Myoneurin regulates BMP signaling by competing with Ppm1a for Smad binding

Highlights

- mynn gene is essential for pharyngeal cartilage development
- mynn is required for the proliferation, differentiation, and survival of the CNCCs
- Mynn has an evolutionarily conserved function in supporting BMP signal
- Mynn maintains BMP signal activity by competing with Ppm1a for Smad binding
Myoneurin regulates BMP signaling by competing with Ppm1a for Smad binding

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SUMMARY
A delicate balance of BMP activity is critical for tissue formation and organogenesis. However, the mechanical molecular details in ensuring the proper duration and intensity of BMP signaling have yet to be fully elucidated. Here, we identified a zebrafish mutant with a disrupted gene encoding for the BTB/POZ and zinc finger protein myoneurin (Mynn). mynn$^{-/-}$ mutants exhibited severe loss of pharyngeal cartilage elements, owing to poor proliferation, blocked differentiation, and low viability of cranial neural crest cells. Depletion of mynn in both zebrafish embryos and mammalian cells led to a reduction of the BMP signal activity. Mechanistically, Mynn interacts with Smad proteins in the nucleus, thereby disrupting the association between Smad protein and the phosphatase Ppm1a. Ultimately, this interaction prevents Smad dephosphorylation. More broadly, our findings may provide a new strategy to balance BMP signal activity via competitive binding of Mynn and Ppm1a to Smad proteins during pharyngeal cartilage formation.

INTRODUCTION
Craniofacial malformations are common congenital birth defects that impact the development of the head, face, and neck. Globally, they account for approximately three-fourths of all human birth defects (Cordero et al., 2011). Aberrant cartilage development is one of the primary causes of craniofacial anomalies (Hamerman, 1989). The neurocranium, which is composed of both the upper and front portions of the skull, arises from both the cranial neural crest cells (CNCCs) and mesoderm. Comparatively, the pharyngeal skeleton—which includes the jaw and branchial arches—is derived exclusively from CNCCs (Yelick and Schilling, 2002). CNCCs originate from the boundary between the epidermal and neural territories during the process of neurulation. Subsequently, these cells undergo an epithelial-to-mesenchymal transition, after which they collectively migrate from the midbrain and hindbrain to pharyngeal arches (Donoghue et al., 2008; Trainor and Nieto, 2003). Once they have reached their destinations, the CNCCs proliferate and differentiate into chondrocytes, which produce a large amount of extracellular matrix consisting of type II collagen (Col2) (Hall and Miyake, 1995).

These dynamic processes that occur during pharyngeal cartilage development are controlled by a diverse set of interacting signals such as BMP, Wnt, and FGF pathways (Crump et al., 2004; LaBonne and Bronner-Fraser, 1999; Nie et al., 2006). In particular, previous studies have demonstrated that BMP signaling is essential for the induction and proper migration of CNCCs to facial primordia (Kanzler et al., 2000; Liem et al., 1995; Tribulo et al., 2003; Wilson et al., 1997). Recent reports from our laboratory have provided evidence that balanced BMP signaling is achieved by the interaction between BMP ligands and their antagonist Noggin3. Moreover, this balanced interaction is necessary to ensure proper proliferation, differentiation, and survival of postmigratory CNCCs (Li et al., 2018; Ning et al., 2013). However, further studies are needed to identify additional genes involved in controlling the delicate balance of BMP signaling during cartilage development.

The BTB/POZ and zinc finger (BTB/POZ-ZF) protein family is classified according to the presence of an N-terminal POZ domain and a C-terminal C2H2 ZF motif. This family comprises a diverse group of transcription factors that either activate or suppress the transcription of distinct genes (Kelly and Daniel, 2006). Various vertebrate POZ-ZF proteins have been isolated and characterized; many are linked either directly or indirectly to human cancers (e.g. B cell lymphoma 6 (Bcl-6), promyelocytic leukemia zinc finger (PLZF),...
Myoneurin (Mynn) gene encodes for a protein with features of a BTB/POZ-ZF protein and was initially cloned during a screening of the human λgt11 testicular library (Alliel et al., 2000). Murine Mynn protein is highly homologous to its human ortholog. Moreover, Mynn transcripts have been identified in both mouse embryos and adult tissues, including cerebellum, skeletal muscle, neuromuscular system, testis, heart, brain, and liver (Alliel et al., 2000; Cifuentes-Diaz et al., 2004). Because its expression is strongly elevated through increased gene copy, Mynn has been identified as one of the potential drivers in ovarian cancer (Ramakrishna et al., 2010). Moreover, genotyping analysis in patients diagnosed with bladder cancer revealed a strong cumulative association between a single nucleotide polymorphism in Mynn and tumorigenesis (Polat et al., 2019). Collectively, these observations suggest a potential role for Mynn throughout development and adulthood. Despite this, the developmental and physiological roles of Mynn have been largely unknown and unexplored.

In this study, we identified a zebrafish mynn mutant line using a Tol2 transposon-mediated gene trapping approach. Homozygous mynn mutant embryos showed multiple developmental defects, including pericardial edema, smaller heads and eyes, curved tails, rough skins, and a near-complete loss of the pharyngeal cartilages. In particular, the cartilage defects are resulted from impaired proliferation, differentiation, and maintenance of CNCCs. We found that the depletion of mynn in embryos and mammalian cells led to a significant decrease in BMP signaling, which partially account for the observed defects in cartilage development. Moreover, biochemical and functional studies revealed that Mynn interacts with Smad proteins. Importantly, this interaction disrupts their association with Ppm1a, which is a serine/threonine phosphatase. This disruption then supports the phosphorylation status of Smad proteins. Hence, these findings uncovered an unexpected role for Mynn in balancing BMP signal activity through competitive binding to Smad proteins with Ppm1a during pharyngeal cartilage development.

RESULTS

T054 mutants possess a marked defect in pharyngeal cartilage formation

To identify the genes essential for zebrafish embryo development, an insertional mutagenesis screen with a Tol2 transposon-mediated gene trap strategy was used. This approach allowed for the isolation of mutants with various developmental defects in specific tissues and organs (Han et al., 2011; Zhao et al., 2008). The Tol2 transposon-based trap vector TSG contained a promoter-less EGFP reporter, with fish carrying TSG insertions showing specific GFP expression patterns under the control of neighboring endogenous promoters (Zhao et al., 2008). One of the trapped lines, T054, was notable as a proportion of the embryos from heterozygous F1 intercrosses exhibited multiple morphological defects, including pericardial edema, smaller heads and eyes, curved tails, rough skins, and severe jaw malformations from 48 to 96 h postfertilization (hpf) (Figure 1A). Such embryos died by approximately 120 hpf. During T054 embryo development, the expression of EGFP was clearly observed at the one-cell stage, suggesting its maternal origin (Figure S1). Then, EGFP was ubiquitously expressed until 24 hpf, becoming more prominent in the head, lens, otic vesicle, and trunk musculature during the pharyngula period (Figure S1). From 48 to 72 hpf, intensive EGFP was detected in the developing pharyngeal arches (Figure 1B). Under close observation of the pharyngeal region at higher magnification, the pharyngeal cartilages were clearly EGFP-positive (Figure 1B). Moreover, immunostaining experiments showed that in T054 embryos at 72 hpf, the EGFP-positive chondrocytes located in the pharyngeal arches had high levels of the secreted primary cartilage matrix protein Col2 (Figure 1C). Taken together, these results suggest that the gene trapped by Tol2 transposon in the T054 line is expressed in the pharyngeal arches. Critically, the interruption of this locus may be responsible for the observed mutant phenotype.
Interestingly, the offspring of T054 heterozygous fish showed a different expression intensity of the EGFP reporter. In agreement with Mendel’s law of segregation, among the 2095 embryos at 24 hpf from 23 pairwise crosses of heterozygous fish, 24% produced strong EGFP expression (homozygous insertion), 51% expressed moderate EGFP (heterozygous insertion), and 25% were negative for EGFP. Heterozygous fish were mated with wild-type fish to generate F2 families. We observed an identical EGFP expression pattern in approximately 50% of their progeny. Simultaneously, no EGFP expression was detected in the other half, implying that a single locus has been trapped in the T054 embryos. The F2 embryos that had GFP expression exhibited normal morphological features and survived to adulthood. Consistently, Alcian blue staining results revealed that, in comparison to wild-type and heterozygous siblings, T054 homozygous mutants derived from F2 fish lost almost all their pharyngeal skeletal elements (Figure 1D). In addition, the dorsal neurocrania were also absent in the homozygous mutants (Figure 1D). Collectively, these data indicate that the insertion of Tol2 element in the T054 embryos causes an inherited disorder associated with severe defects in the craniofacial skeleton.

The interrupted gene in the T054 line is mynn

To identify the trapped gene in the T054 line, we performed 5’-RACE and thermal asymmetric interlaced PCR (TAIL-PCR). Sequence and bioinformatic analyses revealed that the Tol2 transposon inserted into
The homozygous mutants were sorted out at 24 hpf by their EGFP intensity, and their morphology was observed at 72 hpf. The ratio of embryos with mean G mynn expression was evident in the midbrain-hindbrain boundary, pectoral fin, and pharyngeal arches (Figure 2C).

The zebrafish mynn locus consists of 9 exons and 8 introns and has an open reading frame of 2610 base pairs, which encodes a putative peptide of 808 amino acids (Figure 2A). Since the putative translation start codon of mynn was located in the third exon (Figure 2A), we speculated that the expression of the coding sequence and 3’ untranslated region protein of the mynn gene would be disrupted by the transposon inserted in the mutant line. In support of our hypothesis, we found that the expression of mynn in mynn−/− mutants was either dramatically reduced (Figures 2D and 2E). Moreover, microinjection of an antisense mynn morpholino (mynn MO), which efficiently repressed the translation of mynn-EGFP transcripts, resulted in morphological malformations and cartilage phenotypes mimicking the ones observed in mynn−/− mutants (Figures S2A–S2C). Therefore, these results demonstrate that mynn is the interrupted gene in the T054 line.

To further confirm that the loss of pharyngeal skeleton in mynn−/− mutants is a direct result of mynn deficiency, we examined whether injection of mynn mRNA could rescue the mutant phenotype. As expected, the morphological malformations and cartilage defects were successfully rescued after injection of 200 pg of mynn mRNA into one-cell stage mynn−/− embryos (Figures 2F and 2G). Moreover, overexpression of mynn under the control of sox10 promoter, which is well known to drive a tissue-specific expression in the cranial and trunk neural crest cells (NCCs) (Carney et al., 2006), significantly reduced the cartilage loss in mynn−/− mutants (Figure 2G), indicating a cell-autonomous role for mynn in head skeleton development. Interestingly, most morphological defects in mynn−/− mutants, including smaller heads and eyes, rough skin, and curved tails, were also alleviated by such tissue-specific expression of mynn (Figure S3), further implying a non-cell-autonomous function of NCC-expressed mynn during embryogenesis.

Taken together, the above data provide strong evidence that we identified a functional transposon insertion in mynn gene, which is essential for the development of pharyngeal cartilage in zebrafish.

### Loss of mynn compromises the proliferation and chondrogenic differentiation of CNCCs

The formation of pharyngeal skeletons involves multiple steps, including specification of CNCCs, migration of CNCCs ventrally into the pharyngeal arches, preochondrogenic condensation, and chondrogenic
Figure 3. Reduced cell proliferation and differentiation of CNCCs in mynn−/− embryos
(A) Live imaging of the pharyngeal region of mynn−/− embryos in Tg(fli1:EGFP) background. The pharyngeal arches were numbered. The boxed areas in the left panel were enlarged in the right panels. PA: pharyngeal arch. Scale bars, 20 μm.
(B and C) BrdU incorporation experiments showed reduced proliferating CNCCs in mynn−/− mutants. BrdU-treated mynn−/− mutants with fli1:EGFP expression were harvested at indicated stages and stained with anti-BrdU (red) and anti-GFP (green) antibodies. The pharyngeal regions were observed by confocal microscopy (B). Scale bars, 20 μm. The number of BrdU+ and GFP+ cells in the second pharyngeal arch was calculated from six embryos (C). Error bars indicate ± S.D. **, p < 0.01; ***, p < 0.001 (by Student’s t test).
(D and E) Detection of Col2 proteins in the pharyngeal arches. Wild-type and mynn−/− embryos in Tg(fli1:EGFP) background at 60 (E) or 72 (D) hpf were co-immunostained with anti-GFP (green) and anti-Col2 (red) antibodies. The palatoquadrate (pq), ethmoid plate (ep), and ceratohyal (ch) cartilages were shown with anterior to the left. Scale bars, in panel D, 20 μm; in panel E, 50 μm.
(F) Expression pattern of hand2, sox9a, and col2a1 in mynn−/− embryos and their siblings at indicated stages detected by in situ hybridization.

to uncover the cellular defects responsible for the reduction of pharyngeal cartilage induced by mynn inactivation, in vivo imaging was performed on Tg(fli1:EGFP) embryos, which expresses EGFP in CNCC derivatives (Lawson and Weinstein, 2002; McGurk et al., 2014). We observed that, in mynn−/− mutants at 48 hpf, the EGFP-positive CNCCs aggregated as prechondrogenic condensations. However, these condensations became much smaller (Figure 3A), suggesting that the loss of cartilage in the EGFP-positive CNCCs aggregated as prechondrogenic condensations. However, these condensations

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To uncover the cellular defects responsible for the reduction of pharyngeal cartilage induced by mynn inactivation, in vivo imaging was performed on Tg(fli1:EGFP) embryos, which expresses EGFP in CNCC derivatives (Lawson and Weinstein, 2002; McGurk et al., 2014). We observed that, in mynn−/− mutants at 48 hpf, the EGFP-positive CNCCs aggregated as prechondrogenic condensations. However, these condensations became much smaller (Figure 3A), suggesting that the loss of cartilage in mynn−/− mutants may be due to proliferation defects of CNCCs. To test this hypothesis, we performed bromodeoxyuridine (BrdU) incorporation assays in mynn−/− mutants and their siblings at 36, 48, and 60 hpf. We found that the BrdU signal was not obviously changed in the trunk region of mynn−/− mutants, but distinctly reduced in the eye, brain, and pharynx, where mynn was highly expressed (Figure S5A), suggesting a reduction of proliferating cells in multiple head tissues. To further determine whether the proliferation of CNCCs was affected upon mynn-depletion, BrdU incorporation experiments were performed on embryos with the Tg(fli1:EGFP) transgenic reporter. As expected, depletion of mynn resulted in a notable elimination of proliferating BrdU+ CNCCs (Figures 3B and 3C).

Furthermore, immunostaining analysis indicated that Col2 signal, which was clearly seen in the first arch-derived palatoquadrate (pq) and the second arch-derived ceratohyal (ch) cartilages in control animals at 72 hpf, was almost absent in the arches of mynn−/− mutants (Figure 3D). This indicates a failure of chondrogenic differentiation. However, the presence of a large amount of cellular debris in the pharynx of mynn−/− mutants raised a possibility that the loss of Col2 might be due to a loss of CNCCs (Figure 3D). To confirm the differentiation defects in the mutants, the expression of Col2 protein was further examined at 60 hpf, when the majority of CNCCs in mynn−/− mutant were not broken (Figure 3B). Indeed, there was no detectable expression of Col2 in mynn-depleted CNCCs at this developmental stage (Figure 3E). In addition, an apparent diminution in the expression of chondrogenic marker genes, such as hand2, sox9a, and col2a1, was found in mynn−/− embryos at 48 or 56 hpf (Figure 3F), demonstrating a crucial role of mynn in chondrocyte differentiation.

Collectively, these results explicitly demonstrate that mynn is required for the proliferation and chondrogenic differentiation of CNCCs.

**mynn is required for the survival of chondrogenic cells**

As we have described above, loss of mynn impaired the chondrogenic differentiation of CNCCs (Figures 3D and 3E). In siblings, differentiated CNCCs appeared elongated and were organized with a columnar orientation in the arches. Comparatively, no regular chondrocytes and only abundant cellular debris were observed in the pharynx of mynn−/− mutants (Figure 3D), suggesting that increased cell death was occurring in these structures.

To confirm this finding, apoptotic analysis using a TUNEL assay was conducted in embryos on the Tg(fli1:GFP) transgenic background. Before and at 36 hpf, there were no detectable apoptotic cells in
the pharyngeal region in siblings and mynn<sup>−/−</sup> embryos. However, from 42 to 72 hpf, levels of prechondrogenic and chondrogenic cell apoptosis were significantly elevated in mynn<sup>−/−</sup> mutants when compared with control embryos (Figures 4A and 4B). These results indicate an important role of mynn in cell survival during pharyngeal cartilage development. Besides, mynn deficiency-induced cell apoptosis was ubiquitously detected in the whole embryo and occurred mostly in the eye and brain (Figure S5B), suggesting that mynn is broadly required for cell viability during embryo development.

It is well known that the tumor suppressor p53 plays a primary role in inducing apoptosis in many cell types, including chondrocytes (Ito et al., 2014; Komori, 2016; Ning et al., 2013; Vogelstein and Kinzler, 1992).
Figure 5. Mynn negatively regulates BMP signaling

(A) Expression of dGFP reporter in the pharyngeal region. Mynn<sup>-/-</sup> mutants and their siblings in Tg(BRE:dGFP) background were stained with the indicated antibodies. Nuclei were counterstained with DAPI (blue). Lateral views, anterior to the left. Scale bars, 20 μm.

(B and C) Expression of p-Smad1/5/9 was dramatically reduced in mynn<sup>-/-</sup> mutants. mynn<sup>-/-</sup> embryos and their siblings were harvested at 48 hpf, and then stained with anti-p-Smad1/5/9 antibodies (B). Scale bars, 20 μm. The levels of p-Smad1/5/9 in cell lysis of head region were further analyzed by Western blots (C).

(D) Expression levels of BMP target gene id1 were detected by in situ hybridization. Embryos were lateral views with anterior to the left.
produced by excessive BMP signaling (Figure S6A), suggesting that Mynn is not sufficient to activate BMP signaling, but rather maintains it.

pharyngeal arch skeletons of embryos lacking the mynn gene (Figure 5A), suggesting a requirement of cartilage development. To characterize the temporal and spatial differences of BMP signal activity between mynn mutants phenocopied most of the abnormal behaviors of chondrogenic progenitors induced by BMP signaling. To verify this, we generated double mutants by crossing mynn mutants. Moreover, the expression of casmad1 mRNA at the one-cell stage and then immunostained with anti-Col2 antibody at 72 hpf. pq, palatoquadrate; ch, ceratohyal. Scale bars, 20 μm.

(G) Alcian blue staining on embryos from different genotypes. Note that the missing cartilages in mynn-/- mutants could be rescued by specific overexpression of casmad1 in sox10+ NCs. Scale bars, 200 μm.

(H) The effectiveness of Mynn shRNAs. HEK293T cells were transfected with the indicated shRNA plasmids and harvested 48 h after transfection for Western blot analyses.

(i and J) HEK293 cells transfected with indicated shRNA plasmids were treated with TGF-β1 (5 ng/mL) for 2 h (i) or BMP4 (20 ng/mL) for 4 h (J), and then collected for Western blots with the indicated antibodies.

(K) Hep3B cells transfected with plasmids expressing indicated shRNAs and GFP proteins were treated with or without BMP4 for 4 h. The expression levels of p-Smad1/5/9 were determined with immunostaining. As indicated with white arrowheads, the expression of p-Smad1/5/9 was obviously decreased in the cells expressing Mynn shRNAs. Scale bars, 20 μm.

identify the mechanism underlying the effect of mynn deficiency on the apoptosis of chondrogenic cells, we performed WISH experiments to examine whether p53 signaling was elevated in mynn-/- mutants. A drastic increase of p53 expression was observed in mynn-/- embryos at 48 hpf (Figure 4C). Interestingly, the elevated p53 transcripts were mainly restricted to the eye, brain, and pharyngeal region. Furthermore, when compared with control groups, gadd45sa expression—a conventional downstream gene of p53—was concomitantly increased in mynn-depleted embryos (Figure 4C). The upregulation of these pro-apoptotic genes was further validated by quantitative RT-PCR analysis (Figure 4D). These data imply that the p53 apoptotic pathway may be involved in mynn deficiency-induced apoptosis.

Finally, we tested whether p53 inactivation could rescue the cell survival defects observed in mynn-/- mutants. To verify this, we generated double mutants by crossing mynn-/- fish with p53-/- mutants. In the resulting mynn-/-:p53-/- embryos, there was a clear decrease in apoptotic cells in the pharyngeal region. However, these apoptotic cells were not completely eliminated (Figures 4E–4G). Furthermore, depletion of p53 could not recover the loss of pharyngeal cartilages in mynn-/- mutants, indicating mynn is doing more than just cell survival, consistent with its crucial role in the proliferation and differentiation of NCs. Taken together, these results strongly suggest that the absence of mynn induces chondrogenic cell apoptosis. Moreover, this effect is partially caused by activation of the p53 pathway.

**mynn inactivation attenuates BMP signaling**

Our previous study found that the inhibition of BMP signaling resulted in poor proliferation, impaired differentiation, and unsustainable survival of chondrogenic progenitors (Ning et al., 2013). Since mynn-/- mutants phenocopied most of the abnormal behaviors of chondrogenic progenitors induced by BMP signaling deficiency, we next asked whether mynn functions in regulating BMP signaling during pharyngeal cartilage development. To characterize the temporal and spatial differences of BMP signal activity between siblings and mynn-/- mutants, we used a BMP signal reporter transgenic line BRE:dGFP. This transgenic line expressed a destabilized form of GFP (dGFP) driven by the BMP response element (BRE) derived from the mouse promoter Id1 (Collery and Link, 2011). When compared with control animals, mynn-/- mutants showed much lower levels of GFP fluorescence intensity in the ventral arch domains at 36 and 48 hpf (Figure 5A), suggesting a requirement of mynn in BMP signal maintenance in NCs.

We further examined the spatial distribution of phosphorylated Smad1/5/9 (p-Smad1/5/9)—intracellular effectors of BMP signaling—using immunostaining in mynn-/- mutants and their siblings. Consistent with previous reports (Alexander et al., 2011; Ning et al., 2013), in control embryos, a gradient of p-Smad1/5/9 expression was detected in the pharyngeal region. More specifically, high levels were observed ventrally, while low levels were observed dorsally (Figure 5B). In contrast, the phosphorylation of Smad proteins was seriously compromised in the absence of mynn (Figure 5B). Corroborating these findings, Western blotting also showed a marked reduction of p-Smad1/5/9 levels in the head tissues of mynn-/- mutants (Figure 5C). Moreover, the expression of id1—the classic target gene of BMP—was notably reduced in the pharyngeal arch skeletons of embryos lacking the mynn gene (Figure 5D). Taken together, these results indicate that mynn is indispensable for BMP signal activity in pharyngeal chondrogenic progenitors. Surprisingly, injection of mynn mRNA into wild-type embryos did not induce a ventralized phenotype usually produced by excessive BMP signaling (Figure 5E), suggesting that Mynn is not sufficient to activate BMP signaling, but rather maintains it.
We next want to know if an elevated BMP signal can rescue the cartilage defects in *mynn*-/- embryos. To test this, mRNA encoding a constitutively active Smad1 protein (caSmad1)—the phosphomimetic form of Smad1, in which the C-terminal SVS motif was mutated to an aspartic acid-valine-aspartic acid (DVD) sequence—was injected into *mynn*-/- embryos at the one-cell stage (Tsukamoto et al., 2014). To avoid possible effects on embryonic dorsoventral patterning, a small dose of *casmad1* mRNA (10 pg) was injected into each embryo. We found that such injection did not result in obvious ventralized phenotypes, but completely rescued the proliferation defect in *mynn*-defective CNCCs (Figures 5E and S6B). However, although overexpression of *casmad1* in *mynn*-/- mutants partially alleviated the impaired chondrogenic differentiation at 60 hpf (Figure 5F), it did not improve the defects in cartilage formation at 96 hpf. Meanwhile, the morphological malformations in *mynn*-/- mutants were not eliminated in the rescue experiments (Figure S6B). The inefficiency of such rescue experiments might be due to the much lower dose of *casmad1* mRNA injection and the degradation of injected mRNA at later stages. To overcome these drawbacks, *casmad1* was specifically overexpressed in NCCs of *mynn*-/- mutants using the sox10 promoter. As shown in Figure 5G, the NCC-specific overexpressed *casmad1* almost perfectly restored the loss of pharyngeal cartilages in *mynn*-/- mutant embryos. Based on these observations, we proposed that Mynn regulates pharyngeal cartilage development via upholding BMP signaling. In addition, the morphological abnormalities except pericardial edema in *mynn*-/- mutant were eliminated by NCC-specific overexpression of *casmad1* (Figure S6C), suggesting the non-cell-autonomous role of *mynn* also depends on its function in maintaining BMP signal activity.

To next clarify whether Mynn has a role in maintaining BMP signaling in mammalian cells, we designed two DNA constructs expressing independent short hairpin RNAs (shRNAs, termed Mynn shRNA1 and Mynn shRNA2) that targeted human Mynn. Both constructs were able to effectively inhibit Mynn expression (Figure 5H). As shown in Figures 5I and 5J, depletion of Mynn in human embryonic kidney 293 (HEK293) cells had no effect on TGF-β1-induced Smad2 phosphorylation, however, inhibited the phosphorylation of Smad1/5/9 triggered by BMP4, a well-known growth factor that induces the nuclear translocation of Smad proteins (Derynck and Budi, 2019). These results imply a specific function of Mynn in BMP signal transduction. Furthermore, the nuclear accumulation of p-Smad1/5/9, which was induced by BMP4 ligands, was significantly inhibited in Mynn-depleted hepatocellular carcinoma Hep3B cells (Figure 5K), indicating an evolutionarily conserved function of Mynn in BMP/Smad signal regulation.

**Mynn physically interacts with Smad protein**

Mynn is a nuclear protein that functions either directly or indirectly to modulate gene expression (Alliel et al., 2000; Kelly and Daniel, 2006). Given this, we hypothesized that Mynn regulates BMP signaling by interacting with Smad proteins, which constitutively shuttle between the cytoplasm and nucleus (Derynck and Budi, 2019). To test this, Flag-tagged Mynn was coexpressed with Myc-tagged Smad1 in HEK293 cells. After, co-immunoprecipitation (co-IP) experiments were performed using an anti-Flag antibody. Results indicated that Smad1 was co-precipitated with Mynn (Figure 6A). Importantly, we also found that overexpressed Myc-Smad1 was able to interact with endogenous Mynn (Figure 6B).

In general, the phosphorylated Smad proteins migrate more slowly than un-phosphorylated ones. We noted that immunoblotting with anti-Myc antibody revealed two bands that corresponded to a phosphorylated and an un-phosphorylated form of Smad1 in proteins co-precipitated with Mynn, although there was only a single anti-Myc band in the input lysates (Figure 6A). These observations may imply that Mynn prefers to interact with the phosphorylated form of Smad1, resulting in an efficient enrichment of phosphorylated Smad1 in the immunoprecipitated proteins. To address this, the interaction between Mynn and Smad1 was further examined in live cells by a bimolecular fluorescence complementation (BiFC) assay as previously described (Wei et al., 2017). The N-terminal half of yellow fluorescent protein (YFP) was fused to Mynn (YN-Mynn) while the C-terminal half of YFP was fused to Smad1 (YC-Smad1). Immunofluorescence staining revealed that YN-Mynn was localized to the nucleus; comparatively, YC-Smad1 was mainly distributed in the cytoplasm (Figure 6C). Interestingly, the reconstituted YFP signal was specifically observed in the nuclei (Figure 6D). Given this, it is likely that Mynn associates with nuclear Smad1 proteins, most of which are phosphorylated. Consistent with this idea, the association between Mynn and Smad1 was enhanced in the presence of BMP4 (Figure 6E).

Smads are structurally similar proteins containing two conserved polypeptide segments, the N-terminal MH1 and carboxyl-terminal MH2 domains linked by a less conserved linker region (Feng and Derynck, 2005). To determine which domain of the Smad protein is responsible for binding with Mynn, various
truncated mutants of Smad1 were constructed (Figure 6F). Domain mapping revealed that only the MH2 domain—neither the MH1 domain nor the linker region—interacted with Mynn (Figure 6G). Thus, these results indicate that Mynn binds to the MH2 domain of Smad1.

Mynn upholds BMP activity by competitively binding to the Smad protein with Ppm1a

The results presented above suggested that Mynn interacts with Smad proteins and maintains Smad phosphorylation in the nucleus. The next interesting question is how Mynn regulates the level of Smad phosphorylation. The receptor-regulated Smads undergo continuous nucleocytoplasmic shuttling, and their export from the nucleus is controlled by a mechanism involving dephosphorylation (Lin et al., 2006; Xu et al., 2002). In particular, it has been suggested that Ppm1a, a serine/threonine phosphatase, binds to both the MH1 and MH2 domains of Smad1 and dephosphorylates Smad1 in the nucleus (Duan et al., 2006). Given that Mynn and Ppm1a each bind to the MH2 domain of Smad1, Mynn might compete with Ppm1a for this binding motif on Smad1.

This hypothesis prompted us to use co-IP experiments to investigate the effect of Mynn on the binding of Ppm1a to Smad1. Indeed, in HEK293T cells, we found that the presence of Mynn dramatically weakened the interaction between Ppm1a and Smad1 (Figure 7A). Next, we examined the effect of Mynn on Ppm1a-induced dephosphorylation of Smad1. Ectopic expression of ALK1 (Q233D), a constitutively active form of BMP type I receptor (caALK1), led to a dramatic elevation of Smad1 phosphorylation. This phosphorylation was almost abolished by coexpression of Ppm1a (Figure 7B). It is noteworthy that, in the presence of Mynn, the Ppm1a-induced elimination of Smad1 phosphorylation was obviously reduced (Figure 7B).
Figure 7. Mynn promotes BMP signaling through competing with Ppm1a for Smad binding

(A) Presence of Mynn attenuates the interaction of Ppm1a and Smad1. HEK293T cells were co-transfected with Myc-Smad1 and HA-Ppm1a together with or without Flag-Mynn, and then harvested for immunoprecipitation.

(B) Mynn weakens the Ppm1a-mediated dephosphorylation of p-Smad1/5/9. HEK293T cells were transfected with the indicated constructs. CaALK1 is a constitutively active form of BMP type 1 receptor ALK1 (Q233D). The expression levels of p-Smad1/5/9 and total Smad1 were detected by Western blots with appropriate antibodies.

(C) Overexpression of mynn can restore the decrease of BMP signal activity caused by excessive Ppm1a. Tg(BRE:dGFP) embryos were injected with indicated mRNAs at the one-cell stage and subjected to immunostaining for GFP (green) at 48 hpf. Scale bars, 50 μm. Injection dosages: mCherry mRNA, 400 pg; ppm1a mRNA, 400 pg; mynn mRNA, 300 pg.

(D and E) Ppm1a-induced pharyngeal cartilage defects were compromised by overexpression of mynn. Wild-type embryos were injected with 400 pg ppm1a mRNA together with or without 300 pg mynn mRNA at the one-cell stage, and stained at 96 hpf for cartilage with Alcian blue. Representative pictures of different classes of defects in pharyngeal cartilages were shown in (D). The percentage of embryos with different degrees of cartilage defects was calculated (E). Scale bars, 200 μm.
To demonstrate the effect of Mynn on Ppm1a-mediated BMP signal suppression during pharyngeal skeleton formation, Tg(BRE:dGFP) embryos were injected with ppm1a mRNA together with or without mynn mRNA. It has been suggested that Ppm1a can also dephosphorylate Smad2/3 to inhibit Nodal signaling during zebrafish early embryogenesis (Lin et al., 2006). Indeed, a portion of embryos injected with ppm1a mRNA exhibited notochord defects, including discontinuous anterior notochord and thinner posterior notochord (Figures S7A and S7B), which are usually observed in Nodal-deficient mutants (Lin et al., 2006). However, overexpression of mynn in embryos-injected ppm1a mRNA could not obviously improve these morphological defects (Figures S7A and S7B), further confirming our finding that Mynn has a specific role in helping to maintain BMP but not TGF-β/BMP signaling in the pharyngeal region and disrupted cartilage formation (Figures 7C–7E). Notably, both the reduction of BMP activity and the defects of pharyngeal cartilages in Ppm1a-overexpressed embryos were effectively relieved after Mynn coexpression (Figures 7C–7E).

To further explore the role of mynn in pharyngeal cartilage development, we hypothesized that suppression of ppm1a expression in mynn−/− mutants would rescue the CNCC proliferation and differentiation defects. To verify this, we designed a MO that could effectively block ppm1a translation (Figure S7C). We found that in mynn−/− mutant embryos injected with ppm1a MO, the proliferation ability of CNCCs was well recovered and the chondrogenic differentiation was partially rescued (Figures 7F–7H).

Based on the totality of these data, we present a model in which Mynn competes with Ppm1a for the MH2 domain on Smad proteins to uphold BMP signaling during cartilage formation. Mynn absence leads to more Ppm1a proteins associating with and subsequently dephosphorylating Smads. This then turns off BMP signaling, ultimately, resulting in impaired formation of pharyngeal cartilages.

**DISCUSSION**

Members of the BTB/POZ-ZF family are involved in a multitude of developmental events in invertebrates and vertebrates (Adhikary et al., 2003; Barna et al., 2000; Giniger et al., 1994; Pengue et al., 1994; Piazza et al., 2004; Xiong and Montell, 1993). Mynn is a novel member of the BTB/POZ-ZF family and is expressed in various tissues in human and mice, including cerebellum, testis, ovary, placenta, heart, brain, liver, and muscle (Alliel et al., 2000). Recently, Mynn has been suggested to be involved in tumorigenesis (Polat et al., 2019; Ramakrishna et al., 2010). However, its physiological functions during embryonic development remain poorly understood.

In this study, we identified a zebrafish line in which the mynn locus was interrupted by a Tol2 transposon. Zebrafish mynn transcripts are maternally deposited and ubiquitously expressed during the gastrulation stage and segmentation period. From 24 hpf, mynn expression is mainly restricted to the central nervous system, muscles, and pharyngeal arches, suggesting a conserved expression pattern across vertebrate species. mynn mutant embryos show a normal external morphology at early developmental stages, but display a variety of abnormalities at later stages, such as smaller head and eyes, coarse and rough skin, curved tail, and missing pharyngeal skeletons. However, because all of mynn zygotic mutants died at approximately 120 hpf, we were unable to generate maternal-zygotic mynn mutants. The phenotype we observed in mynn mutants was only an indication of loss-of-function effect of zygotic mynn. As such, it is impossible to ignore the possibility that mynn has unknown functions at early stages of embryogenesis, which may be masked by the persistence of maternal transcripts and proteins.

Because the head skeletons were almost completely missing in mynn mutants, we focused on the role of mynn in the development of pharyngeal cartilages. Our study reveals an essential role for mynn in CNCC proliferation, differentiation, and survival. Although expressed in multiple head tissues, mynn plays...
a particularly important and cell-autonomous role during pharyngeal cartilage development. This was evidenced when re-supplying Mynn to NCCs restored cartilage formation in mynn mutants. Interestingly, such NCC-specific expression of mynn in the mutants also alleviated most morphological defects, including smaller heads and eyes, rough skin, and curved tails. Because the cranial and trunk NCCs and their derived mesenchymal cells are required to support the growth and development of various kinds of tissues, such as thymic epithelium, pharyngeal arch artery, and hematopoietic stem cells (Blackburn and Manley, 2004; Damm and Clements, 2017; Ohnemus et al., 2002), it is reasonable to hypothesize that NCC-expressed mynn also has a non-cell-autonomous function during embryogenesis.

Embryos depleted for mynn exhibited similar abnormal behaviors of chondrogenic progenitors induced by deficient BMP signaling as we have previously reported (Ning et al., 2013). Indeed, loss of Mynn in zebrafish embryos and mammalian cells resulted in a dramatic reduction in BMP signal activity, indicating a conserved function of Mynn in regulating BMP signaling. Moreover, NCC-specific overexpressed casmad1 not only restored the loss of pharyngeal cartilages, but also recovered most morphological abnormalities in mynn−/− mutant, suggesting both the cell-autonomous and non-cell-autonomous roles of mynn depend on its function in maintaining BMP signal activity. A number of studies have indicated that single nucleotide polymorphisms in Mynn are associated with both the development and progression in colorectal, ovarian, and bladder cancers (Do et al., 2015; Houlston et al., 2010; Lubbe et al., 2012; Polat et al., 2019; Song et al., 2018). Furthermore, alterations in BMP signaling have been implicated in the pathogenesis of several tumor types including prostate, colorectal, osteosarcomas, myelomas, and breast cancers (Blanco Calvo et al., 2009; Sagorny et al., 2012; Wang et al., 2019). Therefore, our findings also provide meaningful clues for exploring the function of Mynn in tumorigenesis.

It has been reported that BMP signaling is negatively regulated by a few phosphatases, including Smad phosphatase Ppm1a (Duan et al., 2006; Lin et al., 2006). In this study, we find that Mynn physically interacts with Smad1 in the nucleus. The Mynn and Ppm1a proteins compete for binding with the Smad proteins, thus balancing BMP signaling for proper cartilage formation. Interestingly, previous work has also suggested that the BTB/POZ zinc finger family member Kaiso directly represses canonical Wnt signaling by associating with LEF/TCF proteins (Park et al., 2005). Given this, BTB/POZ zinc finger proteins are tightly linked to specific intracellular signaling pathways. However, overexpression of casmad1 in mynn−/− mutants by mRNA injection or using sox10 promoter could not rescue the pericardial defect. One possible explanation for this is that, in addition to regulating BMP signaling, Mynn may also play a role as a transcription factor to control gene expression during embryo development.

In summary, our data support the idea that Mynn is crucial for pharyngeal cartilage development by maintaining an adequate level of canonical BMP pathway activity. Mynn relieves Ppm1a-mediated dephosphorylation through competitive binding to Smad proteins. Therefore, our study provides new insight into the regulatory interactions between Mynn and BMP signaling during vertebrate development.

Limitations of the study
This study mainly explored the role for Mynn in balancing BMP signal activity through competitive binding to Smad proteins with Ppm1a during pharyngeal cartilage development. Mynn is a transcription factor that belongs to the BTB/POZ-ZF family, and mynn mutants exhibit various developmental defects in multiple tissues and organs. Future work is necessary to determine whether Mynn has a more widespread effect by acting as a transcription factor during embryo development as well, potentially independent of BMP signaling.

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AUTHOR CONTRIBUTIONS
S.Y. designed, performed the majority of experiments, analyzed the data, and wrote the original draft. G.N. performed the majority of experiments in the revised paper and involved in data generation and interpretation. Y.H. performed in situ hybridization experiments and raised the zebrafish lines. Critical reagents were provided by Y.C. and J.X. J.W. and T. Z. discussed the results and editing the manuscript. Q.W. designed experiments, interpreted data, and revised the manuscript. All authors contributed to manuscript editing and review.

DECLARATION OF INTERESTS
The authors declare no competing interests.

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## STAR METHODS

### KEY RESOURCES TABLE

| REAGENT or RESOURCE | SOURCE | IDENTIFIER |
|---------------------|--------|------------|
| **Antibodies**      |        |            |
| Rabbit Phospho-Smad1 (Ser463/465)/Smad5 (Ser463/465)/Smad9 (Ser465/467) (DSB10) mAb | Cell Signaling Technology | Cat#13820 |
| Rabbit Smad1 Antibody | Cell Signaling Technology | Cat# 9743 |
| Mouse anti-Collagen type II | DSHB | Cat#II-l683 |
| Mouse anti-Zn5 | Zebrafish International Resource Center (ZIRC) | Cat#111605 |
| Monoclonal Anti-BrdU Mouse | Sigma Aldrich | Cat#B-2531 |
| Rabbit GFP Polyclonal Antibody | Thermo Fisher Scientific | Cat#A11122 |
| Mouse Anti-FLAG® M2 Affinity Gel | Sigma Aldrich | Cat#A2220 |
| Rabbit Anti-c-Myc Agarose Affinity Gel | Sigma Aldrich | Cat#A7470 |
| Mouse DYKDDDDK-Tag(3B9) Antibody | Abmart | Cat#M20008M |
| Mouse Anti-Myc-tag mAb | MBU(Medical & Biological Laboratories) | Cat#M047-3 |
| Mouse anti-Myoneurin (JB-17) | Santa Cruz Biotechnology | Cat# 101082 |
| Sheep Anti-Digoxigenin-AP | Roche | Cat# 11093274910 |
| Donkey anti-Rabbit, Alexa Fluor 488 | Thermo Fisher Scientific | Cat# A-21206 |
| Donkey anti-Mouse, Alexa Fluor 594 | Thermo Fisher Scientific | Cat# A-21125 |
| Donkey anti-Rabbit, Alexa Fluor 594 | Thermo Fisher Scientific | Cat# A-21207 |
| Donkey anti-Mouse, Alexa Fluor 647 | Thermo Fisher Scientific | Cat# A-31571 |

### Chemicals, peptides, and recombinant proteins

|                       | Pathway/Target | Cat#   |
|-----------------------|---------------|--------|
| 4',6-Diamidino-2-phenylindole dihydrochloride (DAPI) | Sigma Aldrich | 10236276001 |
| Alcian Blue 8GX | Sigma Aldrich | 05500 |
| 5-bromo-2'-deoxyuridine (BrdU) | Sigma Aldrich | 85002 |
| Methyl cellulose | Sigma Aldrich | M6385 |
| Trypsin | AMRESCO | 0458 |
| Ethyl 3-aminobenzoate methanesulfonate salt (Tricaine) | Sigma Aldrich | 5040 |
| N-Phenyliothiourea (PTU) | Sigma Aldrich | P7629 |
| Lipofectamine 2000 | Thermo Fisher Scientific | 11668019 |
| RNA Polymerase, T7 | Roche | 10881775001 |
| RNA Polymerase, SP6 | Roche | 11487671001 |
| DIG RNA Labeling Mix | Roche | 11277073910 |
| BMP4 | R&D | 314-BP-010 |

### Critical commercial assays

|                       | Pathway/Target | Cat#   |
|-----------------------|---------------|--------|
| FirstChoice® RLM-RACE Kit | Thermo Fisher Scientific | AM1700 |
| RNeasy Mini Kit | Qiagen |  |
| In Situ Cell Death Detection Kit, TMR red | Roche | 12156792910 |
| ReverTra Ace | TOYOBO | TRT-101 |
| Trelief™ SoSo Cloning Kit | Tsingke Biotechnology | TSV-S2 |
| TB Green® Premix Ex Taq™ II (Tli RNaseH Plus) | Takara | RR820A |
| mMESSAGE mMACHINE™ SP6 | Thermo Fisher Scientific | AM1340 |

### Experimental models: Cell lines

|                       | Pathway/Target | Cat#   |
|-----------------------|---------------|--------|
| HEK293T | ATCC | CRL-3216 |
| HEK293 | ATCC | CRL-1573 |

(Continued on next page)
| REAGENT or RESOURCE | SOURCE | IDENTIFIER |
|---------------------|--------|------------|
| Hep3B               | ATCC   | HB-8064    |
| HeLa                | ATCC   | CCL-2      |

Experimental models: Organisms/strains

- Tg(fli1:EGFP)  
  (Lawson and Weinstein, 2002)  
  ZDB-TGCONSTRCT-070117-94
- Tg(BRE:EGFP)  
  (Laux et al., 2011)  
  ZDB-FISH-150901-20390
- tp53M214K  
  (Berghmans et al., 2005)  
  ZDB-ALT-050428-2
- mynnT054  
  This paper  
  N/A

Oligonucleotides

- mynn-qPCR Forward:  
  5'-CAGTGTCCCTCAACATCCC-3'  
  This paper  
  N/A
- mynn-qPCR Reverse:  
  5'-GGAAAGATCATGGAGGGCTG-3'  
  This paper  
  N/A
- gadd45a-qPCR Forward:  
  5'-CTTGCACTGTACCCTTGCA-3'  
  This paper  
  N/A
- gadd45a-qPCR Reverse:  
  5'-CTCATCGCTCTGGAAGGTTG-3'  
  This paper  
  N/A
- p53-qPCR Forward:  
  5'-CTTCTTCAAGCTACATTACGACCTGAGGGAGC-3'  
  This paper  
  N/A
- p53-qPCR Reverse:  
  5'-GCAGGCACCACATCAGACTG-3'  
  This paper  
  N/A
- gapdh-qPCR Forward:  
  5'-AGGCCGGTGCTAGTATGTC-3'  
  This paper  
  N/A
- gapdh-qPCR Reverse:  
  5'-GCAGGCACCACATCAGACTG-3'  
  This paper  
  N/A
- genotyping-P1:  
  5'-CCGTCAAAGCAAAGCGATTTTAATCT-3'  
  This paper  
  N/A
- genotyping-P2:  
  5'-GGCTTATCCAGAAATGCACCTG-3'  
  This paper  
  N/A
- genotyping-P3:  
  5'-CCTCTCTACCACACTCAACTCCACCTG-3'  
  This paper  
  N/A
- mynn MO:  
  5'-CAGGCCGTGTACAGAGCAGC-3'  
  This paper  
  N/A
- ppm1a MO:  
  5'-GGCTTATCCAGAAATGCACCTG-3'  
  This paper  
  N/A
- control MO  
  5'-CCCTTACCTCCACCTCAAATTTAAT-3'  
  Gene tools
- mynn shRNA NO1:  
  5'-GGAATGTGCAGCTGCCCTCTT-3'  
  This paper  
  N/A
- mynn shRNA NO2:  
  5'-GCACACTGGTGAGTACAGC-3'  
  This paper  
  N/A

Recombinant DNA

- TSG  
  (Han et al., 2011)  
  N/A
- pCS2-flag-mynn  
  This paper  
  N/A
- Pcdna3.0-6xmyc-smad1  
  This paper  
  N/A
- YN-mynn  
  This paper  
  N/A
- YC-smad1  
  This paper  
  N/A
- pCS2-myc-smad1-MH1  
  This paper  
  N/A
- pCS2-myc-smad1-linker  
  This paper  
  N/A
- pCS2-myc-smad1-MH2  
  This paper  
  N/A
- pCS2-HA-ppm1a  
  This paper  
  N/A
- pCS2-caAlk1  
  from Tsinghua University  
  N/A
- pCS2-flag-ppm1a  
  (Duan et al., 2006)  
  N/A
- pCS2-casmad1  
  from Tsinghua University  
  N/A
- pSTG-sox10-casmad1-P2A-mCherry  
  This paper  
  N/A

(Continued on next page)
RESOURCE AVAILABILITY

Lead contact
Further information and requests for resources and reagents should be directed to and will be fulfilled by the Lead Contact Qiang Wang (qiangwang@scut.edu.cn (Q.W.)).

Materials availability
Plasmids and animal models generated in this paper will be shared freely upon request to the lead contact.

Data and code availability
- Data: All data reported in this paper will be shared by the lead contact upon request.
- Code: This paper does not report original code.
- Any additional information required to reanalyze the data in this paper is available from the lead contact upon request.

EXPERIMENTAL MODEL AND SUBJECT DETAILS

Zebrafish strains
Zebrafish strains were maintained in standard laboratory conditions. Embryos were obtained from natural zebrafish matings, raised in Holtfreter’s solution at 28.5°C, and staged by morphology as previously described (Kimmel et al., 1995). The following published strains were used in this study: Tg(fli1:EGFP) (Lawson and Weinstein, 2002); Tg(BRE:EGFP) (Laux et al., 2011) and tp53 mutant (Berghmans et al., 2005). Live embryos or adults were anesthetized in fish water containing 0.4% tricaine (ethyl 3-aminobenzoate methanesulfonate salt; Sigma-Aldrich). Our work involving zebrafish embryo collection and analysis was carried out in accordance with and approved through the Animal Care Committee at the Institute of Zoology, Chinese Academy of Sciences (Permission Number: IOZ-13048).

Cell lines
HEK293 (CRL-1573, ATCC), HEK293T (CRL-3216, ATCC), Hep3B (HB-8064, ATCC) and HeLa cell lines (CCL-2, ATCC) were cultured in Dulbecco’s modified Eagle’s medium (DMEM, HyClone) supplemented with 10% fetal bovine serum (FBS, HyClone) and 1% penicillin-streptomycin (HyClone) at 37 °C in a humidified incubator with 5% CO2. Cell transfections were performed by using Lipofectamine 2000 (11668019, Invitrogen) according to the manufacturer’s instructions.

Generation of zebrafish line T054
For transposon-mediated insertional mutagenesis, the transposon vector TSG was constructed as previously described (Han et al., 2011). 50 ng TSG plasmid and 100 pg tol2 mRNA were injected into embryos at the one-cell stage. The injected founder (F0) embryos were raised to adulthood and outcrossed with wild-type fish, and their progeny (F1) were observed for GFP expression pattern under a fluorescence microscope. The GFP-positive embryos were grown up to establish Tol2 insertion mutant lines. Particularly, the T054 line was identified in such pilot experiments.
METHOD DETAILS

Morpholino, mRNA and microinjection
Capped mRNAs were synthesized in vitro for mynn, casmad1, ppm1a, and Tol2 from the corresponding linearized plasmids using the mMessage mMachine kit (Thermo Fisher Scientific, AM1340). mynn MO (5’-CATGGCGTGTAGAGGAAAAACCTCA-3’) and ppm1a MO (5’-GGCTTATCCAGAAATGCACCCATGT-3’) were designed to target the translational start regions, respectively. A non-targeting standard control MO (5’-CCTCTACCTCAGTTACAATTTATA; Gene Tools) was used as a control for unspecific effects. Both mRNA and MO were injected into the yolk of one-cell stage embryos. In the rescue experiments, the recombinant plasmids pSTG-sox10:mynn-P2A-mcherry or pSTG-sox10:casmad1-P2A-mcherry and the Tol2 mRNA were coinjected into fertilized eggs by microinjection.

Alcian blue staining
Embryos at 96 hpf were fixed in 4% paraformaldehyde overnight. Then embryos were washed in distilled water until the embryo was transparent. After staining with Alcian blue staining buffer (0.015% Alcian Blue, 80% ethanol, and 20% acetic acid) overnight at room temperature, the embryos were de-stained in ethanol of gradient concentrations. Next, the embryos were treated with 0.5% trypsin (0458, AMRESCO) in supersaturated borax at room temperature until the tissues were soft enough to dissect. The embryos were then transferred to 1% KOH and dehydrated with a graded series of glycerol solutions.

5’-RACE, genotyping, and quantitative real-time PCR
5’-RACE was performed using the FirstChoice® RLM-RACE Kit (Thermo Fisher Scientific, AM1700) according to the manufacturer’s instructions.

For genotyping, embryos from T054 intercrosses were separated into three classes according to their fluorescence intensity at 24 hpf. Genomic DNA was extracted from single embryos for genotyping by PCR amplification with sequence-specific primers listed in the key resources table.

For quantitative real-time RT-PCR, total RNA of siblings and T054 embryos were extracted from a pool of 30 embryos at indicated stages by RNeasy Mini Kit (Qiagen, 74104). According to the manufacturer’s instructions (TOYOBO, TRT-101), cDNAs were synthesized from 2 μg RNA using oligo-dT primer, and then quantitative real-time PCR was carried out with TB Green® Premix Ex Taq™ II (Takara, RR820A) in the Mx3000P real-time PCR system (Stratagene). Expression level of β-actin was analyzed as an internal control.

Whole-mount in situ hybridization and immunostaining
For whole mount in situ hybridization, Digoxigenin-UTP-labeled RNA probes were synthesized in vitro from linearized DNA templates using the RNA Polymerase T7/Sp6 system (Roche, Cat#10881775001/11487671001) according to the manufacturer’s instructions. Whole-mount in situ hybridizations were performed following the standard procedure.

For immunostaining, embryos were fixed in 4% paraformaldehyde at 4°C for 24 h, and then dehydrated with methanol. The embryos were re-dehydrated through a series of washings with methanol in PBST. Then embryos were permeabilized with proteinase K (10 mg/mL) for 30 to 60 min and incubated in blocking buffer (2% BSA, 5% normal goat serum, 0.1% Tween-20 in PBS) for 1 h. Embryos were stained with the following affinity-purified antibodies: Rabbit anti-P-Smad1/5/9 (1:500; 18320, Cell Signaling Technology); Mouse anti-Collagen type II (1:100; II-116B3, DSHB); Zn5 (1:50; 111605, Zebrafish International Resource Center). Finally, the embryos were mounted in 1% low melting point agarose and imaged using a Nikon A1R+ confocal microscope.

Proliferation and apoptosis analyses
For all BrdU incorporation experiments, embryos were placed in 10 mM BrdU at specific stages for 20 min and then harvested. Incorporated BrdU and GFP were detected using anti-BrdU (1:1,000; B5002, Sigma) and rabbit anti-GFP antibodies (1:1,000, a gift from Dahua Chen Lab) by whole-mount immunostaining. TUNEL assays were performed using In Situ Cell Death Detection Kit, TMR red (12156792910, Roche) according to the manufacturer’s instruction.
Immunoprecipitation and Western blotting
HEK293T cells were transiently transfected with indicated plasmids. 36 h after transfection, cells were harvested and lysed with TNE buffer (150 mM NaCl, 10 mM Tris-HCl (pH7.5), 2 mM EDTA, and 0.5% NonidetP-40) containing protease inhibitors. Immunoprecipitation and Western blots were performed according to standard protocols. For Western blots, affinity-purified anti-Flag (1:5000; M20008M, Abmart), anti-Myc (1:3000; M047-3, MBL), anti-Smad1 (1:1000; 9743, Cell Signaling Technology), anti-Mynn (1:200; SC-101082, Santa Cruz) and anti-P-Smad1/5/9 (1:500; 18320, Cell Signaling Technology) antibodies were used.

RNA interference
Two shRNAs against Mynn were designed (NO.1, 545-565, 5'-GGAATGTGCTGGCCTCTTTA-3', and NO.2, 1182-1202, 5'-GCACACTGTACATGAAACG-3'). Oligonucleotides were chemically synthesized and subcloned into the U6/GFP/Neo shRNA vector by Gene Pharma (Shanghai, China).

BiFC assay
For the BiFC assay, Mynn was fused to the N-terminal half of YFP with restriction endonuclease cleavage sites EcoRI and SalI (YN-Mynn), while Smad1 was fused to the C-terminal half of YFP with restriction endonuclease cleavage sites BglII and ECoRI (YC-Smad1). YN-Mynn and YC-Smad1 were either individually or collectively transfected into HeLa cells. YFP fluorescence was visualized using a Nikon A1R+ confocal microscope at 48 h after transfection.

Quantification and statistical analysis
For GFP- and BrdU/TUNEL-positive cell counts, images were analyzed with Image J software. Statistical analysis was performed with Graph Pad Prism. All statistical values are displayed as mean ± standard deviation. Comparisons between experimental groups were performed using an unpaired two-tailed t test with unequal variance for parametric samples. Differences are considered significant at a p < 0.05 significance level and marked with "*", very significant at p < 0.01 and labeled with "**", and extremely significant at p < 0.001 and labeled with "***".