0-Cre was different from Wnt1-Cre in labeling NC cells, especially in the midbrain and hindbrain regions. Our data indicate that the P0-Cre mouse line is a valuable model for studies of NC and notochord lineage, and that careful attention needs to be paid in choosing promoters to drive Cre and its reporters for studies on NC lineage in early embryos.

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AUTHOR CONTRIBUTION

Experimental design: GC, MI, JY, SC, YK, YM and HXL. Experiment conduction and data analysis: GC, MI, JY, SK, TF, GS, MKR, CS, SC, YK, YM and HXL. Data interpretation: GC, YK, YM and HXL. Writing manuscript: GC, YM and HXL. Approving final version of manuscript: GC, MI, JY, SK, TF, GS, MKR, CS, SC, YK, YM and HXL. HXL and YM take responsibility for the integrity of the data analysis.

REFERENCE


**SUPPORTING INFORMATION**

Additional Supporting Information may be found in the online version of this article.

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