

Mechanical Coupling Coordinates the Co-elongation of Axial and Paraxial Tissues in Avian Embryos

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SUMMARY

Tissues undergoing morphogenesis impose mechanical effects on one another. How developmental programs adapt to or take advantage of these effects remains poorly explored. Here, using a combination of live imaging, modeling, and microsurgical perturbations, we show that the axial and paraxial tissues in the forming avian embryonic body coordinate their rates of elongation through mechanical interactions. First, a cell motility gradient drives paraxial presomitic mesoderm (PSM) expansion, resulting in compression of the axial neural tube and notochord; second, elongation of axial tissues driven by PSM compression and polarized cell intercalation pushes the caudal progenitor domain posteriorly; finally, the axial push drives the lateral movement of midline PSM cells to maintain PSM growth and cell motility. These interactions form an engine-like positive feedback loop, which sustains a shared elongation rate for coupled tissues. Our results demonstrate a key role of inter-tissue forces in coordinating distinct body axis tissues during their co-elongation.

INTRODUCTION

During development, the morphogenesis of one tissue can produce forces that directly impact the dynamics of its neighboring tissues (Savin et al., 2011; Lye et al., 2015; Morita et al., 2017; Bailles et al., 2019). These interactions may play a role not only in controlling the correct development of each tissue and organ but also in coordinating them into an integrated system. Such role of inter-tissue forces is poorly explored due to the challenge of experimentally detecting and applying forces and the lack of a framework that bridges developmental signals, cell dynamics and tissue mechanics.

Avian embryos are an ideal vertebrate system to address these challenges because of their large size and accessibility. The forming body axis provides a striking example of coordinated morphogenesis of multiple distinct tissues (Bénazéraf and Pourquié, 2013). The axis forms in a head to tail (anterior to posterior, AP) sequence where tissues derived from the three germ layers are progressively added from a posterior growth zone (e.g. primitive streak, tailbud). The posterior movement of the growth zone lays down in its trail a conserved pattern of tissues which include the axial organs: neural tube (NT) and notochord (NC), and the flanking paraxial tissues: presomitic mesoderm (PSM), intermediate mesoderm and lateral plates (Figures 1A-B). While these tissues achieve similar lengths at the same pace, they exhibit distinct cellular organization and elongation dynamics (Keller and Danilchik, 1988; Shih and Keller, 1992; Steventon et al., 2016; Colas and Schoenwolf, 2001; Glickman et al., 2003; Sausedo and Schoenwolf, 1993; Yang, et al. 2002; Bénazéraf et al., 2010; Dray et al., 2013; Bénazéraf et al., 2017). First, the elongation of the NC and NT largely relies on intrinsic cell polarity-driven medio-lateral intercalation promoting convergence and extension along the AP axis. In contrast, the adjacent PSM shows limited convergence in the posterior body axis region (Shih and Keller, 1992; Bénazéraf and Pourquié, 2013). Second, in NC and NT, cells exhibit an epithelial organization, move coherently and only limited cell rearrangements are seen, while in the PSM cells are mesenchymal, show random cell movements and extensive cell mixing (Colas and Schoenwolf, 2001; Bénazéraf et al., 2010). Third, in tailbud stages, both NT and PSM rely on new cell addition from a specific subdomain of the growth zone called the chordo-neural hinge (or PD for progenitor domain) at the posterior end of the axis (Figure 1A). To enter the PSM, newly specified PSM cells on the ventral side of the PD move laterally away from the midline (Denans et al., 2015). In contrast, NC lengthens mainly through cell rearrangement and

differentiation with little addition of new cells (Sausedo and Schoenwolf, 1993; Bénazéraf et al., 2017). These differences in cell behaviors lead to counterintuitive cellular dynamics such as sliding between PSM and NC cells where NC cells appear to move faster posteriorly than PSM cells relative to the somites, despite that the two tissues elongate hand-in-hand (Figure 1B, Yang et al., 2002; Glickman et al., 2003; Bénazéraf et al., 2017). These observations raise the question of how do the PSM, NC and NT, which employ such distinct cellular processes to lengthen, coordinate their elongation during body axis formation?

Mechanical interactions between tissues might offer an answer to this question. The close proximity and lack of interstitial space between the PSM, NC, NT and the PD at the posterior end of the body axis (Schoenwolf 1979) means that the deformation of one would likely have a direct mechanical effect on the others. Indeed, it was found that ablating the posterior PSM (pPSM) greatly reduces the elongation rate of all tissues (Bénazéraf et al., 2010). This finding suggests that the pPSM may mechanically contribute to axial elongation. It was further hypothesized that cell motility in the PSM produces tissue expansion forces (Bénazéraf et al., 2010; Regev et al., 2017). To test these possibilities, tissue and cell behaviors need to be observed *in vivo* with readout of tissue forces.

Here we used soft gels to replace different tissues in the elongating posterior body axis of chicken embryos to track the forces applied by the neighboring tissues. We found that the pPSM compresses the axial tissues. This extrinsic force adds to the intrinsic process of cell intercalation to promote convergence and elongation of the NT and NC. The compression requires FGF dependent cell motility in the pPSM while the NT cell intercalation requires Wnt/PCP (planar cell polarity) mediated cell polarity. Using an agent(cell)-based model that incorporates cell motility and polarity, we faithfully recapitulate the dynamics of cells and tissues under different experimental conditions. The model predicts that the advancing axial tissues exert a pushing force on the PD, which we validated by gel deformation. Surprisingly, the model also predicts that the lateral movement of the PSM cells from the ventral midline requires this axial pushing force. To test this hypothesis, we inserted a magnet-controlled pin in place of the posterior NC/NT (pNC/pNT) region to apply a posteriorly oriented force to the PD. This mechanical signal partially restores PSM cell movements and posterior movement of the PD (i.e. axial elongation). Together our findings reveal mechanical coupling between the PSM and axial tissues: the left and right pPSMs generate a bilateral compression of the axial tissues promoting

their elongation. The axial tissues in turn push on the PD promoting PSM growth by driving cell flow. These interactions form a self-sustaining positive feedback loop that prevents elongation offset from occurring between different tissues. Just like an internal combustion engine that self-refills fuel for the next cycle, expanding pPSM (combustion) forces axial tissues (piston) to drive PSM progenitors (fuel) out of the PD (fuel reserve). Our model thus provides a simple physical mechanism and an intuitive explanation of the striking robustness and coordination between tissues observed in chicken embryonic body axis elongation.

RESULTS

Cellular dynamics of axial and paraxial elongation

To set up a quantitative baseline of tissue and cell behaviors during body axis elongation, we performed live imaging of Tg(CAG:GFP) chicken (McGrew et al., 2008) embryos after labeling the new PSM cells and progenitors in the primitive streak and tailbud with Dil (Figures S1A-B, Movie S1) between Hamburger Hamilton (HH) stages 8 and 14. During these stages, body elongation first follows the regression of the primitive streak and then the posterior movement of the tailbud. These movements result in the progressive formation of the NT, NC and PSM (among other tissues) at the posterior end of the embryo (Hamburger 1992).

Using cell tracking (Figure S1C), we analyzed the movement of cells relative to the Node (Figure 1C). Both axially-localized cells (including NC cells and new PSM cells from the PD) and paraxially-localized PSM cells are left behind by the Node but at increasingly different speeds (Figure 1D, note that while some axial cells move posteriorly with the Node, others are left behind, resulting in a net negative average speed at the Node [position 0]). This speed offset reflects the rate of the posterior movement of the PD (i.e. the elongation rate). In early embryos, at the PD level, cells move away from the midline while in the PSM flanking the NC, cells converge toward the midline (Figure 1E, positive indicates medial to lateral movement). Similar dynamics occur in later stage embryos but at markedly reduced average speeds (Figure 1E). The medial to lateral dispersion of new PSM cells at the PD level results in their joining the PSM on both sides of the axis (Figures S1D). This dispersion speed is an indirect measurement of the PSM cell addition rate as the PD and PSM form a continuum of cell flow. To measure the motility of cells (Figures S1E), we fit the tracks with a diffusion model to separate the directional

and non-directional components (Regev et al., 2017). This allows us to assess the local motility or "cell diffusivity" in the PSM. A posterior to anterior motility gradient is observed in both early and later stages of elongation (Figure 1F). These cellular dynamics are consistent with previous studies (Kulesa and Fraser, 2002; Yang et al., 2002; Manning and Kimelman, 2015; Lawton et al., 2013; Bénazéraf et al., 2010, 2017).

Posterior PSM exerts a lateral to medial compression on the axial tissues

To investigate how the pPSM affects the axial tissues, we performed bilateral ablations of the pPSM (Figure 2A). Interestingly, both pNC and pNT remained wider even after PSM cuts closed, suggesting impaired tissue convergence (Figures 2B-C, S2L). Correspondingly, the elongation rate of the operated embryos reduces drastically (Figures S2A-C). These data show that the pPSM is required for the convergence and elongation of the axial tissues. To test if pPSM supplies a biochemical cue to promote axial elongation, we inserted tailored aluminum foil pieces between pPSM and pNC/NT to cut off signal exchange (Figure 2D). Unlike pPSM ablation, foil insertion does not slow down elongation (Figure 2E), suggesting the requirement of pPSM is not through diffusible signals. To test if the pPSM passively confines pNC/pNT to prevent their lateral expansion, we replaced the pPSM with a stiff alginate gel (1% alginate at 100mM Ca²⁺) as a barrier. The gel-flanked portions of pNC/pNT remained wider (Figure S2D, n=4/4), suggesting that the pPSM actively compresses the axial tissues. To detect this compression, we replaced the pNC with a soft alginate gel containing fluorescent beads (Figure S2F, 1% alginate at 10mM Ca²⁺, order of 100 Pa [Banerjee et al., 2009], comparable to embryonic tissues [Zhou et al., 2009]), which showed ~5% narrowing indicating that pPSM produces a compressive stress in the order of 10s of Pa. We further lowered the gel stiffness by injecting 1% alginate solution into ablation sites allowing gelation *in vivo*. Using live confocal imaging, we found that these implants are more deformable as a paste and round up edges quickly under surface tension (Figure 2F, Movie S2). The pNC/pNT implant narrows significantly over time along the LM axis (Figures 2F-H). Similar deformation dynamics were observed in replacement experiments of the pNC (Figures S2G-I) or the pNT (Figure S2J-K) alone. These results together show that pPSM contributes to axial convergence and elongation through lateral tissue compression.

Cellular basis of PSM compression, axial convergence, and elongation

To investigate how pPSM creates compression on axial tissues, we constructed a 2D agent (i.e. cell)-based computational model (Figure 3A, Movie S3). The model contains a field of PSM cells subdivided by the axial tissue in the midline, and the posterior PD that supplies new PSM cells (for the feasibility of simulation the number of cells is small compared to the real tissues but has a similar paraxial-axial proportion). All cells occupy a volume in the tissue and thus push against each other upon close contact. Cell movement is countered by a resistance mimicking the effect of the viscous extracellular environment (Figures S3A-B, methods). When a motility gradient is added across the field of PSM cells, the high motility region (corresponding to the pPSM) shows a diffusion-like expansion (Figure S3C) on the tissue level that reproduces observations in isolated PSM explants (Palmeirim et al., 1998). When this expansion encounters a lateral boundary, such as the axial tissues, a pressure gradient parallel to the motility gradient is recorded (Figure S3D). To test this pressure gradient *in vivo*, we replaced the aNC with gels (Figures S3E-F). In contrast to the pNC control, gels replacing aNC show minimal deformations (Figures S3G-H). These data are consistent with the pPSM (but not the aPSM) generating compression in the region of maximal cell motility (Figure 1F). Our model also generates the cell density gradient observed in the PSM *in vivo* (Figures S3I-J), which is another consequence of the motility gradient and tissue expansion. To model axial cell behaviors, we implemented a polarity parameter that controls the shape of the axial cells. A polarized axial cell is elongated along the medio-lateral axis and intercalates with other axial cells more easily in the model, mimicking the cell shape and behavior observed *in vivo* such as neural progenitors in the NT (Colas and Schoenwolf, 2001). To model the PD production of new PSM cells, we periodically added cells posterior to the axial end in the simulation iterations. This mimics the *in vivo* configuration where the anterior primitive streak or tailbud adds PSM cells ventrally at the axis end by ingression and proliferation (for simplicity, we only consider the PD placement at the posterior end of the midline, which is similar to the actual geometrical arrangement between different stages and species. In addition, the newly added cells behave identically to PSM cells as their *in vivo* counterparts have undergone PSM specification and epithelial to mesenchymal transition [EMT, Denans et al., 2015] before moving to the ventral midline). A combination of the PSM motility gradient, the polarization of axial cells and the cell addition from PD produces sustained axial convergence and elongation *in silico* (Figure 3B, Movie S3).

To test the key cell behavior assumptions (motility and polarity) of the model, we

considered the molecular signals known to control these cell properties. FGF signaling is known to control the cell motility gradient in the PSM (Bénazéraf et al., 2010). We reduced cell motility by electroporating an inducible dominant-negative FGF receptor (dnFGFR1) construct and a fluorescent reporter in the PD (the anterior primitive streak, Imura and Pourquie, 2008). The electroporation allows PSM-specific expression of the transgenic construct (Figure S3K). Embryos in which FGF signaling is inhibited in the PSM exhibit a wider pNT as seen after pPSM ablation, together with slower pNT convergence and axis elongation (Figures 3C-D, Bénazéraf et al., 2010). Similar results on pNT convergence and elongation were obtained by electroporation of a dominant negative MAP kinase (dnMKK1, Figures S3L-M, Delfini et al., 2005) and by embryo treatment with the FGF inhibitor PD173074 (data not shown). Furthermore, embryos treated with Blebbistatin to directly inhibit cell movement show no medial-lateral compression on implanted gels and delayed axial convergence (Figures S3O,4G). These data support the hypothesis that pPSM cell movement downstream of FGF signaling collectively generates compression on the axial tissues.

pNT and pNC cells exhibit cell polarity which promotes active medio-lateral intercalation driving intrinsic tissue elongation (Keller et al., 2008; Williams et al., 2014). In amphibian embryos, interfering with cell polarity by disrupting the posterior WNT5 gradient by either up- or down-regulating the pathway reduces axial convergence and elongation (Moon et al., 1993; Wallingford and Harland, 2001). We tested whether disrupting medio-lateral cell intercalation by WNT5A also interferes with axis elongation in the chicken embryo. We injected WNT5A protein into the anterior portion of the pPSM or the pNT, where endogenous expression is low (Chapman et al., 2004). This efficiently slowed down elongation while control injections of buffer, WNT10 at the same location, or WNT5A at the Node (where endogenous expression is high) do not have significant impacts (Figures S4A-C). Consistently, the pNT in WNT5A injected embryos maintains a larger width compared to controls (Figures 3E, S4A'-B') indicating impaired convergence. To analyze cell polarity changes following WNT5A perturbation, we electroporated a membrane tagged GFP in the NT which allows mosaic labeling of pNT cells that highlight cell shapes (Figure S5A). We also compared NT cell nuclei shapes following DAPI staining (Figure S5B). This analysis reveals that WNT5A causes a reduction of the lateral-medial polarization of NT cells in terms of cell shape (Figure S5D), while PSM cells do not exhibit changes in cell shape or density in the WNT5A condition (Figures S5C,E). Furthermore, the PSM cell motility gradient

is unaffected (Figures S5F). Using live imaging and cell tracking, we found that the medial-lateral intercalation of pNT cells is greatly reduced following WNT5A injection (Figures S5G-H). These results suggest that WNT5A perturbation delays convergence and elongation by disrupting NT cell polarity, consistent with previous studies. Taken together, our data suggest that FGF signaling in the PSM and WNT/PCP signaling in the NT provide extrinsic and intrinsic forces of axial elongation through motility and polarity, respectively. These results add confidence to our model assumptions, allowing us to utilize the model to further dissect the role of these mechanical interactions between tissues.

Elongating axial tissues produce a pushing force on the progenitor domain

Our perturbations show that posterior movement of the PD (which marks elongation) greatly slows down whenever axial tissues are perturbed (Figures S2B-C, S4), suggesting that the movement of the PD is not autonomous but rather requires pNT/pNC. Inserting a piece of aluminum foil to separate pNT/pNC with the PD does not affect elongation (Figures 2D, S2E), suggesting that the pNT/pNC requirement is not due to a diffusible molecular signal. To understand how axial elongation promotes PD movement, we resorted to our simulation. Counter-intuitively, ablating PD is predicted to slow down elongation more mildly compared to pPSM or axial ablations, which agrees with experiments (Figures 4A-B, Movie S4, compare to Figures S2A-C). We followed the average AP forces that two representative cells experience in the simulations (arrows in Figure 4A): one axial cell in the midline bordering the PD thus detecting the net force from the axial tissue and the PD, the other a paraxial PSM cell that starts off at the same AP level for comparison. The PSM cell is initially pushed posteriorly (by other pPSM cells) but then anteriorly as it transitions to become an aPSM cell, consistent with the model of pPSM expansion (Figure 4C). In contrast, the axial cell (and the PD) is continuously pushed towards the posterior. To test whether this predicted pushing force exists *in vivo*, we replaced the PD with a soft alginate gel. Strikingly, the gel thins along AP especially in the middle where it is in direct contact with the axial structures (Figure 4D-F, Movie S5). This is in sharp contrast to the lateral to medial compressive deformations seen in gels that replace the pNT or pNC (Figures 2F-H, S2F-K, Movie S2), showing that elongating axial tissues indeed push on the PD. This pushing force is also dependent on cell movement and the pPSM (Figures 4G). Notably, as the PD is the source that supplies new cells to the pPSM, the axial push might impact the

dynamics of new PSM cells on the midline. In some PD gel replacement experiments, as the axial tissues advance the gel is seen to move aside from the midline and into the paraxial pPSM ([data not shown](#)), raising the possibility that new PSM cells might respond to the axial push in a similar fashion. To test this, we replaced the PD and midline PSM domain of a transgenic host embryo by a different colored donor ([Figure 4H; Huss et al., 2015](#)). The transplant appears flattened on the anterior similar to gel deformations, suggesting tissues respond to the high stress at the axial-PD interface. Cells at the anterior lateral sides of the transplant start to move medial-laterally into the pPSM. These observations suggest that the axial push may be required for driving the medial-lateral PSM cell flow.

Axial push is required for new PSM cell addition to the posterior PSM

To test the effect of the axial push on the midline PSM cells, we analyzed the average PSM cell movement in the simulation. The model recapitulates the long-term trajectories of PSM cells including their medial to lateral movement from the midline ([Figures 5A-C](#), see also [Figure 1E](#)). In the simulation, there are no additional assumptions beyond cell motility and mechanical pushes between cells. Therefore, the medial to lateral movement is promoted simply as the elongating axial tissue pushes the PD posteriorly and occupies the midline space ([Movie S4](#)). Indeed, our cell tracking (both in simulations and imaging experiments) shows that, following axial ablation, the medio-lateral (ML) speed of PSM cells exiting the midline is reduced along with the elongation rate ([Figures 5B-E, S5I, Movie S6](#)), indicating that the rate of PSM growth is slowed down. Furthermore, the motility gradient is not sustained ([Figure 5F](#)) due to reduced addition of motile cells. These results suggest that the axial tissues not only push the PD posteriorly, but in doing so also promote cell addition into the pPSM thus extending PSM length and sustaining the motility gradient. However, a more definitive test of this hypothesis would require direct application of pushing forces on the PD in place of the axial tissues.

Externally applied axial push rescues PD posterior movement and cell addition to the PSM

To apply an ectopic pushing force on the PD, we first performed an ablation of the pNC and then inserted a steel pin through the NT at the ablation site immediately anterior to the PD ([Figure 6A](#)). The main portion of the pin resides in the semi-liquid culture medium while the remaining tip portion stays at the embryonic tissue level. This tip can rotate in the opposite direction of the

main portion at the vitelline membrane entry point (Figure 6B). In this way, the pin acts as a lever which can be controlled remotely using magnets, allowing a pushing force to be applied on the tissue towards a specified direction (Figure 6C). Using alginate gels (1-10kPa) in place of embryos under the same experimental set-up (Figure S6A), we found that the pin creates strains comparable to those induced by pNC in softer gels (Figure S6B), suggesting the ectopic pin stress is higher than the endogenous forces. Nonetheless, the NT and PD stayed largely intact despite some tear near the pin insertion site (Figure 6C). Embryos with clean and minimally invasively inserted pins were incubated for 4 hours with or without the magnets and then imaged without the pin (live imaging was not feasible with the pin and magnet setup). In the absence of the magnet, the pin stands straight and axis elongation and PSM cell addition are slowed down as expected in pNC ablation experiments, indicated by a reduction of both PD posterior movement and medial to lateral spread of midline PSM cells (Figures 6D-G). Strikingly, when the magnetic field was present to let the pin push along the posterior direction onto the PD (Figure 6C), both the posterior movement of the PD and the medial lateral spread of PSM cells were partially restored (Figures 6D-G). This rescue experiment shows that an external mechanical push can mimic the axial pushing force to drive the PD to move posteriorly and to increase lateral addition of new PSM cells.

Coupling cell dynamics and tissue forces results in a self-sustaining elongation engine

Our findings show two mechanical interactions between tissues in the elongating body axis: the medio-lateral compression of axial tissues by the pPSM and the antero-posterior push on the PD by axial tissues (Figure 7A). In addition, the axial push drives new PSM cell addition to sustain pPSM growth and cell motility, thus forming a positive feedback loop (Figure 7B). While these forces do not represent all processes that drive elongation in these tissues, their simple cross-talk uniquely ensures a tight coordination between axial and paraxial tissues, which would be difficult to achieve with separate tissue-intrinsic elongation programs. A closed loop implies that pPSM will feedback to itself. To test this, we looked at PSM cell addition in pPSM ablated embryos. Indeed, lack of pPSM delays the lateral movement of PSM cells even with intact axial tissues (Figures 7C-E, Movie S7), and the axial pushing force is reduced when pPSM is ablated or loses cell motility (Figure 4G). These results show a tight linkage between inter-tissue mechanics and cell dynamics in this system. Our model also predicts that co-elongation will be

disrupted (i.e. length offsets between tissues will emerge) as a result of breaking the feedback loop (Figure S6C, Movie S4). Indeed, ablating the PD, while itself not causing any immediate tissue length offset and allowing continued elongation (as pPSM compression remains intact in short-term), results in NC and NT that elongate further than the PSM (Figures 4D, S6B). Conversely, ablating axial tissues causes the pPSM from both sides to merge on the midline and continue to grow from the PD. This results in an axial PSM that elongates further than the NC/NT (Figures S6D-F). Coupled together, these mechanical interactions produce local mechanical heterogeneity causing distinct spatial-temporal dynamics of different tissue ablation sites, which we reproduced both *in silico* and *in vivo* (Figure S7).

DISCUSSION

Cellular responses to extrinsic morphogenetic forces

Our study identifies inter-tissue forces as a key element in coordinated morphogenesis. This opens new avenues of investigation of specific cellular mechanisms that respond to these forces, and potential cross-talks with other cellular changes taking place at the same time. For example, for new PSM cells, the exit from the PD and migration into the pPSM are not only cell movement but also accompanied by drastic changes of the cell state. During primitive streak regression, PSM progenitors located in the anterior primitive streak region upregulate Wnt and FGF signaling, undergo an epithelial to mesenchymal transition (EMT) and gain enhanced motility. After tail bud formation, the PD territory becomes located at the tip of the NC and NT (the posterior wall of the chordo-neural hinge) which expresses both *Sox2* and *Brachyury* (a neural and a mesodermal marker, respectively) and continues to produce motile mesodermal precursors at a smaller spatial scale (Sun et al., 1999; Tzouanacou et al., 2009; Olivera-Martinez et al., 2012; Denans et al., 2015; Oginuma et al., 2017). In fish, the equivalent of the PD (called Dorso-Medial Zone) exhibits coherent cell movements (Lawton et al., 2013) suggesting an epithelial-like organization similar to that of the epiblast. More posteriorly in the tailbud, cells move incoherently, become specified to PSM and migrate laterally, causing increased tissue fluidity (Mongera et al., 2018), which may propagate mechanical information to anterior tissues (Das et al., 2019). We showed that the axial pushing force on the PD regulates this lateral migration of new PSM cells in chick. However, whether a mechanistic link exists between the

axial force and the EMT of progenitors remains to be investigated as the movement does not directly measure PSM production. On one hand, we found that in the absence of axial tissues, the PSMs from both sides merge and continue to grow from the PD and eventually form somites along the midline (which are larger and mis-located) indicating normal PSM differentiation, suggesting that the axial force controls morphogenesis but not cell fate specification. On the other hand, we found a decrease of PSM cell motility in axial ablation experiments suggesting that the axial push may contribute to the upregulation FGF signaling in the PD, which is important for both EMT and PSM cell motility (Sun et al., 1999; Bénazéraf et al., 2010). Interestingly, mechanically driven FGF upregulation was recently reported in the context of branching morphogenesis in the embryonic lung explants (Nelson et al., 2017). Our simulations also predict the force experienced by single cells through their movement in the tissue which may influence their differentiation, in particular we predicted PSM cells to sense a reversing net force over time. Future studies involving imaging of force or signaling reporters and *in vivo* mechanical perturbations will be important to test these possibilities.

Force generation by cell dynamics in the posterior PSM

The physical mechanism linking the movements of PSM cells and the tissue-level expansion remains to be clarified. A jamming-transition model relates PSM cell movement to tissue mechanical properties (Mongera et al., 2018) but does not account for the source of internal stress that drives expansion. In our computational model, the force arises as highly motile cells collectively increase the space they occupy through agent repulsion. While this simple assumption proved effective for us to connect the observed compression and its dependency on cell motility, and to subsequently recapitulate tissue shape changes and cell trajectories, the process *in vivo* is likely different and involves the remodeling of the extracellular matrix (ECM) driven by the net sum of forces exerted by single cells. Consistent with this hypothesis, fibronectin fibers in the posterior PSM follow a movement pattern that is identical to the average movement of the cells (Bénazéraf et al., 2010). Unlike the cells, the fibers do not show local random movement. Our 2D model also does not include the surface and interface ECM between tissues, whose mechanical role is supported by the observations that knocking-down fibronectins leads to decoupling of PSM and axial tissues, causing defects in NC and NT morphogenesis in zebrafish (Dray et al., 2013; Guillon et al., 2020). These elements may be

added in a future agent-mesh hybrid model and extended to 3D.

In addition to producing compression on the lateral sides, the pPSM expansion also creates an anterior to posterior pushing force alongside the axial tissues from the underlying random cell motility (Regev et al., 2017). The presence of this force is supported by the observation that PSM tissue can elongate on its own posteriorly without NC (Charrier et al., 1999; this study). The importance of this force in elongation is further evidenced by the ability of PSM to elongate to some extent even when PD is ablated (i.e. no new PSM cells, Bénazéraf et al., 2010; this study). While this pushing force appears weaker compared to the axial force as gel implants deform most in the midline, it is sufficient to continue elongation by pushing the PD posteriorly as a joint midline PSM (this study). To further dissect the contribution of axial and paraxial forces in elongation, it is important to design and adapt quantitative tools of mechanical measurement for very small (orders of 10-100 μ m) live embryonic tissues (Serwane et al., 2017; Mongera et al., 2018).

Robustness and self-maintenance of axis elongation in a coupled system

Our findings reveal a mechanical positive feedback loop coordinating elongation across germ layers. This situation is reminiscent of that observed during elongation of the drosophila embryo where extrinsic forces generated by the mesoderm and the invaginating gut, coupled with intrinsic forces from cell reorganization, lead to AP extension of the epithelial germ band (Butler et al., 2009; Lye et al., 2015; Collinet et al., 2015). Whether similar mechanical coupling occurs in the body axis of other vertebrates remains to be investigated, as there are many tissue and cellular differences between species and stages of elongation (reviewed in Mongera et al., 2019). It is tempting to speculate that our model may also play a role in other contexts because it provides some advantages in addition to coordinating elongation rates. For example, the FGF-dependent PSM cell motility sets a relationship between pPSM expansion and cell density, such that the compression on axial tissues increases when motile cells become more packed. This means that temporarily slower axial convergence will elicit stronger compression leading to correction. Such a mechanism could also play a role in ensuring the robustness of straight elongation (bilateral symmetry), as bending to one side will encounter increasing resistant pressure that increases with curvature, making it difficult to form large curvatures. This might provide a mechanical explanation to the recent observation that disordered cell motion in the

posterior PSM is required for bilateral symmetry of paraxial mesoderm formation ([Das et al., 2017](#)).

Our model also explains the self-maintenance of axis elongation. Transplantation of the Node to an ectopic location such as the area opaca can induce a new embryonic axis which undergoes posterior elongation similar to that of the endogenous AP axis ([Waddington 1932](#)). Our simulation shows that the generation of elongation behavior does not necessarily require either the formed portion of the axis or any global cues that direct cell movement (while they could still contribute to elongation, [Yang et al., 2002](#)), but only a minimal set of tissues (namely pPSM, pNC/pNT, and PD) arranged around and inducible by the Node (the initial symmetry-breaking). As long as a PD (the fuel tank) that produces high motility PSM cells (combusting fuel) continues to function under the corresponding inductive signals, the positive feedback loop (the engine) can kick start locally and drive posterior Node movement, leaving a patterned body axis behind. Consistent with this analogy, elongation can terminate naturally when progenitors run out due to suppression of signals supporting PD proliferation (no more fuel, [Denas et al., 2015](#)), or in situations when signals (e.g., FGF) supporting the “combustion” are depleted ([Bénazéraf et al., 2010](#)).

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AUTHOR CONTRIBUTIONS

F.X., L.M. and O.P. planned the study. F.X. and W.M. built the models. B.B. contributed dnFGFR1 data. F.X. performed the experiments, analyzed the data and prepared the manuscript with O.P. and input from other authors. All authors participated in the interpretation of the data.

The authors declare no competing financial interest.

FIGURE LEGENDS

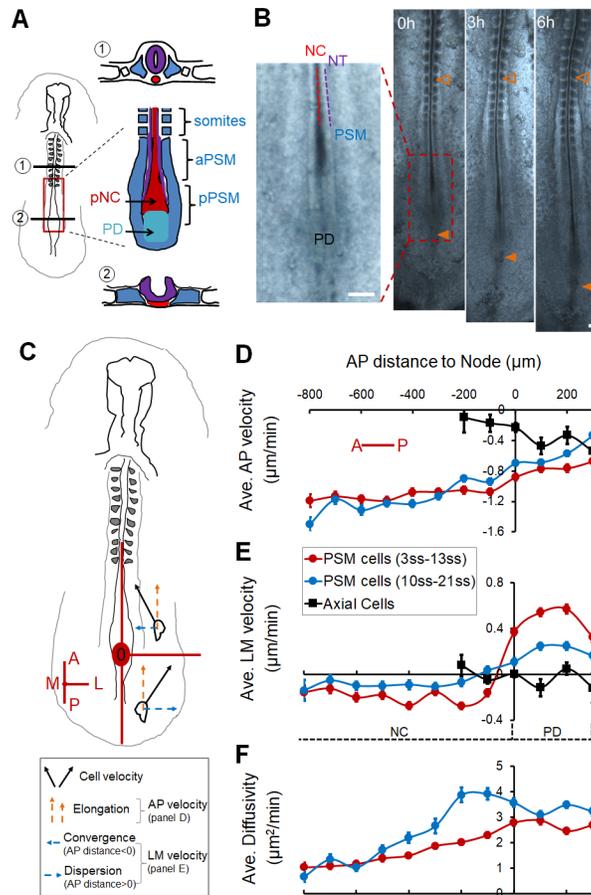


Figure 1. Avian posterior body axis tissues and their cellular dynamics during elongation

(A) Diagram of the avian body axis (Left) and its composing tissues (Right). Cross-section (①) shows the formed section of the body axis including a closed neural tube (NT, purple), rod-like notochord (NC, red) and somites (blue), with significant interstitial space between them. Cross-section (②) shows the newly formed section in the posterior near the progenitor domain (PD) including a folding neural plate (purple), a wider NC (red), and presomitic mesoderm (PSM, blue), which are in close contact. In ①, the uncolored small squares by the somites represent the intermediate mesoderm, the further lateral uncolored blocks represent the lateral plate mesoderm. The red box marks the posterior body axis shown in a detailed ventral view of the tissue arrangement. The NT and NC are in the middle anterior to the PD, flanked by two stripes of PSM and columns of somites. A lowercase letter "p" (or "a") in front of NC/NT/PSM indicates

"posterior" (or "anterior"), same in other figures.

(B) The elongating posterior body axis of a chicken embryo. On the left the magnified area shows the spatial arrangement of the PD and newly formed NC, NT and PSM, as illustrated in (A). On the right axis elongation is observed over time. Empty arrowheads mark the same somite as a reference; filled arrowheads marks the PD. Scale bars: 200 μ m.

(C) Schematic view of the reference axes used for plotting cell dynamics data. The Node area (red circle) is set as the origin (0,0). A, anterior; P, posterior; M, medial; L, lateral. In this reference frame body axis elongation is represented by posterior to anterior movement of PSM cells as they are left behind by the node (orange arrows). The posterior area to the Node (PD) has a positive AP position while the anterior area to the Node (NC) has a negative AP position.

(D-E) Average cell speed along the AP (D) and LM (E) axes. Along AP, negative indicates P to A movement relative to the Node (0,0). Along LM, negative and positive indicate convergent (L to M) and dispersive (M to L) movements, respectively. Each bin integrates 100 μ m along the AP axis averaging 100-200 speed measurements from all time points of the movie. Error bars: \pm SEM. Each movie is repeated at least 3 times. In each embryo, around 50 cells were tracked through ~200 time points.

(F) Cell "Diffusivity" obtained by fitting the mean square displacement (MSD) of single cell tracks with a drifting speed (directional, v) and a diffusion component (random, D) where $MSD=4Dt+v^2t^2$. t is time.

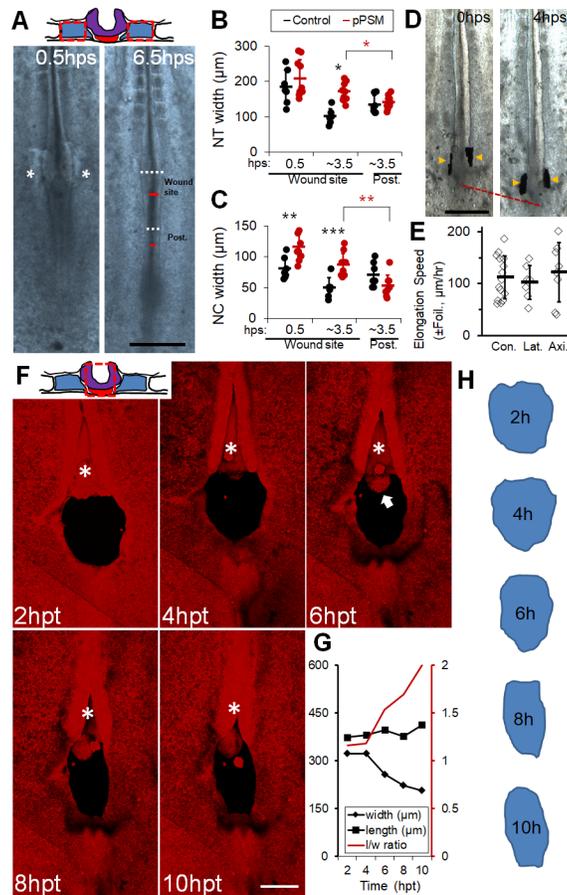


Figure 2. Posterior PSM exerts a lateral compression on axial tissues

(A) Ablation of the pPSM. The top schematic shows cross-section of the operated area. Red dashed boxes mark ablated pPSM (blue). Embryo images show axial tissue dynamics after pPSM ablation. Asterisks indicate the closing PSM ablation sites shortly after ablation (hps, hours-post-surgery). The dark line in the midline of the embryo delineates NC (the short red lines measure NC's width). Lighter regions besides the NC show the neural folds on the further dorsal side (the white dashed lines measure NT's width). The widths were measured for both the wound sites and a control area posterior to the wound sites (Post.). Scale bar: 500µm.

(B-C) NT and NC width after pPSM ablation (Control n=7, pPSM n=10). Asterisks indicate significant difference ($p < 0.01$, t-tests, unpaired (black), paired (red)). "Post." is a control area posterior to the wound site as annotated in panel (A).

(D) Foil insertion between pPSM and pNC/pNT. Foils (arrowheads) are vertically inserted between the tissues and are observed to move along during elongation. Scale bar: 500µm.

(E) Elongation speeds in Foil insertion experiments. Con., Control (no insertion); Lat., Lateral

insertion between pPSM and pNC/NT; Axi., axial cross-sectional insertion between pNC and PD (See [Figure S2E](#)). The average speeds of these groups are not significantly different ($p > 0.05$, t-tests).

(F) Gel implant followed by confocal timelapse. The top schematic shows cross-section of the operated area. Red dashed boxes mark ablated pNC (red) and pNT (purple). Images are maximum projections of GFP (shown in red) z-stacks (dorsal view). Asterisk shows the convergence and closure of NT anterior to the implant site. Arrow marks protruding NC underneath the NT. Scale bar: 200 μ m. See also [Movie S2](#).

(G) Quantified shape changes of the gel implant in (F). Length (l) is measured along the midline, width (w) is measured following the widest line perpendicular to the midline. Consistent narrowing was observed in n=3 movies.

(H) Contours of the gel implant in (G).

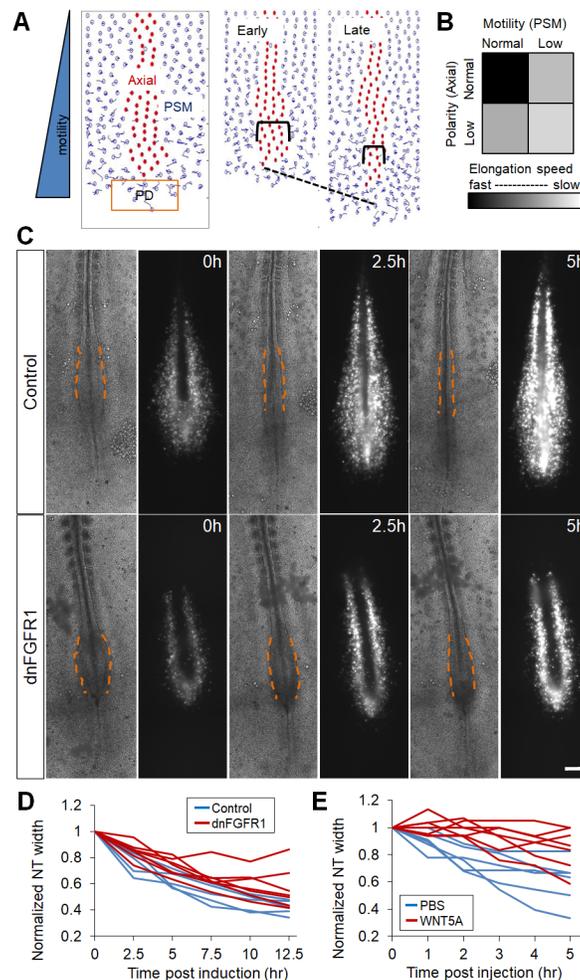


Figure 3. PSM cell motility and axial cell polarity contribute to axial elongation

(A) Agent-based 2D model for posterior body axis elongation. Cell centroids but not cell shapes are shown for the ease of visualization. NC/NT cells (red) are flanked by PSM cells (blue). New PSM cells are added from the PD domain (orange box) periodically on the midline. The simulation recapitulates axis elongation (dashed line) and axial convergence (brackets). See also [Movie S3](#).

(B) Elongation speed as a function of PSM cell motility and axial cell polarity in the model. Results were averaged from 10 simulation runs (4000 time-steps each) for each parameter pair.

(C) Inhibition of FGF signaling by electroporating a dominant negative FGFR1 and a GFP construct into the PSM. Bright field and GFP fluorescence images of representative embryos (ventral view) at three different time points are shown. Dashed lines contour the narrowing pNT behind the pPSM, dark strips in the fluorescent images are the NC tissue flanked by electroporated PSM cells. Scale bars: 200 μ m.

(D-E) Convergence of pNT after dnFGFR1 induction and after WNT5A injection, respectively. Each line represents one embryo. The width is normalized to the width at the first time point. The perturbed convergence rate was significantly lower compared to control in both cases ($p < 0.05$, t-tests).

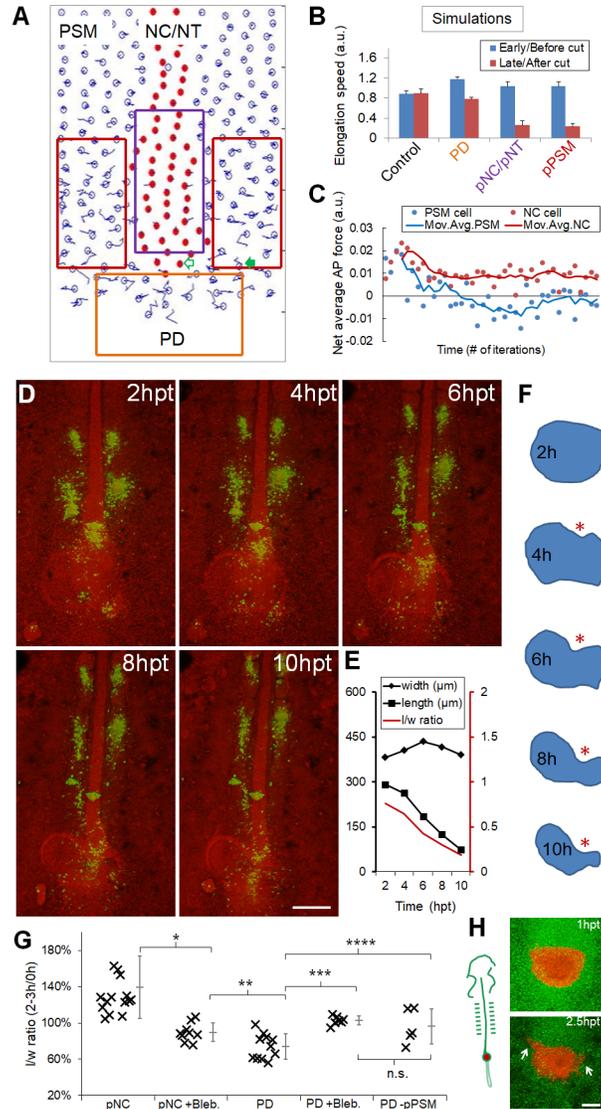


Figure 4. Axial tissues exert a pushing force on the PD

(A) Layout of the model showing simulated ablations (boxes) corresponding to results in (B). Empty arrow, an axial cell; filled arrow, a PSM cell. See also [Movie S4](#).

(B) Axial elongation speeds (\pm SEM) for both before and after tissue ablation. For PD, cell addition was disabled in the beginning of simulation. Speeds were averaged over 2000 time points in each simulation. Each experiment includes $n=40$ simulations. $p=0.93$, control; $p<0.05$, other groups (paired t-tests).

(C) Net forces along AP experienced by an axial cell and a PSM cell. The cells start at the same AP level as marked in panel (A). Positive indicates A to P. Data points are binned by 200 iteration windows. Curves: 4 times moving average.

(D) Gel replacement of PD followed by confocal timelapse. Green indicates spots of Dil injection to follow cell movements. Spots in the pPSM can be seen expanding while the axial tissues narrow. Scale bar: 200 μ m.

(E) Quantified shape changes of the gel implant in (D). Consistent observations in n=3 embryos.

(F) Contours of the gel implant in (D). Asterisk highlights the indentation where the gel is in contact with pNC.

(G) Summary of gel replacement experiments. The l/w ratios after 2-3 hours are plotted as a percentage of the initial aspect ratio. Tissue names indicate implant locations (e.g. pNC, PD), “+/-” indicates further perturbations (e.g. Blebbistatin treatment [+Bleb.], pPSM ablation [-pPSM]). Asterisks: p<0.05, t-tests. n.s.: p=0.4. See also [Figures S3O-P](#).

(H) Shape change and cell dispersion of PD transplant. A group of GFP+ progenitors (shown in red) is transplanted posterior to the pNC of cherry+ host (shown in green). Arrows: cells in the transplant disperse into host pPSM. Anterior flattening of the transplant and cell dispersion were observed in n=4/6 similar experiments. Scale bar: 100 μ m.

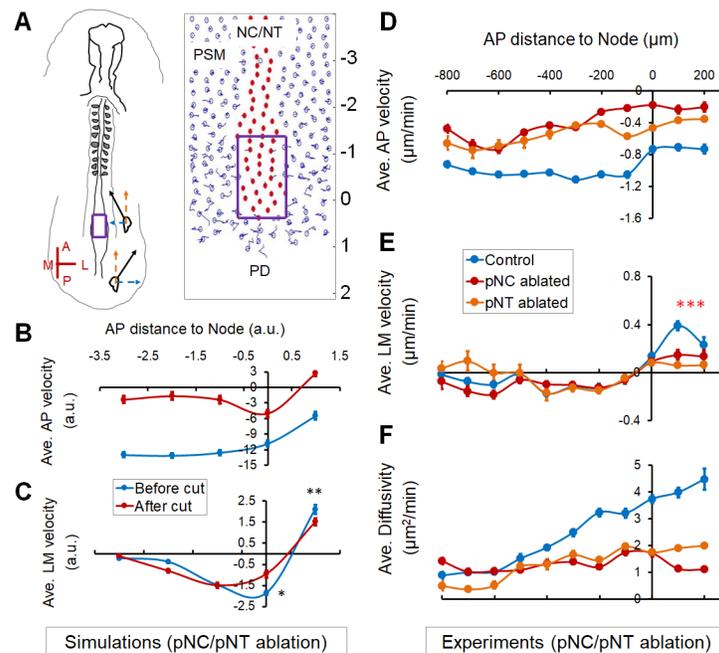


Figure 5. Axial push is required for cell addition from the PD to the PSM

(A) Axial ablation in the embryo and the model. Purple boxes label the surgically and digitally ablated areas, respectively. The AP axis labels of the simulation field (-3 to 2) correspond to the AP distances to Node in (B) and (C). See also [Movies S4](#).

(B) Modeled AP velocity (compare to experimental measurements in [D]), recapitulating

elongation (indicated by a negative AP speed) and slow-down of elongation after axial ablation. (C) Modeled PSM cell LM velocity (compare to experimental measurements in [E]), recapitulating convergence, midline-exit and reduction of midline-exit after axial ablation (n=40 simulations. Asterisks, $p < 0.05$, paired t-tests).

(D-F) Average PSM cell AP (D) and LM (E) velocity and diffusivity (F) under different surgical ablations (n=3 each). Measured and plotted similarly as in [Figures 1D-F](#). See also [Movie S6](#).

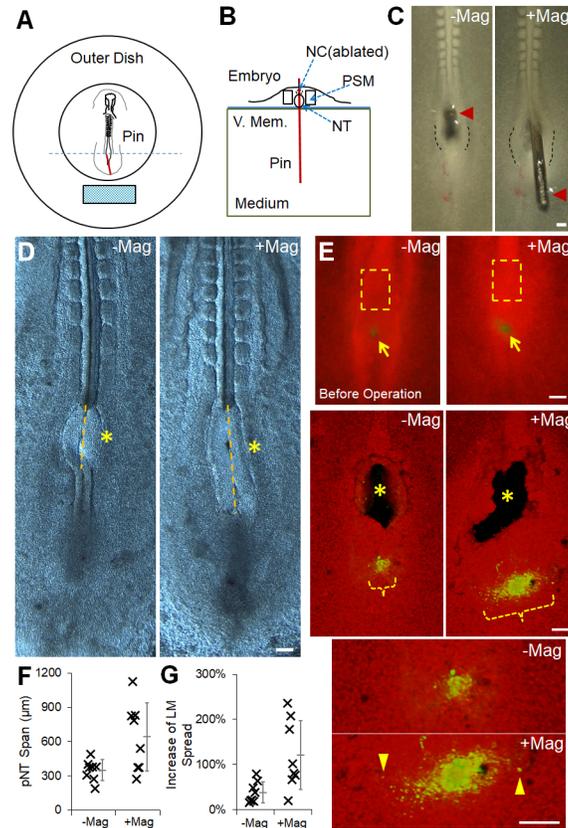


Figure 6. Axial push can drive PD movement and cell addition

(A-B) Schematics of the magnetic pin experiment set up. Top (A) and cross-sectional (B, corresponding to the section marked by the blue dashed line in [A]) views, respectively. Red line indicates the inserted pin and the cyan block shows the magnet (movable). The pin is perpendicularly inserted in the initial position. It rotates around the entry point of the vitelline membrane (V.Mem.). The medium is the semi-liquid albumin agar mix of the embryonic culture allowing pin rotation. See also [Figure S6A](#).

(C) Example of pNC replacement with pin (left) and the push along the posterior direction after magnet is moved in range (right). With magnet, the embryo can be seen stretched and NT walls

(marked by dashed lines) at the insertion site show expansion under stress. Scale bars: 100 μ m.

(D) Elongation after pNC ablation by the pin. Examples of phenotype after 4 hours with or without magnet. Pin was removed and the embryo was cleaned before imaging. Asterisks mark the pNC ablation and pin insertion site. Yellow lines mark the pNT span over the ablation site to measure PD posterior displacement.

(E) Cell dispersion followed by pNC ablation and pin insertion. Top images show the labeled spot (Dil, green) of ventral midline PSM cells (arrows). Before the microsurgery and incubation, they occupy a narrow LM range compared to the pNC (dashed boxes). Middle images are 3D projected confocal images showing the ablation/insertion sites (asterisks) and cell dispersion span (Orange bracket) after 4 hours. Note that the cells have dispersed to wider LM ranges than pre-operation in both experiments. Bottom images are enlarged views showing cells that have migrated significantly laterally (arrowheads).

(F-G) Quantification of elongation (F, $p=0.002$) and cell dispersion (G, $p=0.011$) corresponding to (D) and (E), respectively. t-tests.

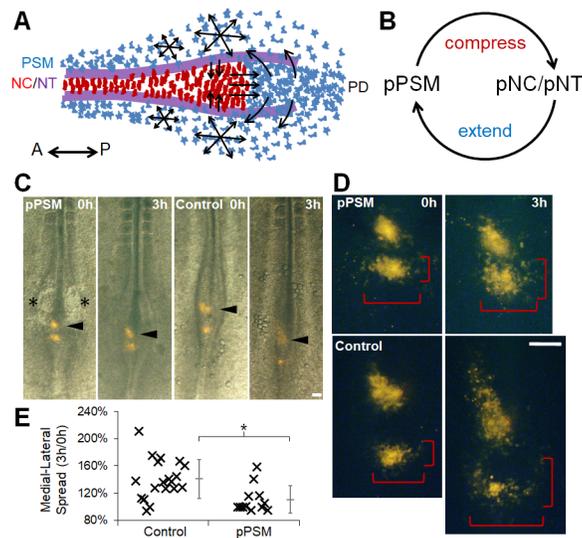


Figure 7. A mechanical feedback loop couples elongating tissues

(A) Schematic ventral view of observed tissue geometry and cell behavior in PSM, NC, NT and PD. Red cells: NC cells (polarized shape). Blue cells: PSM cells (mesenchymal). Purple lines: NT folds on the dorsal side. Arrows show cell movement.

(B) A conceptual mechanical positive feedback loop that couples co-elongation of tissues.

(C) Movement of labeled midline PSM cells. Two Dil-labeled clusters immediately posterior to the pNC are followed in control and pPSM ablation experiments (asterisks indicate the ablation

sites). After pPSM ablation, the labeled clusters move less posteriorly (arrowheads), and remain concentrated (cells did not disperse as much) as compared to the control. Scale bar: 100 μ m.

(D) A close-up view of labeled clusters showing dispersion of single cells. The brackets measure the AP and LM spread of the cells over time. Scale bar: 100 μ m. See also [Movie S7](#).

(E) The LM spreads measured as in (D) are compared between control and pPSM ablation groups as a ratio between 3h and 0h. Asterisk: $p=0.004$. t-test.

STAR METHODS

KEY RESOURCE TABLE

RESOURCE AVAILABILITY

Lead Contact

Further information and requests for resources and reagents should be directed to and will be fulfilled by the Lead Contact, Olivier Pourquié (pourquie@hms.harvard.edu).

Materials Availability

This study did not generate new unique reagents.

Data and Code Availability

The published article includes most datasets generated during this study. Additional replicas and source datasets are available upon request. The computational model is publicly accessible via the link provided in the Key Resource Table and methods.

EXPERIMENTAL MODEL AND SUBJECT DETAILS

Chicken eggs

Wild type chicken eggs were supplied by Charles River Laboratories. Tg(CAG-GFP) ([McGrew et al., 2008](#)) chicken eggs were provided by Clemson university (originally by University of Edinburgh). Tg(PGK1:H2B-chFP) ([Huss et al., 2015](#)) quail eggs were provided by Ozark egg company. Eggs were kept in monitored 15°C fridge for storage and 37.5°C ~60% humidity egg incubators for incubation. HH stage 4-12 embryos were used. Sex as a variable is not relevant at the stages studied. All procedures have been approved by Harvard Medical Area (HMA) Standing Committee on Animals.

METHOD DETAILS

Embryo preparation

Whole embryo explant cultures were prepared using filter paper extraction ([Chapman et al., 2001](#)). Two adjacent 0.5cm holes are punched on a 2cm x 2cm piece of filter paper

(Whatman). Eggs were opened into a petri dish where the embryo will be found near the top of the yolk. The thick albumin is swept aside gently with small filter paper pieces with a tweezer. The holed filter paper is then lowered to attach to the vitelline membrane with the holes allowing embryos to be viewed. The vitelline membrane is then cut around the filter paper to release the embryo. The filter culture embryo is then rinsed in PBS to remove excess yolk. Cleaned embryo is placed on a 3.5cm petri dish containing 2ml of semi-liquid culture media made with the following formula (per 100ml of culture media): Part A: 50ml Albumin (beaten for 15min) then supplement with 0.8ml 20% D-Glucose (Sigma); Part B: 0.3g BactoAgar (Sigma) solved in 50ml water in a microwave then supplement with 1.23ml 5M NaCl. Warm part A and cool part B to 55°C in a water bath. Mix thoroughly and add to petri dishes before gelation. The embryo cultures are maintained in a slide box with wet paper towels in the 37.5°C ~60% humidity egg incubators except when surgeries or snapshot imaging are performed.

Dye labeling, injections, and electroporation

3-14ss embryos were used for cell labeling. 2.5mg/ml Dil in ethanol was diluted in PBS to 0.5mg/ml before injection. The injection solution was loaded into a sharp-tipped glass needle and injected by mouth pipetting from the ventral side of the embryo into spots in different tissues (e.g., PSM, NC, the Node, and primitive streak). For over-expression of transgenes, DNA solutions (pBI-EGFP, pBI-FGFRdn and pCIRX-rtTA-Advanced, [Bénazéraf et al., 2010](#)) were injected from the ventral side into the anterior primitive streak of HH stage 4-5 embryos, followed by electroporation using a NEPA21 type II electroporator (NEPA GENE) in PBS. The pulse setting is at 6V, 150ms interval, 25ms duration per pulse, and 3 pulses per embryo. For PSM expression, the electrode targets the anterior primitive streak (the injection site). For NT expression, the electrode targets the area slightly anterior and flanking the Node. Embryos were checked for fluorescence and development 12hr later. Poorly developed and low/mis-expressed embryos were excluded from imaging and analysis. For the induction of dnFGFR1 expression in the PSM, embryos were incubated on plates with 2µg/ml of doxycyclin (Sigma). Imaging was started 2-3hrs after the incubation started to allow transgene induction. FGF signaling inhibitor PD173074 (Sigma) was solved in DMSO and diluted to 2µM in PBS before use. ~20µL solution or DMSO control was dropped on each cultured embryo on the ventral side. Blebbistatin (Sigma) was diluted to 70µM in PBS from DMSO stock (25µM) before use.

~50 μ L solution or control was dropped on the ventral side. For injections of PBS, WNT5A and WNT10 (R&D systems, recombinant human/mouse), 10 μ g/mL protein solution in PBS was loaded in a glass pipette and injected into the anterior portion of the pPSM close to both sides of the NT/NC, or into the NT apical side directly through the NC and floorplate.

Surgeries, gel implants, and pin experiments

8-12ss embryos were used for surgery experiments. Surgeries were performed under a Leica M90 dissecting scope. Cutting was performed with an electrolysis sharpened steel or tungsten needle from either the ventral or dorsal side, for PSM/NC or NT, respectively. In the ventral surgeries, endoderm was cut open first and gently peeled aside, followed by shallow cutting into the mesoderm while avoiding damaging the neural ectoderm and epiblast. In the dorsal surgeries, the vitelline membrane over the surgery site was first slit in the middle and gently peeled on either side to make a triangular shape opening, followed by cutting around the neural plate/NT area, care was taken to minimize damage to NC and endoderm. The wound areas were gently brushed by the needle to removal cells. For transplant, the comparable region of the donor was cut intact and moved out of wound with a tungsten rod. The tissue was then transferred with a micropipette loaded with PBS to the host. A smaller opening was cut open at the desired location and the transplant was gently pushed in. For gel implant, the cleared cut opening either received an injection of 1% (w/v) alginate sodium (Sigma Aldrich) solution, or a small piece of pre-formed soft alginate gel (40:1 mix with 5% (w/v) calcium chloride solution). Injected alginate solution formed gel with calcium ion in the embryo culture and integrated much better than transplant of pre-formed gel. The deformation results of both methods were qualitatively similar (e.g., transplanted: [Figure S2D](#); injected: [Figure S2E](#)) even though the transplanted gel (higher stiffness, ~100 Pa, [Banerjee et al., 2009](#)) shows less deformation. Gels prepared with higher concentration of calcium ions (20:1 or 10:1 mix with 5% (w/v) calcium chloride solution) or agarose gels do not show significant deformation as tracked by the movement of encapsulated beads (For example, 0.5% agarose with a modulus in the order of 1k Pa [[Markert et al., 2013](#)], [data not shown](#)). Between 20 to 45min of healing and integration was allowed before mounting the host for imaging. For steel pin (MINUCIE, stainless, No.15, ~40 μ m) insertion, the tip of the pin is cut to around 1cm long. The pin is held with a tweezer and inserted between the neural folds into the culture medium from the ventral

side after pNC ablation until only a small head remains level with the embryo. The embryo culture dish is placed in a bigger dish where the magnet is placed. Both distance and angle from the embryo to the magnet are adjusted by hand under a dissecting microscope to get optimal pin movement. The magnet and angle are fixed once proper pin rotation and tissue deformation under the pushing force can be seen and the configuration stays the same through a 4hr incubation.

Imaging

Bright field and fluorescent images of the embryos were taken with a Leica M205 fluorescent microscope. Embryo culture dishes were kept in a slide box with wet paper towel. The box was kept in the egg incubator at all times except image taking (~5 minutes for 10 embryos). For confocal timelapse imaging, the embryo (cultured as explant on filter paper) was mounted ventral or dorsal side down on a 50mm glass bottom dish (MatTek) pre-warmed to 37.5°C. A thin layer of albumin-agarose medium (200µl) was plated on the cover glass before mounting. The sample was then imaged on an inverted Zeiss confocal 780 with heating stage and environmental chamber to maintain temperature and humidity. Typical imaging set up for movies such as [Movie S1](#): Imaging space: 1.4x1.4x0.15mm; voxel size: 1.4x1.4x2.5µm; time coverage: 15-20h; temporal resolution: 2-3min; lasers: 561nm 0.4%, 488nm 3%.

Image Analysis

Dissecting scope images were analyzed by drawing lines over features of interest (e.g., length, wound width, notochord width, etc.) in PowerPoint (Microsoft) or Fiji (ImageJ). For elongation, the AP level of somite 2-3 were chosen as the reference point (except where it is noted differently in Figure legends). The distance increase between the reference point and the Node was used to calculate elongation speed. For confocal timelapse movies, the original .ism (Zeiss) files were rendered into 3D maximum projection movies in FluoRender and compiled to movies in Fiji. Cells were tracked on this movie with the Manual Tracking plugin in Fiji. For cell density measurements, region of interest was selected from PSM z-stack images for fluorescent intensity or nuclei count in Fiji. Tracking results and measurements were processed in Matlab (Mathworks) with custom scripts. A track of the moving Node and a track

of the anterior midline were used to linear transform the coordinates of cells to anterior-posterior / lateral-medial axes before all position and speed calculation.

Modeling

All models were coded in Matlab. Detailed documentation is provided below. The model can be downloaded following the link in the Key Resource Table.

1. Model design and protocol

The goal of the model is to test what types of tissue behavior emerge from cells behaving under simple rules. Tissue geometry and cell activities break the homogeneity and symmetry and underlie these behaviors, although they act with a number of other interactions that should be accounted for in the model. For simplicity and feasibility, the tissue is modeled as a field of cells on a 2D plane with boundaries. The field is seeded with a finite number of cells (much fewer than in actual tissues for feasibility but geometry and proportions are kept) with user defined properties including position, velocity and cell type. Time is introduced as number of iterations for the field to evolve. At each iteration, a cell receives a net "force" defined by its position, previous velocity, type, and motility (eq.1, implementation of each term is explained in section 2). This force causes it to change position and velocity. Collectively the field evolves to a new pattern. Additionally, functionalities are build-in to allow new cells to be added or specific cells to be removed during simulation, to provide tools for assessing progenitor addition and tissue surgery experiments. A chemotaxis gradient generator is also included but was kept off for the present study.

$$\frac{dx}{dt} = \sum I(x) + \sum B(x) - \beta v + R(x) \quad (1)$$

To use the model, copy all provided scripts to the working folder of Matlab. Run `simulation.m`. In this script, a user can define the geometrical and interaction parameters (to be discussed in detail in section 2), as well as number of iterations in one simulation, rounds of independent simulations, and whether a perturbation cut on the tissue is performed (to be discussed in detail in the section after the next). To create a movie, specify a "moviename" string variable and use command:

```
">>makemovie2(Data,N+NewN,chemoN,tipN,NCN,boundary,moviename)"
```

2. Model assumptions and implementation details

The model includes both symmetry breaking and non-symmetry breaking assumptions. The former dictates the emergent patterns of the field. The assumptions and their rationale, impact and implementation are described below:

Non-symmetry breaking:

1. Repulsion between cells: when cells move close to each other, a repulsion force is generated as a function of distance. This assumption shows the basic volume constrain where no two cells could occupy the same position. This assumption provides the basis for force transduction in the system and is key for implementation of several other assumptions. It alone however does not cause any symmetry break or pattern evolution. The strength of repulsion is controlled by parameter **alpha**. The interaction term for a given cell (I_i) is calculated as eq.2:

$$I_i(x) = \alpha \cdot \sum_j \frac{r_{ij}^{-R}}{r_{ij}} \quad (r_{ij} \leq R)$$

$$I_i(x) = 0 \quad (r_{ij} > R) \quad (2)$$

In which $r_{ij} = |x_i - x_j|$. R is the cell diameter.

2. Viscosity/friction: a resistance against the cell's velocity is added at each simulation, to mimic the fact that cells move in a viscous tissue environment where inertia is not important. This force ensures that simulated cells do not project out of the tissues. The strength of viscosity is controlled by parameter **beta**. In eq.1, the calculation gives a force in the opposite direction proportional to the step of movement (v) from the previous time point.

3. Repulsion between cells and boundary: when cells run into a boundary it is bounced back similarly as it runs into a cell (eq.3). This repulsion is not particularly important for the simulation. The parameter controlling which is set to equal **alpha**. In addition, the size of tissue field and cell field can be set by parameters **boundary** and **boundaryc**.

$$B_i(x) = \alpha \cdot \frac{r_{ib}^{-R/2}}{r_{ib}} \quad (r_{ib} \leq R/2)$$

$$B_i(x) = 0 \quad (r_{ib} > R/2) \quad (3)$$

In which $r_{ib} = |x_i - x_b|$. x_b is the location of the boundaries.

Symmetry breaking:

1. Geometrical layout: in the standard configuration, the model places the NT/NC cells (analogously referred to as NC cells in subsequent descriptions and the model codes) in between two flanking PSM cell groups (controlled by parameter **NCN** assigning NC cell type to the designated cell ID numbers). The progenitor domain that adds new PSM cells is placed

posterior to the end of NC cells (controlled by parameter **tipN**). In the standard configuration, the parameter **tipN** restricts a particular NC cell (the **tipN** cell) at the posterior (open) end of the tissue to only move anterior-posteriorly (no free medial lateral movement that could arise due to small number of cells). A medial-lateral line posterior to the **tipN** cell at a specified distance defines the entry point of the new cells, thereby serving as the PD domain in the simulation. This layout is analogous to the actual embryo.

2. New cell addition: in the standard configuration, the model adds 80 new PSM cells to the existing field of 200 cells (controlled by parameters **N**, **NewN**). The cells are added periodically with parameters available to control the frequency, number and pattern of addition in **gastrulation.m**. At each time when new cells are to be added, their AP locations are determined by the location of the **tipN** cell, thereby following the movement of the PD. The **tipN** cell and the new cells otherwise follow the same mechanical interactions as any other cells in the field. This reflects the actual constant PSM cell production in the progenitor zone and the new PSM cells emerge from the dorsal side to the ventral midline. In the current study the cell addition is either stable throughout simulation or shut down completely following removal of the PD.

3. Random motility gradient across the field in the PSM cells: in the standard configuration, the model implements a simple linear motility gradient as a function of the PSM cell's distance to the PD/pNC boundary (as marked by the location of the **tipN** cell). The motility is produced with a random number to control direction and magnitude, with parameter **gamma** controlling overall magnitude (eq.4).

$$R(x) = \gamma \cdot (C - |x_{AP} - x_{AP}(tipN)|) \cdot rdn(1) \quad (4)$$

NC/NT cells are assumed to not have this motility. This assumption is justified by the observation of the exact type of motility gradient in the PSM in previous studies and the current study (Figure 1), and its known regulation by FGF signaling (also validated in this study, Figures 3,S3). This assumption includes cell property and a geometry asymmetry and is the **key assumption** of the model.

4. Polarity of NC cells: in the standard configuration, the model takes into account NC cell's medial-laterally polarized shape by using a parameter **ncp** to bias the intercalation preference, implemented as differential repulsion strengths along different axes. When the value of **ncp** is bigger than 1 it means a medial-laterally elongated cell that would intercalate more easily

medial-laterally than anterior-posteriorly (eq.5). This assumption is justified by the polarized NT cell shapes in previous studies and its known regulation by WNT/PCP signaling (also validated in this study, Figures S4,S5).

$$I_{AP}(x) = ncp \cdot \alpha \cdot \frac{|x_{AP}(i) - x_{AP}(j)|}{r_{ij}}$$

$$I_{LM}(x) = \left(\frac{1}{ncp}\right) \cdot \alpha \cdot \frac{|x_{LM}(i) - x_{LM}(j)|}{r_{ij}} \quad (5)$$

5. Adhesion in the anterior: an adhesion gradient from anterior to posterior in the PSM (implemented as reduced repulsion in the anterior) is used to mimic the same observed properties *in vivo*. The simulations however show that its impact on elongation is minimal. This is consistent with the observation that the main driver of elongation is the posterior PSM where adhesion is low. For this reason, adhesion is not further investigated in the model or tested in wet lab experiments in this instance.

3. Simulations

This section describes a more detailed account for the simulations. After specifying parameters described in the previous section. The user goes on to determine number of iterations (time). Sufficient time is often needed for the field to evolve into a stable state. The sum of parameters **Tcut** and **Tcut2** is the total number of iterations. This is split into two parts as the algorithm provides an opportunity of tissue cutting perturbation in between, so that the model takes the specified removed cells out of the interactions during **Tcut2**. To specify a region to cut, the user can either select one that is pre-specified in the script (cutmatrix) or write a new one. Note that this region should be a non-zero-area box (or boxes) that are within the **boundary** parameter otherwise it will not have an effect. By controlling parameter **sc**, the user can also specify how many independent runs the simulation runs.

4. Result analysis

The output of **simulaiton.m** includes a cell array named "**Data**", in which the cut perturbation info and the whole simulation are recorded. A user can obtain any cell's position and transient velocity as a function of time (iteration number) and use logic index to obtain information of interest. The elongation speeds, time average cell velocities and regional density changes are computed from "**Data**" using simple custom scripts.

QUANTIFICATION AND STATISTICAL ANALYSIS

Statistical analysis summary

Quantitative results generated from the image analysis were sorted into experimental groups and Student's t-test is performed with Excel (Microsoft) and Matlab (Mathworks) to compare the results with $p < 0.05$ considered significant difference. The type of statistical tests, exact value of n , definition of center, and dispersion and precision measures are all precisely defined in each figure legend in the manuscript.

Statistical considerations in study design

Sample size: For qualitative yes/no tests where the distinction is clear (e.g., direction of gel deformation) and no exceptions were observed, sample size is at least 3. For quantitative comparisons (e.g., elongation speed), at least 3 samples per group are used per experiment. ≥ 5 samples are used to in key groups. This sample size is in line with literature and is preferred for consistency and minimizes system errors that might be introduced between groups when total experiment time becomes too long (> 2 hrs), taking into consideration the time consuming procedure of embryo preparation and individual surgical operations. Replications were performed to confirm the results. For cell behavior analysis, > 30 individual cells were tracked through the duration of the movies (hundreds of time points). Before applying t-tests, the measurements were normalized to the mean and pooled for a chi square goodness of fit test. The tests suggest that the variabilities observed in cell speed, elongation speed, convergence speed, tissue width, and cell spread do not come from a distribution significantly different ($p > 0.05$) from a normal distribution. Data exclusions: NO data point exclusions in any analysis. In sample preparation and before data collection, embryos that show unspecific developmental abnormalities or lack of labels (indicating failed electroporation or injection) were discarded. This health and technical screen is applied in an unbiased manner to all experimental and control groups. Replication: All wet experiments reported have been replicated at least once (key experiments 3 or more times) with a different batch of eggs (shipment in a different week) and new preparation of reagents (culture and imaging plates, injection needles and labeling reagents (different aliquots of the same source)). The same dissecting tools, imaging protocols and microscopes were used during replications. Modeling experiments were replicated at least 10 times for each type of simulation run. Randomization

and Blinding: In each embryonic experimental test involving groups of embryos, all embryos were incubated, prepared and handled together except for the particular experimental condition(s) being tested. After an initial health screen of all embryos, the investigators do not use any criteria or variables when allocating embryos into control and experimental groups. Blinding is not possible for surgical perturbations on the embryos as data collection and analysis require the investigator to look at the images where the type of surgical perturbation is clearly visible. For chemical treatment and injections on the embryo, a look-up table (relating operation to sample number) is generated at the operation time and is not referred to during data collection and analysis. Only when analysis is complete is the look-up table linked to sort the results into different experimental groups. Blinding is not applicable in confocal movie experiment or modeling as the test conditions are clearly affiliated with the samples throughout data acquisition and analysis.

SUPPLEMENTAL MOVIE LEGENDS

Movie S1. Cell tracking analysis of paraxial and axial cells, related to Figure 1

The movie on the left is a 3D maximum projection (ventral view) of a Tg(CAG:GFP) (shown in red) chicken embryo imaged for the indicated duration (starting ~3 somite stage). Green (Dil labeled) cells start as clusters at the injection site. This movie is oriented to have the anterior on the left. NC in the center can be seen moving posterior over time, followed by somite formation on both sides from the PSM. Dil labeled PSM cells disperse from the midline progenitor domain (PD) and join pPSM on both sides. On the right side the movie shows the trajectories of tracked cells of the same dataset. The solid red sphere marks the end of NC. The tracks over time are shown in [Figure S1C](#) and two time points of the movie are used for the fate map in [Figure S1D](#). Time stamp: hh:mm. Scale bar: 100 μm .

Movies S2. Tissue and gel dynamics under axial gel replacement, related to Figure 2

The movie is maximum projection (dorsal view) of a Tg(CAG:GFP) (shown in red) chicken embryo imaged for the indicated duration. pNT and pNC were ablated and the opening was filled with alginate gel. Imaging started 20 minutes after the operation. Time stamp: hh:mm. Scale bar: 100 μm .

Movies S3. Simulation of elongation under cellular perturbations, related to Figure 3

Movies show the 2D field (Anterior to the top) of cells (small spheres). Red cells are axial cells and Blue cells are PSM cells. In the standard simulation, new PSM cells are added at the posterior end of the axial tissues (PD). The cell addition grid is determined by one center advancing NC cell, which is allowed to move along the AP axis only to prevent bending from simulation fluctuations due to small cell number. From left to right, the simulations are: Control, Reduced axial cell polarity, Reduced motility gradient, Reduced both motility and polarity. All perturbations last the duration of the simulation (4000 iterations).

Movies S4. Simulation of elongation under surgical perturbations, related to Figure 5

Similar to Movie S3. The cells matching different tissue areas were identified by their coordinates at the time of surgery and removed from the simulation. From left to right, the

simulations are: Control, pNC/pNT ablation, pPSM ablation, No Axial PD. The surgeries take place at iteration 1001 instantly. The No Axial PD condition lasts the duration of the simulation (i.e., no new PSM cells were added over the 4000 iterations).

Movies S5. Tissue and gel dynamics under PD gel replacement, related to Figure 4

The movie is maximum projection (ventral view) of a Tg(CAG:GFP) (shown in red) chicken embryo imaged for the indicated duration. Green (Dil labeled) cells start as clusters at the injection sites in the PSM and can be observed to spread in the pPSM over time. PD was ablated and the opening was filled with alginate gel. Imaging started 30 minutes after the operation. Time stamp: hh:mm. Scale bar: 100 μ m.

Movies S6. Dynamics of midline PSM cells after axial surgeries, related to Figure 5

Maximum projections (ventral view) of Tg(CAG:GFP) (shown in red) chicken embryos imaged for the indicated durations. Green (Dil labeled) cells start as clusters in the PD and can be seen to move laterally and show impaired movement in the axial tissue ablated conditions. Imaging started 25 minutes after the operation for the two ablated conditions (pNC and pNT as labeled). Time stamp: hh:mm. Scale bar: 100 μ m.

Movies S7. Dynamics of midline PSM cells after pPSM ablation, related to Figure 7

The movie is a maximum projection (ventral view) of a Tg(CAG:GFP) (shown in red) chicken embryo imaged for the indicated durations. Similar to Movie S6, green (Dil labeled) cells start as clusters in the PD show impaired movement after pPSM ablations. Imaging started 30 minutes after the operation. Time stamp: hh:mm. Scale bar: 100 μ m.

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SUPPLEMENTAL FIGURES

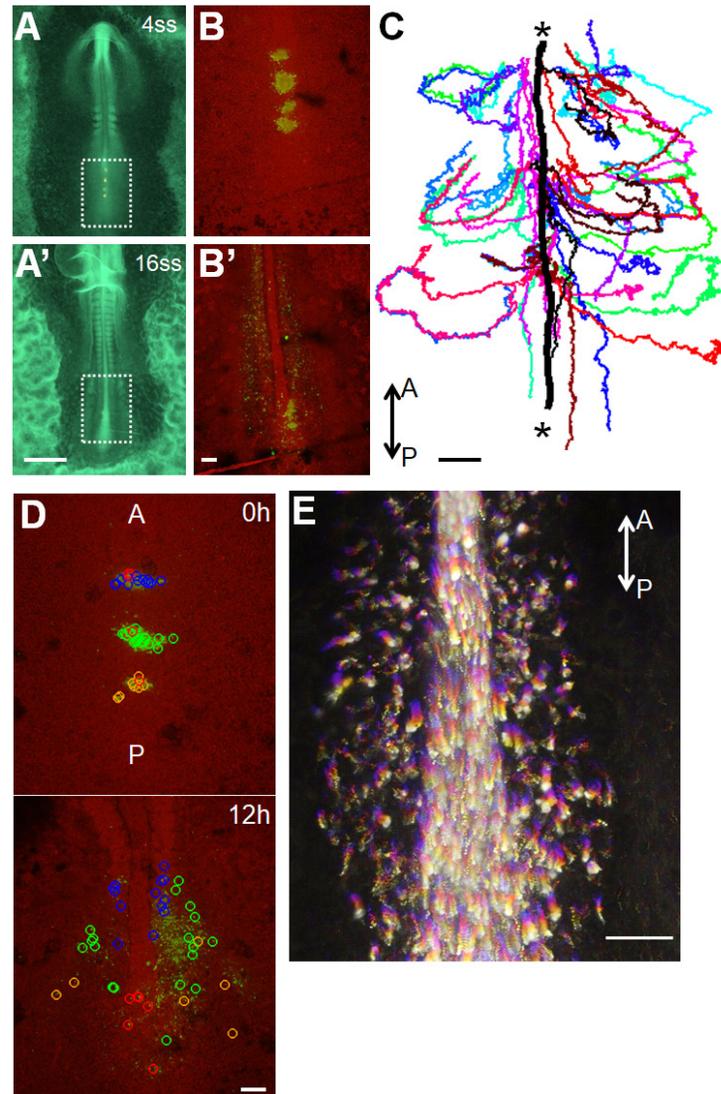


Figure S1. Live imaging and cell tracking in the elongating chicken embryo, related to Figure 1

(A) Examples of live imaging in widefield fluorescence. Whole Tg(CAG:GFP) chicken embryos (ventral view) at indicated somite stages (ss) are shown. Boxes show imaged regions in (B) Scale bar: 1mm.

(B) Examples of confocal cellular live imaging. 3D maximum projections of confocal stacks (ventral view) of the elongating axis are shown. Ubiquitous GFP signal is shown in red, green clusters show Dil labeled sites and cells. These cells were labeled in the PD and many of them later become PSM cells that flank the NC while some remain in the PD (B'). Scale bar: 100 μ m.

(C) Example of cell tracking. Single cell tracks (n=46) obtained by manual tracking ([Movie S1](#)) are plotted over the duration of the movie (~15h). Asterisks mark the trajectory (bold black) of the end of NC/Node area. A, anterior, P, posterior (same below). Scale bar: 100 μ m.

(D) Fate map of labelled cells colored as different groups posterior to NC. Blue circled cells and some Green circled cells are in the PD at 0h. Red circled cells either are NC progenitors or did not yet exit the midline in the duration of the movie (remaining in the PD). Somites and NC are distinguishable in the top and center of the right panel. Scale bar: 100 μ m.

(E) Short-term cell movement analysis using high resolution movie (~1h) overlay. Fire color shows progressing of time (colder is earlier). NC cells in the middle show directional posterior movement whereas PSM cells on both sides do not show directional bias indicating short-term random motility. Scale bar: 100 μ m.

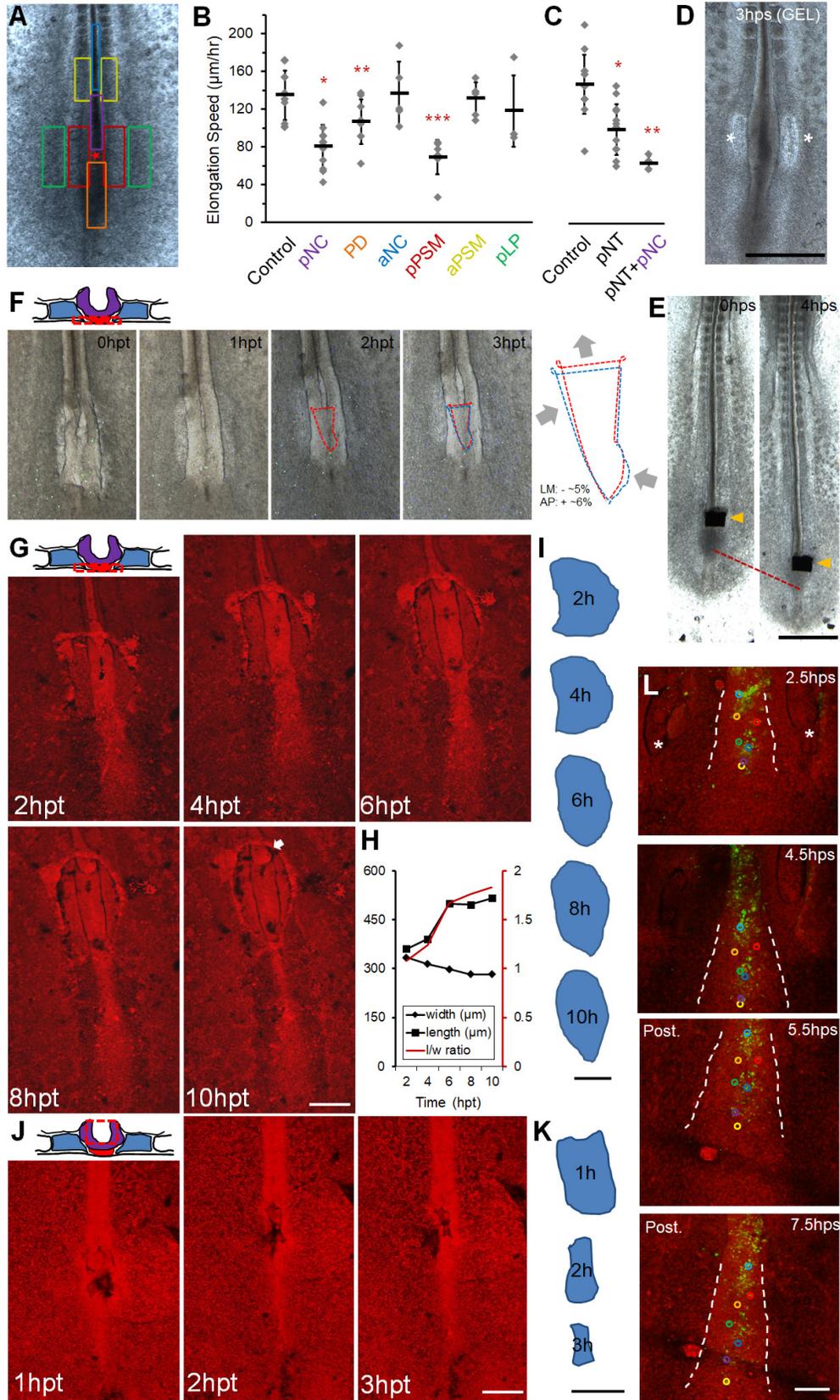


Figure S2. pPSM compresses axial tissues and is important for axial elongation, related to Figure 2

(A-C) Effects of tissue ablation on elongation speed. The embryo image (A) shows different ablation sites (ventral operations) with colors corresponding to elongation speed measurements in (B). (C) shows pNT ablations (dorsal operations) with its own control. Each mark represents an embryo followed over a 6-hour window after surgery. pPSM, pNC, pNT and PD ablations result in significant reduction of elongation speed (asterisks, $p < 0.05$, t-tests). aPSM, aNC and pLP (lateral plate) ablations do not ($p > 0.05$, t-tests). pNT/pNC double ablation results in a stronger reduction than pNT alone ($p = 0.02$, t-test, panel C).

(D) Gel replacement of pPSM. Asterisks, gel implant sites. Widening of the gel flanked portion is observed in 4/4 embryos. Scale bar: $500\mu\text{m}$.

(E) Foil insertion between the pNC and PD. The cross-sectional foils are $\sim 200 \times 200\mu\text{m}$.

(F) Gel replacement of pNC. Cross-section schematic shows the ablation of pNC from the ventral side. Only endoderm and pNC were ablated, NT is visible through the gel. Slight deformation of the gel can be observed through time. Dashed lines collect five fluorescent beads in the gel to a contour. The beads' relative positions change, tracking gel deformation. The blue contour on the 3hpt (hours-post-transplant) image is identical to the red contour on the 2hpt image. The 3hpt contour (red) is narrower laterally than the 2hpt contour (blue) by an average negative strain in LM of $\sim 5\%$, and is longer in AP (arrows) by $\sim 6\%$. Anterior to the top.

(G) Similar to (F) with injected gel. Implant boundary is visible and changes over time. Arrow marks protruding NC from anterior cut site. Scale bar: $200\mu\text{m}$.

(H) Quantified shape changes of the gel implant in (G). Length (l) is measured along the midline, width (w) is measured following the widest line perpendicular to the midline. Consistent narrowing was observed in $n = 7$ experiments.

(I) Contours of the gel implant in (G). Scale bar: $200\mu\text{m}$.

(J-K) Gel replacement of pNT (dorsal view). Consistent narrowing was observed in $n = 3$ experiments. The gel is likely extruded by the tissue in this case. Scale bars: $200\mu\text{m}$.

(L) Tracking of labeled NC cells. In this movie NC cells were labeled with Dil and movement of 7 cells were followed through time (colored circles). White dashed lines mark the lateral boundary of the NC. Asterisks mark pPSM ablation site. As the NC cells pass the ablation site, limited tissue convergence and cell-cell rearrangement are seen. In unperturbed posterior (post.) area, tissue narrowing and medial-lateral cell intercalation are apparent. Scale bar: $100\mu\text{m}$.

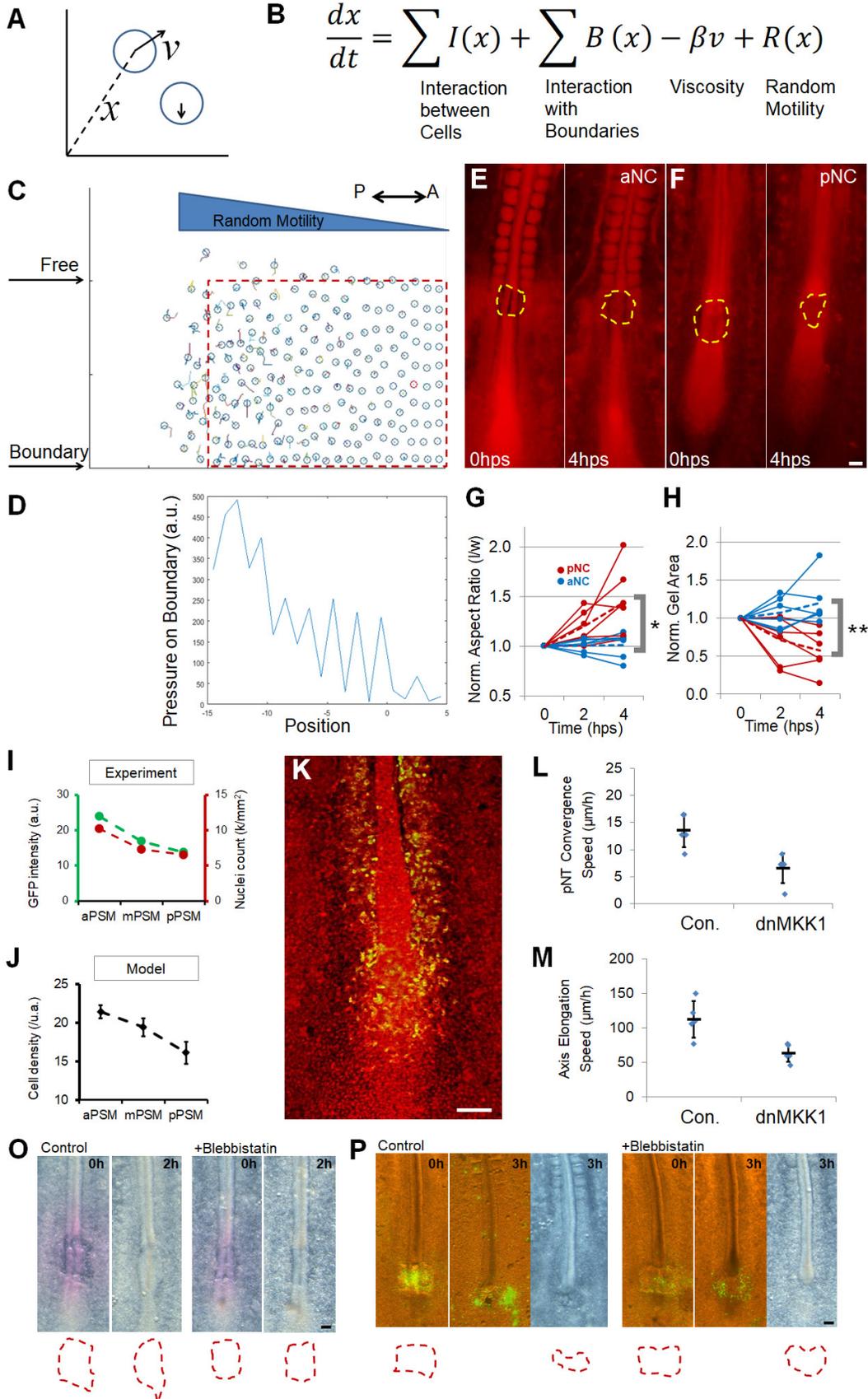


Figure S3. An agent(cell)-based model of co-elongation incorporating the cell motility gradient assumption, related to Figure 3

(A) Cells are defined as circles (with certain radius controlling collision volume) in a plane (tissue space, extracellular matrix). Cells have two main variable attributes: position (x) and velocity (v). Velocity is the time derivative of position.

(B) For each new iteration, a velocity is generated to change the cell's position. The velocity is determined by the forces acting on the cell, which are functions of position and current velocity. The total force is a summary of repulsive interactions between cells (a function of the cell's position and the positions of all other cells) due to collision volume (i.e., cells have volume and cannot occupy the same area), interaction with boundary (such as surface extracellular matrix of the PSM and neighboring tissues such as the lateral plate), effect of viscosity (resistant force against the current movement), and the only active movement component with a random direction and a magnitude as a function of position (i.e., the motility gradient generated by active cell behavior).

(C-D) A random motility gradient causes differential expansion and pressure on confining boundaries. An initial square field of cells (red box) with a gradient of random motility expands over the unconfined (free) space. On a confined boundary, a pressure gradient can be recorded. In the PSM, a similar random motility gradient is observed, and isolated PSM expands preferentially posteriorly. The pressure generated can explain the compression experienced by the pNC/pNT.

(E-F) Deformation of alginate gel implants after surgical ablation of aNC and pNC, respectively. Fluorescent signal (red) is GFP. Yellow dashed lines measure the shape and in-plane area of the gel. Note that the gel volumes are not expected to shrink under compression, the change of area results from gel deformation and flow in the direction perpendicular to the image plane. Scale bar: 100 μ m.

(G) Normalized gel aspect ratio change. "l" measures the length along AP, "w" along LM. l/w ratios for each sample was normalized by the 0 hps data point. Dashed lines are average ratios. Asterisks mark significant difference between averages at 4 hps (*p=0.018, t-test).

(H) Normalized gel top view area change. Similar to (G), l*w was normalized by 0 hps data point. 4 hps difference **p=0.004, t-test.

(I) Cell density measurement by quantifying GFP intensity (green) or nuclei count using Tg(PGK1:H2B-chFP) quail line (red, [Huss et al., 2015](#)) from confocal image z-stacks. a.u., arbitrary units.

(J) Model predictions of cell density along the PSM. Measured areas in the model were adjusted to match measured areas from data. Error bars: \pm SD. /u.a., number per unit area.

(K) Electroporation specificity into PSM cells. Image shows a representative embryo electroporated at stage 5 with a membrane-Venus construct into the anterior primitive streak (PD) of a Tg(CAG:GFP) embryo. Confocal image was taken after 24 hours. Positive cells can be seen in the PD and the PSM but not in the midline NC. Scale bar: 100 μ m.

(L-M) Converging (conv.) speed (\pm SD) of neural folds towards the midline (L) and elongation speed (\pm SD) of the axis (M) measured in 3 hours. n=5 for control and n=6 for dnMKK1, p<0.05, t-tests.

(O-P) Examples of Blebbistatin treatment on gel deformations (see also [Figure 4G](#)). Scale bars: 100 μ m. (O) pNC gel implants. Pink color is from diffused rhodamine-B mixed with the alginate solution to better show initial gel shape. Blebbistatin treatment delays the smoothing of gel shape and deformation (red contours), suggesting a reduction of tissue force. (P) PD gel implants. Fluorescent beads (green) were mixed with the alginate to better show gel deformation. The beads are loosely trapped in the gel (cannot be washed off but can move). Lateral displacement of beads is seen and is delayed in Blebbistatin treatment. Gel is less compressed from the anterior in Blebbistatin treatment.

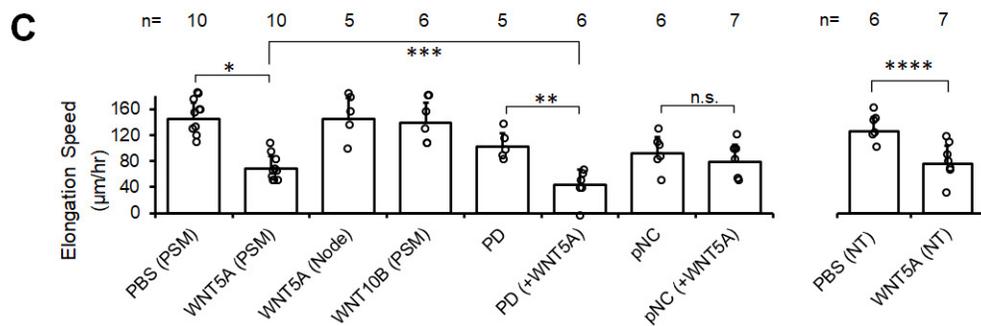
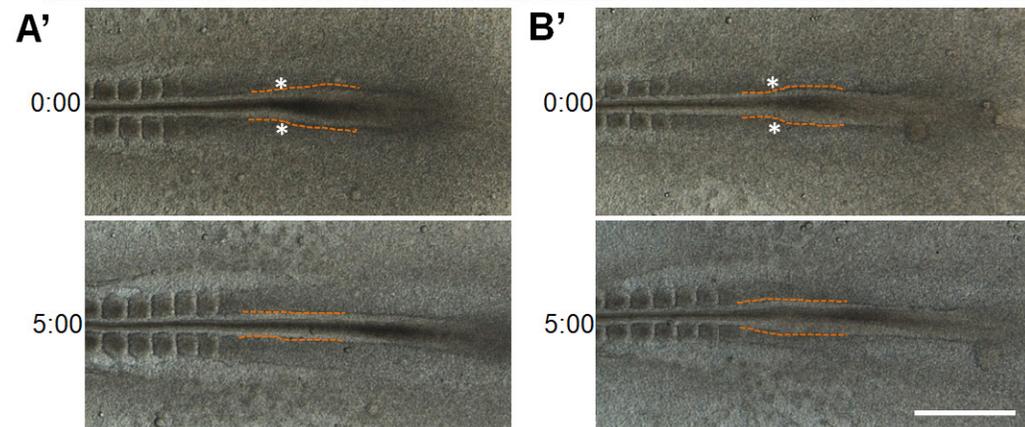
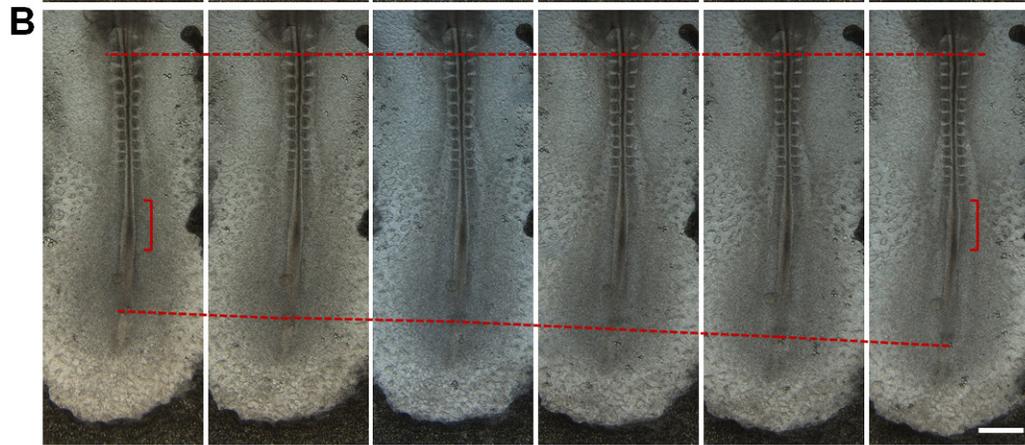
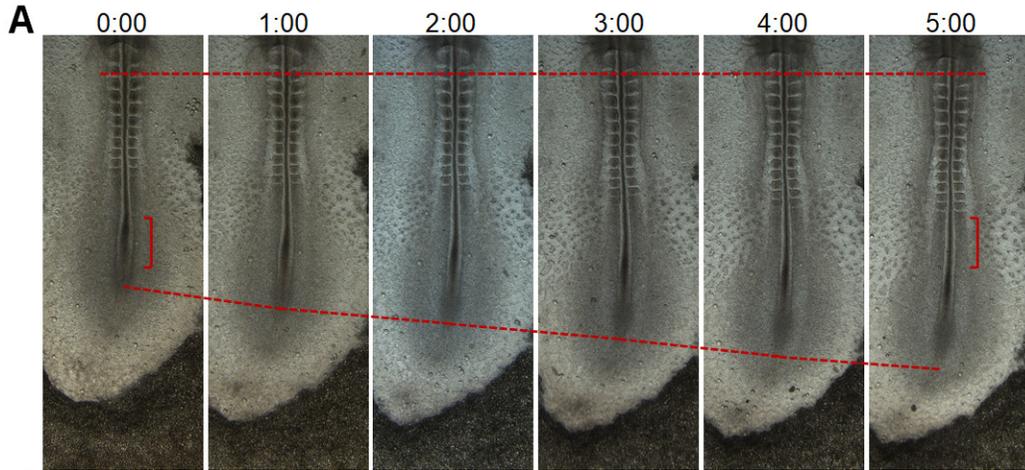


Figure S4. Elongation and tissue dynamics after WNT5A injection, related to Figure 3

(A-B) Time course of embryos after WNT5A injection at the anterior pPSM level (bracket). Top dashed line marks a fixed somite as the reference, bottom dashed line measures elongation. (A'-B') are enlarged views with dashed lines showing the width of the NT. Asterisks mark the approximate injection location. Scale bars: 500 μ m.

(C) Elongation speeds after WNT5A injections. The first 4 columns (from left) are injections only, at the injected tissue level indicated in the parenthesis (e.g., PSM, Node). The second 4 columns couple tissue ablation (e.g., PD, pNC) with injections. The NT data (injections only) is shown in a separate panel on the right as the controls were performed separately. Asterisks indicate significant difference ($p < 0.05$, t-tests).

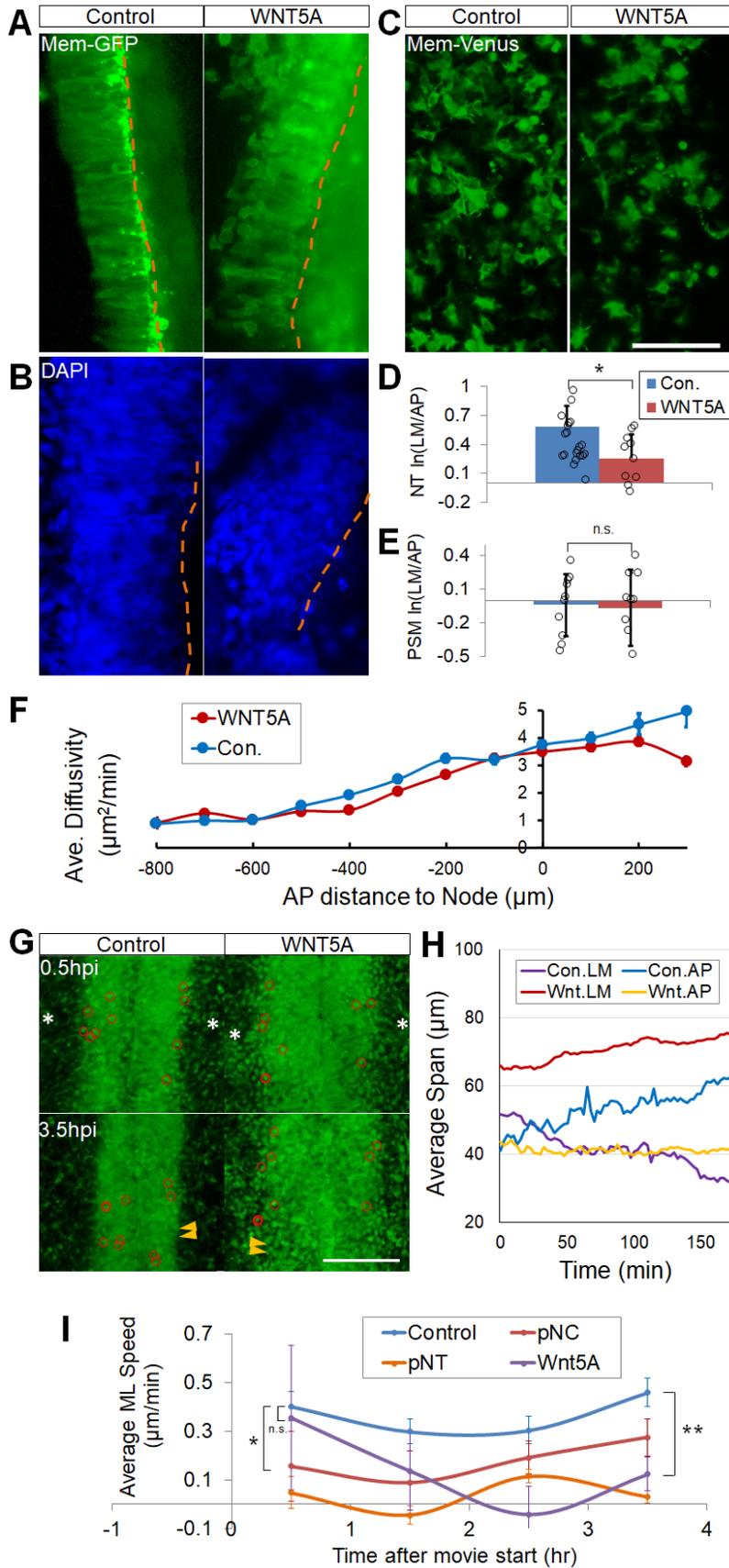


Figure S5. pNT cell polarity and dynamics under WNT5A perturbation, related to Figure 3

(A-C) Cell shape analysis using confocal images of cells in live embryos electroporated with a membrane-GFP construct in the NT (A, dorsal view), fixed embryos stained with DAPI in the NT (B, dorsal view), and live embryos electroporated with a membrane-Venus construct in the PSM (C, ventral view), respectively. Yellow dashed lines mark the apical/medial side of the neural fold. Scale bar: 100 μ m.

(D-E) Quantification of cell shapes by the logarithm of the cell's aspect ratio (lateral-medial length over anterior-posterior length, LM/AP). In (D), NT cells show LM polarization which is reduced after WNT5A injection (n=20, *p=0.004, t-test). In (E), PSM cells show no shape polarization in either condition (n=10, p=0.8, t-test). Similar cell shape changes near the injected area were observed in the NT of all 4 injected embryos but none of 3 controls.

(F) PSM cell motility gradient in a WNT5A injected embryo.

(G) Distribution of tracked cells in a WNT5A injected embryo. The images are 3D maximum projections (dorsal view) of Tg(CAG:GFP) (shown in green) chicken embryos after injection of PBS (control) or WNT5A. Red circles highlighted randomly selected lateral cells at the beginning of the movies at the injected level (marked by asterisks). The pNT is seen in the middle undergoing convergence and elongation through NT cell intercalations. Mesenchymal pPSM cells can be seen flanking the NT. Yellow arrowheads mark the organized vs. disorganized NT areas between control and injected. Scale bars: 100 μ m.

(H) Tracking of NT cell intercalation dynamics. 20 cells in each condition were tracked. Average span is calculated using the average position of all cells along either LM or AP direction. Reduction of LM span and a corresponding increase of AP span (as in illustration) indicate cell intercalation.

(I) Timing of slowdown of medial-lateral movement of midline PSM cells after different perturbations. >100 cell movement measurements were taken in each group. Mean speeds \pm SD are plotted over time. The WNT5A injection group is similar to control at 0.5hr (n.s., p>0.05, t-test) and significantly lower at 3.5hr (**p<0.05, t-test). The surgery groups (pNT and pNC) are significantly lower already at 0.5hr (*p<0.05, t-tests).

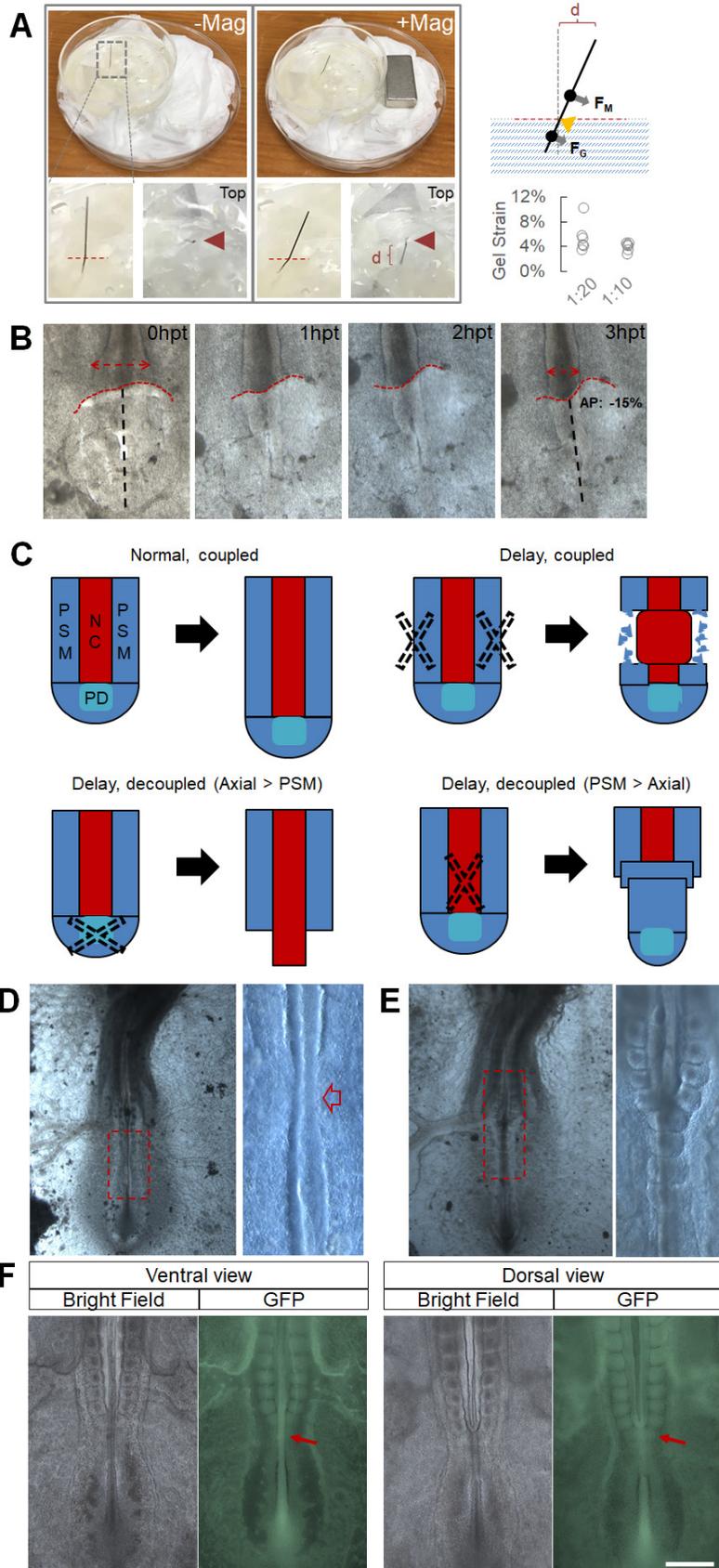


Figure S6. Decoupled elongation as a result of breaking the mechanical feedback loop, related to Figure 7

(A) Magnetic pin stress estimation using alginate gels. The left images show the double dish set up used in the experiments. The sample dish (35mm) with the gel or embryo is placed on one end of the 60mm dish that holds the magnet. The steel pin is inserted vertically into a layer of alginate gels on top of the culture medium. Upon magnet addition, the pin rotates and causes a local compression on the gel. The amount of rotation is related to the gel strain. The force from the gel elastic deformation (F_G) and the magnetic attraction force on the pin (F_M) produce torques that balance to determine the rotation angle. The gels were made with indicated volume ratios (1:20 and 1:10) by mixing 5% (w/v) calcium chloride solution with 1% (w/v) alginate sodium solution. Given the observed strains on these gels that are much stiffer than the PD tissue, the estimated stress produced by the pin on the interface point is estimated to be much higher than the endogenous axial push (compare to [B]).

(B) Gel implant to replace the progenitor domain (PD). Double headed arrows show the narrowing of the NT. Red dashed lines show the outline of the gel in contact with pNC and pPSM. Here NC becomes longer than PSM as the gel deforms to a heart shape. The dark dashed lines measure a change of the gel length in AP at the midline by ~15%. The gel is made by 1:40 mixing of 5% (w/v) calcium chloride solution with 1% (w/v) alginate sodium solution.

(C) Summary illustration of microsurgery ablations that impact elongation rate. pPSM ablation causes a local delay but does not disrupt coupling. PD ablation results in longer axial tissues and axial ablation results in longer and centered PSM.

(D) Left and right PSM merged at the site where pNC was ablated and elongation continued in the long term (>15 hours-post-surgery). Right panel is a zoomed in view of the red box area in the left panel. Red arrow points to the region of PSM merging. The surgery ablation of pNC usually does not completely remove all NC progenitors, thus restoration of NC further posterior is usually observed.

(E) Joined PSM formed large axial somites posterior to the pNC ablation site. Right panel is a zoomed in view of the red box area in the left panel. Here PSM becomes longer than NC. Note that the NT still co-elongated with the PSM.

(F) Joined PSM on the dorsal side at the pNT ablation site (arrow). Ventral view shows an undamaged NC through the ablation site. Dorsal view show joint somites forming at the NT gap. In this case PSM becomes longer than NT while NC still co-elongated with PSM. Scale bar: 500 μ m.

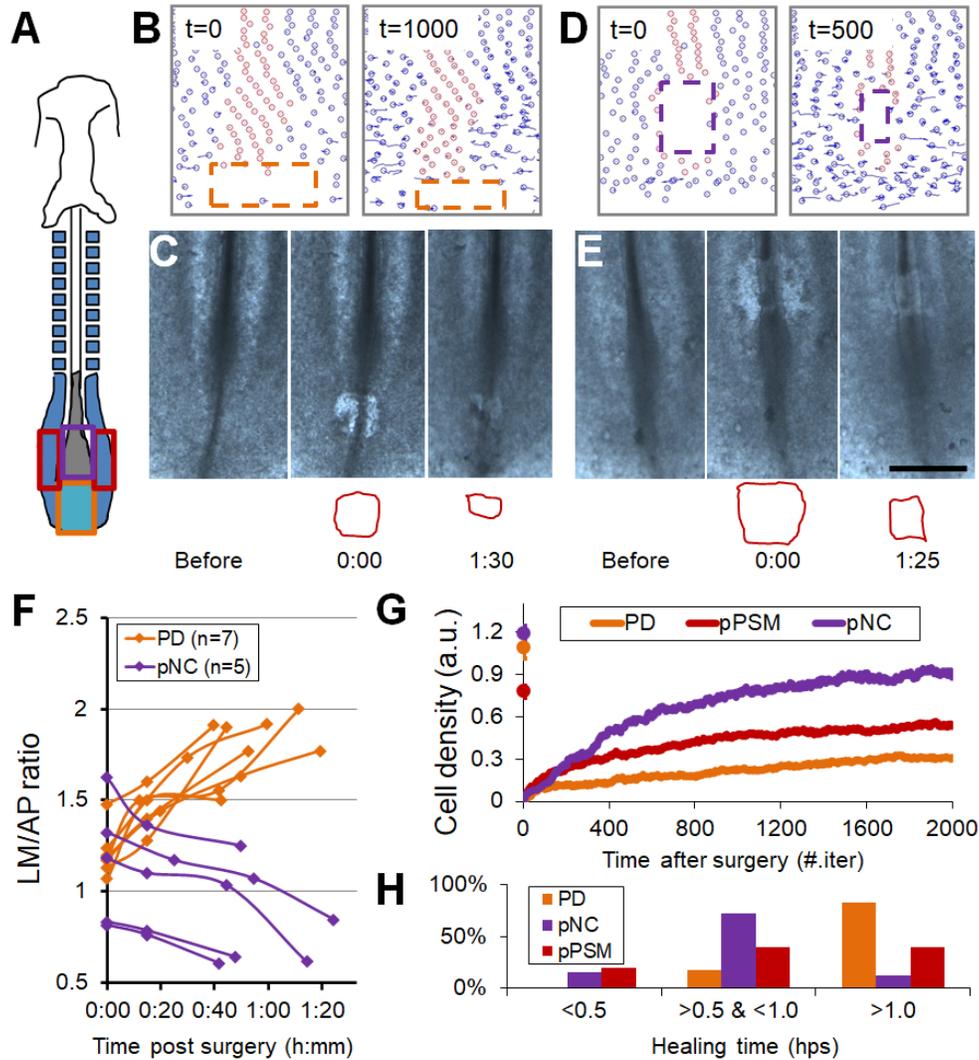


Figure S7. Model predictions of tissue mechanical dynamics, related to Figure 7

(A) Schematic of surgery sites on the embryo corresponding to areas in the model.

(B,D) Simulations of wound closure in response to ablation of PD and pNC. Time indicates number of iterations post-ablation.

(C,E) Closure of rectangle wounds introduced on embryos as shown in (B) and (D), respectively. Red contours outline the shape of the openings. Note that these closing dynamics are much faster than when there is a gel implant. Scale bar: $500\mu\text{m}$.

(F) Summary of wound aspect ratio change. The ratios are plotted as LM over AP lengths, indicating spatially differential tissue forces.

(G) Simulated cell density recovery (an approximate measure of wound closure) after cell deletion in the same area. Curves show the average of 30 simulations in each case. Markers $\pm\text{SD}$ at time 0 indicate cell density before surgery. pNC area is predicted to have the fastest recovery.

(H) Time of complete wound closure by ablated tissues. PD $n=17$, pNC $n=25$, pPSM $n=15$. The majority of pNC wounds close under 1 hps, while the majority of PD wounds remain open.