A gene regulation network controlled by Celf1 protein–rbpj mRNA interaction in Xenopus somite segmentation

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Summary

Somite segmentation is impaired in Xenopus celf1 morphant embryos. The Celf1 RNA-binding protein targets bound mRNAs for rapid degradation, and antisense approaches demonstrated that segmentation defects in celf1 morphants were due to a derepression of rbpj mRNA. Rbpj protein is a key player of Notch signalling. Because segmentation involves complex cross-talk between several signalling pathways, we analysed how rbpj derepression impacted these pathways. We found that rbpj derepression stimulated the Notch pathway. Notch positively controlled the expression of cyp26a, which encodes a retinoic acid (RA)-degrading enzyme. Thus, rbpj derepression led to cyp26a overexpression and RA attenuation. It also repressed fgf8, consistent with an inhibition of FGF signalling. Pharmacological inhibition of the FGF pathway repressed cyp26a, but rbpj derepression was sufficient to restore cyp26a expression. Hence, while it was known that the FGF pathway antagonized RA signalling through expression of cyp26a, our results suggest that Rbpj mediates this antagonism. Furthermore, they show that the post-transcriptional repression exerted by Celf1 on rbpj mRNA is required to keep cyp26a expression under the control of FGF signalling. We conclude that rbpj repression by Celf1 is important to couple the FGF and RA pathways in Xenopus segmentation.

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Key words: RNA-binding protein, Notch, FGF, retinoic acid, Cyp26a

Introduction

In vertebrates, somites are arranged along the anteroposterior axis of the embryo in an organisation known as segmentation. Somite segmentation is a blueprint for vertebral segmentation in adults, and vertebral disorders, such as congenital scoliosis, may arise from defective somite segmentation (Pourquie, 2011). Segmentation results from the periodic emergence of somites from the presomitic mesoderm (PSM). It depends on cross-talk between a clock and a determination front (Mara and Holley, 2007; Dequenat and Pourquie, 2008; Aulehla and Pourquie, 2010; Gibb et al., 2010). The oscillatory expression, in the posterior PSM, of tens of genes encoding factors of the Notch, FGF, and Wnt signalling pathways supports the clock. The determination front is set by the antagonistic activities of the FGF pathway in the posterior PSM and the retinoic acid (RA) pathway in the anterior PSM (Moreno and Kintner, 2004; Goldbeter et al., 2007). It moves toward the posterior extremity during embryo elongation. A prospective somite consists of the cells left behind the determination front during one oscillation of the clock. Oscillations cease in these cells and the expression of certain genes changes from a cyclic to a stable pattern, restricted to part of the future somite. Furthermore, these cells express new segmentation genes, in either the anterior or the posterior half of the future somites. This contributes to their antero-posterior polarity and prefigures morphological segmentation (Mara and Holley, 2007; Dequenat and Pourquie, 2008; Aulehla and Pourquie, 2010; Gibb et al., 2010).

The players in somite segmentation include components of the Notch, RA, FGF and Wnt signalling pathways, and molecules involved in the post-transcriptional control of gene expression. Indeed, the stability of Fgf8 mRNA shapes the postero-anterior FGF gradient, but the factors controlling Fgf8 mRNA degradation remain unknown (Dubrule and Pourquie, 2004), and the oscillations imply a rapid decay of clock mRNAs (Cibois et al., 2010a). One key post-transcriptional regulator of somite segmentation is Celf1 (previously known as EDEN-BP or Cugbp1), a multifunctional RNA-binding protein (Barreau et al., 2006). The knockdown of celf1 expression impairs Xenopus segmentation. Celf1 protein interacts directly with rbpj mRNA [also known as suppressor of hairless, su(h)] (Gautier-Courteille et al., 2004), and two lines of arguments indicate that the loss of the interaction between Celf1 protein and rbpj mRNA is the main cause of impaired segmentation of celf1 morphants. First, rbpj is overexpressed in Celf1-deficient embryos, consistent with the capacity of Celf1 to target bound mRNAs to rapid deadenylation and decay. Furthermore, rbpj overexpression (driven by mRNA...
injection) is sufficient to strongly alter segmentation (Gautier-Courteille et al., 2004). Second, impairing the interaction between Celf1 protein and rbpj mRNA with a “target protector” antisense morpholino yielded a moderate rbpj overexpression, both at the mRNA and protein levels, and recapitulated the segmentation defects. A minute amount of a second morpholino blocking rbpj translation, which drove Rbpj protein abundance back to its initial level, restored segmentation, strongly supporting the specificity of the target protector morpholino (Cibois et al., 2010b). These experiments have revealed the phenotypic consequences of rbpj overexpression for the first time in vertebrates, and have highlighted a post-transcriptional mechanism that prevents rbpj overexpression.

rbpj encodes a DNA-binding protein that plays a key role in the Notch signalling pathway. Upon binding to its ligand, the Notch transmembrane receptor is cleaved, releasing its intracellular domain (NICD). This is subsequently transllocated to the nucleus, where it associates with Rbpj. The Rbpj–NICD complex activates the transcription of target genes, but, in the absence of NICD, Rbpj represses them (Kovall and Blacklow, 2010). Somite segmentation is impaired in both embryos overexpressing rbpj (Cibois et al., 2010b) and in rbpj mutants and morphants (Oka et al., 1995; Echeverri and Oates, 2007). Segmentation therefore requires the presence of optimal amounts of Rbpj protein. In this study, we investigated the consequences of Rbpj overproduction for the Notch pathway and for other signalling pathways involved in Xenopus segmentation.

**Results**

**rbpj overexpression modulates Notch signalling in the presomitic mesoderm**

A target-protector morpholino (TP MO) causes rbpj overexpression by abolishing the repressive interaction between Celf1 protein and rbpj mRNA (Cibois et al., 2010b). In Drosophila, some phenotypes associated with rbpj [Su(H)] overexpression are similar to a Notch gain-of-function while other phenotypes are similar to a Notch loss-of-function. This is probably a consequence of the dual capacity of this protein to activate and repress transcription (Furriols and Bray, 2000). Similarly, in Xenopus segmentation, TP MO-mediated rbpj overexpression could either enhance or attenuate Notch signalling. To discriminate between these possibilities, we injected the TP MO unilaterally with a lineage tracer and analysed the expression of dlc (delta-2). dlc encodes the main Notch ligand in the PSM. It is expressed in the posterior PSM and as 3–4 chevrons in the anterior PSM. The TP MO repressed the posterior expression of dlc, and transformed the stripes in the anterior PSM to a more continuous pattern by filling the gaps (Fig. 1). The expression of dlc in the PSM is controlled by complex feedback loops. Stimulating the Notch pathway with a constitutively active mutant of Rbpj repressed dlc in the posterior PSM, and to a weaker extent in the anterior PSM. Conversely, repressing the Notch pathway with a dominant negative mutant of Rbpj filled the gaps in the anterior PSM and weakly stimulated its expression in the posterior PSM (Jen et al., 1997; Sparrow et al., 1998; Jen et al., 1999). Hence, the TP MO mimics the effect on dlc of a Notch gain-of-function and a Notch loss-of-function, respectively, in the posterior and the anterior PSM. This differential sensitivity of Notch signalling to Rbpj abundance may reflect the differential amounts of NICD in the two compartments of the PSM, but we were unable to accurately compare the amounts of NICD in these two compartments.

**rbpj overexpression represses retinoic acid signalling in the presomitic mesoderm through cyp26a upregulation**

We next analysed the effects of the TP MO on the retinoic acid (RA) signalling pathway. We injected a plasmid carrying the luciferase reporter under the control of RA response elements into embryos (Blumberg et al., 1997). The addition of exogenous RA stimulated the luciferase activity, whereas the dominant negative RA receptor dnRAR repressed it (Fig. 2A). This indicates that luciferase activity adequately reflects RA signalling. Importantly, the TP MO lowered the luciferase activity (Fig. 2A), suggesting that rbpj upregulation represses the RA signalling pathway.

To confirm these data, we analysed the expression of mespa (mesoderm posterior homologue A, thylacin-1, Mesp2 in mice). This gene marks the determination front. Because the position of the front is contributed by an antero-posterior gradient of RA signalling, stimulating the RA pathway shifts posteriorly its expression while repressing the RA pathway shifts it anteriorly (Moreno and Kintner, 2004; Oginuma et al., 2008). The TP MO shifted anteriorly mespa domain of expression, pretty much like dnRAR-mediated inhibition of RA signalling (Fig. 2B). This confirms that the RA pathway is repressed in embryos overexpressing rbpj.

Another gene that positively responds to RA signalling is cyp26a (Moreno and Kintner, 2004). However, we found that the TP MO, unlike dnRAR, stimulated cyp26a (Fig. 2C). It is worth noting that cyp26a is overexpressed in the posterior PSM, at the place where the TP MO stimulates the Notch pathway. We think that the upregulation of cyp26a is the main reason of RA attenuation in TP MO-injected embryos for three reasons. First, cyp26a encodes an RA-degrading enzyme. Second, in zebrafish, the morpholino-mediated knockdown of rbpj repressed cyp26a expression, whereas a dominant activated mutant of rbpj activated it (Echeverri and Oates, 2007). Third, if Cyp26a did not link the stimulation of Notch to the repression of RA signalling, then cyp26a would only be a downstream target of RA signalling, and we would have observed a repression, rather than a stimulation, of cyp26a expression in TP MO-injected embryos. We conclude that overexpressed rbpj stimulates cyp26a, which represses the RA pathway.
**rbpj overexpression represses FGF signalling in the presomitic mesoderm**

We next investigated the effects of *rbpj* overexpression on the FGF pathway. The TP MO repressed *fgf8* expression (Fig. 3A). Because *rbpj* overexpression attenuates RA signalling (see above), and dnRAR repressed *fgf8* (Fig. 3A) as previously described (Moreno and Kintner, 2004), the repression of *fgf8* in TP MO-injected embryos may be due to RA signalling inhibition. *fgf8* encodes the relevant FGF ligand in the PSM (Dubrulle et al., 2001), and the downregulation of *fgf8* expression was expected to translate into attenuation of FGF signalling. We checked this point by analysing the expression of *msgn1* (*mesogenin-1*, *mespo*) and *dusp6* (*mkp3*). The FGF pathway controls these two genes, in conjunction with the Wnt [msgn1 (Wang et al., 2007)], or the RA

**Fig. 3. Impact of *rbpj* overexpression on the FGF signalling pathways.** We injected nLacZ and control or TP MO, or dnRAR mRNA, as indicated, into one blastomere of two-cell embryos, which we allowed to develop until the tailbud stage. Where indicated, we treated the embryos for 2 hours with SU5402 (embryos injected with control MO were treated with DMSO) and we stained them for β-galactosidase activity (red dots) and *fgf8* (A), *msgn1* (B), or *dusp6* (C) mRNA by ISH. We sorted DMSO and SU5402-treated embryos into 5 classes depending on their staining intensities. The right panels show the percentages of embryos within each of these classes, and, for injected embryos, the percentage of embryos with a staining intensity in the injected side above, equal to, and below (respectively green, yellow and red) that in the control side. We compared the distributions, between these three classes, of the control and the other conditions by a chi-square test and we show the p-values. All the photographs are dorsal views, anterior left, injected-side up.
[dusp6 (Moreno and Kintner, 2004)] pathways. Accordingly, the FGF pathway inhibitor SU5402 repressed these two genes (Fig. 3B,C). These two genes were also downregulated in TP MO-injected embryos (Fig. 3B,C), and this supports our interpretation that rbpj overexpression represses FGF signalling.

**rbpj overexpression takes over FGF inhibition in the control of the RA pathway**

Moreno and Kintner showed that the FGF and RA pathways were mutually antagonistic, so that FGF repression was associated with enhanced RA signalling (Moreno and Kintner, 2004). However, our results demonstrate that rbpj upregulation represses both RA and FGF signalling. To solve this apparent discrepancy, we analysed embryos injected with the TP MO and exposed to SU5402. While SU5402 shifted posteriorly the expression of mespa, consistent with an RA gain-of-function, the TP MO reversed this effect (Fig. 4A). Similarly, SU5402 repressed cyp26a, but the TP MO restored a high level of expression of cyp26a in SU5402-treated embryos (Fig. 4B). Consequently, SU5402 and the TP MO have opposite consequences on mespa and cyp26a expression, but the consequences of exerting these two treatments simultaneously are similar to the consequences of TP MO injection. By contrast, the TP MO had no effect on dlc in SU5402-challenged embryos (Fig. 4C). Together, these data show that rbpj overexpression takes over FGF repression in the control of RA signalling, as deduced from cyp26a and mespa expression, but not in the control of dlc expression.

**Discussion**

Pioneering work in Xenopus revealed that cyp26a encodes an RA-degrading enzyme and is positively controlled by the FGF pathway, while dusp6 encodes a phosphatase that antagonizes FGF signalling and is positively controlled by RA. This results in a mutual antagonism between the RA and FGF pathways that defines two compartments of the PSM, a posterior one where FGF predominates and an anterior one where RA predominates (Moreno and Kintner, 2004). The present article extends this work in two directions.

First, we observed that inhibiting the RA pathway either directly (by injecting dnRAR mRNA) or indirectly (by injecting the TP MO) repressed msgn1, probably owing to fgf8 repression (Fig. 3). Conversely, it had previously been reported that dnRAR activated msgn1 expression owing to dusp6 repression (Moreno and Kintner, 2004). A plausible explanation for this discrepancy is the age of the embryos used. Indeed, we examined the effects of dnRAR on msgn1 expression in tailbuds, whereas neurulae were previously analyzed. Nevertheless, taking msgn1 expression as a proxy for FGF signalling, our results suggest that the relationships between the RA and FGF pathways can not be only summarized as a mutual antagonism but that, at least under certain circumstances, RA signalling is able to positively control the FGF pathway (Fig. 5). Cell migration is a major contributor to embryo elongation in fish (Zhang et al., 2008) and, in chick, FGFinduced in the PSM directs cell motions leading to the antero-posterior elongation of the embryo (Bénazéraf et al., 2010). The dependence of posterior FGF signalling on anteriorly produced RA may thus control embryo elongation rate. It is therefore tempting to attribute embryo curvature (e.g. Fig. 2B, Fig. 4A) to a decreased elongation of the TP MO-injected side, due to the repression of RA and consequently FGF signalling.

Second, our results prompt us to reconsider the relationships between the FGF, Notch and RA pathways in segmentation. TP MO-mediated overexpression of rbpj repressed dlc and stimulated cyp26a. Because the Notch pathway represses dlc (Jen et al., 1997; Sparrow et al., 1998; Jen et al., 1999) and cyp26a is under Rbpj control (Echeverri and Oates, 2007), these observations indicate that rbpj overexpression stimulates Notch signalling in the posterior PSM. TP MO-mediated stimulation of Notch signalling repressed RA signalling, while attenuating the FGF pathway by SU5402 treatment stimulated the RA pathway. Importantly, simultaneously activating the Notch pathway and repressing the FGF pathway led to RA repression, as deduced from cyp26a overexpression and mespa anterior shift. This shows that rbpj overexpression takes over FGF repression in the control of RA signalling. Furthermore, FGF may positively control the Notch pathway, since FGF repression downregulated dlc. This suggests that the Notch pathway is an intermediate by which the FGF pathway represses RA signalling. A demonstration of this model would be to show that Rbpj is required for the repression of RA induced by FGF signals. Unfortunately, we were unable to make FGF gain-of-function experiments, because premature FGF activation strongly alters gastrulation. Taking this caveat into account, we propose that, in a control situation, Celf1 minimises Rbpj protein abundance to keep Notch signalling at a level compatible with FGF regulation, ensuring a coupling between the
FGF and RA pathways (Fig. 5A). When the repression of 
rbpj by Celf1 is abolished by the TP MO, 
rbpj overexpression leads to high, FGF-independent Notch signalling that results in 
cyp26a overexpression and RA attenuation (Fig. 5B). Hence, the post-
transcriptional control exerted by Celf1 protein on 
rbpj mRNA is required to couple the FGF with the Notch and RA pathways in somite 
segmentation. 

The Notch pathway plays a central role in the clock (Mara and 
Holley, 2007; Dequenat and Pourquié, 2008; Aulehla and 
Pourquié, 2010; Gibb et al., 2010), and we propose here that it 
also contributes to the cross-talk between the FGF and RA pathways to determine the determination front. Somite 
segmentation is impaired in 
rbpj mutants and morphants (Oka et al., 1995; Echeverri and Oates, 2007), but also in embryos 
overexpressing 
rbpj (Cibois et al., 2010b). Segmentation therefore requires the presence of optimal amounts of 
Rbpj protein. This requirement for such a tight control of Rbpj abundance was not expected given the ubiquitous expression of the 
rbpj gene (Wettstein et al., 1997). celf1 is also required for zebrafish segmentation (Matsui et al., 2012) and is expressed in 
nice PSM (Kress et al., 2007), and it will be important to test if 
the function of Celf1 to control 
rbpj is conserved in the segmentation process of other vertebrates.

Materials and Methods

Plasmids and probes

The plasmids encoding 
dlc, nuclear β-galactosidase (nLacZ, pCS2+nBGal vector) 
and dnRAR have been described (Blumberg et al., 1997; Gautier-Courteille et al., 2004). 
mespa ORF was PCR-amplified using forward (ATGATTTTCCTCCA-
ACAAAC) and reverse (TGAATAGCGAAGTAGCTGAAGTG) primers and 
Xenopus embryo cDNA, and cloned in pGEM-T (Promega). Other plasmids were 
purchased from Imagene (Xenopus embryo cDNA, and cloned in pGEM-T (Promega)). Other plasmids were 

Xenopus laevis embryos procedures

We injected embryos at the two-cell stage in one blastomere with one or several of the following: 2 pmol of control or target-protector morpholino (Cibois et al., 2010b), 0.5 fmol of nLacZ mRNA, 5 fmol of dnRAR RNA, 1.5 fmol of hGR-
NICD or hGR-Su(H)-Ank mRNA. They were allowed to develop at 16–22°C. 
When required, tailbud (stage 24-26) embryos were incubated for 2 hours at 22°C with 
RA (1 µM, Sigma), cycloheximide (10 µg/ml, Sigma), SU5402 (400 µM), dexamethasone (20 µM), or a concentration of DMSO or ethanol corresponding to the highest amount in the drug solutions. For ISH, embryos were fixed 45 min in 
MEMFA, rinsed in PBS and preincubated in PBS supplemented with 2 mM 
MgCl2. They were revealed for β-galactosidase activity by incubation in X-Gal 

Fig. 5. Model for the relationships between the RA, FGF 
and Notch pathway in Xenopus PSM. Three intra-pathway 
feedback loops attenuate each pathway (RA stimulate 
cyp26a and FGF stimulates 
dusp6 while Notch represses 
dlc). In addition, the RA pathway either stimulates (through 
fgf8) or represses (through 
dusp6) the FGF pathway, the FGF pathway 
stimulates the Notch pathway (through 
dlc), and the Notch pathway 
represses the RA pathway (through 
cyp26a). (A) The 
Celf1 RNA-binding protein attenuates the Notch pathway by 
limiting the abundance of Rbpj protein through a post-
transcriptional control. Notch signalling depends on FGF 
stimulating 
and RA pathways. (B) When the TP MO impairs 
the repressive interaction between Celf1 and 
rbpj mRNA, 
rbpj is 
overexpressed, which stimulates the Notch pathway.

Consequently, the RA pathway is repressed (due to 
cyp26a overexpression) and the FGF pathway is repressed (due to 
fgf8 repression). FGF repression does not lead to Notch repression 
owing to 
rbpj overexpression, and relieving the repressive 
interaction between Celf1 protein and 
rbpj mRNA uncouples the 
Notch and RA pathways from the FGF pathway.

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Competing Interests

The authors have no competing interests to declare.

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mesoderm 

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