Unc5B Interacts with FLRT3 and Rnd1 to Modulate Cell Adhesion in *Xenopus* Embryos

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Abstract

The FLRT family of transmembrane proteins has been implicated in the regulation of FGF signalling, neurite outgrowth, homeotypic cell sorting and cadherin-mediated adhesion. In an expression screen we identified the Netrins receptors Unc5B and Unc5D as high-affinity FLRT3 interactors. Upon overexpression, Unc5B phenocopies FLRT3 and both proteins synergize in inducing cell deadhesion in *Xenopus* embryos. Morpholino knock-downs of Unc5B and FLRT3 synergistically affect *Xenopus* development and induce morphogenetic defects. The small GTPase Rnd1, which transmits FLRT3 deadhesion activity, physically and functionally interacts with Unc5B, and mediates its effect on cell adhesion. The results suggest that FLRT3, Unc5B and Rnd1 proteins interact to modulate cell adhesion in early *Xenopus* development.

Introduction

FLRT proteins comprise a small family of fibronectin type III domain (FNIII) and leucine-rich repeats (LRR) transmembrane proteins in vertebrates [1]. FLRT3, the best-characterized member of the family, can physically interact with FGF receptors and modulate FGF-ERK signalling [2]. In addition, FLRT3 induces homotypic cell sorting in cultured cells and in *Xenopus* embryos [3]. FGF signalling is dependent on the cytoplasmic tail, whereas FGF receptor binding is mediated by the FNIII domain of FLRT3 [2]. The extracellular LRR domains are dispensable for FGF signal activation [2], but are essential for the FLRT3-mediated cell sorting activity [3]. FLRT3 was also identified as a target gene of Nodal signalling, inhibiting cadherin adhesion in *Xenopus* early development through interaction with the Rho family GTPase Rnd1 [4]. In the mouse, FLRT3 knockout embryos display defects in ventral closure, headfold fusion and definitive endoderm migration [3], as well as disorganization of the basement membrane which leads to rupture of the anterior visceral endoderm [6], suggesting that cell adhesion is affected upon FLRT3 ablation. Furthermore, FLRT3 has been implicated in neurite outgrowth [7,8].

The Netrin receptor Unc5 of the immunoglobulin (Ig) superfamily is involved in axon guidance, vasculogenesis and apoptosis regulation. Unc5 was originally identified as a mutation resulting in an uncoordinated movement phenotype [9]. Four Unc5 paralogs (A to D) are present in vertebrates, where they regulate axon migration together with the Netrin coreceptor DCC [10]. In addition, Unc5 proteins show properties of dependence receptors and can induce apoptosis through their Death domain in the absence of Netrin binding [11,12]. Unc5B also plays a role in vascular morphogenesis [13,14]. Despite the important roles ascribed to the Unc5 receptors, currently there is limited knowledge of their binding partners, modulators and downstream targets.

Through a cell surface binding screen for FLRT3 partners, we identified Unc5B and Unc5D as high-affinity FLRT3 interactors. We show that FLRT3 and Unc5B functionally interact in modulating cell adhesion during early *Xenopus* development, and provide evidence that the Unc5B effect on adhesion is mediated by Rnd1.

Results

FLRT3 binds to Unc5

To identify interacting partners of the *Xenopus* FLRT3 ectodomain, we performed a cell surface binding screen [15]. Pools of about 250 clones prepared from a mouse embryonic cDNA expression library were transfected in HEK293T cells. Two days later, the cells were incubated with soluble alkaline phosphatase-FLRT3 ectodomain fusion protein (AP-FLRT3) and screened for FLRT3 interactors by a chromogenic assay for cell-surface bound AP activity. In this screen, a splice variant of the mouse *Unc5D* gene was isolated. FLRT3 was found to also interact with *Xenopus* Unc5B. FLRT3 interaction with Unc5B and -D is specific, as no binding was observed to other receptors of the Ig superfamily, such as Robo2 and -3 (Figure 1A). Additional in vitro binding and coimmunopre-
Figure 1. Unc5 proteins are high affinity interactors of FLRT3. (A) FLRT3 ectodomain binds to cells expressing Unc5. HEK293T cells were transfected (T) with the indicated expression constructs, incubated with AP-tagged FLRT3ΔTM conditioned medium (M) and stained for bound alkaline phosphatase (AP) activity. Schematic representation of the FLRT and Unc5 family domain structures is shown on top. (B) FLRT3 and Unc5D ectodomains interact in solution. Equal amounts of V5-tagged FLRT3ΔTM conditioned medium were mixed with the indicated FLAG-tagged condition media and subjected to immunoprecipitation (IP) and immunoblotting (IB). (C,D) Binding curves and Scatchard analyses of AP-FLRT3ΔTM binding to Unc5B or Unc5D transfected cells. The dissociation constants (K_d) are indicated. (E) The LRR domains of FLRT3 mediate binding to Unc5D. Unc5B Interacts with FLRT3
Whole cell lysates of HEK293T cells transfected with the indicated Renilla- (RL-) and V5-tagged constructs were immunoprecipitated with anti-V5 antibody. RL activity of individual samples after IP was normalized to the total activity of the respective lysate. (f) Both the IG and the TSP domains of Unc5D are required for FLRT3 binding. HEK293T cells were transfected with the indicated V5-tagged Unc5D expression constructs, incubated with recombinant AP-FLRT3ΔTM and stained for bound AP activity. Similar levels of protein production were confirmed by V5 immunoblotting (data not shown).

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Despite these efforts, it was also observed that FLRT3 and Unc5D ectodomains interact in solution with high specificity (Figure 1B). Next, the binding affinity of the FLRT3 ectodomain to Unc5 was determined in equilibrium binding experiments. The dissociation constant \( K_d \) of AP-FLRT3ΔTM was 50 nM with Unc5B and 26 nM with Unc5D (Figure 1C,D). These values are comparable to the Netrin-Unc5 \( K_d \) [16]. We conclude that Unc5 and FLRT3 ectodomains interact specifically and with high affinity.

To further characterize the FLRT3-Unc5 interaction, a series of FLRT3 deletion constructs was used in communoprecipitation experiments with Unc5-ΔN-RENilla fusion protein to determine the FLRT3 domain required for Unc5 binding. The interaction depends mostly on the LRR region of FLRT3 (Figure 1E), which is in contrast to the binding of FLRT3 to FGF receptors for which the FNIII domain is essential (Figure 1E and [2]). On the other hand, deletion of either the Ig- or the TSP-domains of Unc5D abolished FLRT3 interaction in cell surface binding experiments (Figure 1F), revealing that both domains are required for efficient FLRT3 binding.

Unc5B and FLRT3 induce embryonic cell deadhesion

To analyse if FLRT3 and Unc5 proteins interact not only biochemically but also functionally, we carried out overexpression experiments in Xenopus laevis embryos, where mRNA injection of FLRT3 has been shown to induce cell deadhesion [4]. Similar to FLRT3, Unc5B mRNA injection induced loss of cell adhesion and inhibited wound healing (Figure 2A). Deletion of the intracellular part of Unc5B abolished the deadhesion phenotype (Figure 2A, Unc5BAC), despite the fact that this construct was expressed at higher protein levels (Figure 2B). Expression of C-cadherin, which is the major cadherin in early Xenopus development [17] was not affected by Unc5B, as was also the case for FLRT3 (Figure 2B and data not shown). Unc5B effect on adhesion was specific and not due to cell toxicity or apoptosis, since (i) the detached cells remained viable and proliferative when cultured ex vivo (data not shown), (ii) deletion of the pro-apoptotic Death domain did not abolish cell deadhesion (Figure 2A, Unc5BΔD) and (iii) the phenotype could be rescued by C-cadherin overexpression or by Rnd1 depletion (see below). Nevertheless, to avoid potential toxic effects associated with the Death domain [10], subsequent experiments were performed with Unc5BAD. We conclude that Unc5B phenocopies FLRT3 in inducing cell deadhesion and that this effect requires its cytoplasmic part (excluding the Death domain).

To test for their functional interaction, limiting mRNA doses of FLRT3 and Unc5BAD were injected, which individually yielded no significant deadhesion (Figure 2C). Upon co-injection, however, a synergistic cell dissociation effect was observed in isolated animal caps and in whole embryos, indicating that both proteins can act together to interfere with cell adhesion (Figure 2C).

FLRT3-induced deadhesion was proposed to involve C-cadherin endocytosis and could be rescued by C-cadherin overexpression [4]. We therefore tested whether Unc5B operates through a similar mechanism. Indeed, C-cadherin was able to substantially rescue Unc5BAD-induced deadhesion (Figure 2D,E), suggesting that Unc5B, like FLRT3, can interfere with cadherin-mediated cell adhesion in Xenopus embryos. However, we failed to observe significant C-cadherin internalization upon either Unc5BAD or FLRT3 overexpression (data not shown).

Unc5B-FLRT3 interaction in Xenopus morphogenesis

Since multiple FLRT and Unc5 paralogs are expressed in early X. laevis development [2,4] and data not shown), and because of the existence of pseudoalleles in this tetraploid species, we chose the related diploid species X. tropicalis for loss-of-function experiments. Of the four Unc5 paralogs, Unc5B proved the most abundantly expressed gene throughout X. tropicalis development (Figure 3A and data not shown). Comparison of the spatio-temporal expression patterns of Unc5B and FLRT3 reveals overlapping expression domains in the mesoderm and the sensorial ectoderm at blastula stage, and later in the mesoderm at gastrula stage, tissues which undergo intense morphogenetic movements during gastrulation (Figure 3B,C). Also in later development the two genes show partially overlapping expression in certain tissues (Figure S1).

In order to test for functional interaction, we injected X. tropicalis embryos with Morpholino (MO) antisense oligos targeting FLRT3 and Unc5B individually or in combination (Figure 3D). Because of a lack of suitable antibodies to assess endogenous protein depletion and in order to achieve efficient knock-down effects [18], we used a mixture of two non-overlapping MOs for FLRT3 and Unc5B, targeting their ATG and UTR regions. While individual injection of FLRT3 or Unc5B MOs did not significantly affect early development, combined FLRT3/Unc5B morphant embryos had shorter bodies and smaller heads, suggesting morphogenetic defects (Figure 3D). The expression domain of the dorsal mesodermal marker Xnot in gastrula stage embryos was shorter and broader, a hallmark of impaired convergent extension movements during gastrulation (Figure 3D). Quantitative RT-PCR analysis at gastrula and neurula stages revealed no major changes in the expression of epidermal, neuroectodermal, neural crest, paraxial, dorsal and ventral mesodermal marker genes (Figure 4). This indicates that cell fate determination and tissue differentiation were not significantly affected, consistent with morphogenetic rather than differentiation defects. The fact that (i) only combined FLRT3/Unc5B MO injection yields phenotypic defects, which (ii) concern the dorsal mesoderm where both genes are coexpressed, and (iii) leaves cell differentiation markers largely unaffected, supports the specificity of the Morpholino knock-downs. We conclude that Unc5B and FLRT3 functionally interact during early Xenopus development.

Unc5B acts upstream of the small GTPase Rnd1

Both Unc5B and FLRT3, while interacting via their ectodomains, modulate cell adhesion through their cytoplasmic tails. The small GTPase Rnd1 was shown to bind to FLRT3 and to mediate its effect on cell adhesion [4]. We therefore tested whether Rnd1 can also interact with Unc5B.

In communoprecipitation assays Unc5BAD (which causes deadhesion) interacted robustly with Rnd1, whereas Unc5BAC
Figure 2. Unc5B and FLRT3 induce cell deadhesion in *Xenopus laevis* embryos. (A) Embryos were injected with the indicated mRNAs (1.2 ng FLRT3, 4 ng Unc5B variants) and at blastula stage 9 were photographed 10 min after partial excision of the animal caps. (B) V5 and C-cadherin immunoblots for protein production of sibling embryos from panel A. (C) FLRT3 and Unc5B functionally synergize. Embryos were injected with the indicated mRNAs (0.2 ng FLRT3, 1.6 ng Unc5BΔD) and cell deadhesion was assessed in animal caps of blastula stage 8.5 embryos (upper row), as well as in whole embryos at neurula stage 13, where ectodermal thinning and blastocoel edema were observed (lower row, indicated is the fraction of affected embryos). PPL, preprolactin (control mRNA). (D) C-cadherin rescue of Unc5B-induced deadhesion. Embryos were injected with the indicated
Rnd1 acts downstream of Unc5B to mediate its cell deadhesion by Rnd1 MO. These data indicate an epistatic relationship where the results showed substantial rescue of the deadhesion phenotype MO and monitored cell deadhesion at blastula stage (Figure 5E,F).

Discussion

In this study we discover a novel function of the Netrin receptor Unc5B to physically and functionally interact with FLRT3. We also find that Unc5B interacts with and signals via the small GTPase RhoA (Figure 5B).

To examine whether Unc5B-mediated loss of cell adhesion requires Rnd1, as was demonstrated for FLRT3 [4], we injected Unc5BAD mRNA in X. laevis embryos pre-injected with Rnd1 MO and monitored cell deadhesion at blastula stage (Figure 5E,F). The results showed substantial rescue of the deadhesion phenotype by Rnd1 MO. These data indicate an epistatic relationship where Rnd1 acts downstream of Unc5B to mediate its cell deadhesion activity.

Methods

Cell surface binding assays

Cells plated in poly-L-lysine-treated (Sigma, 100 μg/ml) 24-well culture plates were transiently transfected with expression constructs using FuGENE 6 (Roche). After 48 h, concentrated AP-FLRT3ATM conditioned media (3 to 8 U/ml AP activity) in AP-binding buffer (DMEM, 10% FCS, 50 mM HEPES) was added to the cells. After 1 h incubation at RT, cells were washed three times with PBS and stained for bound AP with Fast Red substrate (Roche). For binding curves, different amounts of concentrated AP or AP-FLRT3ATM CM in AP-binding buffer were added to transfected cells in 48-well plates (always at least in triplicates) and incubated for 1 h at RT. Cells were then harvested by pipetting, counted, washed 4 times with PBS containing 0.2 M NaCl, lysed in 70 μl per well 1% Triton X-100, 10 mM Tris (pH 8.0) and cleared by centrifugation. The supernatants were added to transfected cells in 96-well microtiter plates and measuring the absorbance at 405 nm in an ELISA reader (Multiskan RC, Labsystems). Background counts were subtracted from each data point and the absorbance normalized to cell number was calculated.

Embyrological assays

Xenopus embryo culture and manipulations were carried out following standard protocols and details are described in the Supplement (Text S1). Morpholino (MO) antisense oligonucleotides (Gene Tools) were as following: Control MO, tccttaactcacttataata (Gene Tools); Rnd1 MO, tgggtctctgtctctaggg [4]; FLRT3-ATG MO, atctcaagttcgg-
Figure 3. Unc5B-FLRT3 interaction in *Xenopus tropicalis* morphogenesis. (A) Developmental quantitative RT-PCR analysis of FLRT3 and Unc5A-D expression. (B–C) Expression patterns of FLRT3 and Unc5B at blastula stage 9 and gastrula stage 10.5. The dorsal blastopore lip is indicated with an arrow. DM, dorsal mesoderm. (D) Representative phenotypes of embryos injected with FLRT3 and/or Unc5B Morpholinos (MOs). A mixture of two MOs per gene targeting the ATG and the UTR regions was injected (10 ng each). A stunted axis was observed in 0% (n = 29), 3% (n = 34), 7% (n = 28) and 100% (n = 31) of the injected embryos, respectively. Lower row: whole mount in situ hybridization of *Xnot* at late gastrula stage 12. doi:10.1371/journal.pone.0005742.g003
gacattgc [4]; FLRT3-UTR MO, gatttcgagctatagagctct; Unc5B-ATG MO, gatataaatgcatcgcttagc; gacattgcagctct. FLRT3-UTR MO, gacattgcagctct. FLRT3-ATG and Unc5B-UTR MOs have one single terminal mismatch each in X. tropicalis, which should not significantly affect their activity [18].

Additional experimental methods and reagents are described in the Supplement (Text S1).

Supporting Information

Figure S1 Spatio-temporal expression patterns of FLRT3 and Unc5B in Xenopus tropicalis development

Text S1 Supplementary Materials and Methods

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Author Contributions

Conceived and designed the experiments: EK RTB CN. Performed the experiments: EK RTB PS WW. Analyzed the data: EK RTB CN. Contributed reagents/materials/analysis tools: EK RTB WW MR SO KC. Wrote the paper: EK CN.
Figure 5. Unc5B acts upstream of the small GTPase Rnd1. (A,B) Rnd1 binds to Unc5BΔD. Immunoprecipitation (IP) assays followed by immunoblotting (IB) with lysates of HEK293T cells overexpressing the indicated V5- and HA-tagged proteins. (C) Unc5B and Rnd1 synergize in cell deadhesion. *X. laevis* embryos were injected with the indicated mRNAs (0.8 ng Unc5BΔD, 0.16 ng Rnd1). Loss of adhesion was observed as ectodermal thinning and blastocoel edema at neurula stage 13 (upper row), as well as in paraffin sections of blastula stage 9 embryos (lower row). PPL, preprolactin (control mRNA). (D) Quantification of the deadhesion phenotypes from panel C (stage 9 embryos). (E) Rnd1 Morpholino (MO) injection rescues Unc5B-induced deadhesion. *X. laevis* embryos were injected sequentially with the indicated MO (40 ng) and mRNA (4 ng), and grown until blastula stage 9 in high-salt medium, which promotes ectodermal lesions when adhesion is inhibited (upper row) or in normal medium (lower row, paraffin sections). (F) Quantification of the Rnd1 MO rescue. Two embryo batches (exp.1,2) were injected as in panel E, grown in normal medium and cell deadhesion was assessed at blastula stage 9.

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References


