



## TGF- $\beta$ mediated FGF10 signaling in cranial neural crest cells controls development of myogenic progenitor cells through tissue–tissue interactions during tongue morphogenesis

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### ABSTRACT

Skeletal muscles are formed from two cell lineages, myogenic and fibroblastic. Mesoderm-derived myogenic progenitors form muscle cells whereas fibroblastic cells give rise to the supportive connective tissue of skeletal muscles, such as the tendons and perimysium. It remains unknown how myogenic and fibroblastic cell–cell interactions affect cell fate determination and the organization of skeletal muscle. In the present study, we investigated the functional significance of cell–cell interactions in regulating skeletal muscle development. Our study shows that cranial neural crest (CNC) cells give rise to the fibroblastic cells of the tongue skeletal muscle in mice. Loss of *Tgfb2* in CNC cells (*Wnt1-Cre;Tgfb2<sup>fllox/fllox</sup>*) results in microglossia with reduced *Scleraxis* and *Fgf10* expression as well as decreased myogenic cell proliferation, reduced cell number and disorganized tongue muscles. Furthermore, TGF- $\beta$ 2 beads induced the expression of *Scleraxis* in tongue explant cultures. The addition of FGF10 rescued the muscle cell number in *Wnt1-Cre;Tgfb2<sup>fllox/fllox</sup>* mice. Thus, TGF- $\beta$  induced FGF10 signaling has a critical function in regulating tissue–tissue interaction during tongue skeletal muscle development.

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### Introduction

During skeletal muscle development, two mesenchymal cell lineages are required. Myogenic cells give rise to mature muscle cells, and fibroblastic cells give rise to the surrounding connective tissue such as perimysium and tendon. In the craniofacial region, skeletal muscle cells are derived from unsegmented paraxial mesoderm, and the surrounding connective tissue comes from cranial neural crest (CNC)-derived cells (Noden, 1988). In the trunk region, on the other hand, skeletal muscle cells are derived from somite-derived cells and segmented paraxial mesoderm cells, and the surrounding connective tissue comes from lateral mesoderm cells (Chevallier et al., 1977). The cell origins of the tongue are essentially a hybrid. Myogenic cells in the tongue are derived from occipital somites and connective tissue is derived from CNC cells (Noden, 1983; Noden and Francis-West, 2006). To date, these conclusions have been based on chick/quail recombination experiments (Noden, 1983), and

the extent to which mammalian development conserves these mechanisms has remained unknown.

The Cre-LoxP system provides a tool for mapping cell fate and deleting specific genes with spatial and temporal specificity. We previously reported CNC cell fate analysis using *Wnt1-Cre;R26R* mice (Chai et al., 2000). During tongue skeletal muscle development, myogenic cells and the surrounding CNC cells can be distinguished using the Cre-LoxP system. In the present study, we investigate the relative contribution of CNC and myogenic cell lineages in the developing tongue primordium using *Wnt1-Cre* to target CNC cells and *Myf5-Cre* to target myogenic cells (Chai et al., 2000; Tallquist et al., 2000).

Chick/quail recombination experiments have previously demonstrated that CNC cells surround the myogenic cell lineage at an early stage, but do not penetrate into the myogenic core (Bogusch, 1986; Noden, 1986; Noden and Francis-West, 2006). Early in development, CNC cells secrete BMP and Wnt inhibitors, which induce myogenic differentiation in the branchial arch (Tzahor et al., 2003). Borue and Noden proposed a passive displacement model based on the interface between CNC and myogenic cells in later developmental stages (Borue and Noden, 2004). Finally, CNC cells give rise to tissue surrounding skeletal muscles such as perimysium, epimysium, endomysium, and tendon (Couly et al., 1992; Evans and Noden,

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2006), however, the molecular mechanism involved in regulating their development is still unknown.

Transforming Growth Factor- $\beta$  (TGF- $\beta$ ) is composed of three isoforms in mammals, TGF- $\beta$ 1, - $\beta$ 2, and - $\beta$ 3. TGF- $\beta$  ligands bind to a TGF- $\beta$  type II receptor (TGF $\beta$ RII) and then Type II and Type I receptors form a hetero-tetramer. Subsequently, Smad2/3 are phosphorylated by the receptor complex and bind to Smad4, the common Smad. This Smad complex then translocates into the nucleus to regulate downstream target genes (Massague, 1998; Wu and Hill, 2009). TGF- $\beta$  signaling is involved in multiple biological functions, such as cell proliferation, differentiation, extracellular matrix synthesis, and cell migration during embryonic development, wound healing, and carcinogenesis (Hosokawa et al., 2005; Massague, 1998; Massague and Gomis, 2006). Previous studies indicate that TGF- $\beta$ 1 and TGF $\beta$ RII are co-expressed in undifferentiated mesenchymal cells (Lawler et al., 1994). Furthermore, TGF- $\beta$ 1 is expressed in the surrounding tissue at late stages of skeletal muscle development (McLennan, 1993). The function of TGF- $\beta$  signaling during tongue muscle formation *in vivo* is still unknown. In the present study, we investigate the function of TGF- $\beta$  signaling in CNC cells during tongue muscle development.

Loss of *Tgfb2* (which encodes for TGF $\beta$ RII) in CNC cells results in microglossia and disorganized tongue muscles. Specifically, there is compromised FGF10 signaling in CNC-derived cells and retardation of myogenic cell proliferation activity. Our data suggests that TGF- $\beta$  induced FGF signaling regulates tissue–tissue interactions to control tongue muscle development.

## Materials and methods

### Generation of *Wnt1-Cre;Tgfb2<sup>fllox/fllox</sup>* mutant mice

*Wnt1-Cre* transgenic mice have been described previously (Chai et al., 2000). We crossed *Wnt1-Cre;Tgfb2<sup>fllox/+</sup>* with *Tgfb2<sup>fllox/fllox</sup>* mice to generate *Wnt1-Cre;Tgfb2<sup>fllox/fllox</sup>* mice, which were genotyped using PCR primers as previously described (Ito et al., 2003).

### Two-component genetic system for marking myogenic and CNC-derived cells

The R26R reporter allele has been described previously (Soriano, 1999). We mated *Wnt1-Cre* or *Myf5-Cre* mice with R26R mice to generate *Wnt1-Cre;R26R* or *Myf5-Cre;R26R* embryos in which CNC- or myogenic-derived cells could be detected, respectively. Detection of  $\beta$ -galactosidase activity in sections was carried out as previously described (Chai et al., 2000). *Wnt1-Cre;Tgfb2<sup>fllox/+</sup>* mice were crossed with *Tgfb2<sup>fllox/fllox</sup>;R26R<sup>fllox/fllox</sup>* mice to produce embryos with the genotype of *Wnt1-Cre;Tgfb2<sup>fllox/fllox</sup>;R26R<sup>fllox/+</sup>*.  $\beta$ -galactosidase analysis was carried out as previously described (Chai et al., 2000).

### Histological analysis

Tissues were fixed in 4% paraformaldehyde in phosphate-buffered saline, paraffin embedded, sectioned at 7  $\mu$ m and stained with hematoxylin/eosin. For immunostaining, tissue sections were incubated with anti-TGF- $\beta$  IIR (1:100, Santa Cruz biotechnology Inc, Santa Cruz, CA) and anti-Myogenin [MF 20, 1:100, DSHB (Developmental Studies Hybridoma Bank), Iowa City, IA] following standard procedures. For LacZ and BrdU/MHC double labeling, bromodeoxyuridine (BrdU; Sigma, St Louis, MO) was injected intraperitoneally into mice at E12.5 and E13.5 at a dose of 100  $\mu$ g BrdU per gram of body weight. Mice were sacrificed 1 h after injection and pups were harvested. Tissues were fixed in 0.05% glutaraldehyde and processed into cryosections. We first performed LacZ staining and then anti-BrdU or MHC (myosin heavy chain) (1:100, DSHB, Iowa City, IA) staining, following the manufacturer's directions (BrdU antibody: Zymed, South San Francisco, CA) or standard procedure (MHC immunostaining).

*FDG (Fluorescence di- $\beta$ -D-Galactopyranoside) staining for  $\beta$ -galactosidase and Fluorescence Activated Cell Sorting (FACS)*

At E12.5, we digested the tongue primordium with TrypLE Express (Invitrogen, Grand Island, NY) and collected the cells. FDG staining and FACS analysis were performed as reported previously (Zhao et al., 2006).

### *In situ hybridization*

Sectioned *in situ* hybridization was performed according to standard procedure (Wilkinson, 1998). Several negative controls (sense probe and no probe) were run in parallel with the experimental reaction. *Fgf10* RNA probe was generated as reported previously (Bellusci et al., 1997).

### Cell culture system

Timed-pregnant mice were sacrificed on postcoital day 12.5 and staged according to somite age. Tongue primordium was removed from the first branchial arch and cut into tissue blocks. These tissue blocks were seeded on 35 mm culture plates (BD biology, San Jose, CA) and cultured with 0.5 ml of growth medium (DMEM with 40% FBS) (Invitrogen, Carlsbad, CA) at 37 °C overnight. The next day, an additional 1.5 ml of growth medium was added (Oh et al., 2004). After 2 days, the tissue blocks were removed and the remaining primary cells were cultured under the growth medium for 2 more days. After that, the medium was switched to differentiation medium (DMEM with 5% horse serum) and 10 ng/ml FGF10 (R&D Systems Inc, Minneapolis, MN) was added (Harada et al., 2002), followed by cell culture for ten days. Eight fields (20 $\times$ ) from each genotype were used for quantification of muscle cell number (Doherty et al., 2005).

### *Organ culture of wild type and Wnt1-Cre;Tgfb2<sup>fllox/fllox</sup> mutant tongue explants and bead implantation experiments*

Timed-pregnant mice were sacrificed on postcoital day 12.5. Tongue explants were cultured in serum-less, chemically-defined BGJB medium according to standard methods (Chai et al., 1994). For delivery of TGF- $\beta$ 2, we used affi-gel blue beads (Bio-Rad, Hercules, CA), diameter 50–80  $\mu$ m. The beads were washed in phosphate-buffered saline (PBS) and then incubated for 1 h at room temperature in 10  $\mu$ g/ml TGF- $\beta$ 2 (R&D, Minneapolis, MN). Control beads were incubated in 0.1% BSA. TGF- $\beta$  or BSA-containing beads were placed in the tongue explant.

### Real-time quantitative RT-PCR

The mRNA level of *Fgf10* was analyzed by real-time quantitative RT-PCR (Bio-Rad iCycler system). Tongue primordium was dissected at E12.5 and total RNA was extracted. The mRNAs were reverse-transcribed into cDNAs by using SuperScript™ First-Strand (Invitrogen). The real-time PCR was performed using a SYBR super mix kit (Bio-Rad) as reported previously (Oka et al., 2007). *Fgf10* and *Tgfb2* primers were generated as reported previously (Oka et al., 2007; Sakaue et al., 2002).

### *In vivo transplantation of cultured cells*

We isolated myogenic and cranial neural crest cells by FACS as described above. CNC and myoblast cells were combined as depicted schematically in Figs. S3A–C. Cultured cells ( $1.0 \times 10^6$ ) were mixed with 3D scaffold collagen composition (BD, Bedford, MA), and the scaffold with cell mixture was then transplanted subcutaneously into the dorsal surface of 4-week-old female immunocompromised beige mice (NIH-bg-nu/*ν*-xid, Harlan Sprague Dawley, Indianapolis, IN, <http://www.harlan.com>).

## Results

### *The relationship of mesoderm- and CNC-derived cells and expression of TGF- $\beta$ type II receptor in the tongue primordium*

To investigate cell lineages in the tongue primordium, we examined myogenic or CNC-derived cells in *Myf5-Cre;R26R* and *Wnt1-Cre;R26R* mice, respectively. In *Myf5-Cre;R26R* mice, LacZ positive cells were detected in the center of the tongue primordium (Fig. 1A). In contrast, the LacZ expression pattern in *Wnt1-Cre;R26R* mice was the reverse of that of *Myf5-Cre;R26R* mice (Fig. 1B). This pattern suggests that CNC cells circumscribe somite-derived myogenic progenitor cells at E13.5 (Figs. 1A and B). Furthermore, the expression pattern of myosin heavy chain (MHC), a marker for myogenic differentiation, coincided with LacZ positive cells in *Myf5-Cre;R26R* mice and LacZ negative cells in *Wnt1-Cre;R26R* mice (Figs. 1A–C). We also found that tongue skeletal muscle cells in *Myf5-Cre;R26R* mice appeared to form bundle structure, whereas CNC-derived connective tissue cells in *Wnt1-Cre;R26R* mice populated the region between the presumptive *Myf5* positive bundles at E14.5 (Figs. 1D and E). Furthermore, we assessed the expression pattern of TGF- $\beta$  type II receptor during tongue development. We detected TGF- $\beta$  IIR expression in CNC-derived cells (Fig. 1G, red) and myogenic progenitors (Fig. 1G, green).

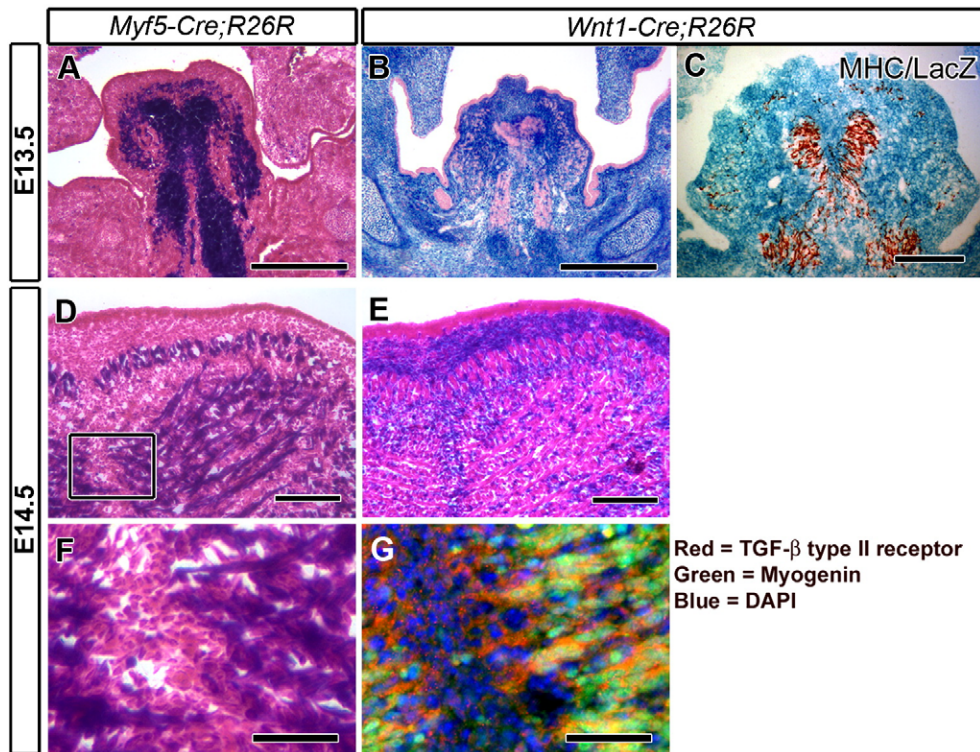
### *Loss of TGF- $\beta$ signaling in CNC cells produces microglossia and disorganization of muscle cell alignment*

The loss of TGF- $\beta$  signaling in CNC cells resulted in a severe defect in tongue development. *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>* mice exhibited microglossia when compared with *Wnt1-Cre;Tgfb2<sup>flox/+</sup>* (control) mice

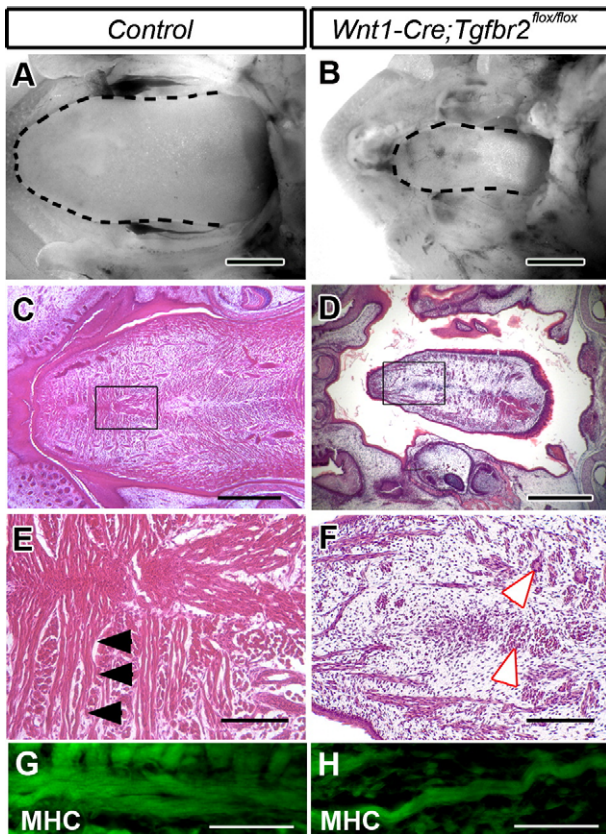
(Figs. 2A and B), which were indistinguishable from wild type mice (data not shown). We detected well-organized muscle cells in control mice by histological analysis (Figs. 2C and E). In contrast, muscle cells were disorganized and present in low density in *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>* mice (Figs. 2D and F). We detected that MHC expression in muscle cells of *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>* mice was similar per cell to that of control mice, although muscle cells were decreased in number and curved (Figs. 2G and H). We previously reported that the oral space of *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>* mice was smaller than that of control mice because of their small mandible and cleft palate (Ito et al., 2003; Oka et al., 2007). To test the hypothesis that these physiological differences affected the tongue development process, we performed tongue organ culture to remove any physical constraints on tongue growth. After isolation of the tongue primordium at E12.0, there was no structure inhibiting tongue growth via physiological force. Following seven days of culture, we found that the size of the tongue primordium from mutant mice ( $n=6$ ) was smaller than that from control mice ( $n=6$ ) (Figs. S1A and B). The maximal length and width of tongues from *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>* mice were reduced by  $21.1 \pm 2.5\%$  and  $31.7 \pm 1.7\%$ , respectively. Thus, we conclude that the tongue phenotype of *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>* mice was a primary defect and not a secondary defect due to constraints such as a small mandible. Because TGF- $\beta$  signaling is lost only in CNC cells, our data suggest that TGF- $\beta$  signaling regulates tissue–tissue interaction during skeletal muscle development.

### *Loss of TGF- $\beta$ signaling in CNC cells results in reduced myogenic cell proliferation*

During tongue skeletal muscle development, myogenic cells migrate from the occipital somite to the tongue primordium through



**Fig. 1.** The interface between cranial neural crest and myogenic cells in the tongue primordium. LacZ staining and immunostaining of frontal sections from *Myf5-Cre;R26R* (A, D, and F) and *Wnt1-Cre;R26R* (B, C, E, and G) mice. (A) LacZ positive cells (*Myf5-Cre;R26R*) form the core of the tongue primordium at E13.5. (B) LacZ positive cells (*Wnt1-Cre;R26R*) circumscribe the LacZ negative core at E13.5. (C) Myosin heavy chain (MHC) immunostaining and LacZ staining of E13.5 *Wnt1-Cre;R26R* mice. MHC positive cells are detectable in the center of the tongue primordium. LacZ positive cells (*Wnt1-Cre;R26R*) circumscribe the MHC positive cells. (D, F) LacZ positive cells (*Myf5-Cre;R26R*) form myofibril-like structures at E14.5. F is enlarged from the box in D. (E) LacZ positive cells are detectable between muscle cell-like structures at E14.5. (G) Immunofluorescence of TGF- $\beta$  type II receptor (red) and myogenin (green). *Myf5* negative cells, which are CNC-derived cells, are immunopositive for TGF- $\beta$  type II receptor (red). Myogenin positive cells (green) coincide with *Myf5* positive cells (see F), and some are immunopositive for TGF- $\beta$  type II receptor (yellow). Scar bars: 300  $\mu$ m in A, B, 100  $\mu$ m in C–E, and 50  $\mu$ m in F, G.



**Fig. 2.** *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>* mice exhibit microglossia and muscle cell disorganization. Macroscopic appearance (A and B), hematoxylin and eosin staining (C–F), and myosin heavy chain (MHC) immunohistochemistry (G and H) of tongues from *Wnt1-Cre;Tgfb2<sup>flox/+</sup>* (control) (A, C, E, and G) and *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>* (B, D, F, and H) newborn mice. (A and B) *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>* mice exhibit microglossia (dashed lines indicate the outline of tongue). (C–F) Hematoxylin and eosin staining of control mice shows well-organized myofibrils (C, E; black arrowheads), but muscle cells are disorganized and scattered in *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>* mice (D, F white arrowheads indicate the diminished muscle cells). E and F are enlarged from the boxes in C and D, respectively. (G and H) Muscle cells are positive for MHC expression (green) in both control and *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>* mice. Scale bars: 1 mm in A, B, 500  $\mu$ m in C, D, 100  $\mu$ m in E, F, and 50  $\mu$ m in G, H.

the hypoglossal cord (Huang et al., 1999). We found that myogenic precursor cells started to invade the tongue primordium around E12.0 (data not shown). To compare the cell population in *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>;R26R* and *Wnt1-Cre;Tgfb2<sup>flox/+</sup>;R26R* (control) mice, we performed Fluorescence Activated Cell Sorting (FACS) using FDG staining for  $\beta$ -galactosidase. Through the sorting process, we separated cranial neural crest derived cells (FDG-positive) and non-cranial neural crest cells (FDG-negative) (Fig. 3A). We found that the cell numbers of both FDG-positive and -negative cells in *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>* mice were indistinguishable from those of control mice. After FACS, we assessed *Tgfb2* gene expression in CNC cells and non-CNC cells in *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>;R26R* samples. Real-time PCR data demonstrated that *Tgfb2* gene expression was deleted only in CNC cells of *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>* mice (Fig. 3B). Furthermore, to assess myogenic cell migration in *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>* mice, we analyzed MHC expression in the tongue primordium at E12.5. We detected MHC positive cells throughout the anterior to posterior regions of the tongue primordium in *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>* and control mice (Figs. 3C and D). Thus, myogenic cell migration into the tongue primordium in *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>* mice was indistinguishable from that of control mice. In order to investigate the cellular mechanism responsible for microglossia, we examined cell proliferation activity in *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>* mice. To determine whether CNC or myogenic cell lineages exhibited reduced cell proliferation, we

performed anti-BrdU and LacZ double staining. In *Wnt1-Cre;Tgfb2<sup>flox/+</sup>;R26R* (control) mice, both CNC cells (Fig. 3E, open arrow) and myogenic cells (Figs. 3E and G, black arrows) were positive for cell proliferation activity. In contrast, proliferation activity was reduced in myogenic cells of *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>;R26R* mice (Figs. 3F and H, black arrows), although CNC cells maintained proliferation activity similar to that of control mice (Fig. 3F, open arrow). We also analyzed the number of BrdU positive and negative CNC and myogenic cells of the tongue primordium (Figs. 3I and J). This quantitative analysis confirmed that cell proliferation activity was significantly decreased in the myogenic cell lineage but not in the CNC cell lineage of *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>* mice.

#### *Skeletal muscle organization is compromised in Wnt1-Cre;Tgfb2<sup>flox/flox</sup> mice*

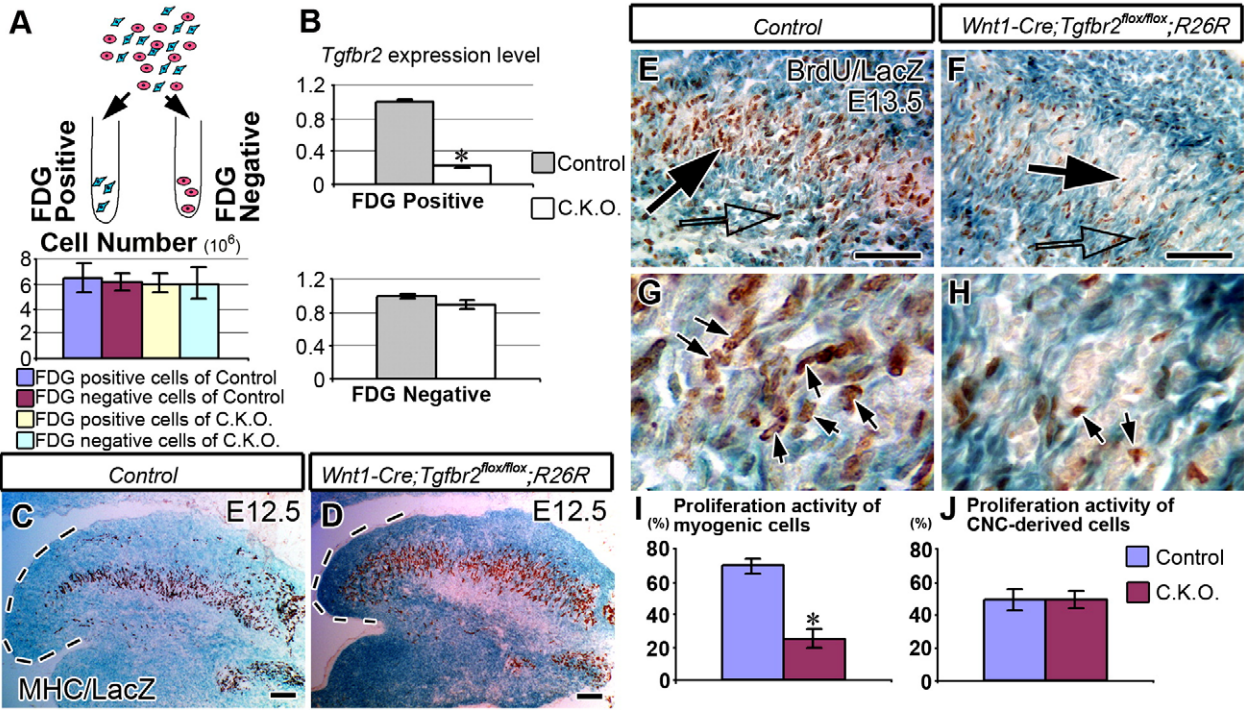
We assessed whether skeletal muscle organization was altered in *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>* mice. At E12.5, myogenic cells derived from occipital somites formed a myogenic core in the center of the tongue primordium of *Wnt1-Cre;Tgfb2<sup>flox/+</sup>;R26R* (control) mice (Fig. 4A, delineated by the dashed line). The myogenic core was also detectable in the tongue primordium of *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>;R26R* mice (Fig. 4B), implying that the early stage of myogenic formation in *Wnt1-Cre;Tgfb2<sup>flox/+</sup>* mice was unaffected. Division of the myogenic core had occurred in control mice at E13.5, based on the expression pattern of *p57*, a marker for myogenic cell differentiation (Zhang et al., 1999) (Fig. 4C), whereas division of myogenic core failed to occur in *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>* mice (Fig. 4D). Thus, we conclude that although myogenic differentiation at early stages was normal as indicated by MHC staining (See Fig. 3D), the division process was compromised in *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>* mice.

#### *Cell autonomous requirement for TGF- $\beta$ signaling during the differentiation of CNC-derived cells in the tongue primordium*

We hypothesized that TGF- $\beta$  signaling controls the fate of CNC-derived cells and, therefore, analyzed cell differentiation status in *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>* mice. *Type I collagen*, the main component of connective tissue, was expressed in the dense lamina propria and septum of the tongue in control mice (Figs. 5A and C), whereas this staining appeared diminished in *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>* mice (Figs. 5B and D). In contrast, expression of *type I collagen* in the mandible was indistinguishable between control and *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>* mice (Figs. 5A and B, open arrows). *Scleraxis (Scx)*, a bHLH transcription factor, is expressed in the mature tendons of limbs and trunk as well as their progenitors, and plays an important role in regulating tendon differentiation (Brent et al., 2003). *Scx* was expressed in the perimysium, epimysium, and septum that surround tongue muscle fibers of control mice (Fig. 5E, white arrowheads). In *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>* mice, *Scx* staining was diminished in the tendon of the tongue muscle (Fig. 5F). We hypothesized that TGF- $\beta$  signaling induces *Scx* expression during tongue development. Our study showed that TGF- $\beta$ 2 soaked beads induced *Scx* expression in the tongue primordium of control mice (Fig. 5G), but BSA soaked beads had no effect (Fig. 5H). In contrast, TGF- $\beta$ 2 soaked beads failed to induce *Scx* expression in *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>* mice (Fig. 5I). These data suggest that TGF- $\beta$  signaling is required to play a cell autonomous role in inducing *Scx* expression and in regulating CNC differentiation during tongue development.

#### *Loss of TGF- $\beta$ signaling in CNC cells results in a reduction of muscle cell number in vitro*

To investigate the relationship between cranial neural crest and myogenic cells, we analyzed a primary cell culture of tongue primordium from *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>* mice. The cells harvested



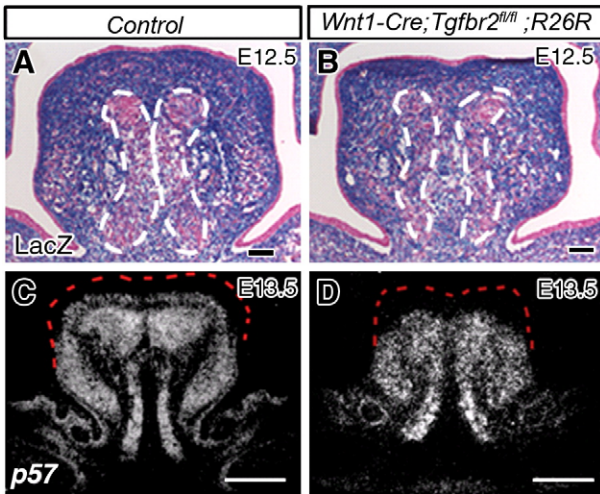
**Fig. 3.** Loss of *Tgfr2* adversely affects myogenic cell proliferation. (A) Schematic diagram of Fluorescence Activated Cell Sorting (FACS) by Fluorescence di-β-d-Galactopyranoside (FDG) staining. The graph indicates the number of FDG-positive or -negative cells from *Wnt1-Cre;Tgfr2<sup>flx/flx</sup>* (Control) or *Wnt1-Cre;Tgfr2<sup>flx/flx</sup>* mice. (B) *Tgfr2* gene expression level in either FDG-positive or -negative cells. Student *t*-tests were used for statistical analysis. \*: *P* < 0.05. (N = 2). (C and D) MHC immunostaining and LacZ staining of E12.5 sagittal sections from *Wnt1-Cre;Tgfr2<sup>flx/flx</sup>;R26R* (Control) and *Wnt1-Cre;Tgfr2<sup>flx/flx</sup>;R26R* mice. MHC positive cells are detectable at the tip of the tongue primordium in Control and *Wnt1-Cre;Tgfr2<sup>flx/flx</sup>;R26R* mice. (E–H) BrdU staining with LacZ staining of E13.5 sections from *Wnt1-Cre;Tgfr2<sup>flx/flx</sup>;R26R* (E, G, Control) and *Wnt1-Cre;Tgfr2<sup>flx/flx</sup>;R26R* (F, H, conditional knockout mice). G and H are enlarged from E and F, respectively. (E and G) BrdU positive cells in control mice are distributed between CNC cells (blue staining, white arrow) and myogenic cells (non-blue cells, black arrow). (F and H) BrdU positive myogenic cells are scattered in *Wnt1-Cre;Tgfr2<sup>flx/flx</sup>;R26R* mice (black arrows). (I and J) BrdU labeling index in each myogenic cell (G) and CNC cells (H). Five randomly selected, non-overlapping samples were used from each experimental group. Student *t*-tests were used for statistical analysis. \*: *P* < 0.05. (N = 5). Scale bars: 200 μm in C, D, E, and F.

from *Wnt1-Cre;Tgfr2<sup>flx/flx</sup>* mice were indistinguishable from those of control mice when cultured in growth medium, which inhibited myogenic differentiation (data not shown), and after changing from growth medium to differentiation medium (total 6 days of culture) (Figs. 6B and C). Moreover, *Wnt1-Cre;Tgfr2<sup>flx/flx</sup>* and control cells

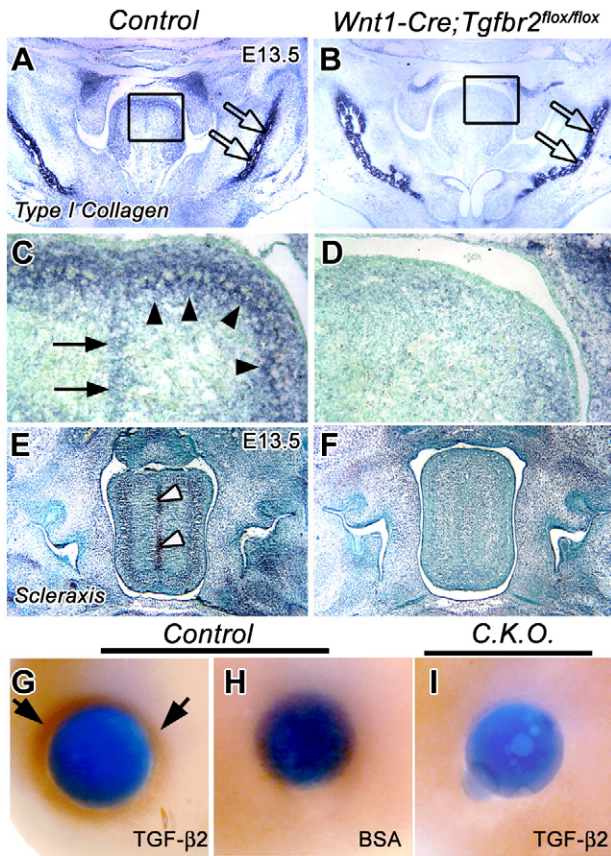
both expressed myosin heavy chain (MHC), a marker of myogenic differentiation, at this stage (Figs. 6B and C, inserts). Thus, primary myogenic differentiation appears normal in *Wnt1-Cre;Tgfr2<sup>flx/flx</sup>* mice. After 14 days of culture, the number of muscle cells in *Wnt1-Cre;Tgfr2<sup>flx/flx</sup>* mice was decreased when compared with that from control mice (Figs. 6D, E and G). These observations suggest that CNC cells might have a critical function in muscle cell formation. To test this hypothesis, we applied conditioned medium from the control culture to the *Wnt1-Cre;Tgfr2<sup>flx/flx</sup>* primary culture. This addition resulted in an increased number of muscle cells in the *Wnt1-Cre;Tgfr2<sup>flx/flx</sup>* primary cell culture (Figs. 6F and G). This stimulation indicates that there may be secreted factors from cranial neural crest cells that are required for myogenic development.

*FGF10 functions downstream of TGF-β signaling to regulate tongue development*

To elucidate the molecular mechanism of TGF-β signaling during tongue development, we performed microarray analyses to compare gene expression profiles between wild type and *Tgfr2* mutant tongue at E12.5. We decided to focus on E12.5 because we detected cell proliferation defects in the developing tongue of *Wnt1-Cre;Tgfr2<sup>flx/flx</sup>* mutant mice beginning at E13.5 (see Figs. S2 and 3D). In our initial set of three microarray experiments, we detected a significant reduction in *Fgf10* expression in the tongue primordium of *Wnt1-Cre;Tgfr2<sup>flx/flx</sup>* mice (data not shown). Both FGFs and their receptors are expressed during tongue development (Nie, 2005). Furthermore, FGF signaling has been shown to be critical for skeletal myogenesis during *Xenopus* and chick embryonic development (Itoh et al., 1996; Standley et al., 2001). We detected *Fgf10* expression in CNC-derived cells in tongue primordium of control mice at E12.5 (Fig. 7A). In the *Wnt1-Cre;Tgfr2<sup>flx/flx</sup>*



**Fig. 4.** Compromised patterning of myogenic cells in *Wnt1-Cre;Tgfr2<sup>flx/flx</sup>* mice. (A and B) LacZ staining of *Wnt1-Cre;Tgfr2<sup>flx/flx</sup>;R26R* mice (A) and *Wnt1-Cre;Tgfr2<sup>flx/flx</sup>;R26R* mice (B) at E12.5. White dashed line indicates the margin of myogenic core. (C and D) In situ hybridization of p57 from *Wnt1-Cre;Tgfr2<sup>flx/flx</sup>* mice (C) and *Wnt1-Cre;Tgfr2<sup>flx/flx</sup>* mice (D). Red dashed line indicates the margin of tongue primordium. Scale bars: 200 μm in A–D.



**Fig. 5.** Scleraxis (*Scx*) and type I collagen expression in CNC-derived cells in the tongue. In situ hybridization of type I collagen and Scleraxis (*Scx*) in control (A,C,E) and *Wnt1-Cre;Tgfr2<sup>lox/lox</sup>* (B,D,F) mice at E13.5. (A–D) Type I collagen is expressed in the dense lamina propria (arrowhead) and septum (arrow) but its expression is greatly reduced in the *Tgfr2* mutant sample. Open arrow points to type I collagen expression in the developing mandible. (E and F) *Scx* is expressed in the tongue septum of control mice (white arrowheads), but not in the tongue of *Tgfr2* mutant mice. (G–I) Explants from tongue primordia of E12.5 control and *Wnt1-Cre;Tgfr2<sup>lox/lox</sup>;R26R* mice treated with beads for 12 h. The control sample treated with TGF- $\beta$ 2 beads (G) is positive for *Scx* expression (black arrows), but BSA beads do not induce *Scx* expression (H). The conditional knockout sample treated with TGF- $\beta$ 2 beads is negative for *Scx* expression (I).

mutant, there was a reduction in *Fgf10* expression in the CNC-derived cells surrounding the myogenic core (Fig. 7B). In order to test whether there was a significant reduction in the expression level of *Fgf10*, we performed real-time RT-PCR analysis. We found that there was a significant reduction in *Fgf10* expression in the tongue primordium and in primary culture of tongue primordium in *Wnt1-Cre;Tgfr2<sup>lox/lox</sup>* mice (Figs. 7C and D). Thus, our data suggest that there is a cell autonomous requirement for TGF- $\beta$  signaling to induce *Fgf10* expression in CNC-derived cells during tongue development.

#### Exogenous FGF10 rescues myogenic cell proliferation in *Tgfr2* mutant mice

Mutation in *Tbx1* results in down-regulation of FGF10 expression, which affects the patterning of CNC cells in the mandibular arch and results in defects in branchiomeric myogenesis in mice (Kelly et al., 2004). To test the hypothesis that FGF10 acts downstream of TGF- $\beta$  to control myogenic cell proliferation, we performed rescue experiments using the primary tongue cell culture model. Following four days of culture in growth medium, we switched the cultured tongue cells into differentiation medium for another ten days. Cells from the control sample expanded their population and differentiated into myogenic cells, as validated by positive MHC staining (Fig. 8A). In *Wnt1-Cre;Tgfr2<sup>lox/lox</sup>* mutant samples, there was a significant reduction

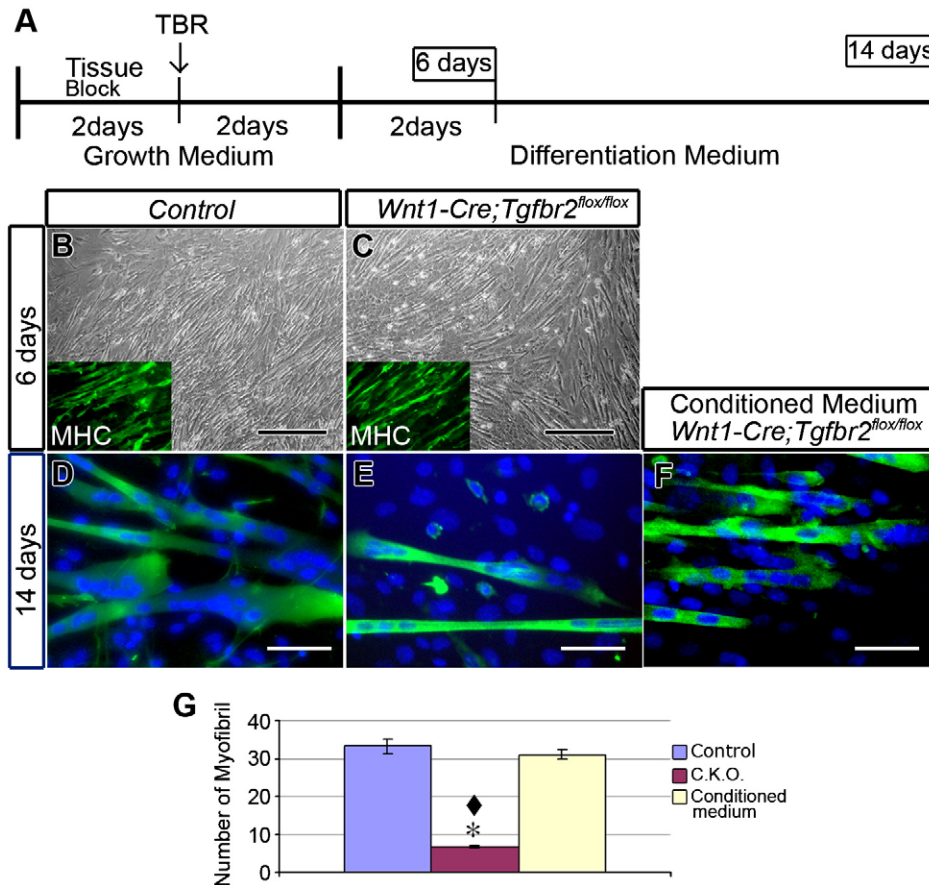
( $P < 0.05$ ) in the number of muscle cells. Significantly, the addition of exogenous FGF10 in the differentiation medium was able to increase the number of muscle cells (Fig. 8B). In contrast, addition of BSA failed to restore the number of muscle cells in the *Wnt1-Cre;Tgfr2<sup>lox/lox</sup>* mutant tongue cell culture (Fig. 8C). Statistical analyses revealed that exogenous FGF10 was able to restore the number of muscle cells in the *Wnt1-Cre;Tgfr2<sup>lox/lox</sup>* cell culture substantially, but not completely to wild type levels, although addition of FGF10 had no effect on control samples (Fig. 8D). These results suggest that additional members of the FGF family, or other factors, may also mediate TGF- $\beta$  signaling to control the proliferation and survival of myogenic cells during tongue development. In a parallel approach, we placed beads carrying FGF10 or BSA in cultured tongue explants. The number of muscle cells in *Wnt1-Cre;Tgfr2<sup>lox/lox</sup>* mutant tongue explants treated with FGF10 beads increased close to that of control mice explants, whereas BSA beads failed to restore the number of muscle cells in mutant tongue explants (data not shown). Finally, to test whether TGF- $\beta$ 2 could induce *Fgf10* expression, we applied TGF- $\beta$ 2 ligand or BSA into primary tongue cell cultures of control mice, immediately following the switch from growth medium to differentiation medium. *Fgf10* expression was induced two hours after the addition of TGF- $\beta$ 2, whereas BSA addition had no effect (Fig. 8E). Taken together, our data suggest that TGF- $\beta$  induced FGF signaling plays an important role in regulating tissue–tissue interaction and myogenic cell proliferation during tongue development.

#### Neural crest cells are required for organizing tongue muscle cells

To test the reciprocal relationship between muscle cells and tendon cells (derived from CNC cells), we used a tissue regeneration approach. We separated myogenic and CNC cells from E12.5 control and *Wnt1-Cre;Tgfr2<sup>lox/lox</sup>* tongue primordia using cell sorting and combined cells as depicted in Figs. S3A–C. The first sample contained CNC cells and myoblasts from control mice, the second was CNC cells from control mice and myoblasts from conditional knockout mice, and the third was CNC cells from conditional knockout mice and myoblasts from control mice. These cell mixtures were seeded in a collagen scaffold and the scaffold was inserted beneath the back skin of host mice for 2 weeks. We detected muscle cells along the scaffold in the first sample (Figs. S3D and G). The second sample also resulted in muscle cell formation in which the myoblasts had also been derived from conditional knockout mice (Figs. S3E and H). In contrast, we observed disorganized MHC (myosin heavy chain) positive cells in the third sample, even though the myoblasts were derived from control mice. Our results suggest that CNC cells play a crucial role in organizing myogenic cells. Furthermore we examined the transplanted CNC cells to eliminate the possibility that the phenotype we observed was due to a difference in survival. We detected CNC cells from control mice around muscle cells in the first and second groups (Figs. S3J and K) and the CNC cells from conditional knockout mice in the scaffold (third group, Fig. S3L), scattered but not organized around the myogenic cells. These observations suggest that CNC cells possess a function important for muscle cell formation and organization.

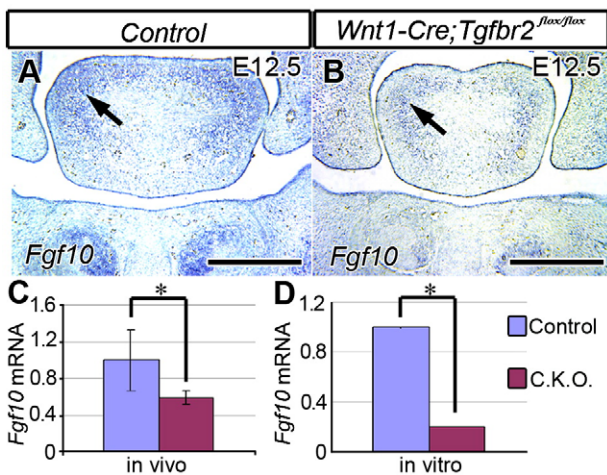
#### Discussion

In this present study, we provide the first evidence that TGF- $\beta$  signaling has a critical function in regulating skeletal muscle development through tissue–tissue interactions. We find that loss of TGF- $\beta$  signaling in CNC cells results in microglossia with defects in CNC-derived connective tissue and tongue muscle development. Specifically, TGF- $\beta$  is required for Scleraxis and *Fgf10* expression in CNC-derived cells and cell proliferation and organization in their neighboring myogenic progenitors. Thus, we demonstrate that the CNC-derived cells surrounding the myogenic progenitors have a critical function in regulating the development of tongue muscle.



**Fig. 6.** Compromised muscle cell formation in primary cell culture of *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>* mice. (A) Schematic diagram of the timetable of primary cell culture. TBR: Tissue Block Remove. (B and C) After 6 days of culture, the appearance of cells from *Wnt1-Cre;Tgfb2<sup>flox/+</sup>* (control) and *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>* mice is indistinguishable under the phase microscope. Inserts show immunostaining of myosin heavy chain (MHC). (D–F) MHC immunostaining (MHC appears green, DAPI appears blue) after 14 days of *Wnt1-Cre;Tgfb2<sup>flox/+</sup>* (control) and *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>* primary cell culture and *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>* cells treated with conditioned medium from the control culture. After 14 days of culture, the number of muscle cells in the *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>* culture (E) is reduced compared to control (D). The addition of conditioned medium to the *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>* culture (F) increased the number of muscle cells. (G) Quantitation of muscle cell number from D to F. Scale bars: 300  $\mu$ m in A–F.

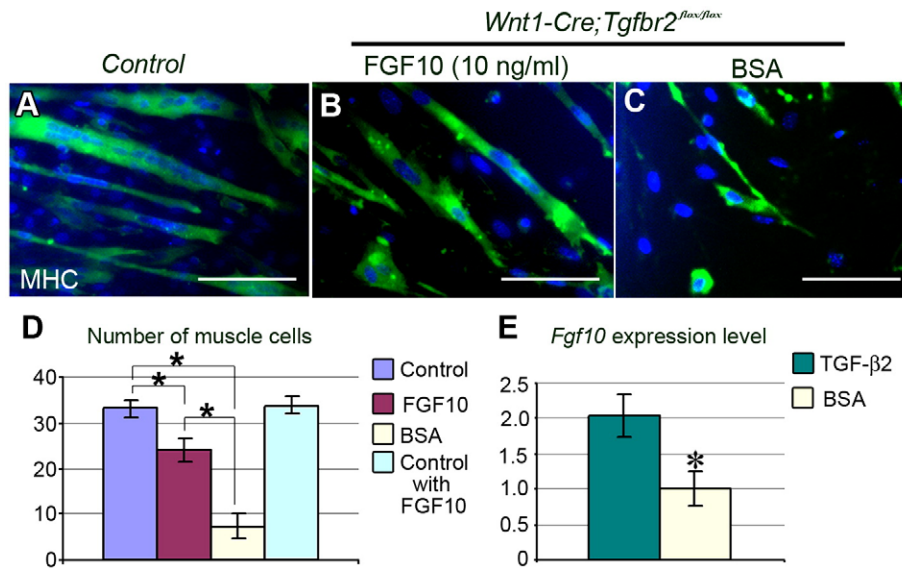
#### Tissue–tissue interactions in regulating tongue muscle development



**Fig. 7.** *Fgf10* expression analysis in the developing tongue. (A and B) In situ hybridization of *Fgf10* mRNA in E12.5 *Wnt1-Cre;Tgfb2<sup>flox/+</sup>* (control) and *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>* mice. *Fgf10* mRNA is expressed in cranial neural crest cells around a myogenic core in the tongue primordium (arrows) of control and *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>* mice. (C and D) Quantitative analysis of *Fgf10* expression in *Wnt1-Cre;Tgfb2<sup>flox/+</sup>* and *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>* mice using real-time PCR. The expression level of *Fgf10* in *Wnt1-Cre;Tgfb2<sup>flox/flox</sup>* mice at E12.5 is reduced in vivo (C). *Fgf10* expression is also reduced in primary cell culture samples in vitro (D). Student *t*-tests were used for statistical analysis. \*:  $P < 0.05$ . (N = 5). Scale bars: 300  $\mu$ m in A, B.

Craniofacial organs offer unique opportunities for the investigation of reciprocal tissue–tissue interactions in regulating embryonic development. For instance, tooth development involves reciprocal interactions between the oral epithelium and the CNC-derived ectomesenchyme tissues (Kettunen et al., 2000). Muscles in the craniofacial region derive from two mesenchymal cell lineages; one consists of mesoderm-derived myogenic progenitors and their surrounding tissues derive from cranial neural crest cells. The intimate relationship between these two cell lineages suggests reciprocal interactions between CNC and myogenic cells in the process of skeletal muscle development. Furthermore, because of our ability to generate CNC or myogenic cell specific inactivation of a gene of interest, we have a unique opportunity to investigate the functional significance of tissue–tissue interactions in regulating craniofacial skeletal muscle development.

The tongue is a muscular organ and performs important physiological functions in our daily lives. Using our two component genetic model for indelibly marking the progenies of CNC cells, we show in this study that tendons and connective tissues are derived from CNC cells in the tongue and are critical for the organization and function of tongue. Previous studies have shown that, in invertebrates such as *Drosophila*, the ectoderm is the origin for tendons (Schnorre and Dickson, 2004). Chick/quail experiments indicates that the tendon and surrounding tissue of craniofacial muscles are composed of CNC cells in the craniofacial region (Noden, 1988). In vertebrates,



**Fig. 8.** Exogenous FGF10 rescues the reduction of muscle cells in *Wnt1-Cre;Tgfb2<sup>flax/flax</sup>* mice. (A–C) Immunostaining of MHC (green) in *Wnt1-Cre;Tgfb2<sup>flax/+</sup>* primary cell culture (Control, A) and *Wnt1-Cre;Tgfb2<sup>flax/flax</sup>* primary cell cultures with FGF10 (B) or BSA (C) added into the medium. (D) Quantitative analysis of the number of muscle cells detectable in sample fields (20×). Student *t*-tests were used for statistical analysis. \*:  $P < 0.05$ . ( $N = 5$ ). (E) *Fgf10* gene expression level analyzed by real-time PCR. Exogenous TGF-β2 ligand induces *Fgf10* expression. Student *t*-tests were used for statistical analysis. \*:  $P < 0.05$ . ( $N = 2$ ). Scale bar: 100 μm in A–C.

tendons are derived from the lateral mesoderm, except in the craniofacial region where CNC-derived cells give rise to tendon and connective tissue in the skeletal muscles.

In the mouse model, *Scleraxis* (*Scx*) is considered a marker for tendon cells (Brent et al., 2003). Tendon formation is diminished in *Scx* knockout mice (Murchison et al., 2007). *Scx* is a basic helix–loop–helix transcription factor involved in controlling collagen expression. Pro-α1(I) collagen (*COL1a1*) is one of the chains composed of Type I collagen (van der Rest and Garrone, 1991). There are TSE1 (Tendon-specific elements) and TSE2 regions in the promoter region of *COL1a1* (Terraz et al., 2002). *SCX* binds to TSE1 and TSE2 and controls the expression of *COL1a1* (Lejard et al., 2007). Studies of TSE1 and TSE2 LacZ transgenic mice indicate that these regions express in the fibroblast cells giving rise to tendon cells (Terraz et al., 2002). The LacZ staining in these mice is detectable not only in the tendons of the trunk region but also in the tendons of craniofacial muscles such as the masseter. Our present study reveals that *Scx* and *Type I collagen* expression is compromised following the loss of TGF-β signaling and that TGF-β ligand can induce *Scx* expression. Taken together, we conclude that TGF-β mediated *Scleraxis* and *Type I collagen* expression is critical in regulating the fate of CNC cells during craniofacial skeletal muscle development.

Skeletal muscle development involves cell migration, proliferation, differentiation, and organization stages (Birchmeier and Brohmann, 2000; Buckingham et al., 2003; Mackenzie et al., 1998; Noden and Francis-West, 2006). During the migration stage, myoblast precursor cells can migrate into their presumptive location even if the CNC cells are ablated (Huang et al., 1999; von Scheven et al., 2006). This result implies that CNC cells are not required for the migration of myogenic precursors towards their presumptive destinations, including the branchial arch and tongue primordium. At the differentiation stage, CNC cells in the branchial arch of chicks secrete BMP and Wnt inhibitors, which induce myogenic differentiation (Tzahor et al., 2003). However, it is still unclear how CNC cells control the organization of myogenic cells. During skeletal muscle development, muscle cells and surrounding tissue, such as tendon cells, both appear to be involved in the formation of the attachment to the skeletal muscle. For instance, the loss of *stripe*, which is involved in early steps of tendon differentiation, results in the disruption of the entire

somatic muscle pattern (Frommer et al., 1996). A defect in tendon formation causes compromised muscle formation in *Drosophila* and zebrafish (Frommer et al., 1996; Kudo et al., 2004). The loss of *Periostin*, an adhesion molecule, in the myoseptum causes a differentiation defect in myoblasts (Kudo et al., 2004). These results imply that the reciprocal interaction between myogenic and surrounding cells, including tendon cells, is critical as a regulatory mechanism of skeletal muscle development.

#### TGF-β mediated FGF signaling controls tissue–tissue interactions during tongue development

Previous studies have shown that FGF ligands and their receptors are required for skeletal muscle development (de Alvaro et al., 2005; Flanagan-Steet et al., 2000). For instance, the viral expression of truncated *Fgf type I* receptor causes a skeletal muscle defect in the chick limb (Flanagan-Steet et al., 2000). FGF signaling controls myogenic cell differentiation status through MAP kinase (de Alvaro et al., 2005; Gonzalez et al., 2004; Naya and Olson, 1999). We demonstrate here that secreted factor(s) from CNC cells are critical for myogenic development, because conditioned cell culture medium can rescue the reduction of muscle cell number in the *Wnt1-Cre;Tgfb2<sup>flax/flax</sup>* sample. *Fgf10* is a candidate for this secreted factor. *Fgf10* expression is reduced in *Wnt1-Cre;Tgfb2<sup>flax/flax</sup>* mice, and exogenous FGF10 reverses the reduction of tongue muscle cell number in vitro. The *Fgf10* conventional knockout mouse is another model that has revealed reciprocal interactions between two cell lineages. *Fgf10* null mice exhibited a reduction of cell proliferation activity in the dental epithelium, even though *Fgf10* expression is detected only in the dental mesenchyme. Exogenous FGF10 reverses the reduction of cell proliferation in *Fgf10* null mice (Harada et al., 2002). FGF10 has also been shown to be critical for mediating tissue–tissue interactions during lung and palate development (Bellusci et al., 1997; Bhushan et al., 2001; Rice et al., 2004). Taken together, our study and others show that a paracrine function of FGF10 is critical for embryonic organ development.

Our conditional knockout mouse model helped us to explore the reciprocal relationship among tissue layers during organogenesis. Specifically, this study demonstrates that the loss of *Tgfb2* in non-



myogenic cells results in the disorganization of muscle cells. In contrast, conditional inactivation of *Tgfb2* in myogenic specific cell lineages exhibit organized muscle cells (our unpublished data), although myogenic differentiation is compromised as reported in a previous cell culture study (Filvaroff et al., 1994). We have also investigated how the cells surrounding the myoblast contribute to muscle cell organization during their formation using a tissue regeneration experimental approach. Our data indicate that TGF- $\beta$  signaling is crucial in CNC-derived cells, but not in myoblasts, for organizing muscle cells during tongue muscle development.

We have presented the first study investigating the function of CNC cells during skeletal muscle development using a genetic, rather than surgical or tissue transplantation approach. Although cell line experiments have revealed that many genes are critical for skeletal muscle development in a cell autonomous manner (de Alvaro et al., 2005; Deng et al., 1994), they have yet to provide information about the functional significance of tissue–tissue interactions during muscle development. Previously, *Drosophila* or zebrafish studies have been used to reveal the mechanism of skeletal muscle organization (Frommer et al., 1996; Kudo et al., 2004). Our in vivo system allows us to investigate the functional significance of genes in the spatial complexity of the mouse muscular organ and has provided evidence for the importance of TGF- $\beta$  signaling in regulating tissue–tissue interactions during tongue muscle development. In summary, TGF- $\beta$  signaling regulates fibroblast differentiation by controlling *Scleraxis* expression in a cell autonomous manner. TGF- $\beta$  signaling controls myogenic cell proliferation via secreted factors from CNC cells and tissue–tissue interactions during tongue morphogenesis. Disruption of this interaction may lead to dysmorphogenesis.

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## Appendix A. Supplementary data

Supplementary data associated with this article can be found, in the online version, at doi:10.1016/j.ydbio.2010.02.030.

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