Oscillatory links of Fgf signaling and Hes7 in the segmentation clock

Yukiko Harima[1,2,3] and Ryoichiro Kageyama[1,3]*

[1]Institute for Virus Research, Kyoto University, Kyoto 606-8507, Japan.

[2]Kyoto University Graduate School of Biostudies, Kyoto 606-8502, Japan.

[3]Japan Science and Technology Agency, CREST, Shogoin-Kawahara, Sakyo-ku, Kyoto 606-8507, Japan

*Corresponding author: Ryoichiro Kageyama

Institute for Virus Research

Kyoto University

Shogoin-Kawahara, Sakyo-ku

Kyoto 606-8507

Japan

Tel: 81-75-751-4011

Fax: 81-75-751-4807

E-mail: rkageyam@virus.kyoto-u.ac.jp

Summary

Somitogenesis is controlled by the segmentation clock, where the oscillatory expression of cyclic genes such as *Hes7* leads to the periodic expression of *Mesp2*, a master gene for somite formation. Fgf signaling induces the oscillatory expression of Hes7 while Hes7 drives coupled oscillations in Fgf and Notch signaling, which inhibits and activates *Mesp2* expression, respectively. Because of different oscillatory dynamics, oscillation in Fgf signaling dissociates from oscillation in Notch signaling in S−1, a prospective somite region, where Notch signaling induces *Mesp2* expression when Fgf signaling becomes off. Thus, oscillation in Fgf signaling regulates the timing of *Mesp2* expression and the pace of somitogenesis. In addition, Fgf signaling was found to be a primary target for hypoxia, which causes phenotypic variations of heterozygous mutations in *Hes7* or *Mesp2*, suggesting gene-environment interaction through this signaling.

Introduction

Somites, metameric structures that later give rise to vertebrae, ribs, skeletal muscles, and subcutaneous tissues, are formed by periodic segmentation of the anterior parts of the presomitic mesoderm (PSM). This periodic event is regulated by a biological clock called the segmentation clock, which involves the oscillatory expression of cyclic genes such as the basic helix-loop-helix (bHLH) gene *Hes7* (Figure 1) [1-5]. Genes in the Notch, Fgf, and Wnt pathways are expressed in an oscillatory manner in the mouse PSM [6-9]. One major outcome of such oscillatory expression is periodic activation of the bHLH gene *Mesp2*, a master gene for somite formation, in the prospective somite region termed S−1 (note that S0 is the next forming somite while S−1 is the PSM region that will form a somite after S0) (see Figure 3) [10,11]. So, the question is how cyclic genes periodically induce *Mesp2* expression in S−1.

Oscillatory expression is halted in the anterior region of the PSM, where overt somite differentiation begins, and the interface between this anterior region and the posterior oscillatory region is called the wavefront. It was previously shown that the wavefront is established by morphogens such as Fgf8 [12,13]. The *Fgf8* gene is transcribed only in the posterior end of the PSM, but due to slow degradation, the amount of *Fgf8* mRNA gradually decreases in PSM cells, which move anteriorly as the PSM grows posteriorly, forming the posterior to anterior *Fgf8* gradient [14]. The

anterior border of the Fgf gradient corresponds to the wavefront, and after passing this line, PSM cells express *Mesp2* because *Mesp2* expression is repressed by Fgf8 [11]. Fgf signaling seems to sweep back at a steady speed as the PSM grows, suggesting that the wavefront might regress steadily. However, *Mesp2* expression initiates synchronously in the whole S−1 region, and this synchronous expression seems to be important for the subsequent somitogenic processes [11]. These results raise the alternative possibility that the wavefront periodically jumps. Recent studies revealed that oscillators in the Fgf and Notch pathways are essential for periodic induction of *Mesp2* expression in S−1.

Fgf signaling induces traveling waves

The Notch intracellular domain (NICD), an active form of Notch, activates the *Hes7* promoter both in in-vitro culture and in-vivo transgenic mouse studies, indicating that Hes7 expression is controlled by Notch signaling [8,15-17]. In agreement with this notion, Hes7 expression is severely down-regulated in the absence of Rbpj, an essential mediator of Notch signaling [8,17]. However, a low level of *Hes7* expression still oscillates in the PSM of *Rbpj*-null mice, suggesting that another signaling may be responsible for oscillatory expression of *Hes7* [8,17]. Indeed, Fgf signaling is required for Hes7 expression because Hes7 expression totally disappears in the presence of Fgf inhibitors or in the absence of *Fgfr1*, an Fgf receptor gene essential for somitogenesis [8,18,19]. Thus, Hes7 expression is cooperatively regulated by both Fgf and Notch signaling. In zebrafish, it was previously proposed that non-synchronous oscillatory expression remains in embryos mutant for Notch signaling [20], and it was recently shown that expression of *her1*, a zebrafish *Hes7* homologue, still oscillates non-synchronously in such mutants [21•]. In contrast, treatment with an inhibitor of Fgf signaling abolished *her1* expression, whereas transplanted Fgf8-soaked beads induced ectopic traveling waves of *her1* expression in zebrafish embryos [22••]. These results indicate that Fgf signaling is essential for *Hes7*/*her1* oscillations in the PSM.

The stripes of both *Hes7* and *her1* become narrower as they move anteriorly in the PSM, and this narrowing is owing to slowing oscillations (longer periods) [23,24]. The *Fgf8* gradient could be involved in the slowing oscillations because the Fgf level affects the length (or narrowing) of *her1* stripes [22••]. However, pERK, an effector of Fgf signaling [13,25], is expressed in an oscillatory manner and does not form any apparent gradients in the mouse PSM (see below) [8,26••], indicating that the Fgf8

gradient is not translated into pERK levels. It was shown that Notch signaling-mediated intercellular coupling regulates the pace of the segmentation clock [27]. Fgf signaling could cross-talk with Notch signaling and thereby control the period of oscillatory expression. Further study will be required to understand the mechanism of how *Hes7*/*her1* oscillations slow in the anterior PSM and whether Fgf signaling is involved in this slowing.

Coupled and dissociating oscillations of Notch and Fgf signaling

In the mouse PSM, expression of the bHLH repressor gene *Hes7* oscillates by negative feedback [15]. Hes7 oscillation drives the oscillatory expression of the Notch modulator *Lunatic fringe* (*Lfng*), which inhibits Notch signaling [5,17,28-30•]. When Notch signaling is activated by its ligands such as Delta-like1, the transmembrane protein Notch is processed, releasing NICD. Lfng oscillation periodically inhibits Notch signaling, resulting in cyclic formation of NICD in the PSM (Figure 2) [26••,31,32]. It has been shown that NICD activates *Mesp2* expression in collaboration with Tbx6, a T-box protein required for PSM differentiation and segmentation [11,33]. Hes7 also drives the oscillatory expression of Dusp4, a phosphatase of phophorylated ERK (pERK), an effector of Fgf signaling. Owing to Dusp4 oscillation, pERK is periodically dephosphorylated, and the amount of pERK also oscillates in the PSM (Figure 2) [8,26••]. It was shown that Hes7 induces oscillatory expression of another Fgf inhibitor gene, *Sprouty4*, in the mouse PSM, which could also contribute to pERK oscillation [34]. Thus, *Hes7* induces oscillations of both Notch and Fgf signaling in the PSM. Conversely, Notch and Fgf signaling cooperatively up-regulate *Hes7* expression (Figure 2) [7]. These results indicate that Notch and Fgf signaling molecules and Hes7 form the oscillatory gene network. The next question is how the oscillatory expression of NICD, an activator of *Mesp2*, and pERK, a repressor of *Mesp2*, induces *Mesp2* expression in S−1.

Detailed expression analysis showed that the dynamics is different between NICD and pERK oscillations: NICD expression travels like a wave, narrowing down to a near somite size, while pERK exhibits an On-Off pattern (Figure 3). Narrowing of the NICD expression domain is due to slower oscillation in the anterior PSM compared to the posterior PSM, whereas the pERK oscillation does not slow. When NICD is expressed in the posterior to middle PSM, pERK is also present, inhibiting NICD from

inducing *Mesp2* expression (phases II and III, Figure 3). However, when the NICD expression domain moves and narrows down to the region around S−1, pERK suddenly disappears, which allows NICD to induce *Mesp2* expression synchronously in the whole S−1 region (Figure 3, panel furthest to the right) [26••]. Thus, two types of oscillations in Notch and Fgf signaling, which are coupled by Hes7 in the posterior PSM, dissociate in the anterior PSM, and this dissociation allows a group of cells ($NICD^{+}pERK^{-}$) to express *Mesp2* simultaneously in the whole S−1 region (Figure 3).

Retinoic acid (RA) signaling forms an opposite gradient to and antagonizes the Fgf8 signaling [35], and it has been proposed that the bistability between antagonistic Fgf and RA gradients leads to jumps of the wavefront [36]. However, this notion remains to be experimentally analyzed. A recent study with the monolayer PSM culture system shows that the phase of Lfng oscillation is delayed in the anterior PSM compared to the posterior PSM [37•]. Interestingly, the phase difference between anterior and posterior cells (phase gradient) was found to be inversely proportional to the sizes of formed segments, suggesting that the segment size or the wavefront jump is predetermined by the phase gradient [37•]. Strikingly, the phase gradient in monolayer PSM culture tissues is maintained even in the absence of an anterior opposing gradient [37•], suggesting that the segment size is controlled independently of an anterior opposing gradient.

Oscillation in Fgf signaling is important for the timing of somite formation

It was previously thought that Notch signaling function as a pacemaker of the segmentation, while the distance that the wavefront (the anterior boarder of Fgf signaling) travels during one oscillation cycle defines the somite size [1]. However, recent data on the oscillator networks needs revision of this view. Cyclic down-regulation of pERK seems to be important for the S−1 cells to express *Mesp2* periodically, suggesting that pERK oscillation in Fgf signaling regulates the pace of segmentation (Figures 3 and 4A) [26••]. In contrast, the NICD expression domain narrows in the anterior PSM, and this narrowing is important for the size of somites, because the $NICD^{+}$ region that co-expresses Tbx6 induces *Mesp2* expression after being freed from Fgf-pERK regulation (Figures 3 and 4A) [26••,31]. Thus, NICD oscillation plays an important role in the spatial regulation of somitogenesis. It was previously shown that increased Fgf activity reduces the somite size without affecting the pace of

segmentation [12]. Fgf was shown to caudalize the PSM and thereby shift the wavefront anteriorly [12]. In addition, as discussed above, Fgf signaling could be involved in narrowing of Hes7 stripes, which would lead to narrowing of NICD stripes via Lfng. These effects could secondarily affect the somite size. Further analysis will be required to determine the role of Fgf signaling in somite formation to differentiate between direct and indirect activities.

In *Hes7* KO mice, although segmentation is severely defective, *Mesp2* is expressed in the S−1 region, as observed in wild-type mice [5,26••]. In the absence of *Hes7*, Lfng and Dusp4 expression becomes non-oscillatory, and therefore NICD and pERK expression become steady (Figure 4B) [26••]. In *Hes7* KO mice, NICD expression continues longer than pERK, forming the $NICD^{+}pERK^{-}$ region around S−1 where *Mesp2* expression is induced. However, time-lapse imaging analysis showed that this *Mesp2* expression domain moves steadily as the PSM grows (Figure 4B). As a result, in the absence of *Hes7*, the onset of *Mesp2* expression does not occur simultaneously in the whole S−1 region but gradually proceeds from the anterior to posterior even in the same S−1 region (Figure 4B), which may cause the segmentation defects. Thus, although a snapshot of *Mesp2* expression is not significantly different between the wild-type and *Hes7* KO embryos (green signals in the left panels of Figure 4A,B), the spatiotemporal profiles of *Mesp2* are totally different (green signals in the middle panels of Figure 4A,B) [26••].

The notion that Hes7-induced NICD and pERK oscillations regulate periodic expression of *Mesp2* raises the possibility that Hes7 is a key pacemaker of the segmentation clock. It has been suggested that the negative feedback with a delayed timing is essential for sustained oscillation with an appropriate period [38-42], and that negative feedback with shorter delays accelerates the tempo and dampens or abolishes the oscillation. It was recently shown that intronic delays, which include the time required for transcription and splicing of intron sequences, constitute an important part of such proper delays. Due to rapid transcription, splicing events rather than the intron length may be more important for the intronic delay [43]. The *Hes7* gene has three introns, and deletion of all three introns reduces the delay by 19 min and completely abolishes oscillatory expression, leading to steady Hes7 expression and fusion of all somites [44]. In contrast, deletion of two introns of the *Hes7* gene reduces the delay by 5 min and accelerates the pace of the segmentation clock, although the oscillation is

eventually halted [45•]. Interestingly, these mutant mice have nine cervical vertebrae whereas the wild-type mice have seven, indicating that two more pulses of Hes7 oscillation occur during formation of cervical vertebrae [45•]. It is likely that oscillations in both Notch and Fgf signaling are also accelerated in these mutant mice, suggesting that *Hes7* is a fundamental pacemaker of the segmentation clock.

Hypoxia and Fgf signaling

Congenital scoliosis, which is caused by vertebral defects, occurs about 1 in 1,000 human live births, and it has been shown that heterozygous mutations in *HES7* or *MESP2* cause this disease [46••]. Interestingly, some of those who have the same mutations do not have any abnormal vertebrae, indicating partial penetrance. Partial penetrance of vertebral defects is also observed in mice carrying heterozygous mutations in *Hes7* or *Mesp2* [46••]. Notably, short-term hypoxia induces significantly higher rates and severity of vertebral defects in *Hes7* or *Mesp2* heterozygous mutant mice, suggesting that the phenotypes of genetic disorders are affected by environmental conditions such as hypoxia [46••,47]. Hypoxia does not alter *Fgf8* and *Fgfr1* expression in the PSM, but pERK expression becomes absent or significantly reduced (Figure 2). Furthermore, *Hes7* expression becomes absent or significantly reduced, which is similar to the phenotype caused by treatment with Fgf signaling inhibitors [46••]. These results suggest that hypoxia primarily affects Fgf signaling, although it may also affect other pathways like Wnt and Notch signaling (Figure 2). The mechanism of how hypoxia affects pERK formation remains to be determined.

Perspectives

In the mouse PSM, NICD makes a traveling wave, while pERK makes an On-Off pattern, and the mechanism by which the dynamics of NICD and pERK oscillations are different remains to be determined. The kinetics of the cell-cell communication may be different between Notch and Fgf signaling. One possibility is that Notch signaling travels relatively slowly because the ligand activates Notch only in adjacent cells, while Fgf is secreted and may rapidly reach distant cells, making the whole population respond simultaneously. Further analysis will be required to determine the parameter values of each signaling communication speed.

Another important aspect of Fgf signaling is its role in elongation of the

embryonic axis [48-50]. Fgf signaling activates random motility of PSM cells, particularly in the posterior region, but not the proliferation of these cells, and this graded motility (high in the posterior and low in the anterior) seems to contribute to the axis elongation [48]. It remains to be determined how Fgf signaling coordinates the axis elongation with the pace of segmentation.

Recent studies showed that Fgf signaling induces the oscillatory formation of pERK in non-PSM cells such as fibroblasts, and pERK pulses are shown to affect cell cycle progression [51,52]. Thus, the oscillation in Fgf signaling is not specific to the PSM but is involved in other biological events of many cell types. Further analyses will be required to understand the significance and mechanism of oscillation in Fgf signaling activity not only in the segmentation clock but also in other biological events.

References and recommended reading

1. Pourquié O: **Vertebrate segmentation: from cyclic gene networks to scoliosis.** *Cell* 2011, **145:**650-663.

2. Oates AC, Morelli LG, Ares S: **Patterning embryos with oscillations: structure, function and dynamics of the vertebrate segmentation clock.** *Development* 2012, **139:**625-639.

3. Saga Y: **The mechanism of somite formation in mice.** *Curr Opin Genet Dev* 2012, **22:**331-338.

4. Palmeirim I, Henrique D, Ish-Horowicz D, Pourquié O: **Avian *hairy* gene expression identifies a molecular clock linked to vertebrate segmentation and somitogenesis.** *Cell* 1997, **91:**639-648.

5. Bessho Y, Sakata R, Komatsu S, Shiota K, Yamada S, Kageyama R: **Dynamic expression and essential functions of *Hes7* in somite segmentation.** *Genes Dev* 2001, **15:**2642-2647.

6. Aulehla A, Wehrle C, Brand-Saberi B, Kemler R, Gossler A, Kanzler B, Herrmann BG: ***Wnt3a* plays a major role in the segmentation clock controlling somitogenesis.** *Dev Cell* 2003, **4:**395-406.

7. Dequéant M-L, Glynn E, Gaudenz K, Wahl M, Chen J, Mushegian A, Pourquié O: **A complex oscillating network of signaling genes underlies the mouse segmentation clock.** *Science* 2006, **314:**1595-1598.

8. Niwa Y, Masamizu Y, Liu T, Nakayama R, Deng C-X, Kageyama R: **The initiation and propagation of Hes7 oscillation are cooperatively regulated by Fgf and Notch signaling in the somite segmentation clock.** *Dev Cell* 2007, **13:**298-304.

9. Krol AJ, Roellig D, Dequéant ML, Tassy O, Glynn E, Hattem G, Mushegian A, Oates AC, Pourquié O: **Evolutionary plasticity of segmentation clock networks.** *Development* 2011, **138:**2783-2792.

10. Saga Y, Hata N, Koseki H, Taketo MM: **Mesp2: a novel mouse gene expressed in the presegmented mesoderm and essential for segmentation initiation.** *Genes Dev* 1997, **11:**1827-1839.

11. Oginuma M, Niwa Y, Chapman DL, Saga Y: **Mesp2 and Tbx6 cooperatively create periodic patterns coupled with the clock machinery during mouse somitogenesis.** *Development* 2008, **135:**2555-2562.

12. Dubrulle J, McGrew MJ, Pourquié O: **FGF signaling controls somite boundary position and regulates segmentation clock control of spatiotemporal *Hox* gene activation.** *Cell* 2001, **106:**219-232.

13. Sawada A, Shinya M, Jiang, Y-J, Kawakami A, Kuroiwa A, Takeda H: **Fgf/MAPK signalling is a crucial positional cue in somite boundary formation.** *Development* 2001, **128:**4873-4880.

14. Dubrulle J, Pourquié O: ***fgf8* mRNA decay establishes a gradient that couples**

axial elongation to patterning in the vertebrate embryo. *Nature* 2004, **427:**419-422.

15. Bessho Y, Hirata H, Masamizu Y, Kageyama R: **Periodic repression by the bHLH factor Hes7 is an essential mechanism for the somite segmentation clock.** *Genes Dev* 2003, **17:**1451-1456.

16. Chen J, Kang L, Zhang N: **Negative feedback loop formed by Lunatic fringe and Hes7 controls their oscillatory expression during somitogenesis.** *Genesis* 2005, **43:**196-204.

17. Ferjentsik Z, Hayashi S, Dale JK, Bessho Y, Herreman A, De Strooper B, del Monte G, de la Pompa JL, Maroto M: **Notch is a critical component of the mouse somitogenesis oscillator and is essential for the formation of the somites.** *PLoS Genet* 2009, **5:**e1000662.

18. Yamaguchi T, Harpal K, Henkemeyer M, Rossant J: ***fgfr-1* is required for embryonic growth and mesodermal patterning during mouse gastrulation.** *Genes Dev* 1994, **8:**3032-3044.

19. Wahl MB, Deng C, Lewandoski M, Pourquié O: **FGF signaling acts upstream of the NOTCH and WNT signaling pathways to control segmentation clock oscillations in mouse somitogenesis.** *Development* 2007, **134:**4033-4041.

20. Jiang Y-J, Aerne BL, Smithers L, Haddon C, Ish-Horowicz D, Lewis J: **Notch signaling and the synchronization of the somite segmentation clock.** *Nature* 2000, **408:**475-479.

21•. Delaune EA, François P, Shih NP, Amacher SL: **Single-cell-resolution imaging of the impact of Notch signaing and mitosis on segmentation clock dynamics.** *Dev Cell* 2012, **23:** 995-1005.
This study showed oscillatory expression of her1 at a single-cell resolution in the zebrafish segmentation clock. It beautifully demonstrated that her1 expression oscillates asynchronously in PSM cells of Notch signaling-mutant zebrafish. Interestingly, her1

oscillation seems to be linked with mitosis so that cell division makes less noise than previously thought.

22••. Ishimatsu K, Takamatsu A, Takeda H: **Emergence of traveling waves in the zebrafish segmentation clock.** *Development* 2010, **137:**1595-1599.
The authors showed that Fgf signaling induces *her1* oscillation, which forms a traveling wave, in zebrafish embryos. Fgf signaling levels affect the length of *her1* stripes, indicating that Fgf regulates the period of *her1* oscillation. The mechanism of how Fgf signaling regulates the period of *her1* oscillation remains to be determined.

23. Giudicelli F, Ozbudak EM, Wright GJ, Lewis J: **Setting the tempo in development: an investigation of the zebrafish somite clock mechanism.** *PLoS Biol* 2007, **5:**e150.

24. Masamizu Y, Ohtsuka T, Takashima Y, Nagahara H, Takenaka Y, Yoshikawa K, Okamura H, Kageyama R: **Real-time imaging of the somite segmentation clock: revelation of unstable oscillators in the individual presomitic mesoderm cells.** *Proc Natl Acad Sci USA* 2006, **103:**1313-1318.

25. Delfini M-C, Dubrulle J, Malapert P, Chal J, Pourquié O: **Control of the segmentation process by graded MAPK/ERK activation in the chick embryo.** *Proc Natl Acad Sci USA* 2005, **102:**11343-11348.

26••. Niwa Y, Shimojo H, Isomura A, González A, Miyachi H, Kageyama R: **Different types of oscillations in Notch and Fgf signaling regulate the spatiotemporal periodicity of somitogenesis.** *Genes Dev* 2011, **25:**1115-1120.
This study showed that Hes7 drives coupled oscillations in Fgf and Notch signaling, which inhibits and activates *Mesp2* expression, respectively. Oscillation in Fgf signaling dissociates from oscillation in Notch signaling in a prospective somite region, where Notch signaling induces *Mesp2* expression in the region where Fgf signaling is off. Thus, oscillation in Fgf signaling regulates the timing of *Mesp2* expression and the pace of somitogenesis.

27. Herrgen L, Ares S, Morelli LG, Schröter C, Jülicher F, Oates AC: **Intercellular coupling regulates the period of the segmentation clock.** *Curr Biol* 2010, **20:**1244-1253.

28. Dale JK, Maroto M, Dequeant ML, Malapert P, McGrew M, Pourquié O: **Periodic Notch inhibition by Lunatic Fringe underlies the chick segmentation clock**. *Nature* 2003, **421:**275-278.

29. Stauber M, Sachidanandan C, Morgenstern C, Ish-Horowicz D: **Differential axial requirements for lunatic fringe and Hes7 transcription during mouse somitogenesis.** *PLoS One* 2009, **4:**e7996.

30•. Okubo Y, Sugawara T, Abe-Koduka N, Kanno J, Kimura A, Saga Y: **Lfng regulates the synchronized oscillation of the mouse segmentation clock via *trans*-repression of Notch signalling.** *Nat Commun* 2012, **3:**1141.
This study sought to answer how Lfng regulates synchronized oscillation in the mouse PSM. The authors showed that Lfng represses Notch activity in neighboring cells by modulating Dll1 function, and proposed that this *trans*-repression is important for synchronized oscillation.

31. Huppert SS, Ilagan MXG, De Strooper B, Kopan R: **Analysis of Notch function in presomitic mesoderm suggests a γ-secretase-independent role for presenilins in somite differentiation.** *Dev Cell* 2005, **8:**677-688.

32. Morimoto M, Takahashi Y, Endo M, Saga Y: **The Mesp2 transcription factor establishes segmental borders by suppressing Notch activity.** *Nature* 2005, **435:**354-359.

33. Yasuhiko Y, Haraguchi S, Kitajima S, Takahashi Y, Kanno J, Saga Y: **Tbx6-mediated Notch signaling controls somite-specific *Mesp2* expression.** *Proc Natl Acad Sci USA* 2006, **103:**3651-3656.

34. Hayashi S, Shimoda T, Nakajima M, Tsukada Y, Sakumura Y, Dale JK, Maroto M,

Kohno K, Matsui T, Bessho Y: ***Sprouty4*, an FGF inhibitor, displays cyclic gene expression under the control of the Notch segmentation clock in the mouse PSM.** *PLoS One* 2009, **4:**e5603.

35. Diez del Corral R, Olivera-Martinez I, Goriely A, Gale E, Maden M, Storey K: **Opposing FGF and retinoid pathways control ventral neural pattern, neuronal differentiation, and segmentation during body axis extension.** *Neuron* 2003, **40:**65-79.

36. Goldbeter A, Gonze D, Pourquié O: **Sharp developmental thresholds defined through bistability by antagonistic gradients of retinoic acid and FGF signaling.** *Dev Dyn* 2007, **236:**1495-1508.

37•. Lauschke VM, Tsiairis CD, François P, Aulehla A: **Scaling of embryonic patterning based on phase-gradient encoding.** *Nature* 2013, **493:**101-105.
The authors developed a real-time imaging system of *Lfng* expression in monolayer PSM cultures and showed that the size of formed segments is adjusted (scaled) with that of the remaining PSM tissues. This study also indicated that the segment size is controlled independently of opposing gradients.

38. Lewis J: **Autoinhibition with transcriptional delay: a simple mechanism for the zebrafish somitogenesis oscillator.** *Curr Biol* 2003, **13:**1398-1408.

39. Monk NAM: **Oscillatory expression of Hes1, p53, and NF-kB driven by transcriptional time delays.** *Curr Biol* 2003, **13:**1409-1413.

40. Jensen MH, Sneppen K, Tiana G: **Sustained oscillations and time delays in gene expression of protein Hes1.** *FEBS Lett* 2003, **541:**176-177.

41. Hirata H, Bessho Y, Kokubu H, Masamizu Y, Yamada S, Lewis J, Kageyama R: **Instability of Hes7 protein is crucial for the somite segmentation clock.** *Nat Genet* 2004, **36:**750-754.

42. Zeiser S, Rivera O, Kuttler C, Hense B, Lasser R, Winkler G: **Oscillations of Hes7 caused by negative autoregulation and ubiquitination.** *Comput Biol Chem* 2008, **32:**47-51.

43. Hanisch A, Holder MV, Choorapoikayil S, Gajewski M, Ózbudak EM, Lewis J: **The elongation rate of RNA polymerase II in zebrafish and its significance in the somite segmentation clock.** *Development* 2013, **140:**444-453,

44. Takashima Y, Ohtsuka T, González A, Miyachi H, Kageyama R: **Intronic delay is essential for oscillatory expression in the segmentation clock.** *Proc Natl Acad Sci USA* 2011, **108:**3300-3305.

45•. Harima Y, Takashima Y, Ueda Y, Ohtsuka T, Kageyama R: **Accelerating the tempo of the segmentation clock by reducing the number of introns in the *Hes7* gene.** *Cell Rep* 2013, **3:**in press.
The authors showed that deletion of two introns in the *Hes7* gene shortens the transcriptional delay and accelerates the tempo of both *Hes7* oscillation and somite segmentation. This study indicated that *Hes7* is a key regulator of the pace of the segmentation clock.

46••. Sparrow DB, Chapman G, Smith AJ, Mattar MZ, Major JA, O'Reilly VC, Saga Y, Zackai EH, Dormans JP, Alman BA, McGregor L, Kageyama R, Kusumi K, Dunwoodie SL: **A mechanism for gene-environment interaction in the etiology of congenital scoliosis.** *Cell* 2012, **149:**295-306.
The authors showed that short-term hypoxia induces significantly higher rates and severity of vertebral defects in *Hes7* or *Mesp2* heterozygous mutant mice. It seems that hypoxia primarily affects Fgf signaling. This study indicated that the phenotypes of genetic disorders are affected by environmental conditions such as hypoxia.

47. Dunwoodie SL: **The role of hypoxia in development of the mammalian embryo.** *Dev Cell* 2009, **17:**755-773.

48. Bénazéraf B, François P, Baker RE, Denans N, Little CD, Pourquié O: **A random**

cell motility gradient downstream of FGF controls elongation of an amniote embryo. *Nature* 2010, **466:**248-252.

49. Stulberg MJ, Lin A, Zhao H, Holley SA: **Crosstalk between Fgf and Wnt signaling in the zebrafish tailbud.** *Dev Biol* 2012, **369:**298-307.

50. Boulet AM, Capecchi MR: **Signaling by FGF4 and FGF8 is required for axial elongation of the mouse embryo.** *Dev Biol* 2012, **371:**235-245.

51. Nakayama K, Satoh T, Igari A, Kageyama R, Nishida E: **FGF induces oscillations of Hes1 expression and Ras/ERK activation.** *Curr Biol* 2008, **18:**R332-R334.

52. Albeck JG, Mills GB, Brugge JS: **Frequency-modulated pulses of ERK activity transmit quantitative proliferation signals.** *Mol Cell* 2013, **49:**1-13.

Figure legends

Figure 1. Oscillatory expression of Hes7 in the PSM during somite segmentation. Spatiotemporal profiles of Hes7 expression are shown in the lower middle. The x axis represents time, while the y axis represents space. Anterior is top, and posterior is bottom. The spatial patterns of Hes7 expression at different time points are shown at both sides and on top. Note that the posterior end of the PSM grows posteriorly (downward).

Figure 2. Hes7-mediated coupled oscillations in Fgf and Notch signaling in the segmentation clock. Hes7 drives oscillatory expression of Dusp4, leading to pERK oscillation. Hes7 also drives oscillatory expression of Lfng, leading to oscillatory formation of NICD. Thus, pERK-Dusp4 and Lfng-NICD oscillations are coupled by Hes7 oscillations. Conversely, Hes7 oscillations are cooperatively regulated by Fgf and Notch signaling in the PSM.

Figure 3. Dynamic expression of NICD, pERK and Mesp2 (adapted from [26••]). The

posterior NICD domain moves anteriorly and narrows, while the pERK domain expands anteriorly, covering the NICD domain (middle two panels). After segmentation, pERK expression is down-regulated, and NICD now induces *Mesp2* expression in the S−1 region (panel furthest to the right). Thus, oscillations in Notch signaling periodically segregate a group of synchronized cells, and oscillations in Fgf signaling release these synchronized cells for somitogenesis at the same time.

Figure 4. Spatiotemporal profiles of NICD, pERK and Mesp2 (adapted from [26••]). (A) Spatiotemporal patterns of NICD, pERK and *Mesp2* expression in the wild type. *Mesp2* expression is induced periodically by NICD in the whole S−1 region after pERK expression disappears. (B) Spatiotemporal patterns of NICD, pERK and *Mesp2* expression in *Hes7* KO mice. In *Hes7* KO mice, pERK expression steadily regresses, and *Mesp2* expression also steadily regresses in the anterior PSM after Fgf/ERK signaling is turned off. Thus, *Mesp2* expression occurs at different time between the anterior and posterior cells even in the same prospective somites.

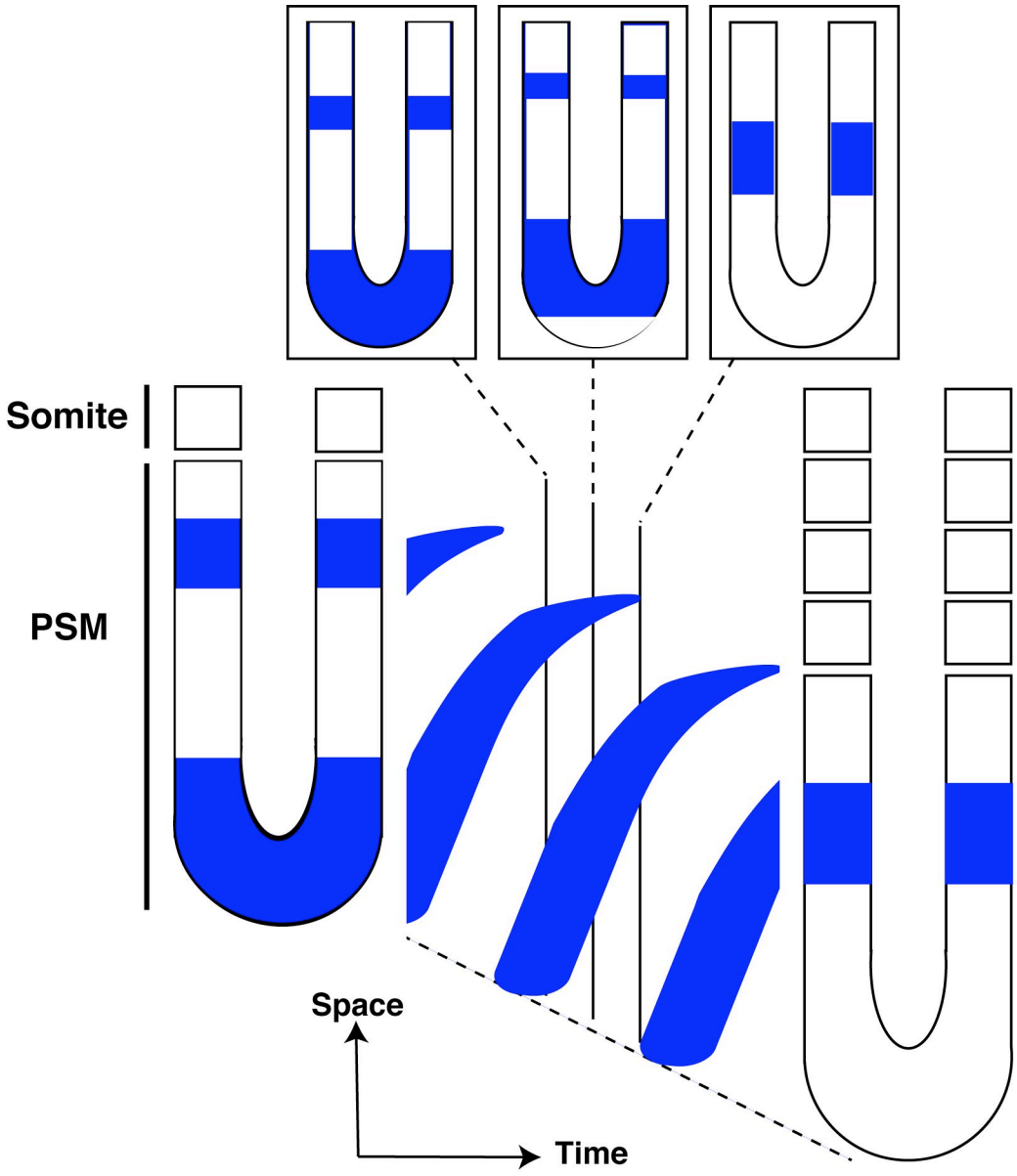

Figure 1

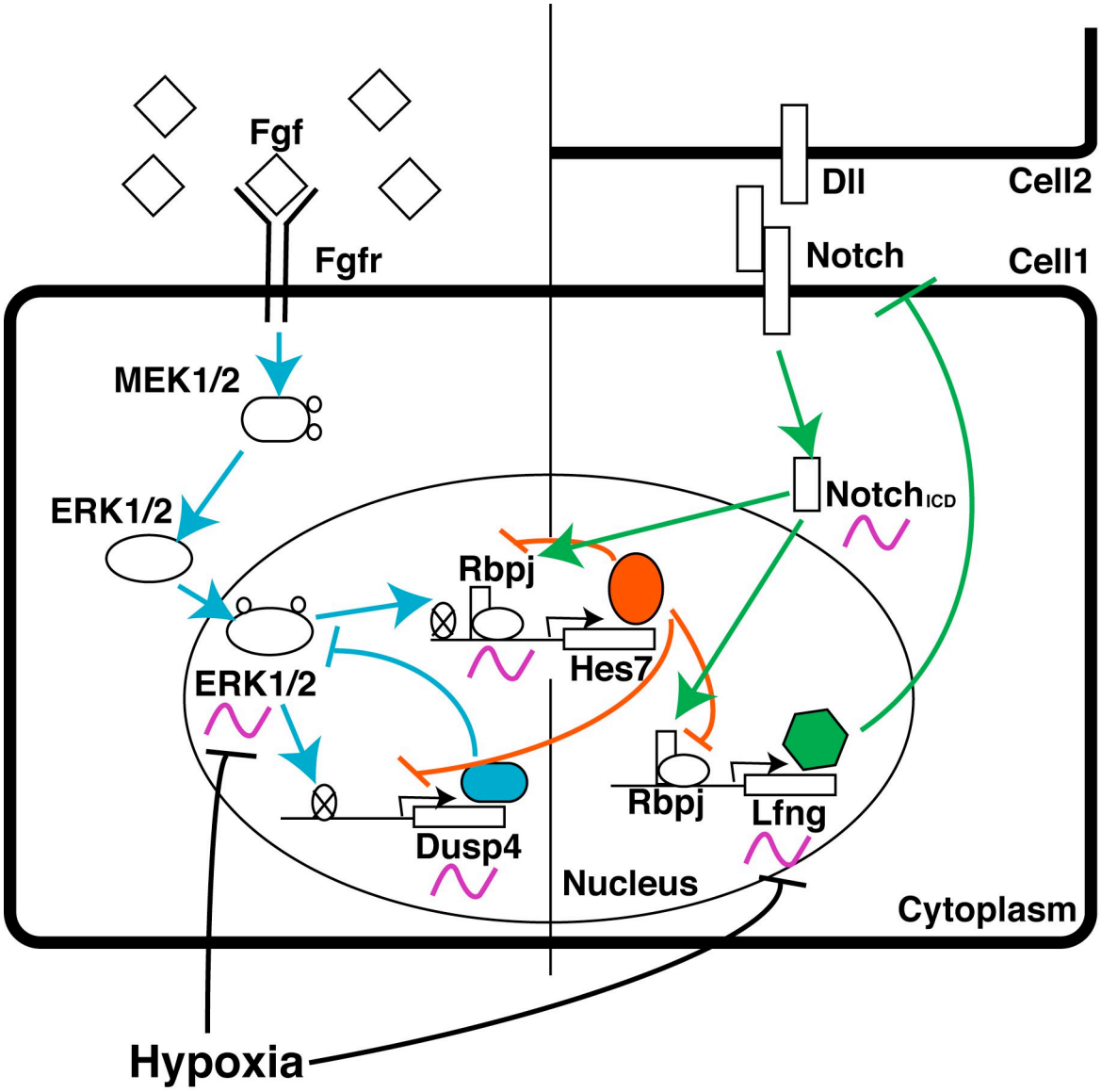
Fgf
Fgfr
Dll
Cell2
Notch
Cell1
MEK1/2
ERK1/2
Notch ICD
Rbpj
Hes7
ERK1/2
Rbpj
Lfng
Dusp4
Nucleus
Cytoplasm
Hypoxia

Figure 2

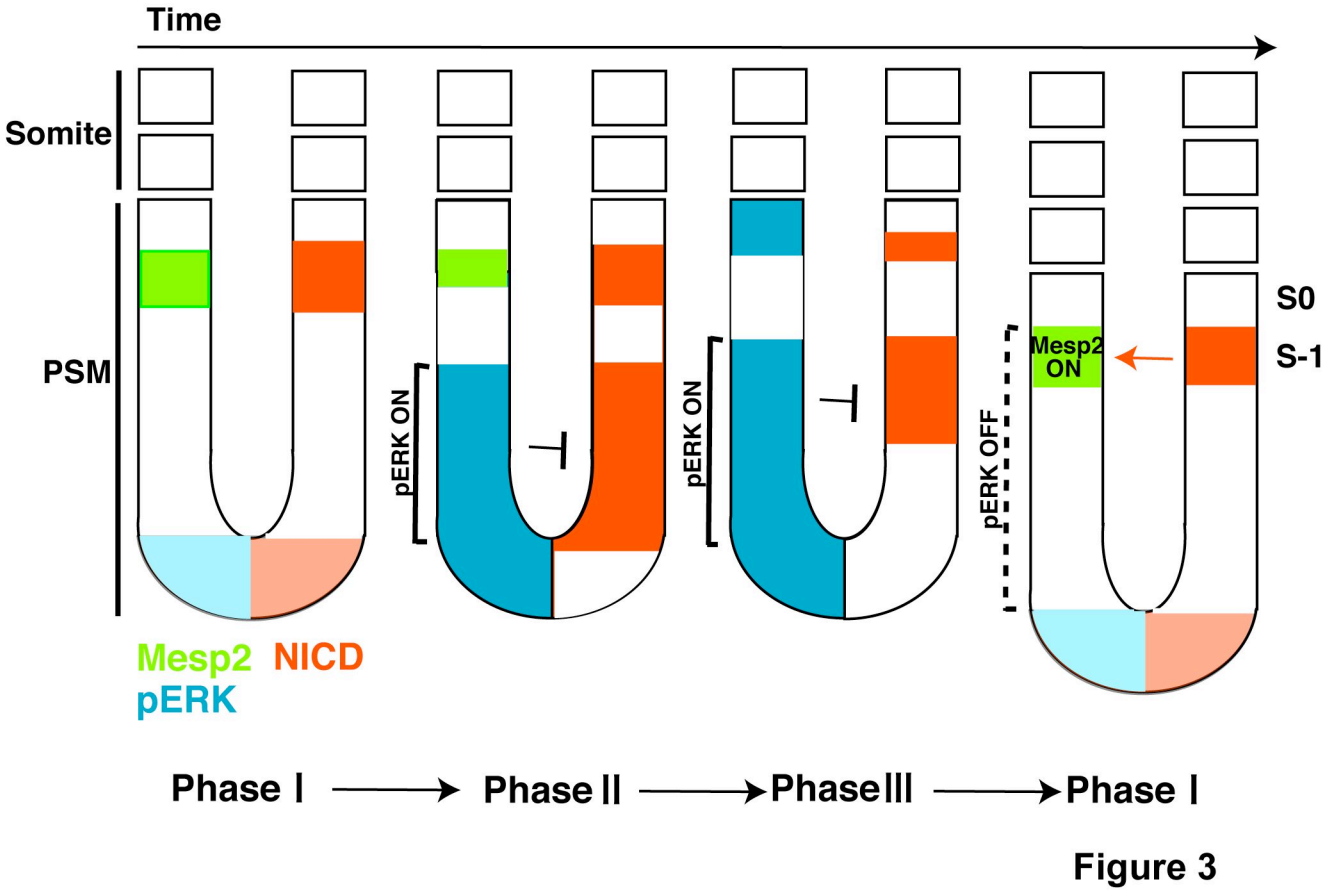
Time
Somite
PSM
pERK ON
pERK ON
pERK OFF
Mesp2 ON
S0
S-1
Mesp2
NICD
pERK
Phase I
Phase II
Phase III
Phase I

Figure 3

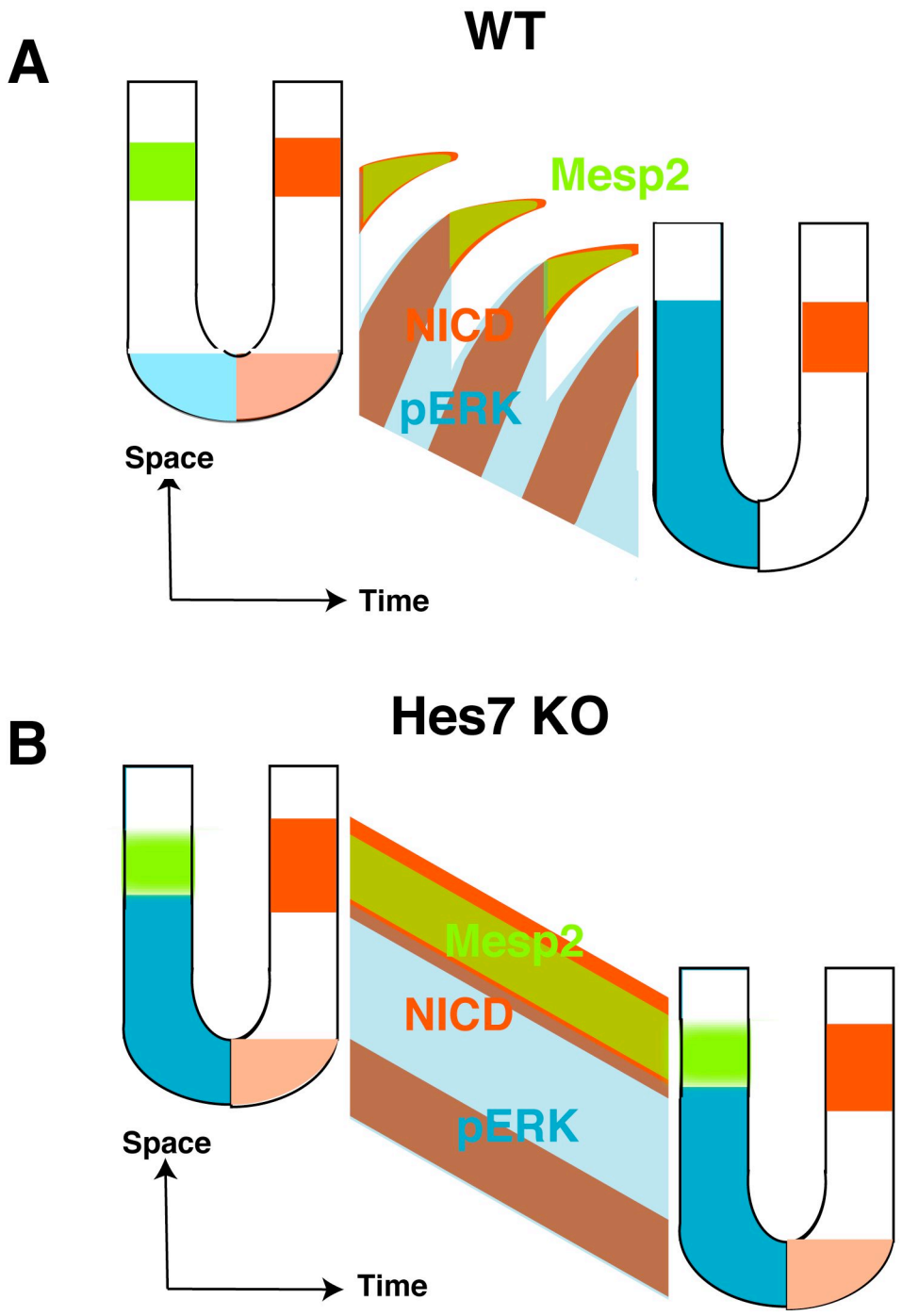
A
WT
Mesp2
NICD
pERK
Space
Time
B
Hes7 KO
Mesp2
NICD
pERK
Space
Time

Figure 4