RESEARCH ARTICLE

Semaphorin 6B acts as a receptor in post-crossing commissural axon guidance

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ABSTRACT

Semaphorins are a large family of axon guidance molecules that are known primarily as ligands for plexins and neuropilins. Although class-6 semaphorins are transmembrane proteins, they have been implicated as ligands in different aspects of neural development, including neural crest cell migration, axon guidance and cerebellar development. However, the specific spatial and temporal expression of semaphorin 6B (Sema6B) in chick commissural neurons suggested a receptor role in axon guidance at the spinal cord midline. Indeed, in the absence of Sema6B, post-crossing commissural axons lacked an instructive signal directing them rostrally along the contralateral floorplate border, resulting in stalling at the exit site or even caudal turns. Truncated Sema6B lacking the intracellular domain was unable to rescue the loss-of-function phenotype, confirming a receptor function of Sema6B. In support of this, we demonstrate that Sema6B binds to floorplate-derived plexin A2 (PlxnA2) for navigation at the midline, whereas a cis-interaction between PlxnA2 and Sema6B on pre-crossing commissural axons may regulate the responsiveness of axons to floorplate-derived cues.

KEY WORDS: Spinal cord development, In ovo RNAi, Cis-interaction, PlexinA2, Chick

INTRODUCTION

During development, a large number of neurons must connect with their target cells to establish functional neural circuits. A well-orchestrated set of axon guidance cues and their receptors on growth cones directs axonal navigation through the pre-existing tissue. On their journey, axons contact one or several intermediate targets before they reach their final destination. The floorplate at the CNS midline is one such intermediate target, where commissural axons cross to the contralateral side and grow along the rostrocaudal axis (Nawabi and Castellani, 2011; Chédotal, 2011). Owing to their stereotypic trajectory, dI1 commissural axons are a well-studied model for axon guidance. A number of long-range guidance cues directing these axons ventrally toward the floorplate have been identified (Kennedy et al., 1994; Augsburger et al., 1999; Charron et al., 2003; Islam et al., 2009). Midline crossing is regulated by a shift from attraction to repulsion (Stoeckli and Landmesser, 1995; Stoeckli et al., 1997; Long et al., 2004; Sabatier et al., 2004; Mametisawa et al., 2005; Chen et al., 2008). Axonal repulsion is induced by the upregulation of Robo1 on the growth cone surface, which allows for the detection of Slit, a negative floorplate-derived signal (Philipp et al., 2012). Axonal navigation at the floorplate exit site is subsequently directed by F-spondin (Burston-Cohen et al., 1999), SynCAMs/Nectin-like molecules (Niederkofler et al., 2010), MDGA2 (Joset et al., 2011), and the morphogens Wnts (Lyuksyutova et al., 2003; Domanitskaya et al., 2010) and sonic hedgehog (Shh) (Bourikas et al., 2005; Wilson and Stoeckli, 2013).

Semaphorin-plexin signaling also has been implicated in guiding commissural axons (Zou et al., 2000; Nawabi et al., 2010). GDNF- and NrCAM-mediated inhibition of the protease calpain 1 stabilizes plexin A1 (PlxnA1) on the growth cone (Nawabi et al., 2010; Charoy et al., 2012). The receptor complex neuropilin 2-PlxnA1 triggers sensitivity to the repulsive function of Sema3B, thus expelling axons from the midline area. Interestingly, Shh is also involved in this switch by downregulating protein kinase A (PKA) activity and thus inducing growth cone sensitivity to midline-derived class-3 semaphorins (Porra and Zou, 2010).

In contrast to the well-studied class-3 semaphorins, much less is known about class-6 semaphorins (Pasterkamp and Kolodkin, 2003; Kolodkin and Tessier-Lavigne, 2011; Pasterkamp, 2012). Class-6 semaphorins are a family of four type-I transmembrane proteins that bind directly to class-A plexins (plexinAs) in a neuropilin-independent manner. They have mainly been implicated as ligands for plexinAs. For example, Sema6A and Sema6B are required in the navigation of cardiac neural crest cells. They interact with PlxnA2 for proper cardiac outflow tract formation (Toyofuku et al., 2008). In axon guidance, Sema6A and Sema6B cooperate in the regulation of hippocampal mossy fiber targeting via interactions with PlxnA2 and PlxnA4 (Suto et al., 2007; Tawarayama et al., 2010). Sema6A directs the growth of sensory and sympathetic neurons (Xu et al., 2000; Haklai-Topper et al., 2010), and Sema6A/PlxnA4 signaling regulates the dendritic development of motoneurons (Zhuang et al., 2009). Sema6A also regulates cerebellar granule cell migration by controlling nucleus-centrosome coupling (Kerjan et al., 2005; Renaud et al., 2008). At the optic chiasm, Sema6D interacts with PlxnA1 and NrCAM to promote midline crossing of retinal axons (Kuwajima et al., 2012).

Although class-6 semaphorins act as ligands in all these processes, their intracellular domains, which include Src homology-3 (SH3) and zyxin-like domains, also suggest receptor functions (Eckhardt et al., 1997; Klostermann et al., 2000; Toyofuku et al., 2004a; Kolodkin and Tessier-Lavigne, 2011; Pasterkamp, 2012). This idea is supported by the analysis of corticospinal tract formation in Sema6a knockout mice, in which Sema6A may guide axons in a cell-autonomous manner. However, Sema6A is also expressed in the surrounding area, making it impossible to distinguish between cell-autonomous versus non-autonomous functions (Leighton et al., 2001; Rünker et al., 2008). A receptor function for Sema6A was also suggested by our previous
studies on boundary cap cell clustering (Mauti et al., 2007). Boundary cap cell clusters act as gate keepers, allowing axons, but not cell bodies, to cross the CNS/PNS boundary. After knocking down Sema6A in boundary cap cells, motoneurons migrated out of the spinal cord along the ventral roots. This phenotype could be explained by a model in which Sema6A is a receptor on migrating boundary cap cells that recognizes PlxnA1 on motor axons as a stop signal. The Sema6A-PlxnA1 interaction would lead to the accumulation of boundary cap cells and indirectly initiate their clustering at the ventral motor exit point (Mauti et al., 2007).

So far, the only direct evidence for a receptor function of class-6 semaphorins in vivo comes from heart development, where Sema6D was shown to be a ligand and receptor for PlxnA1 (Toyofuku et al., 2004a,b).

Here, we demonstrate in chick that Sema6B is an axon guidance receptor. We show that Sema6B guides post-crossing commissural axons by binding to PlxnA2 expressed by floorplate cells. In the absence of axonal Sema6B or its ligand PlxnA2 in the floorplate, post-crossing commissural axons fail to turn rostrally along the longitudinal axis of the spinal cord.

RESULTS

Sema6b is transiently expressed in dorsal commissural neurons

Sema6b mRNA was readily detectable in dI1 commissural neurons of the chicken neural tube during the time window when their axons cross and exit the floorplate (Fig. 1A-F). Importantly, Sema6b mRNA was only detectable at HH22 (Fig. 1C), when most axons have reached the ipsilateral floorplate border. No Sema6b expression was detectable in commissural neurons at HH21, that is, shortly before their axons reach the floorplate (Fig. 1B,E). Highest expression levels were found at HH24, when commissural axons exit the floorplate and turn rostrally along the contralateral floorplate border (Fig. 1D,E). At later stages (not shown), Sema6b was found at lower levels throughout the gray matter, consistent with findings in mouse (Suto et al., 2005). The identity of dI1 neurons was confirmed by labeling adjacent sections with probes for the marker Lhx2/9 (Helms and Johnson, 2003) (Fig. 1F,G).

As we described previously, Sema6a was not expressed in commissural neurons at the lumbar level (Mauti et al., 2007; but see Fig. 1.

Fig. 1. Sema6b expression in dI1 commissural neurons peaks at HH24 and is required for post-crossing commissural axon guidance. (A-E) Expression of Sema6b mRNA in transverse sections of the chicken lumbar spinal cord at the indicated developmental stages (A-D) with schematic representations of the corresponding growth of dI1 axons (E). Sema6b is first detected in dI1 neurons at HH22, when their axons have reached the ipsilateral floorplate border (C, arrowhead). Expression of Sema6b in dI1 neurons is strongest at HH24 (D, arrowheads). (H,J) No staining is seen with a Sema6b sense probe. (J) Diffuse expression of Sema6d at HH25, with higher levels in motoneurons. Neither Sema6b nor Sema6d is expressed in the floorplate (D,J, open arrowheads). (K) Schematic of an open-book preparation and application of Dil for axonal tracing (red). D, dorsal; V, ventral; R, rostral; C, caudal. (L) Commissural axons cross the midline and extend along the contralateral floorplate border (arrows) in untreated control embryos. (M,N) After silencing Sema6B by injection and electroporation of dsRNA (dsSema6b), axons stall at the contralateral floorplate border (closed arrowheads) or even turn caudally (open arrowhead). (O) Silencing Sema6D by injection and electroporation of dsRNA (dsSema6d) did not affect axon guidance (arrows). (P) Expression of a co-electroporated EGFP reporter confirmed the exclusively dorsal targeting of dsRNA (asterisks). Arrows mark axons of contraterally projecting commissural neurons. The floorplate is indicated by dashed lines. (Q) Quantification of Dil injection sites with aberrant axonal pathfinding. ***P<0.001; error bars indicate s.e.m. Scale bars: 50 µm in A-D,F-J; 100 µm in L-P.
dorsal expression at the brachial level described by Toyofuku et al., 2008). *Semat6d* was strongly expressed in motor neurons and at lower levels in the spinal cord gray matter (Fig. 1J) (Mauti et al., 2007). No class-6 semaphorins were expressed in the floorplate (Fig. 1) (Mauti et al., 2007). No ortholog of *Semat6c* is found in the chicken genome.

**Loss of Sema6B causes defects in commissural axon guidance**

The transient expression of *Sema6b* in dI1 neurons during the window of axonal midline crossing and turning suggested a role for Sema6b in axonal pathfinding. To test this hypothesis, we used *in ovo* RNAi to knock down Sema6B in commissural neurons (Fig. 1K-Q). Long dsRNA derived from *Sema6b* or *Sema6d*, together with an EGFP reporter plasmid, was injected into the central canal at HH18 and electroporated into the dorsal spinal cord (Fig. 1P). Commissural axon pathfinding was assessed 2 days later by axonal tracing with lipophilic DiI applied to the cell bodies of dI1 neurons (Fig. 1K-O).

In untreated control embryos, commissural axons crossed the floorplate and turned rostrally along the contralateral floorplate border (Fig. 1L,Q; 95.3±2.5% of DiI sites were normal; N=19 embryos, N=164 DiI injection sites). When Sema6B was downregulated, axons failed to turn rostrally and instead stalled or even turned caudally at the floorplate exit site in 50.1±5.4% of the DiI sites (n=22, N=149; Fig. 1M,N,Q). By contrast, loss of Sema6D did not affect the guidance of post-crossing axons (abnormalities at only 10.2±5.7% of injection sites; n=13, N=84; Fig. 1O,Q). Taken together, these experiments indicated that downregulation of Sema6B specifically interfered with commissural axon guidance upon floorplate exit.

**Sema6B acts as a receptor in commissural axons**

Semaphorins are typically known as ligands for plexin receptors. Sema6B acts as a receptor in commissural axons and that Sema6B functions cell-autonomously as an axon guidance receptor.

**PlxnA2 is a potential interaction partner for Sema6B in commissural axon guidance**

We next sought a potential floorplate-derived ligand for the Sema6B receptor. Based on the known interactions of class-6 semaphorins with plexinAs (Suto et al., 2005; Mauti et al., 2007; Toyofuku et al., 2008), we assessed the expression patterns of plexinAs during commissural axon pathfinding. As we described previously (Mauti et al., 2006), plexins have dynamic and distinct expression patterns in the chicken lumbar spinal cord (Fig. 3). At HH21, just before the first dI1 commissural axons reach the floorplate at lumbar levels of the spinal cord, *Plxna1* was strongly expressed in the ventral spinal cord (including the floorplate), and at lower levels throughout the rest of the neural tube (Fig. 3A). By HH25, *Plxna1* expression was high in dI1 neurons and in motoneurons (Fig. 3A’), but was no longer expressed in the floorplate (inset in Fig. 3A’). *Plxna2* was expressed strongly in dI1 neurons and the floorplate at both HH21 and HH25 (Fig. 3B,B’). *Plxna4* was expressed in dI1 neurons, but was not found in the floorplate (Fig. 3C,C’). By contrast, *Plxnc1* was found in the floorplate (Fig. 3D,D’). No ortholog of *Plxna3* is found in the chicken genome.

Consistent with their potential role as interaction partners for Sema6B, unilateral downregulation of plexinAs interfered with commissural axon pathfinding (Fig. 3E-H). The loss of *Plxna1* caused post-crossing defects at 43.8±7.3% of DiI sites (n=16, N=106; Fig. 3E,M). After downregulation of *Plxna2*, post-crossing commissural axons failed to turn rostrally at 54.1±9.1% of injection sites (n=15, N=92; Fig. 3F,M). The same phenotype was observed at 46.3±11.3% of injection sites after downregulation of *Plxna4* (n=11, N=66; Fig. 3G,M).

Based on these results, we analyzed the requirement for plexinAs in post-crossing commissural axon guidance more specifically by targeted electroporation of the floorplate (Fig. 3I-L). Because *Plxna1* expression disappeared from the floorplate during axonal crossing and turning, and based on biochemical interaction assays that indicated that PlxnA1 did not bind to Sema6B (Toyofuku et al., 2008; and see below), we instead considered PlxnA2 as the most likely binding partner for axonal Sema6B in the floorplate. Indeed, after knocking down PlxnA2 in the ventral spinal cord, we found contralateral stalling phenotypes comparable to those observed in the absence of Sema6B at 36.2±5.2% of injection sites (n=18, N=149; Fig. 3I,N). By contrast, ventral electroporation of *Plxna4* dsRNA did not interfere with commissural axon guidance (Fig. 3J,N). As *Plxna4* was never found in the floorplate (Fig. 3C,C’), this experiment was a negative control. Aberrant axonal behavior was detected at only 5.0±2.6% of injection sites (n=7, N=56). Downregulation of PlxnC1, which is expressed in the floorplate but interacts with class-7 semaphorins (Pasterkamp, 2012), did not interfere with commissural axon navigation: aberrant behavior was detected at only 5.8±4.1% of injection sites (n=12, N=75; Fig. 3K,N). Together, the targeted electroporation experiments supported a specific axon guidance role of floorplate-derived PlxnA2.
The intracellular domain of PlxnA2 is dispensable for its axon guidance function in the floorplate

If PlxnA2 was acting as a floorplate-derived ligand for Sema6B during commissural axon guidance, then the cytoplasmic domain of PlxnA2 could be dispensable for its axon guidance activity in the floorplate. To test this idea, we downregulated PlxnA2 specifically in the floorplate using an miRNA (miPA2), then attempted to rescue the phenotype by expressing miRNA-resistant constructs encoding either the full-length protein (PlexinA2ΔmiR) or a truncated version containing the extracellular and the transmembrane domains but lacking the cytoplasmic tail (PlexinA2ΔCTΔmiR). The miRNA and rescue constructs were verified in vivo and in vitro (supplementary material Fig. S2). All constructs were under the control of a Hoxa1 enhancer in order to drive specific expression in the floorplate (Fig. 4A) (Wilson and Stoeckli, 2011).

As expected, the miRNA-based knockdown of PlxnA2 specifically in the floorplate caused axon pathfinding errors. Whereas the control group injected and electroporated with miLuc and pMES showed normal axon guidance at 60.9±8.5% of the injection sites per embryo (n=12 embryos, N=107 injection sites), downregulation of PlxnA2 specifically in the floorplate with miPA2 reduced normal axon guidance to only 30.6±6.1% of the injection sites (n=14, N=97). Axon guidance was rescued by co-injection and electroporation of knockdown-resistant full-length PlxnA2 (PlexinA2ΔmiR; normal axon guidance at 63.2±5.5% of the injection sites per embryo; n=14, N=97) and truncated PlxnA2 (PlexinA2ΔCTΔmiR; normal axon guidance at 54.2±5.1% of the injection sites per embryo; n=17, N=119). The integrity of the floorplate was not affected in these experiments, as the expression of neither markers (Hnf3β and Nkx2.2) nor other axon guidance molecules (Shh and Slit2) was perturbed (supplementary material Fig. S4).

Together, these results suggest that PlxnA2 acts directly as an axon guidance molecule in the floorplate. The extracellular domain of PlxnA2, and not its intracellular domain, is crucial for this function. This finding was consistent with our hypothesis that PlxnA2 is a floorplate-derived ligand for Sema6B during commissural axon guidance.
Sema6B interacts with PlxnA2

To support the above hypothesis, we next confirmed a physical interaction between Sema6B and PlxnA2 in co-immunoprecipitation (co-IP) and binding assays. The co-IP of Sema6A with PlxnA2 was used as positive control, as this interaction has been described previously (Suto et al., 2005, 2007; Toyofuku et al., 2008; Janssen et al., 2010; Nogi et al., 2010). Immunoprecipitation of myc-tagged PlxnA2 ectodomains from conditioned medium was found to pull down both Sema6B-Fc and Sema6A-Fc ectodomains (Fig. 4F-I).

We confirmed the interaction between Sema6B and PlxnA2 by cell-binding assays. HEK293 cells expressing different class-6 semaphorins were incubated with PlxnA2 ectodomains fused to alkaline phosphatase (PlexinA2<sup>ecto</sup>-AP; Fig. 4J-N). As expected, binding of PlexinA2<sup>ecto</sup>-AP to Sema6A-expressing cells was very strong (Fig. 4J). PlexinA2<sup>ecto</sup>-AP binding to Sema6B-expressing cells was clearly detectable, but weaker than to Sema6A-expressing cells (Fig. 4K). PlexinA2<sup>ecto</sup>-AP also bound to Sema6BCT (Fig. 4M). However, no binding of PlexinA2<sup>ecto</sup>-AP was found to cells transfected with Sema6D (Fig. 4L) or a control EGFP plasmid (Fig. 4N). By contrast, PlexinA1<sup>ecto</sup>-AP bound to cells expressing Sema6A (Fig. 4O) and Sema6D (Fig. 4Q) but not to cells expressing Sema6B (Fig. 4P,R). Thus, we could exclude PlxnA1 as a Sema6B interaction partner. Taken together, our results confirmed a previously reported interaction between Sema6B and PlxnA2 (Toyofuku et al., 2008), and strongly supported our hypothesis that PlxnA2 is a floorplate-derived ligand for Sema6B during commissural axon guidance.

Sema6B mediates an outgrowth response of commissural neurons to PlxnA2

We next investigated whether the response of commissural axons to PlxnA2 substrate was altered by a lack of Sema6B (Fig. 5). On coverslips coated with Albumax, Laminin or concentrated conditioned medium from cells expressing either AP alone or PlexinA2<sup>ecto</sup>-AP, axons failed to turn rostrally at the contralateral floorplate border (arrowheads). (H) Only injection sites in the electroporated area (verified by EGFP expression) were included in the analysis. (I-K) Analysis of commissural axon pathfinding after downregulation of PlxnA2 (l, arrowheads), post-crossing axons failed to turn into the longitudinal axis or even turned caudally. Axon guidance was unaffected after ventral downregulation of PlxnA4 (J, arrow) or PlxnC1 (K) exclusively in the ventral spinal cord. After ventral downregulation of PlxnA2 (l, arrowheads), post-crossing axons failed to turn into the longitudinal axis or even turned caudally. Axon guidance was unaffected after ventral downregulation of PlxnA4 (l, arrow) or PlxnC1 (K, arrow).

(M,N) Quantification of injection sites with aberrant axonal pathfinding after (M) unilateral or (N) ventral electroporation of dsRNA. ***<i>P</i> < 0.001; **<i>P</i> < 0.01; error bars indicate s.e.m. Scale bars: 50 µm in A-D’; 100 µm in E-L.
comprised both wild-type and miRNA-expressing (EBFP2-positive) neurons from the same embryos. Thus, the wild-type neurons provided an internal control. Cultures were allowed to grow for 48 h before fixation and immunolabeling for the axonal marker axonin 1 (contactin 2).

As expected, we observed modest outgrowth of commissural axons on Albumax (Fig. 5G; data not shown), whereas the Laminin and AP substrates encouraged slightly longer axons (Fig. 5A-D,G). By far the longest axons were found on PlxnA2-coated coverslips (Fig. 5E-G). However, the expression of miS6B (but not miLuc) significantly dampened this outgrowth response to PlxnA2 (Fig. 5E-H). These results indicate that PlxnA2 promotes the outgrowth of commissural axons in a pathway that is mediated by Sema6B.

**Axonal PlxnA2 contributes to post-crossing commissural axon guidance**

Our functional analyses suggested that PlxnA2 is a floorplate-derived ligand for Sema6B during commissural axon guidance. However, Plxna2 was expressed not only in the floorplate but also in commissural neurons (Fig. 3B). Because PlxnA1 has previously

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**Fig. 4.** The extracellular domain of PlxnA2 mediates its axon guidance activity in the floorplate and binds to Sema6B.

(A) Schematics of the constructs used in B-E. Hoxa1 drives floorplate-specific expression.
(B-D) Open-book analysis of embryos co-electroporated with: (B) miPA2 and Hox-EGFP; (C) miPA2 and Hox-PA2ΔmiR; or (D) miPA2 and Hox-PA2ΔCTmiR. (B’) The successful and exclusive targeting of floorplate cells was confirmed by EGFP and EBFP2 fluorescence from the co-electroporated constructs. Post-crossing axons that failed to turn correctly at the contralateral floorplate border are indicated by arrowheads (B). Arrows (C,D) indicate normal crossing and turning of commissural axons.
(E) Quantification of injection sites with normal axon pathfinding after floorplate-specific manipulations of PlxnA2. *P<0.05, **P<0.01; n.s., not significant; error bars indicate s.e.m.
(F) Schematics of the proteins used for co-IPs. (G) Sema6BΔε, or Sema6AΔε ectodomains were incubated with either PA2Myc ectodomains or APMyc and immunoprecipitated with anti-myc agarose beads. Co-IP of Sema6B and Sema6A was detected with anti-myc antibodies on western blots (upper panel). Blots stained with anti-myc antibodies, to demonstrate the successful immunoprecipitation, are shown below. As controls, the input proteins from conditioned media were analyzed on blots developed with (H) anti-Fc antibodies to detect Sema6B and Sema6A ectodomains and (I) anti-myc antibodies to detect PA2Myc and APMyc. (J-S) HEK293T cells were transfected with chicken Sema6A (J), Sema6B (K), Sema6D (L), Sema6BΔCT (M) or EGFP (N, S) and incubated with conditioned medium containing AP-tagged ectodomains of PlxnA2 (PlxnA2ΔCT-AP, J-N) or PlxnA1 (PlxnA1ΔCT-AP, O, S). Scale bars: 100 μm.
been identified as an axonally expressed receptor with important roles in commissural axon guidance (Nawabi et al., 2010; Charoy et al., 2012), we hypothesized that other plexinAs might function similarly. To test this idea, we knocked down PlxnA2 and PlxnA4 only in the dorsal spinal cord by targeted electroporation of long dsRNA. In line with results from unilateral electroporations, knockdown of PlxnA2 in commissural neurons perturbed axon guidance at 44.3±8.1% of injection sites (n=15 embryos, N=115 injection sites; Fig. 6A,D). Similarly, dorsal downregulation of PlxnA4 interfered with post-crossing axon guidance at 38.9±6.0% of injection sites (n=23, N=153; Fig. 6D). Based on these results, both PlxnA2 and PlxnA4 could act as guidance receptors on commissural axons after crossing the midline. Furthermore, these findings suggested the possibility of axonal cis-interactions with Sema6B.

To investigate this idea, we examined the localization patterns of Sema6B and plexinAs in more detail. In vivo, the expression of PlxnA2 was strongest on pre-crossing and commissural segments (Fig. 6E). Unfortunately, staining of Sema6B on sections was not possible with the available antibody, but our earlier in situ results (Fig. 1) suggested that Sema6b was only expressed in pre-crossing axons shortly before they contacted the floorplate. In cultured commissural neurons obtained from HH25 embryos, we found expression of Sema6B all along the axons and on growth cones (Fig. 6F). PlxnA2 (Fig. 6H,K) was expressed similarly. By contrast, PlxnA4 was found in a punctate pattern along the axons (Fig. 6I), suggesting an intracellular, vesicular localization. Additionally, only very low levels of PlxnA4 were found in growth cones (Fig. 6L), making the formation of a cis-complex with Sema6B on the growth cone or axonal surface unlikely.

Taken together, the co-expression of PlxnA2 and Sema6B in similar subcellular locations, together with their ability to interact with each other (Fig. 4; supplementary material Fig. S5), suggested a possible cis-interaction between these molecules on commissural axons as they enter and cross the floorplate.

**PlxnA2 misexpression in commissural neurons leads to axon stalling in the dorsal neural tube**

As knockdown of PlxnA2 in commissural neurons impaired axon guidance (Fig. 6), we tested whether it was required cell-autonomously for growth cone turning (Fig. 7; supplementary material Fig. S2). We used the previously described rescue constructs encoding either full-length PlxnA2 or PlexinA2ΔCT (supplementary material Fig. S2). This time, the miRNA and rescue constructs were driven by a dI1 neuron-specific enhancer of Math1 (Atoh1) (Fig. 7A; supplementary material Fig. S2) (Helms and Johnson, 2003; Wilson and Stoeckli, 2011). However, the...
pre-crossing commissural axons is prevented from mediating a floorplate, where levels of class-3 semaphorins are high, PlxnA2 on derived from the ventral spinal cord, and axons extend readily activated by the low levels of repulsive class-3 semaphorins PlxnA2 expressed on pre-crossing commissural axons is not receptor or co-receptor, when expressed on commissural axons, we interaction between Sema6B and PlxnA2. As PlxnA2 is required semaphorins and failed to extend into the ventral spinal cord. experimentally (Fig. 7B-E), axons were hypersensitive to class-3 axons extend ventrally. However, when PlxnA2 levels were increased are low, as are the levels of repulsive class-3 semaphorins. Therefore, the floorplate. In the dorsal spinal cord, endogenous levels of PlxnA2 spinal cord before floorplate contact to allow growth cone entry into signaling is normally inhibited in pre-crossing axons in the ventral spinal cord, whereas too little PlxnA2 prevents the proper pathfinding of post-crossing axons. Although the overexpression of PlxnA2ACT did not affect the ventral axonal trajectory (in all embryos analyzed, n=7), it nevertheless failed to rescue the post-crossing axon guidance defects (supplementary material Fig. S2D,E).

Taken together, these findings suggest that: (1) the PlxnA2 intracellular domain transduces axonal signals that are required for post-crossing axon pathfinding; and (2) PlxnA2 intracellular signaling is normally inhibited in pre-crossing axons in the ventral spinal cord before floorplate contact to allow growth cone entry into the floorplate. In the dorsal spinal cord, endogenous levels of PlxnA2 are low, as are the levels of repulsive class-3 semaphorins. Therefore, axons extend ventrally. However, when PlxnA2 levels were increased experimentally (Fig. 7B-E), axons were hypersensitive to class-3 semaphorins and failed to extend into the ventral spinal cord.

In summary, our in vivo analyses suggest a complex mode of interaction between Sema6B and PlxnA2. As PlxnA2 is required both as a ligand, when expressed on floorplate cells, and as a receptor or co-receptor, when expressed on commissural axons, we suggest the following model (Fig. 8). In the dorsal spinal cord PlxnA2 expressed on pre-crossing commissural axons is not activated by the low levels of repulsive class-3 semaphorins derived from the ventral spinal cord, and axons extend readily towards the floorplate (Fig. 8A). However, in the vicinity of the floorplate, where levels of class-3 semaphorins are high, PlxnA2 on pre-crossing commissural axons is prevented from mediating a repulsive signal by cis-interaction with Sema6B, which is now expressed on pre-crossing axons of d1l neurons (Fig. 8B). Upon floorplate contact (Fig. 8C), the cis-complex between axonal PlxnA2 and Sema6B is replaced by a trans-interaction between Sema6B and floorplate PlxnA2. The change from Sema6B-PlxnA2 cis- to trans-interaction mediates a Sema6B-dependent turning signal. Axonal PlxnA2 is available for cis-interaction with neuropilin, resulting in a repulsive signal on post-crossing axons (Navabi et al., 2010; Parra and Zou, 2010; Zou et al., 2000).

**DISCUSSION**

Our analyses of Sema6B function in post-crossing commissural axon guidance provide the first report of a receptor function of class-6 semaphorins in vertebrate axon guidance. Our evidence strongly supports the conclusion that Sema6B on commissural axons is the receptor that binds PlxnA2 expressed at the intermediate target: (1) Sema6B is expressed by commissural neurons, PlxnA2 is expressed by floorplate cells, and a direct interaction between Sema6B and PlxnA2 was shown in binding studies and by co-IP; (2) our loss-of-function studies revealed similar defects in post-crossing commissural axon guidance after silencing Sema6B or PlxnA2; (3) the axon guidance effects of Sema6B were dependent on the presence of its intracellular domain; (4) the extracellular domain of PlxnA2 was sufficient to rescue the effects of PlxnA2 knockdown in the floorplate; and (5) a substrate of PlxnA2 ectodomains enhanced the outgrowth of commissural axons in a Sema6B-dependent manner. Although our in vitro assay (Fig. 5) did not elucidate the guidance activities of PlxnA2 because the axons were not faced with a choice of substrate), our results suggest that PlxnA2 is a floorplate-derived ligand that affects axonal behavior in a Sema6B-mediated manner.

A receptor function for class-6 semaphorins is not without precedent. In vertebrates, our previous studies suggested a receptor role of Sema6A in boundary cap cell clustering (Mauti et al., 2007). In addition, Sema6D acts as both ligand and receptor in heart development (Toyofuku et al., 2004a,b). In invertebrates, there is direct evidence that transmembrane semaphorins are axon guidance receptors. Sema1a, which is the closest *Drosophila* homolog of
class-6 semaphorins, was shown to act as a receptor in R-cell axon guidance in the visual system (Cafferty et al., 2006; Yu et al., 2010), with repulsive class-3 semaphorin activity high (large blue diamond). Pre-crossing axons along the contralateral floorplate border. Note that we have drawn PlxnA2 (green) arbitrarily as dimer or monomer based on reports on the crystal structures of Sema6A and PlxnA2 (for details see Janssen et al., 2010; Nogi et al., 2010). For simplicity, we omitted neuropilins, which would form complexes with plexins. (B) In the ventral spinal cord, close to the floorplate, where repulsive class-3 semaphorin activity is high (large blue diamond), PlxnA2 (green) is prevented from mediating a repulsive signal due to the cis-interaction of axonal PlxnA2 with Sema6B, which is not involved in any cis-interaction with Sema6B, may bind to growth cone-derived Sema6B in trans, thus inducing a growth/turning signal in the axons. Furthermore, axonal PlxnA2 is freed to interact with class-3 semaphorins derived from the midline. Together, these pathways facilitate the post-crossing trajectory (Fig. 8).

At present, it is not known how the turning signals are mediated intracellularly. However, we propose that the axonal PlxnA2 and Sema6B pathways are separated by the Sema6B-PlxnA2 trans-interaction, resulting in distinct Sema6B- and PlxnA2-mediated parallel signaling in post-crossing axons. This scenario would explain why commissural axons acquire responsiveness to class-3 semaphorins only when crossing the floorplate, but not before. Based on its precise temporal and spatial coincidence with axonal midline crossing, the switch from Sema6B-PlxnA2 cis-interaction to trans-interaction constitutes an excellent regulatory mechanism that contributes to and strengthens the previously described mechanisms: the Shh-induced regulation of PKA activity (Parra and Zou, 2010) and the GDNF-mediated and
Nrcam-dependent inhibition of calpain, which in turn stabilizes PlxnA1 on the growth cone (Nawabi et al., 2010; Charoy et al., 2012).

MATERIALS AND METHODS

Plasmids and miRNAs

Primers for cloning and mutagenesis are listed in supplementary material Table S1; for cloning details see supplementary material methods. miRNA target sequences are listed in supplementary material Table S2. Full-length Sema6B (GenBank accession number KJ201030) and a truncated form maintaining the transmembrane region and the five adjacent C-terminal amino acids, but lacking the rest of the intracellular domain, were cloned into pMES (Wilson and Stoeckli, 2011) for in vivo studies and into pcDNA3.1 (Invitrogen) for in vitro studies. Site-directed mutagenesis (Zheng et al., 2004; Wilson and Stoeckli, 2011) was used to introduce six silent mutations into the miS6B target site of Sema6b to make it resistant to knockdown (supplementary material Fig. S1). The same strategy was used to synthesize a knockdown-resistant version of Plxna2 (supplementary material Fig. S2). Constructs for the expression of miRNAs against Sema6b and PlxnA2 were synthesized as described (Wilson and Stoeckli, 2011). GenScript Target Finder was used to predict miRNA target sequences.

In situ probes, dsRNA and immunohistochemistry

Expressed sequence tags (cESTs) were used to generate dsRNA and in situ probes (supplementary material Table S3). In situ hybridization and dsRNA synthesis were performed as described (Mauti et al., 2006; Pekarik et al., 2003). Antibodies used for expression analyses on 20 µm thick cryosections, synthesis were performed as described (Mauti et al., 2006; Pekarik et al., 2011, 2012). Embryos were staged according to Hamburger and Hamilton (1951). To assess binding of soluble Plexinecto-AP fusion proteins, HEK293T cells were transfected with pcDNA3.1-Sema6A-myc/his, pcAGG-Sema6B-ha or pcAGG-Sema6D-ha for expression of full-length class-6 semaphorins. Live cells were incubated for 1 hour with Plexinecto-AP fusion proteins at 4°C before fixation and staining (Flanagan et al., 2000).

For co-IPs, the soluble fusion proteins described above were loaded on Handee spin columns (Thermoscientific) and incubated for 2 hours at 4°C with 10 µl anti-c-Myc agarose, according to the manufacturer’s instructions (ProFound IP-Kit; Thermoscientific, 23620). Immunoprecipitates were analyzed on western blots using the antibodies listed in supplementary material Table S4.

In vitro axon growth assay

Chick embryos at HH17-18 were electroporated with constructs encoding EBFP2 and miLuc or miS6B, as described above. At HH25, dissociated commissural neurons from the electroporated side were collected (pooled from three embryos in each condition) and grown on coverslips precoated with poly-lysine, which were incubated for 45 min at 37°C with Albumax (50 µg/ml), Laminin (10 µg/ml), AP only or PlexinA2 ecto-AP (both 50 ng/ml) for 24 hours before fixation and immunolabeling (Niederkofler et al., 2010) (supplementary material Table S4).

For quantification, at least 20 images per condition were taken from random positions on each coverslip. Each image typically contained both wild-type and miRNA-expressing neurons, which were distinguishable by the expression of EBFP2. At least 34 axons were measured per condition (ImageJ) using a Wacom DU-1931 tablet and pen tool.

Statistics

For open-books, multiple comparisons by one-way ANOVA followed by Tukey (homoedasticity) or Tamhane’s T2 (heteroscedasticity) post-hoc tests were used to calculate P-values, using Statistics-20 Software (SPSS). For the in vitro assays we performed two-sample t-tests for independent samples using the VassarStats website (http://vassarstats.net). P<0.05 was regarded as significant.

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Competing interests

The authors declare no competing financial interests.

Author contributions

I.A., N.H.W. and T.B. performed the experiments; I.A., N.H.W., O.M., M.G. and E.T.S. developed the concepts and approach; S.S. provided unpublished studies and into supplementation to I.A. Work in the lab of S.S. was supported by the National Center Zurich (ZNZ) to I.A. Work in the lab of E.T.S. and by fellowships of the Roche Research Foundation and the Neuroscience Center Zurich (ZNZ) to I.A. Work in the lab of S.S. was supported by the National Institutes of Health (NS046336).

Supplementary material

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References


