Activation of the WNT-BMP-FGF Regulatory Network Induces the Onset of Cell Death in Anterior Mesodermal Cells to Establish the ANZ

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The spatiotemporal control of programmed cell death (PCD) plays a significant role in sculpting the limb. In the early avian limb bud, the anterior necrotic zone (ANZ) and the posterior necrotic zone are two cell death regions associated with digit number reduction. In this study, we evaluated the first events triggered by the FGF, BMP, and WNT signaling interactions to initiate cell death in the anterior margin of the limb to establish the ANZ. This study demonstrates that in a period of two to 8 h after the inhibition of WNT or FGF signaling or the activation of BMP signaling, cell death was induced in the anterior margin of the limb concomitantly with the regulation of Dkk, Fgf8, and Bmp4 expression. Comparing the gene expression profile between the ANZ and the undifferentiated zone at 22HH and 25HH and between the ANZ of 22HH and 25HH stages correlates with functional programs controlled by the regulatory network FGF, BMP, and WNT signaling in the anterior margin of the limb. This work provides novel insights to recognize a negative feedback loop between FGF8, BMP4, and DKK to control the onset of cell death in the anterior margin of the limb to the establishment of the ANZ.

Keywords: limb development, ANZ, Fgf signaling, Wnt signaling, BMP signaling, programmed cell death, morphogenesis, Anterior Necrotic Zone

INTRODUCTION

Programmed cell death (PCD) is essential to regulate the final morphology and sculpting limbs (Pajni-Underwood et al., 2007; Montero et al., 2020). PCD participates in separating digits and zeugopodial elements (Hinchliffe and Thorogood, 1974; Zuzarte-Luis and Hurle, 2005). The anterior and posterior margins of the avian limb are associated with digit reduction (Saunders and Gasseling, 1962; Zuzarte-Luis and Hurle, 2002; 2005). Those regions are called the anterior necrotic zone (ANZ) and posterior necrotic zone (PNZ) (Saunders and Gasseling, 1962). Meanwhile, PCD in the interdigital regions takes part in species with free digits.

The process of cell death is under the control of the apical ectodermal ridge (AER); it is well known that limb truncation and massive cell death occur after the elimination of the AER, demonstrating that cell survival of mesodermal cells depends on the signals from this epithelium. The molecular analysis of AER indicates that Fibroblast Growth Factor (FGF) family...
members expressed in the AER govern cell proliferation and cell survival. FGF8 signaling protects the mesodermal cells from cell death (Niswander et al., 1993; Sun et al., 2002; Mariani et al., 2008; Ten Berge et al., 2008; Mariani et al., 2017). Mutant mice for the Fgf8 gene or the blockade of the function of FGF signaling induces cell death in the mesodermal cells (Montero et al., 2001; Sun et al., 2002; Mariani et al., 2008; Mariani et al., 2017). FGF8 participates promoting a positive feedback loop, inducing the expression of Fgfl0 in the mesoderm whose interaction with the FGF receptor 2b (FGFR2b) in the AER promotes the expression of Wnt3a (Xu et al., 1998; Danopoulos et al., 2013; Haro et al., 2014; Jin et al., 2018). WNT3A induces the expression of transcription factors Sp6 and Sp8 that induce Fgf8 expression in the AER (Kawakami et al., 2001; Kawakami et al., 2004). WNT signaling from the AER participates in maintaining the undifferentiated state of mesodermal cells under AER (Ten Berge et al., 2008). The function of WNT-ß catenin signaling may be blocked by DKK, an antagonist of this signaling pathway. Gene deletion of DKK results in cell death inhibition, and Dkk expression is observed in cell death regions during limb development (Grotewold et al., 1999; Mukhopadhyay et al., 2001; Grotewold and Ruther, 2002a; b).

During limb development, the bone morphogenetic proteins (BMP) signaling is another pathway controlling cell death. Bmp2, Bmp4, and Bmp7 expression are observed in the interdigital regions and the anterior and posterior margins of the limb (Hogan, 1996; Yokouchi et al., 1996; Macias et al., 1997; Chen and Zhao, 1998; Merino et al., 1999; Salas-Vidal et al., 2001; Zuzarte-Luis et al., 2004). The implantation of BMP-soaked beads promotes cell death in the interdigital tissue, whereas the blockade of BMP signaling with NOGGIN or GREMLIN inhibits it (Ganan et al., 1996; Macias et al., 1997; Zuzarte-Luis et al., 2004). BMP stimulates the SMAD1/5/8 signaling pathway in blood vessels and mesodermal cells, promoting cell death in mesodermal cells, and inhibiting Fgf8 expression in the AER (Zuzarte-Luis et al., 2004; Monteiro et al., 2008; Abarca-Buis et al., 2011).

High levels of FGF signaling in the interdigital mesoderm downregulate Bmp genes, inhibiting cell death (Montero et al., 2001; Hernandez-Martinez et al., 2009). However, FGF2 signaling promoted cell death induced by BMP proteins, suggesting that FGF also works in a feedback loop with BMP signaling (Montero et al., 2001). Mesodermal cells become competent to signaling pathways that control cell fate when WNT and FGF signals are depleted underneath AER (Ten Berge et al., 2008). If cells receive cell death-promoting factors, presumably BMPs, cells enter the cell death program (Montero et al., 2001; Hernandez-Martinez et al., 2009). Otherwise, the cell differentiation program begins if mesodermal cells receive chondrogenic signals (Chimal-Monroy et al., 2003; Monteiro et al., 2008; Marin-Ilera et al., 2019). Thus, FGF and WNT signaling together with BMP signaling establish a well-known regulatory network to control the undifferentiated state, cell proliferation, and cell survival during limb development (Niswander et al., 1993; Montero et al., 2001; Sun et al., 2002; Mariani et al., 2008; Ten Berge et al., 2008; Hernandez-Martinez et al., 2009; Mariani et al., 2017). The participation of this regulatory network is better known during the PCD in the interdigital regions than in the ANZ or PNZ. A previous study demonstrated that a BMP-pulse of 4 h was sufficient to induce cell death in the anterior margin of the limb (Abarca-Buis et al., 2011). Notably, TUNEL-positive cells show no co-localization of nuclear phosphorylated SMAD1/5/8 proteins suggesting that BMP signaling participates in a molecular cascade in the ANZ, culminating in cell death (Abarca-Buis et al., 2011).

Because a short pulse of BMP is sufficient to induce cell death in the anterior margin of the limb, this work aimed to determine how the regulatory network integrated by FGF, BMP, and WNT signaling pathways participate in the establishment of the ANZ. The results presented here demonstrated that inhibition of WNT or FGF signaling or the activation of BMP signaling during a short period is sufficient to induce cell death in the anterior margin of the limb and to regulate Dkk, Fgf8, and Bmp4 expression. Thus, the regulatory network of the FGF-BMP-WNT signaling pathway induces cell death in the anterior margin of the limb to establishing the ANZ.

MATERIAL AND METHODS

Ethics

This protocol was reviewed and approved by the Institutional Ethical Review Board for the Care and Use of Laboratory Animals of the Instituto de Investigaciones Biomédicas, Universidad Nacional Autónoma de México (UNAM, Mexico City, Mexico).

Eggs and Embryo Manipulations

Fertilized White Leghorn chicken eggs (ALPES, Puebla, Mexico) were incubated at 38°C and staged according to Hamburger and Hamilton (1951). Eggs were windowed to expose the right limb at developing stages from 22 to 25 HH for experimental procedures. Heparin beads (Cat. H6508, Sigma-Aldrich, St. Louis, MO, United States) or in Affigel (Bio-Rad Laboratories, Hercules, CA) were soaked in 1 mg/ml FGF8 (cat. 100-25A), FGF10 (100-26), 1 mg/ml DKK (cat. 120-30), BMP4 (cat. 120-05), BMP7 (cat. 120-30P) or 2 mg/ml NOGGIN (cat. 120-10C) from Peprotech, Mexico City, Mexico. Agl-X2 ionic exchange beads (Cat. 1401231, Sigma-Aldrich, Mexico) were soaked in 4 mg/ml SU502 and placed in the ANZ of embryonic limbs. Manipulated embryos were incubated for short times according to the experiments and processed for lysotracker staining, in situ hybridization, or both.

RNA Probes and in Situ Hybridization

RNA antisense probes were labeled with UTP-digoxigenin (11209256910, Roche Applied Science, Indianapolis, IN, United States) and used for whole-mount in situ hybridization (ISH) as described previously (Merino et al., 1998). Samples were treated with 60 μg/ml proteinase K for 25 min at 21°C for Bmp7, Fgf1, Fgf2, Fgf3, Mkp3, Mxs2, and Wif. Bambi required 70 μg/ml proteinase K for 28 min at 25°C; 60 μg/ml was used for 22 min at 21°C for Bmp4 and Dkk. Fgf8 was treated with 15 μg/ml for 20 min at 21°C. The hybridization temperature was 68°C, and post-hybridization washes were at 70°C for all genes. The signal of ISH was visualized with BM-Purple substrate for alkaline phosphatase (Roche Applied Science). Images were acquired...
with the Nikon StereoScope Fluorescence Microscope SMZ1500 (Nikon Corporation, NY, United States) or in AxioZoom V.16 microscope (Carl Zeiss, Oberkochen, Germany) using Zen lite software (Carl Zeiss, Oberkochen, Germany).

Detection of Cell Death With Lysotracker and Neutral Red Staining
Embryonic limbs were incubated in 1 μM LysoTracker Red DND-99 (Cat. L7528, Molecular Probes) at 37°C for 15 min. Samples were rinsed twice in PBS and fixed in 4% PFA overnight at 4°C. Some samples were also processed for ISH as described previously (Abarca-Buis et al., 2011). After ISH, limb buds were dehydrated in increasing methanol-PBS-Tween series and cleared with 2:1 benzyl alcohol: benzyl benzoate solution for 1 h each (Parish, 1999). For Neutral Red staining, limbs were isolated, washed in PBS, and stained with 2% Neutral Red in PBS time at 37°C. Images were acquired with a Nikon StereoScope Fluorescence Microscope SMZ1500 (Nikon Corporation, NY, United States) and Fluorescence Microscope Axio Zoom. V16 Carl Zeiss, Göttingen, Germany.

Real-Time RT-PCR
RNA extractions were performed with NucleoSpin RNA (Macherey-Nagel, cat. no. 740955, Düren, Germany), and retrotranscription of total RNA was achieved using the RevertAid RT kit (Thermo Fisher Scientific, cat. no. K1691, Waltham, MA, United States) according to the manufacturer’s instructions. Expression levels were analyzed using a real-time PCR system and quantified with SYBR green (Thermo Fisher). The Rpl13 gene was used as a normalizer. The expression level was evaluated relative to a calibrator according to the 2^(-ΔΔCt) equation. Each value represented the mean ± SD of three independent experiments and was analyzed using Student’s t-test. Statistical significance was set at p < 0.05.

RESULTS

Anterior Necrotic Zone Appears by the Progressive Loss of the Apical Ectodermal Ridge
To associate the gene expression pattern of Fgf8, a survival signal, and the presence of cell death in the ANZ during limb outgrowth, embryonic limbs from 22HH to 25HH were stained with l lysotracker and hybridized for Fgf8 (Figure 1). The results indicated that cell death was observed in the anterior margin of the limb from stage 23 HH to 25 HH (Figure 1). Concurrently as the limb outgrowth occurs, the Fgf8 expression is gradually downregulated in the AER from the proximal to the distal region, near the ANZ. Thus, these results exhibited a boundary between the downregulation of Fgf8 from the AER and the appearance of the ANZ while progressive cell death in mesodermal cells is observed (Figure 1).

FGF-, BMP-, and WNT-Signaling Pathways Are Active in the Anterior Margin of the Limb Bud When the ANZ Is Established
The association of the expression patterns of the genes related to FGF-, BMP-, and WNT-signaling pathways with the boundary observed between Fgf8, and the appearance of progressive cell death was studied at the 24 HH stage. At this developing stage, cell death and Fgf8 expression are precisely located at neighboring positions (Figure 2), allowing us to study how FGF-, BMP-, and WNT-signaling pathways regulate the onset of cell death and the appearance of ANZ in the developing limb.

The genes related to FGF signaling, such as Mkp3, a target of this signaling pathway, and the three receptors of FGF, Fgfr1, Fgfr2, and Fgfr3 (Figure 2A) were evaluated. Mkp3 and Fgfr1 were expressed in the distal part of the anterior margin and in the anterior half of the distal position. Besides, Mkp3 was expressed in the central part of the proximal region. Fgfr2 was mainly expressed in both anterior and posterior margins, whereas Fgfr3 was slightly expressed in an extended way in the mesoderm (Figure 2A).

Regarding BMP signaling, the expression of Bmp4, Bmp7, and Bambi was evaluated. Bmp4 was expressed in both anterior and posterior margins of the limb (Figure 2B). In contrast, Bmp7 expression was mainly localized in the most distal region of the limb riming undifferentiated region underneath AER and the anterior margin of the limb (Figure 2B). Bambi expression was observed preferentially in the proximal region of the anterior margin and the most distal region of the posterior margin of the limb (Figure 2B). In addition, as a marker of undifferentiated cells and regulated by BMP signaling, we evaluated the Msx2 gene expression pattern.

![FIGURE 1](image-url) Progressive loss of the apical ectodermal coincides with the apparition of cell death in the anterior margin of the limb. In situ hybridization of Fgf8 gene expression and cell death pattern evidenced by lysotracker stain at limb development stages from 22 to 25 HH. Notice that cell death occurs in regions in which Fgf8 expression is disappearing.
Results showed that it is expressed in the anterior margin of the limb (Figure 2B).

On the other hand, Dkk and Wif gene expression was analyzed as genes related to Wnt signaling (Figure 2C). Dkk gene expression was localized in the anterior and posterior margins of the limb (Figure 2). Wif gene expression was observed in all anterior and posterior margins and the distal region of the limb (Figure 2C). These results showed a dynamic gene expression pattern suggesting that all three signaling pathways are active during the process of cell death in the anterior and posterior limb margins.

FGF, WNT, and BMP Signaling Control the Induction and Maintenance of Cell Death in the ANZ

Although the gene expression pattern is quite similar in the anterior and posterior margin of the limb, here was decided to determine the minimum time to promote cell death in the ANZ modulating the function of FGF, WNT, and BMP signaling pathways. As a first approach, FGF8-soaked beads or FGF signaling inhibitor (SU5402)-soaked beads were placed in the anterior margin of the limb at the 24 HH stage (Figure 3A). The results showed that FGF8 treatments did not inhibit cell death. In contrast, inhibiting FGF signaling for 6 h was sufficient to increase cell death in this region (Figure 3A). On the other hand, the minimal time to induce cell death after inhibiting WNT signaling with DKK-soaked beads in the anterior margin of the limb was 8 h. In contrast, as expected, WNT3A soaked beads did not inhibit cell death (Figure 3B).

It is known that a short pulse of BMP induces cell death in the ANZ (Abarca-Buis et al., 2011) and because the expression of the Bmp4 and Bmp7 is observed in the anterior margin. Here the cell death promoted by both proteins was evaluated. The results showed that cell death in the ANZ stimulated by BMP4 and BMP7 occurs after 6 h (Figure 3C) and is inhibited by NOGGIN after 6 h of treatment (Figure 3C). Taking together, these results suggest that the minimum time to trigger cell death in the ANZ is...
6 h after inhibition of FGF and activation of BMP signaling or 8 h after inhibiting WNT signaling.

**FGF- and BMP-Signaling Are Coordinated to Regulate Cell Death in the ANZ**

The inhibition of FGF or activation of BMP signaling triggers cell death after 6 h. Thus, to determine the relation between FGF and Bmp4 and Dkk expression during the induction of cell death, the gene expression of Bmp4 and Dkk was evaluated after inhibiting or activating FGF signaling (Figure 4). The results demonstrated that Dkk expression was inhibited after 2 h of treatment, whereas Bmp4 inhibition occurred after 4 h (Figure 4A). However, after the administration of proteins FGF8 or FGF10 to activate FGF signaling, only FGF10 induced moderately Bmp4 expression without important changes in the expression of Dkk and Bmp4 by FGF8 (Figures 4 B, C). Thus, because the inhibition of FGF signaling for two or 4 h, downregulated Dkk and Bmp4,
**FIGURE 4** | Inhibition of Dkk and Bmp4 gene expression by treatment with an inhibitor on FGF signaling. Gene expression pattern of Bmp4 and Dkk in the anterior margin of the limb at developing stage 24HH. (A) Dkk expression inhibition was observed after 2 h of treatment with SU5402, while Bmp4 was inhibited after 4 h. The gene expression pattern of Dkk and Bmp4 was not modified at least 4 h after (B) FGFR- and (C) FGFR10-treatment.

**FIGURE 5** | BMP signaling regulates Fgf8, Bmp4, Dkk, Msx2, and Bambi gene expression. Regulation of (A–C) Fgf8, (D–F) Bmp4, (G–I) Dkk, (J–L) Msx2, (M–O) Bambi gene expression in the anterior limb margin after BMP4, BMP7, or Noggin. Fgf8 (B, C) and Bmp4 (D, E) are downregulated after 4 h of BMP4 or BMP7. Notice the stronger effect of BMP4 than BMP7. In contrast, NOGGIN-treatment maintained the expression of both genes after treatments. (G–I) Upregulation of Dkk gene expression after 4 h of BMP4- or BMP7-treatment and downregulation of Dkk after 4 h of NOGGIN treatment. (J–L) Msx2 and (M–O) Bambi gene expression were upregulated by BMP4 and BMP7. In contrast, NOGGIN inhibited Bambi expression during the first 4 h of treatment but not Msx2 [Panel (L, O)].
respectively, it is possible to suggest the expression of Dkk and Bmp4 in the anterior margin of the limb depends on FGF signaling.

The next step was to determine the minimum time required for BMP signaling to regulate Fgf8, Bmp4, and Dkk gene expression in the anterior margin of the limb for promoting cell death (Figures 5A–I). The results showed that BMP4 or BMP7 proteins had regulated Fgf8, Bmp4, and Dkk gene expression differentially (Figures 5A,B,D,E,G,H). BMP4 protein inhibited Fgf8 in the AER and Bmp4 in the anterior margin of the limb after 4 h of treatment (Figures 5A,D). In contrast, BMP7 slightly affected Fgf8 gene expression, and it faintly inhibited Bmp4 gene expression (Figures 5B,E). Furthermore, NOGGIN-treatment effects on Fgf8 expression were minor (Figure 5C), while Bmp4 gene expression was not affected after 4 h (Figure 5F). Regarding Dkk gene expression, the treatment with BMP4 showed a more significant effect than BMP7 (Figures 5G,H). Blocking BMP signaling with NOGGIN demonstrated that Dkk gene expression depended on BMP signaling (Figure 5I). Msx2 and Bambi are regulated by BMP signaling (Figures 5J–O). The results showed that Msx2 was regulated by BMP or NOGGIN treatment (Figures 5J–L). In contrast, NOGGIN treatment inhibited Bambi expression entirely, but BMP4 or BMP7 induced it slightly at least 4 h (Figures 5M–O). These results showed that BMP signaling self-regulate Bmp4 and regulates Fgf8, Dkk, Msx2, and Bambi gene expression at short times in the ANZ.

Inhibition of FGF Signaling Is the Last Step in the Molecular Cascade of Cell Death

To determine the hierarchy of FGF and BMP signaling to promote cell death and regulate Dkk gene expression, we performed double treatments to promote or block FGF and BMP signaling for 6 h. Under these conditions, the dual treatment of FGF8 and BMP4 promoted Dkk expression inducing cell death closer to the BMP bead (Figure 6A). Nevertheless, if FGF signaling is inhibited in the presence of BMP4, Dkk is still expressed (Figure 6B). In contrast, if BMP signaling is blocked and FGF8 is present, the expression of Dkk

![Figure 6](https://example.com/figure6.png)

**FIGURE 6** | The last step to induce cell death is the inhibition of FGF signaling. Double treatments were done to promote or block FGF and BMP signaling for 6 h. Under these conditions, lysotracker staining and Dkk gene expression were evaluated. (A) Double FGF8 and BMP4 treatment promoted Dkk expression, and cell death was observed closer to the BMP bead. (B) Inhibiting FGF signaling in the presence of BMP4, Dkk was still expressed. (C) Treatment with NOGGIN and in the presence of FGF8 significantly inhibited Dkk expression, but it slightly diminished the area of cell death. (D) Double treatment inhibiting FGF and BMP signaling inhibited Dkk gene expression, but no cell death. (E) Fgf8 gene expression is inhibited in response to DKK protein. Notice that the blue line of the right limb (DKK treatment)—representing the expression of Fgf8 from the limb posterior margin to the anterior limb margin—is shorter than the contralateral limb.
was inhibited, and the area of cell death was slightly diminished (Figure 6C). The double blockade of FGF and BMP signaling inhibited Dkk gene expression. Under these conditions, cell death is still induced (Figure 6D). Finally, we evaluated the expression of Fgf8 in the AER in the response of DKK protein that inhibited Fgf8 (Figure 6E). These results suggest that FGF signaling must be inhibited in the ANZ and is probably the last step in the molecular cascade to trigger PCD by BMP signaling.

**Dynamic of Gene Expression in Presumptive ANZ and ANZ**

Once it was established that the onset of cell death requires the negative loop triggered by FGF signaling, a comparison of an expression profile of genes related to the maintenance of the undifferentiated stage and the commitment to cell death was performed (Figure 7). The tissue of limb primordia from 22 HH and the ANZ from 25 HH were dissected in two regions: 1) the mesodermal cells of the anterior margin at stage 22 HH (here called presumptive ANZ) or the ANZ at stage 25 HH, and 2) cells from the undifferentiated zone from both developing stages. It allowed us to clearly distinguish the location of genes expressed before establishing ANZ (stage 22 HH) and once established the ANZ (stage 25 HH). In both stages, the profile expression of this zone was compared with its respective undifferentiated zone.

Regarding genes related to the FGF signaling (Figures 7A,B), results showed that Fgf10 expression was lower in the presumptive ANZ and ANZ than in the undifferentiated zone in both developing stages. It tends to be upregulated in the presumptive ANZ than in the undifferentiated zone, but this study found no difference at both stages (Figure 7A). In the presumptive ANZ, Fgfr2 expression was lower than in the undifferentiated zone, but at stage 25HH, the expression levels were similar in the ANZ and the undifferentiated zone (Figure 7A). However, comparing the levels of fgf10, Fgfr1 and Fgfr2 expressed between the presumptive ANZ at 22 HH and ANZ at 25 HH, the Fgf10 expression did not
show significant changes (Figure 7B). In contrast, the expression levels of both receptors increased at stage 25HH than at stage 22HH, although only Fgfr2 increased significatively (Figure 7A). Next, the expression of Mkp3 and Churchill was evaluated, two target genes of FGF signaling (Kawakami et al., 2003; Sheng et al., 2003). In the presumptive ANZ and ANZ, the expression levels of Mkp3 decrease compared to the undifferentiated zone, while Churchill has no significant changes (Figure 7A). Moreover, comparing the expression levels of both genes, they had a higher level in the ANZ than in the presumptive ANZ (Figure 7B). These data showed an interesting dynamic of FGF signaling.

The next group of genes analyzed was Bmp4, Bmp7, Bambi, and those regulated by BMP signaling (Figures 7C,D). At stage 22HH, the Bmp4 expression is higher in the presumptive ANZ than in the undifferentiated zone. In contrast, the Bmp7 showed lower expression while Bambi did not show significant differences (Figure 7C). At stage 25HH, the expression of these genes in the ANZ was like the observed at stage 22HH (Figure 7D). We also evaluated the expression of Msx2 and Id2, two genes that are regulated by BMP signaling. In the presumptive ANZ, the expression level of Msx2 was lower than the undifferentiated zone, whereas Id2 did not show changes in both regions. In contrast, the expression of Msx2 was elevated in the ANZ regarding the undifferentiated zone (Figure 7C). Id2 expression levels increased at stage 25HH in the ANZ compared with the undifferentiated zone, but at stage 22HH, no differences were found (Figure 7C). Comparing the presumptive ANZ with the ANZ, Msx2 and Id2 presented higher expression levels in the ANZ (Figure 7D). In contrast, the levels of Bmp4 and Bmp7 were lower, while Bambi is similar in both developing stages (Figure 7D).

The next group of genes evaluated is either regulated by BMP and FGF signaling or both (Figures 7E,F). Mario is a gene associated with the formation of digit 2, and it is induced by FGF and inhibited by BMP (Amano and Tamura, 2005). Snail is a transcription factor related to areas of undifferentiated mesenchyme and cell death; BMP and FGF signaling induce both genes (Ros et al., 1997; Montero et al., 2001). Results showed that the expression levels of both genes in the presumptive ANZ and ANZ are higher than in the undifferentiated zone (Figure 7E). Comparing the expression levels between the presumptive ANZ and ANZ, it was found that Snail expression is higher than Mario (Figure 7F).

**DISCUSSION**

The AER is a signaling center where FGF, WNT, and BMP signaling pathways play an essential role in controlling cell proliferation, cell survival, and cell differentiation (Fernández-Teran and Ros, 2008; Ten Berge et al., 2008; Mariani et al., 2017). During sculpturing of the limb, mesodermal cells underneath AER in the anterior margin of the limb undergo cell death giving rise to the ANZ that appears gradually, in coordination with the progressive loss of the AER (Todt and Fallon, 1987; Fernandez-Teran et al., 2006; Fernandez-Teran and Ros, 2008).

The intricate regulatory network between BMP, FGF, and WNT signaling that controls cell death in interdigital tissue is well known (Ganan et al., 1996; Pizette and Niswander, 1999; Danopoulos et al., 2013; Haro et al., 2014; Jin et al., 2018). However, the first events triggered by this regulatory network to initiate cell death in the anterior limb undergoing ANZ formation are not well established. The present study aimed to elucidate the earliest events triggered by the regulatory network of FGF, WNT, and BMP signaling in the control of cell death to induce the ANZ formation in the anterior margin of the limb. Previously, it was reported that a short pulse of BMP is sufficient to trigger cell death in the anterior margin of the limb (Abarca-Buis et al., 2011). An interval of two to 8 h is sufficient to induce cell death after the inhibition of WNT or FGF or the activation of BMP signaling. Besides, cell death induction is coordinated with the regulation of Dkk, Fgf8, and Bmp4 expression. Inhibition of FGF signaling inhibited Dkk expression after 2 h of treatment, demonstrating that FGF is necessary to induce Dkk gene expression. In contrast, DKK treatment-induced cell death after 8 h. Likewise, DKK and BMP4 inhibit Fgf8 expression; thus, it is possible to postulate that the negative feedback loop between Bmp4, Fgf8, and Dkk controls the onset of cell death in the ANZ.

It has been demonstrated that the expression of Fgf8 in the anterior AER is not redundant with other Fgf genes expressed in posterior AER (Moon and Capecchi, 2000; Moon et al., 2000; Delgado et al., 2008). Massive cell death in the anterior margin results from BMP activation (Yokouchi et al., 1996). The mutant mouse for Bmpr1a demonstrates that this receptor mediates BMP signaling in controlling Fgf8 expression (Pajni-Underwood et al., 2007). Interestingly in the posterior region of the limb deprived of SHH, signaling massive cell death occurs concomitantly with up-regulation of Bmp4 (Sanz-Ezquerro and Tickle, 2000). The absence of SHH signaling increases the repressor form of GlI3 (GLI3R), which regulates Bmp expression as observed in the anterior margin; GLI3R is abundant and correlates with an increase in Bmp4 expression (Bastida et al., 2004).

In this work, comparing the expression profile of Fgfr1 and Fgfr2 between the presumptive ANZ with the undifferentiated region at stage 22 HH showed differential expression. Fgfr1 expression is higher than Fgfr2 in the presumptive ANZ. In addition, the lower levels of expression of Churchill or Mkp3 that are FGF signaling targets might suggest that minimal amounts of Fgfr1 or Fgfr2 are enough to avoid cell death. Furthermore, FGF signaling may be active at low levels before ANZ formation. However, the levels of Bmp4 might be the result of a regulation of GLI3R (Bastida et al., 2004), and it is possible that although higher expression of Id2 and Msx2 together with the low level of Mkp3 expression observed in the anterior margin of the limb might be not sufficient to induce cell death. Thus, the balance of FGF and BMP signaling may favor FGF signaling.

As development progress, the ANZ is established. At stage 25HH, the expression levels of Fgfr1 and Fgfr2 and Churchill increase. However, the Mkp3 level remains slightly higher in the well-established ANZ than in the undifferentiated region. Interestingly, levels of Churchill are higher, which might indicate that FGF signaling is being inhibited (Kok et al., 2010). Other genes such as Mario and Wif (data not shown) are higher expressed in ANZ, BMP, whereas FGF signaling induces Mario. It has been involved in the boundary between non-digit digit two in the chick embryo (Amano and
Tamura, 2005), whereas Wif is an inhibitor of WNT/β catenin. Wif together with DKK may inhibit WNT signaling and, consequently, FGF signaling, promoting cell death. The levels of expression of Bmp7 and Bambi in both stages are lower than the undifferentiated zone indicating this minor participation in establishing ANZ. Finally, comparing gene expression levels between the presumptive ANZ at stage 22HH and the ANZ of 25HH demonstrates that FGF and BMP signaling is more active in stage 25HH when the ANZ is well-established. These data suggest that the control of FGF and BMP signaling is necessary to regulate cell death.

Based on data obtained in the present study and from the literature, we propose the following model to explain the onset of cell death in the anterior margin of the limb to give rise to ANZ (Figure 8). First, it is known that WNT3A mediated by Sp6/Sp8 induces the Fgf8 expression in the AER (Haro et al., 2014). On the other hand, BMP signaling inhibited Fgf8 expression; meanwhile, DKK inhibits the function of WNT/β catenin signaling, and consequently, the expression of Fgf8 is inhibited. In the presumptive ANZ, the high levels of FGF8 from AER are necessary for cell survival, but FGF8 is also required for promoting cell death regulating Bmp4 and Dkk expression. The establishment of ANZ occurs when the levels of FGF signaling are reduced and BMP signaling increases leading to the inhibition of Fgf8 in the AER inducing cell death. Fgf1, Fgf2, Mkp3, Churchill, Mox2, Id2, and Bambi are expressed differentially in the presumptive ANZ and the established ANZ.

**DATA AVAILABILITY STATEMENT**

The original contributions presented in the study are included in the article/Supplementary Material, further inquiries can be directed to the corresponding author.

**AUTHOR CONTRIBUTIONS**

JC-M, MED-H, and JCM-L conceived and designed the experiments. MED-H, CG-H, KC-S, MB, JCM-L, and SW performed the experiments. All authors analyzed the data. JC-M and JCM-L wrote the final version of the manuscript. The manuscript was revised and approved by all authors.

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