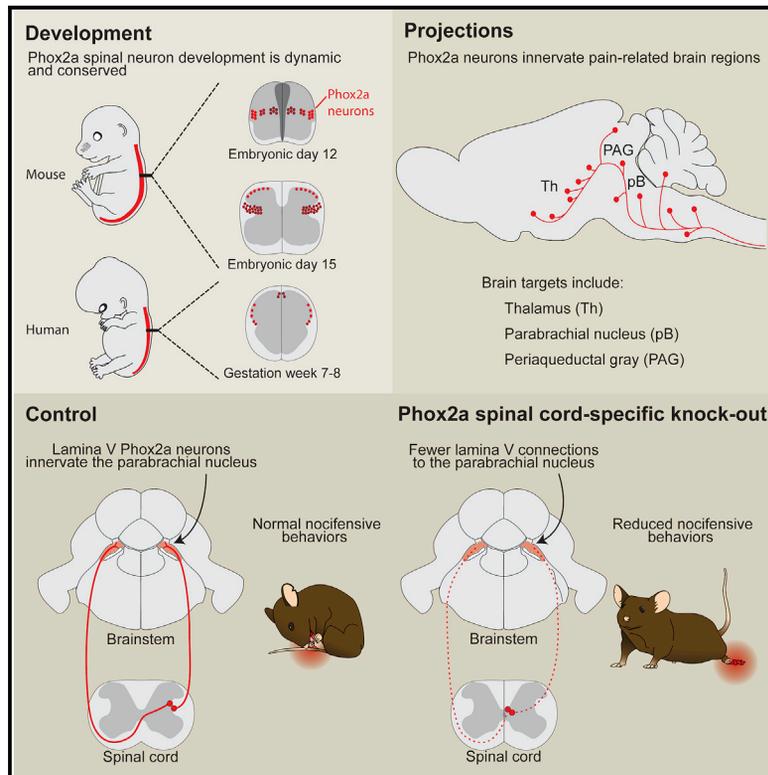


Phox2a Defines a Developmental Origin of the Anterolateral System in Mice and Humans

Graphical Abstract



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In Brief

Roome et al. generate a *Phox2a*^{Cre} mouse that labels anterolateral system neurons during development, revealing their developmental dynamics as well as their molecular conservation in humans. Developmental loss of *Phox2a* results in deficient spinoparabrachial connections and a loss of sensitivity to noxious stimuli.

Highlights

- *Phox2a* is transiently expressed in embryonic anterolateral system (AS) spinal neurons
- *Phox2a* AS neuron development reflects AS neuron diversity
- Spinal *Phox2a* knockout causes aberrant AS connectivity and nociceptive defects
- Human and mouse embryonic spinal *Phox2a* neurons are similar



Article

Phox2a Defines a Developmental Origin of the Anterolateral System in Mice and Humans

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SUMMARY

Anterolateral system neurons relay pain, itch, and temperature information from the spinal cord to pain-related brain regions, but the differentiation of these neurons and their specific contribution to pain perception remain poorly defined. Here, we show that most mouse spinal neurons that embryonically express the autonomic-system-associated Paired-like homeobox 2A (Phox2a) transcription factor innervate nociceptive brain targets, including the parabrachial nucleus and the thalamus. We define the Phox2a anterolateral system neuron birth order, migration, and differentiation and uncover an essential role for Phox2a in the development of relay of nociceptive signals from the spinal cord to the brain. Finally, we also demonstrate that the molecular identity of Phox2a neurons is conserved in the human fetal spinal cord, arguing that the developmental expression of Phox2a is a prominent feature of anterolateral system neurons.

INTRODUCTION

In vertebrates, somatosensory information about noxious stimuli is carried from peripheral nociceptors to the brain via spinal projection neurons collectively known as the anterolateral system (AS), which also carries temperature and itch information. The brain regions innervated by nociceptive AS neurons interpret the transmitted signals as pain—a sensation endowed with discriminative and affective components that, respectively, convey the identity, location, and intensity of the stimulus as well as elicit behavioral responses driven by arousal and aversion (Melzack and Casey, 1968). Since the molecular identity of AS neurons remains unknown, insights into the functional logic of nociceptive information relay from the periphery to the brain remain limited.

Prominent AS targets include the ventroposterolateral thalamus (VPL) (Gauriau and Bernard, 2004; Willis et al., 1979), which relays somatotopically organized nociceptive information (Guilbaud et al., 1980) to the primary somatosensory cortices, and the parabrachial nucleus (pB) (Bernard et al., 1995), which is considered to mediate affective components of pain by relaying noxious information to the amygdala (Han et al., 2015), and via the medial thalamus, to the prefrontal cortex (Bourgeois

et al., 2001). Clinical evidence supports the division between discriminative and affective dimensions of pain, as prefrontal lobotomy (Freeman and Watts, 1948) and insular cortex-related pain asymbolia (Berthier et al., 1988; Rubins and Friedman, 1948) result in the discriminatory nature of noxious stimuli being appreciated in the absence of the negative affect. The critical role of the AS in relaying both discriminative and affective components of nociception to its brain targets is suggested by the effects of lesions to the spinal anterolateral tract (Spiller and Martin, 1912).

The anatomy of AS neurons is well known in rodents, where they are found principally in laminae I and V and the lateral spinal nucleus (LSN) of the spinal dorsal horn (Davidson et al., 2010; Kitamura et al., 1993). Lamina I AS neurons have small receptive fields (Willis et al., 1974) and respond to specific classes of stimuli and their modalities (e.g., temperature, itch, mechanical versus thermal pain) (Andrew and Craig, 2001; Craig and Serrano, 1994), which are relayed to targets thought to mediate discriminatory responses such as the VPL thalamus. Lamina V/LSN AS neurons, in contrast, have broad receptive fields and wide dynamic ranges of receptivity (Craig, 2003b), and their physiology corresponds poorly with the qualitative descriptions of pain (Craig, 2004). Based on their prominent projections to



the dorsal pB (Feil and Herbert, 1995) and medial thalamus (Gauriau and Bernard, 2004), lamina V/LSN neurons likely transmit the affective and motivational dimensions of pain. These AS neuron functions are in line with substance-P receptor (NK1R)-directed AS neuron ablation, resulting in analgesia (Cameron et al., 2015; Mantyh et al., 1997). Recently developed genetic tools have uncovered the identity of interneurons that gate transmission of innocuous sensations to AS neurons (Duan et al., 2014; Petitjean et al., 2019), but genetic access to AS neurons has been elusive. Ablation of *Tachykinin1* (*Tac1*)-expressing spinal interneurons and pB-innervating AS neurons (Huang et al., 2019) produced behavioral deficits consistent with the loss of supraspinal transmission of nociceptive information without affecting the function of spinal nocifensive reflexes. Despite these advances, the genes expressed selectively in AS neurons remain unknown.

Developmental gene expression has been instrumental in studying locomotor circuits of the ventral spinal cord (Arber, 2012; Goulding, 2009) and may also be useful in accessing dorsal spinal cord somatosensory circuits. The dorsal spinal cord is divided into molecularly distinct neural precursor domains, whose link to adult neuronal classes remains obscure (Lai et al., 2016). Spinothalamic neurons express the transcription factor *Lmx1b* (Szabo et al., 2015), a marker of the putative projection neuron domain dl5, also expressed in other dorsal interneuron classes. In contrast, the Paired-like homeobox 2a (*Phox2a*) transcription factor is a more selective, albeit transient, marker of developing dl5 neurons (Ding et al., 2004) and, thus, a potential selective label of AS neurons.

Here, we report that transient embryonic expression of *Phox2a* in spinal neurons defines the identity of several AS projection neuron classes. We also reveal a developmental diversity of AS neurons and show that a loss of *Phox2a* impairs AS neuron innervation of their brain targets, resulting in attenuated supraspinal responses to noxious stimuli. Furthermore, we show that the molecular identity of *Phox2a* AS neurons is conserved in the developing human spinal cord.

RESULTS

Spinal *Phox2a*^{Cre} Neurons Reside in Lamina I, Lamina V, and LSN

Mouse *Phox2a* and its proxy, bacterial artificial chromosome (BAC) transgene *Phox2a*^{GFP}, are expressed embryonically and perinatally in the superficial and deep dorsal horn, where many AS neurons reside (Allen Institute for Brain Science, 2008; GENSAT, 2008). In order to label these neurons in adults, we created the transgenic *Phox2a*^{Cre} mouse line by inserting a Cre-poly(A) minigene into the BAC RP23-333J21 (GENSAT, 2008), at the *Phox2a* ATG codon (Figure 1A), and assessed Cre expression via the Cre-dependent tdTomato (tdT) reporter *R26*^{LSL-tdT} (Ai14). Adult *Phox2a*^{Cre}; *R26*^{LSL-tdT} mice showed tdT expression throughout the rostrocaudal length of the spinal cord in dorsal horn neurons, principally in lamina I (Figure 1B) and lamina V/LSN (Figures 1B and S1A), as well as in spinal accessory nerve (mXI) motor neurons (Figure S1A). Although rare, large “antenna”-like neurons were also found in laminae III/IV (Figure S3) (Marshall et al., 1996). *Phox2a* and tdT are expressed throughout

embryogenesis, but *Phox2a* expression is absent in adults (Figures 1C–E). While tdT expression is specific to cells expressing *Phox2a*, only 33% of lamina V/LSN *Phox2a* cells (*Phox2a*^{Deep}) express tdT at embryonic day (E)16.5, while 82% of lamina I cells (*Phox2a*^{Lami}) do. Similar proportions were observed at E18.5 (Figure S1B), suggesting that the *Phox2a* BAC may be missing some enhancer sequences necessary for *Phox2a* expression. Together, these data constitute evidence that *Phox2a*^{Cre} can be used to trace the fate of *Phox2a*-expressing spinal neurons.

Spinal *Phox2a*^{Cre} Neurons Innervate AS Targets

To reveal the connectivity of spinal *Phox2a*^{Cre} neurons, we restricted *Phox2a*^{Cre}-driven reporter expression to the spinal cord using the Cre-Flp recombinase-dependent reporter *R26*^{FSF-LSL-tdT} (Ai65) combined with the caudal neural tube-specific Flp recombinase mouse line *Cdx2*^{FlpO} (Britz et al., 2015) to generate *Phox2a*^{Cre}; *Cdx2*^{FlpO}; *R26*^{FSF-LSL-tdT} mice (Figure 2A). To validate this genetic intersection, we compared cellular tdT reporter expression between adult *Phox2a*^{Cre}; *R26*^{LSL-tdT} mice (Figures 2B–2F and S2E–S2H) and *Phox2a*^{Cre}; *Cdx2*^{FlpO}; *R26*^{FSF-LSL-tdT} mice (Figures 2B'–2F' and S2E'–S2H'). In the brain, *Phox2a*^{Cre} drove cellular tdT expression in motor and autonomic nuclei (Figures 2B–2E and S2A–S2H), which was not observed in *Phox2a*^{Cre}; *Cdx2*^{FlpO}; *R26*^{FSF-LSL-tdT} mice (Figures 2B'–2E' and S2E'–S2H'). In the caudal spinal cord of *Phox2a*^{Cre}; *Cdx2*^{FlpO}; *R26*^{FSF-LSL-tdT} mice, however, the cellular expression of tdT+ expression was preserved (Figures 2F and 2F'), allowing us to map axonal trajectories and brain targets of spinal *Phox2a*^{Cre} neurons.

In *Phox2a*^{Cre}; *Cdx2*^{FlpO}; *R26*^{FSF-LSL-tdT} mice, tdT+ axons were observed in the lateral funiculus in a distribution similar to previous reports of lamina I spinofugal axon locations (Apkarian et al., 1985; McMahon and Wall, 1983) (Figure 2E'). We observed tdT+ axons in known AS targets such as the globus pallidus (GP; Figure 2G); VPL and posterior (Po) thalamus (Figures 2H and S2I); mediodorsal thalamus (MD; Figure 2I); the posterior triangular thalamus (PoT) and anterior pretectal nucleus (Figure S2K); the deep layers of the superior colliculus, possibly within the orientation barrels (Masullo et al., 2019) (Figure 2J, indicated by white arrowheads); periaqueductal gray (PAG; Figure 2K); the pB (Figures 2M and 2N); the nucleus of the solitary tract (NTS; Figure 2P); the locus coeruleus (LC; Figure 2C'); and the caudal ventrolateral medulla (CVLM; Figure 2D'). These termini colocalized with the presynaptic marker vGluT2, suggesting that they were glutamatergic synapses (Figures 2Q and 2Q'). Within the pB, the dorsal-lateral (pBdl), central-lateral (pBcl), and internal-lateral (pBil) subnuclei and regions surrounding the external-lateral (pBel) contained many tdT+ axons, while the superior-lateral (pBs) and medial (pBm) subnuclei contained fewer axons (Figures 2M, 2N, S2M, and S2N). Consistent with previous reports, the pBel received very limited spinal innervation (Figures 2N, S2E, and S2N) (Bernard et al., 1995). Additionally, spinal *Phox2a*^{Cre} axons were also seen in brain regions not previously thought to receive direct AS innervation, such as the granular layers of the cerebellum (“gr” in Figure 2L), the vestibular nuclei (Figure 2O), the posterior hypothalamus near the A11 dopaminergic cell group (Figure S2J), and a region of the retrorubral area/dorsal-medial substantia nigra (Figure S2L). Thus, spinal *Phox2a*^{Cre} neurons

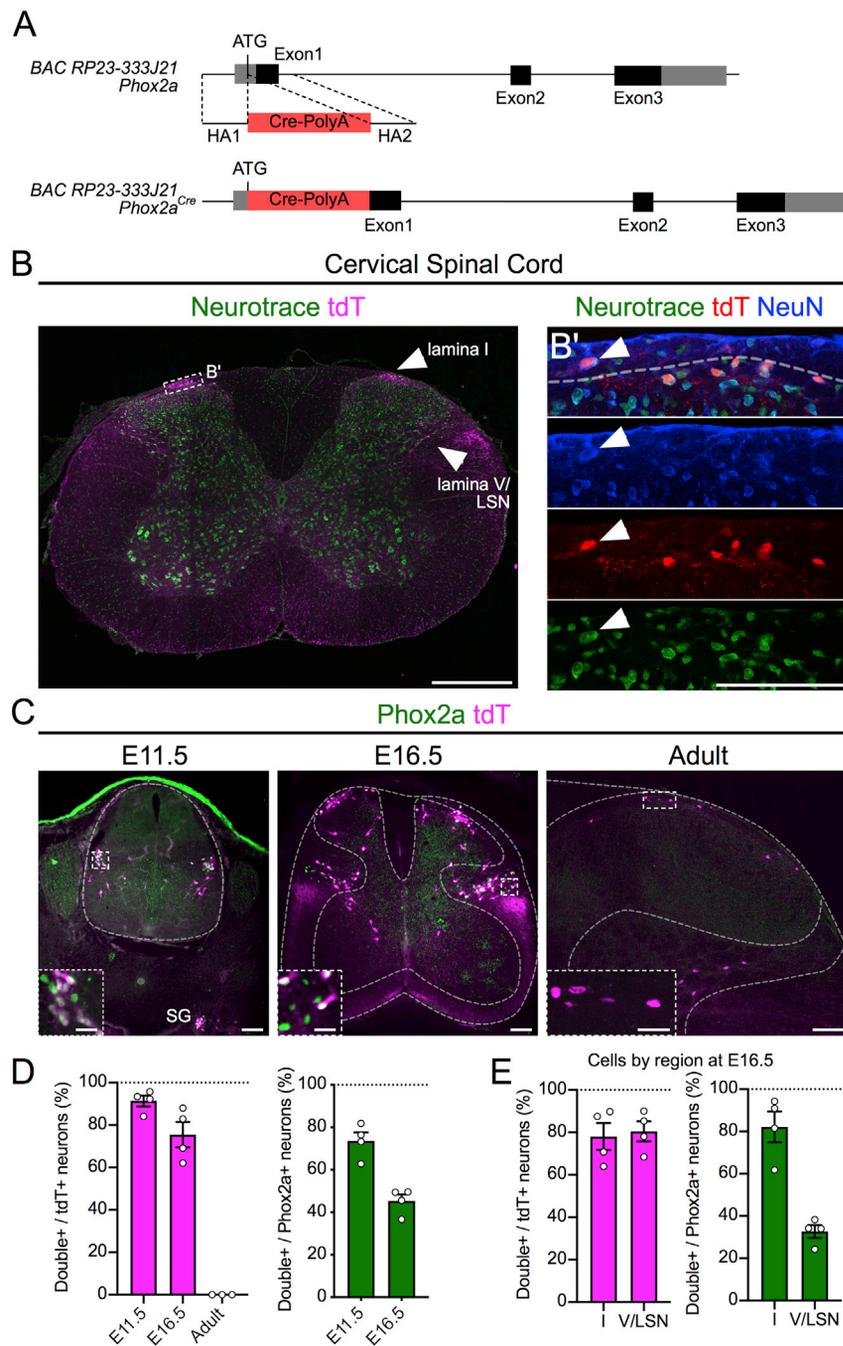


Figure 1. Spinal *Phox2a^{Cre}* Neurons Reside in Lamina I, Lamina V, and LSN

All images are of *Phox2a^{Cre}; R26^{LSL-tdT/+}* mice.

(A) BAC recombination strategy: *Cre-PolyA* insertion 3' to the *Phox2a* ATG codon in the BAC RP23-333J21.

(B) tdTomato (tdT)+ neurons in lamina I, lamina V, and LSN of the cervical spinal cord of adult mice.

(B') Magnified box in (B) showing lamina I Neurotrace, tdT, and NeuN co-labeling.

(C) Expression of tdT and *Phox2a* in E11.5, E16.5, and adult mouse spinal cord.

(D) Percentage of tdT+ neurons that express *Phox2a*, as well as percentage of *Phox2a*+ neurons that express tdT at E11.5 and E16.5 and in adult mice.

(E) Percentage of tdT+ neurons that express *Phox2a*, as well as percentage of *Phox2a*+ neurons that express tdT in the superficial and deep dorsal horn of E16.5 mouse spinal cords.

n = 4 E11.5 mice, n = 4 E16.5 mice, and n = 3 adult *Phox2a^{Cre}; R26^{LSL-tdT/+}* mice. Data are represented as mean ± SEM. Scale bars: 500 μm in (B), 100 μm in (B'), 100 μm in (C), and 25 μm in insets. SG, sympathetic ganglia.

tracers (Tracer+) that were also tdT+, sampled at all spinal cord levels (1,023 FG+, 6,620 CTb-488+, and 3,345 tdT+ cells from 7 mice). We focused on the cervical spinal cord, as spinothalamic neurons are relatively sparse in the mouse caudal spinal cord (Davidson et al., 2010). Overall, *Phox2a^{Cre}* labeled similar ratios of AS neurons traced from the VPL and the pB (26.9% ± 5.0% and 19.7% ± 4.3%, respectively; Figures 3C and 3D; n = 7). Consistent with the commissural nature of lamina I AS neurons, many *Phox2a^{Cre}* AS neurons were localized to the contralateral lamina I (Figures 3E, 3F, 3J, and 3K), in contrast to lamina V/LSN (Figures 3E, 3G, 3H, and 3I) where neurons were frequently seen labeled only with retrograde tracer. At least 20% of lamina V/LSN and approximately half of lamina I AS neurons, therefore, express *Phox2a^{Cre}*. Additionally, a bias in tdT expression to contralaterally projecting lamina I AS neurons suggests that *Phox2a^{Cre}* neurons may be involved in the localization of noxious stimuli (Figure 3K).

innervate brain regions predominantly involved in autonomic regulation and homeostasis such as the pBdl, NTS, and CVLM, as well as nociceptive areas VPL, PAG, and pBil.

Spinal *Phox2a* Neurons Are Predominantly AS Neurons

Next, we determined the fraction of *Phox2a* neurons that are AS neurons. Adult *Phox2a^{Cre}; R26^{LSL-tdT}* mice were injected unilaterally with Fluoro-Gold (FG) in the VPL thalamus (Figure 3A) and with CTb-488 in the pB (Figure 3B). After 7 days, we examined the proportion of spinal neurons labeled with either or both

that *Phox2a^{Cre}* neurons may be involved in the localization of noxious stimuli (Figure 3K).

Contrarily, most tdT+ neurons are AS neurons, regardless of type: in our most comprehensive injections of tracer into the pB and VPL, we reached a labeling ceiling of ~80% of lamina V/LSN tdT+ neurons bilaterally and as much as 100% of lamina I tdT+ neurons, suggesting that all spinal *Phox2a^{Cre}* neurons contribute to the AS (Figures 3L–3N). Smaller fractions of tdT+ neurons were labeled by tracer injection into the pB or VPL, suggesting that *Phox2a^{Cre}* neurons represent multiple AS neuron

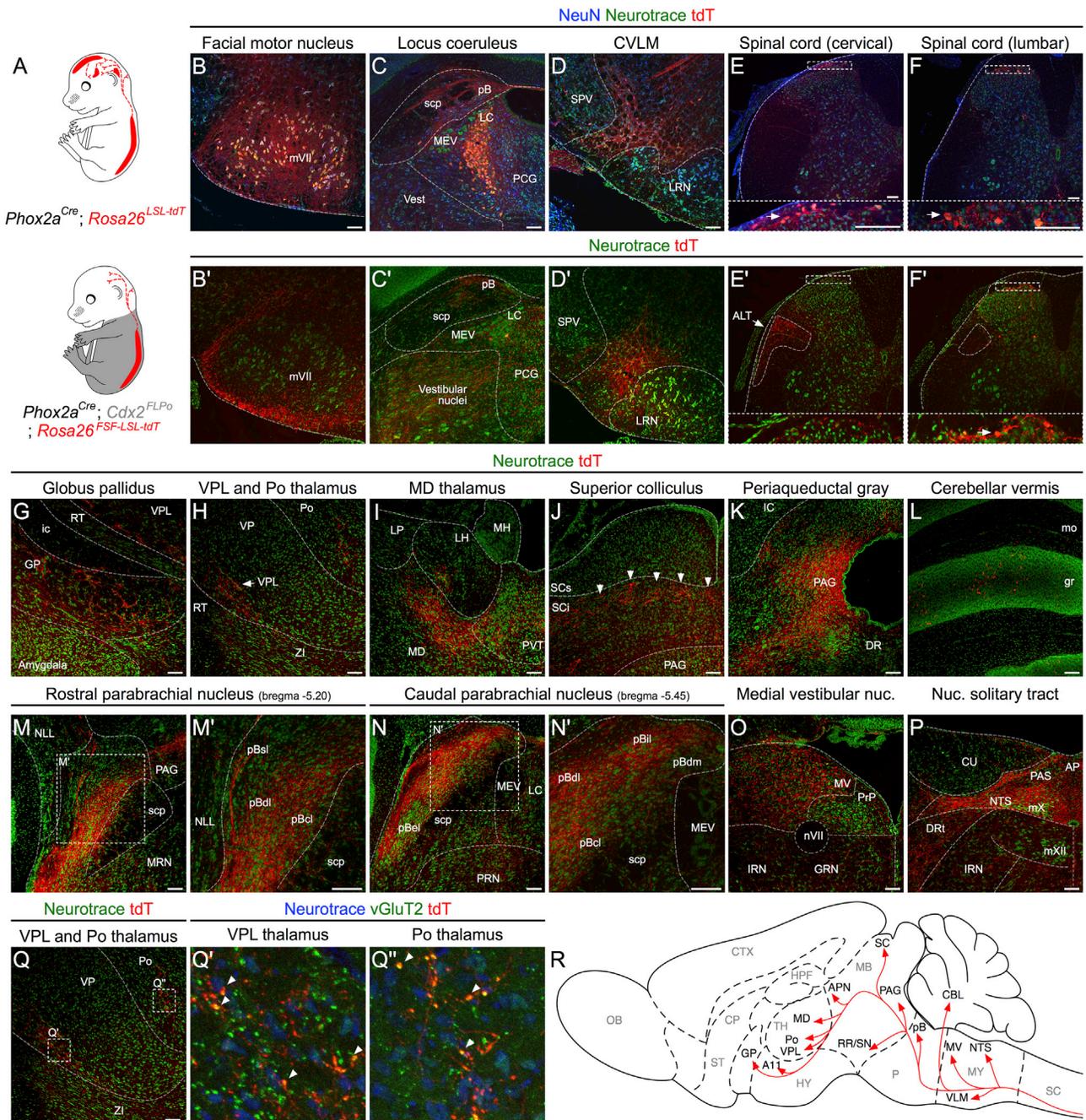


Figure 2. Spinal *Phox2a*^{Cre} Neurons Innervate AS Targets

(A) Intersectional genetic strategy to visualize spinofugal axons with tdT. *Phox2a*^{Cre}; *R26*^{LSL-tdT/+} mice have tdT cellular expression in the brain and spinal cord, while *Phox2a*^{Cre}; *Cdx2*^{FLPO}; *R26*^{FSF-LSL-tdT/+} mice have cellular tdT expression only in spinal *Phox2a* neurons.

(B–F) In (B)–(F): expression of cellular tdT in the brain and spinal cord of *Phox2a*^{Cre}; *R26*^{LSL-tdT/+} mice in comparison to (B'–F') the lack of tdT expression in the brain and spinal cord of *Phox2a*^{Cre}; *Cdx2*^{FLPO}; *R26*^{FSF-LSL-tdT/+} mice, except caudal to the cervical level. In (E'), arrow indicates presumptive anterolateral tract (ALT) axons in white matter. Insets in (E), (F), (E'), and (F') correspond to stippled boxes and show tdT+ cell bodies (white arrows).

(G–P) Prominent targets of tdT+ spinofugal axons. Higher magnifications are shown in (M') and (N').

(Q–Q'') vGluT2 and tdT immunohistochemistry in the thalamus demonstrates putative excitatory synaptic termini arising from spinofugal axons. The image in (Q) is duplicated from (H) and is used as a reference for (Q') and (Q'').

(R) Diagram summarizing the termination sites of tdT+ spinofugal axons.

n = 3 *Phox2a*^{Cre}; *R26*^{LSL-tdT/+} adult mice, and *n* = 3 *Phox2a*^{Cre}; *Cdx2*^{FLPO}; *R26*^{FSF-LSL-tdT/+} adult mice. Scale bars: 100 μm, except 25 μm for (Q') and (Q''). AP, area postrema; CU, cuneate nucleus; DR, dorsal raphe; DRt, dorsal reticular nucleus; GRN, gigantocellular reticular nucleus; ic, internal capsule; IRN, intermediate reticular nucleus; LH, lateral habenula; LP, lateral posterior thalamus; LRN, lateral reticular nucleus; MEV, midbrain trigeminal nucleus; MH, medial habenula; mo, medial olive; nVII, nucleus VII; NTS, nucleus tractus solitarius; PAS, parabrachial nucleus; PrP, parabrachial nucleus; P, pons; RR/SN, reticular raphe nucleus; SC, superior colliculus; VLM, ventral lateral midline; VPL, ventral posterior lateral nucleus; VP, ventral posterior nucleus; ZI, zona incerta.

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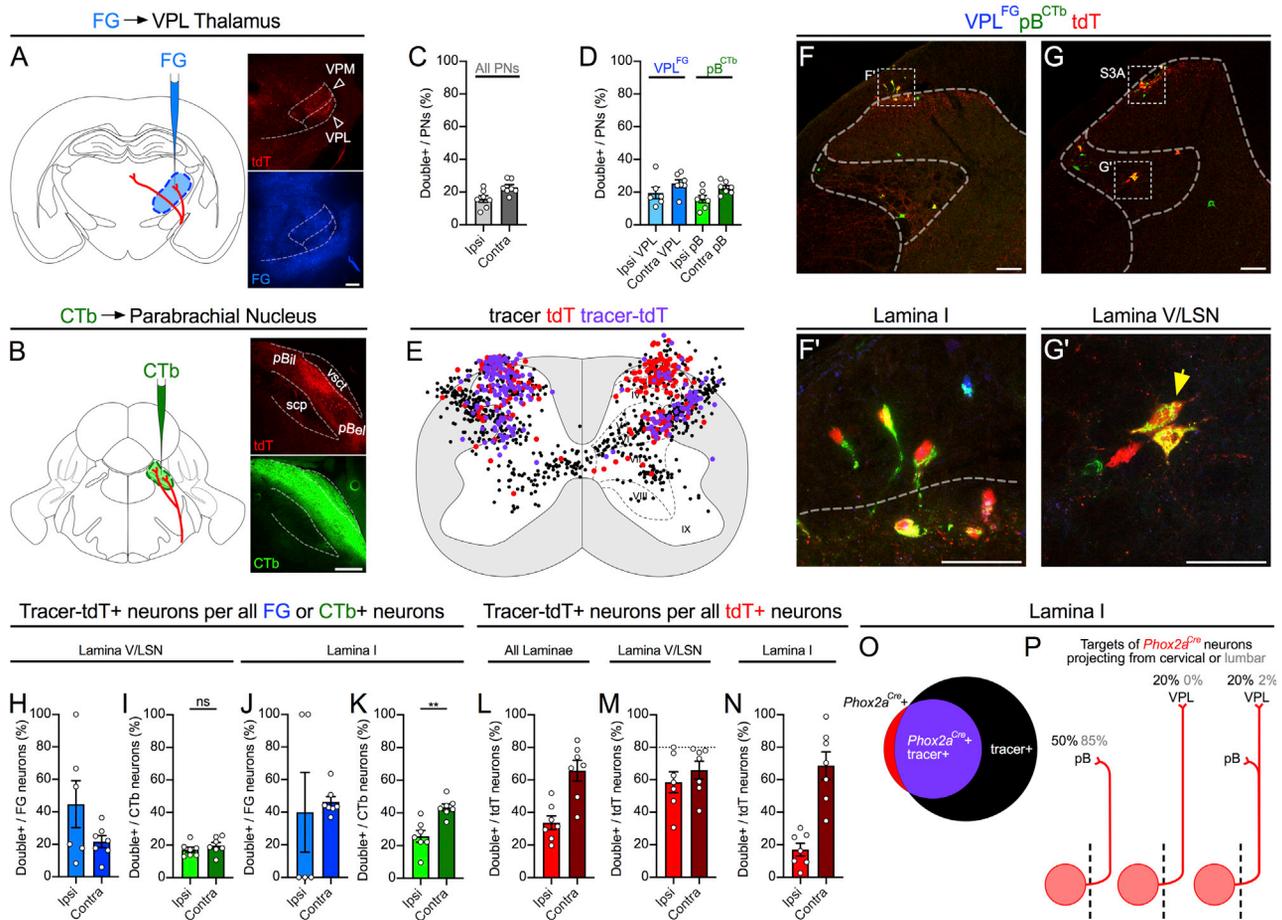


Figure 3. Spinal *Phox2a^{Cre}* Neurons Are Predominantly AS Neurons

(A and B) Adult *Phox2a^{Cre}; R26^{LSL-tdT/+}* mice were injected with FG in the VPL thalamus (A) and CTb-488 in the parabrachial nucleus (B).

(C and D) Percentage of cervical spinal cord dorsal horn projection neurons expressing tdT, classified as those labeled with either tracer (C; All PN) or selectively with FG or CTb (D).

(E) Diagram of the locations of tdT+ only, retrograde-labeled only (FG or CTb), or tdT+ and tracer-labeled (violet) neurons, in 5 non-sequential 25- μ m sections of the cervical spinal cord of one representative animal.

(F–G') In (F) and (G): representative images of the cervical spinal cord demonstrating tdT+ neuron labeling by retrograde tracers. See Figures S3A and S3B for more examples. (F' and G') High magnification of boxed areas in (F) and (G), depicting retrograde-labeled lamina I (F') or lamina V/LSN (G') tdT+ neurons (indicated by yellow arrows).

(H–N) Laminar analysis of neuron location in the cervical spinal cord ipsilateral or contralateral to tracer injection.

(H–K) Percentage of tracer-labeled neurons in lamina I (H and I) or lamina V/LSN (J and K) also expressing tdT, separated by tracer type.

(L–N) Percentage of tdT-labeled neurons labeled with one or both tracers in all laminae (L), in lamina V/LSN (M), or in lamina I (N).

(O) Diagram depicting overlap between tdT and retrograde tracer in lamina I of a representative mouse.

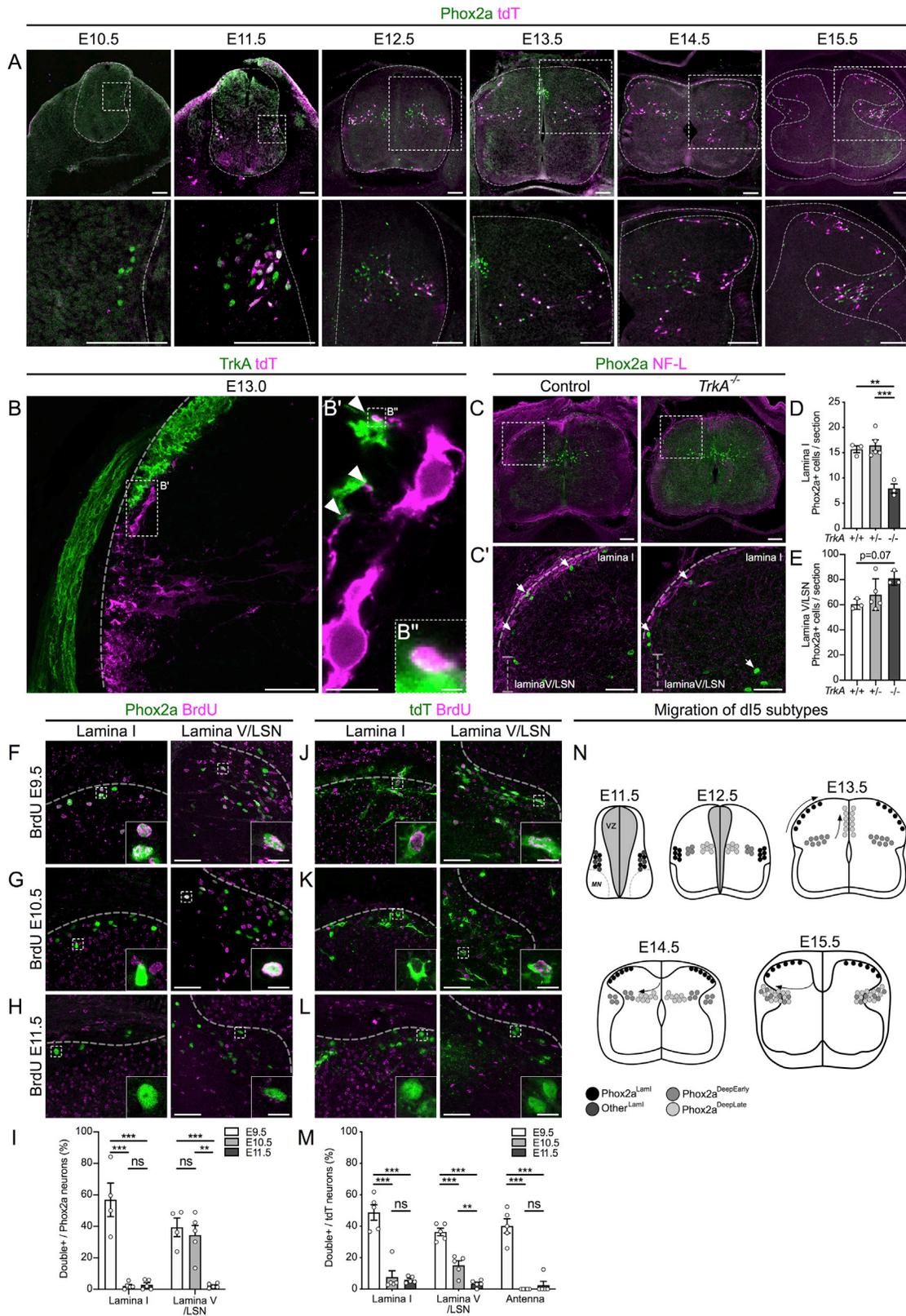
(P) Diagrams illustrating the estimated percentages of cervical and lumbar lamina I *Phox2a^{Cre}* neurons projecting to mouse pB, VPL, or both. Stippled line represents spinal midline.

$n = 7$ *Phox2a^{Cre}; R26^{LSL-tdT/+}* adult mice (4 male, 3 female). Mann-Whitney test in (I) and (K); ** $p < 0.01$; ns, non-significant. Data are represented as mean \pm SEM. Scale bars: 250 μ m in (A) and (B), 100 μ m in (F) and (G), and 50 μ m in (F') and (G').

types (Figure 3L versus Figures S3H–S3K). Although rare, antenna and lamina X tdT+ neurons were also predominantly AS neurons and tended to project contralaterally (Figures S3C–S3G). We also examined spinal projections to the MD thalamus (Figures S3L–S3R) as well as the cerebellar vermis (Figures

S3S–S3Y) via retrograde tracer injection, which labeled fewer neurons than pB/VPL injections but also included tdT+ neurons. Clarke's column and ventral horn spinocerebellar neurons did not express tdT, suggesting that these canonical spinocerebellar neuron types are not derived from *Phox2a*-expressing neurons.

molecular layer of the cerebellum; MV, medial vestibular nucleus; mVII, facial motor nucleus; mX, vagal motor nucleus; mXII, hypoglossal motor nucleus; NLL, nucleus of the lateral lemniscus; nVII, facial motor nerve; PAS, parasolitary nucleus; pBdm, dorsal-medial parabrachial nucleus; PCG, pontine central gray; PRN, pontine reticular nucleus; PRP, nucleus prepositus; PVT, paraventricular thalamus; RT, reticular thalamic nucleus; SCI, superior colliculus, intermediate laminae; scp, superior cerebellar peduncle; SCs, superior colliculus, superficial laminae; SPV, spinal trigeminal nucleus; ZI, zona incerta.



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In the hindbrain, pB, VPL, and MD tracer injections also labeled tdT+ neurons in the CVLM, parvocellular reticular nucleus (PARN), and spinal trigeminal lamina I/paratrigeminal region (Figures S3Z–S3CC), suggesting shared functions with spinal *Phox2a^{Cre}* neurons.

Heterogeneity of Spinal *Phox2a* Neuron Migration, Sensory Afferent Interaction, and Birth Time

We next studied the cellular and molecular events underlying *Phox2a* neuron development. First, we followed their migration via *Phox2a* and tdT expression in *Phox2a^{Cre}; R26^{LSL-tdT}* spinal cords throughout embryonic development. *Phox2a* neurons first appear at E10.5 in the cervical region and begin expressing tdT 1 day later (Figure 4A). At E12.5, three *Phox2a* populations are evident: *Phox2a*+/tdT+ (*Phox2a^{LamiI}*) neurons ventrolateral to the nascent dorsal horn, and two medial populations consisting of *Phox2a*+/tdT+ and those expressing only tdT. At E13.5, *Phox2a^{LamiI}* neurons disperse on the surface of the nascent superficial dorsal horn tangentially, while deeper *Phox2a* neurons (*Phox2a^{Deep}*) acquire positions that correlate with tdT expression: *Phox2a^{Deep}* tdT+ neurons remained ventrolateral to the dorsal horn, while *Phox2a^{Deep}* tdT– neurons accumulated above the central canal. At E14.5, *Phox2a^{Deep}* tdT– neurons migrate laterally and eventually become intermingled with *Phox2a^{Deep}* tdT+ neurons at E15.5, achieving their final configuration (Figure S4A). These results suggest the existence of at least three distinct migratory paths of *Phox2a* neurons.

As the tangential dispersal of *Phox2a^{LamiI}* neurons within the dorsal horn occurs at the time of primary afferent innervation, we asked how these two events are related. Prior to their entry into lamina I, *Phox2a^{Cre}* neurons project processes toward the dorsal root entry zone (Figures S4C and S4D) and form appositions with TrkA+ primary afferents at E13.0 as *Phox2a^{LamiI}* neurons begin migrating dorsally (Figures 4B and S4B), resulting in E13.5 *Phox2a^{LamiI}* neurons becoming encased in TrkA+ afferents (Figures S4E and S4F). To determine whether TrkA+ axons contribute to *Phox2a^{LamiI}* neuron positioning, we examined the location of *Phox2a* neurons in *TrkA* null (*TrkA^{-/-}*) mouse embryos, in which most TrkA+ afferents are absent (Smeyne et al., 1994). Compared to controls, the number of *Phox2a^{LamiI}* neurons in *TrkA^{-/-}* embryos was approximately halved (Figures

4C–4E), suggesting that TrkA+ afferents may interact with migrating *Phox2a* neurons. Although we found no significant effects of TrkA axonal loss on *Phox2a^{Deep}* neurons, their count tended to increase, suggesting that *Phox2a^{LamiI}* neurons fail to migrate and stall at the base of the forming dorsal horn (Figures 4C–4E).

To determine whether spinal *Phox2a* neuron diversity and migration patterns correlate with the time of their birth, we injected pregnant *Phox2a^{Cre}; R26^{LSL-tdT}* mice with bromodeoxyuridine (BrdU) at E9.5, E10.5, and E11.5 and examined strong BrdU co-staining with *Phox2a* or tdT in E16.5 *Phox2a^{Cre}; R26^{LSL-tdT}* embryos (Figures 4F–4M and S4R–S4T). Nearly all *Phox2a^{LamiI}* neurons were born at E9.5, while *Phox2a^{Deep}* neurons were born between E9.5 and E10.5, with very few born later. Furthermore, E11.5 *Phox2a* neurons that give rise to *Phox2a^{LamiI}* neurons (Figure 4A) are also born at E9.5 (Figures S4G–S4M). Given the differential expression of *Phox2a* and tdT in *Phox2a^{Deep}* cells, we developed a model in which *Phox2a^{LamiI}* and antenna neurons are born first, followed by *Phox2a^{Deep}* tdT+ (*Phox2a^{DeepEarly}*), while *Phox2a^{Deep}* tdT– neurons are born last (*Phox2a^{DeepLate}*) but not beyond E11.5 (Figures 4N and S4N–S4Q). More generally, our data show that AS neurons constitute one of the earliest born spinal neuron populations.

The Molecular Identity and Specification of Spinal *Phox2a* Neurons

To uncover the molecular pathways controlling *Phox2a* AS neuron specification, we studied their expression of neuronal identity determinant genes, identified transcription factor programs that specify them, and sought molecular markers that subdivide them. Spinal *Phox2a* expression begins at E9.5 and is restricted to spinal accessory motor neurons (Figures S5A and S5B). Non-motor neuron *Phox2a* expression is first visible at E10.5 in *Lmx1b*+ (dl5) post-mitotic neurons, which do not express the progenitor markers *Ascl1* or *Pax7* or the *dl1*, *dl3*, or *dl4/6* transcription factors *Lhx2*, *Isl1*, or *Pax2*, respectively, but which do express *dl5* transcription factors *Lbx1*, *Tlx3*, and *Brn3b/Pou4F2* and the commissural neuron guidance receptors *Robo3* and *DCC* (Figures 5A, 5B, S5C, and S5D). These findings demonstrate that non-motor neuron spinal *Phox2a* cells are predominantly commissural *dl5* neurons.

Figure 4. Heterogeneity of Spinal *Phox2a* Neuron Migration, Sensory Afferent Interaction, and Birth Time

(A) Migration of *Phox2a*+ and tdT+ neurons in embryonic spinal cords of *Phox2a^{Cre}; R26^{LSL-tdT/+}* mice between E10.5 and E15.5. Boxed regions in upper panels are magnified below.

(B–B'') Location of tdT+ neurons in the dorsal horn of *Phox2a^{Cre}; R26^{LSL-tdT/+}* spinal cords at E13.0, highlighting contacts (B' and B'') between lamina I neurons and TrkA+ sensory afferents (white arrowheads).

(C–E) Spinal *Phox2a* neuron (white arrows) location in E14.5 *TrkA^{+/+}*, *TrkA^{+/-}*, and *TrkA^{-/-}* mouse embryos. Boxed regions in (C) are magnified in (C').

(D and E) Counts of *Phox2a* neurons in lamina I (D) and lamina V/LSN (E).

(F–M) Birthdating of spinal *Phox2a* (F–I) or tdT (J–M) neurons in E16.5 *Phox2a^{Cre}; R26^{LSL-tdT/+}* mouse embryos, exposed to BrdU at E9.5 (F and J), E10.5 (G and K) or E11.5 (H and L).

(I and M) *Phox2a*+/BrdU+ (I) or tdT+/BrdU+ (M) neurons as a percentage of all *Phox2a*+ or tdT+ neurons in either lamina I, lamina V/LSN, or laminae II/III ("Antenna"-like neurons) and compared between groups.

(N) Diagram of migration of spinal *Phox2a* neuron subpopulations.

Phox2a^{Cre}; R26^{LSL-tdT/+} embryos: (A) n = 3 E10.5, n = 3 E11.5, n = 3 E12.5, n = 3 E13.5, n = 3 E14.5, and n = 3 E15.5; (B–B'') n = 3 E13.0 embryos; (F–M) n = 4–5 E16.5 embryos per condition. (C–E) n = 3 *TrkA^{+/+}*, n = 5 *TrkA^{+/-}*, and n = 3 *TrkA^{-/-}* E14.5 embryos.

In (D) and (E), one-way ANOVA, with Tukey's multiple comparisons test; in (I) and (M), individual one-way ANOVAs for each neuron type with Tukey's multiple comparisons test. **p < 0.01; ***p < 0.001. Data are represented as mean ± SEM. Scale bars: 100 μm in (A) and (C); 50 μm in (B), (C'), (F)–(H), and (J)–(L); 10 μm in (B') and insets in (F)–(H) and (J)–(L); and 1 μm in (B''). NF-L: Neurofilament-L.

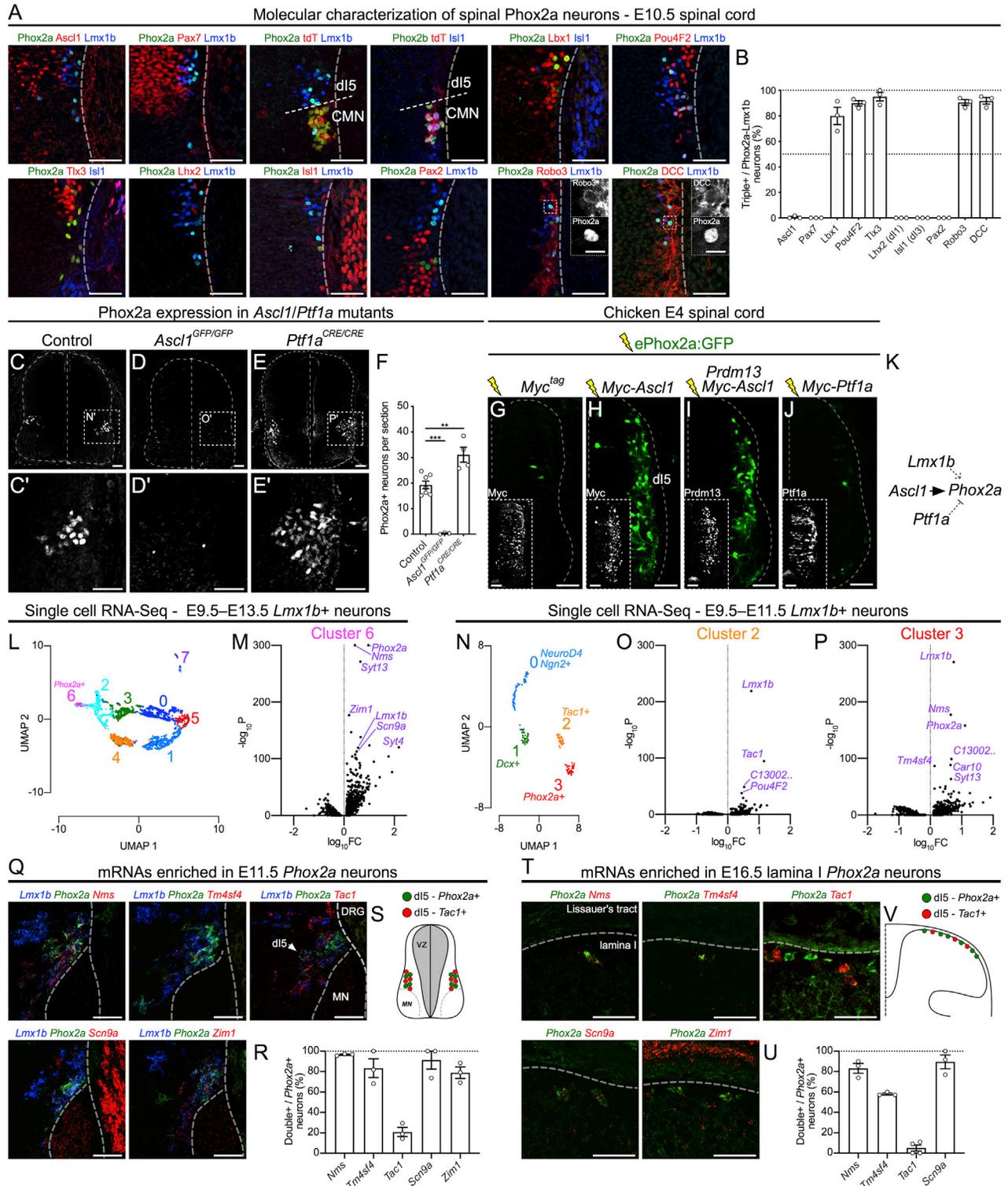


Figure 5. The Molecular Identity and Specification of Spinal Phox2a Neurons

(A) Co-expression of Phox2a with embryonic spinal neuron markers in the E10.5 *Phox2a*^{Cre}; *R26*^{LSL-tdT/+} spinal cord using immunohistochemistry.

(B) Quantification of each marker as a percentage of Phox2a+/Lmx1b+ neurons.

(C–F) Phox2a expression in control (C), *Ascl1* null (D), and *Ptf1a* null (E) E11.5 spinal cords, with the average number of Phox2a+ cells per section quantified in (F). (C'), (D'), and (E') show high magnification of boxed regions in (C), (D), and (E), respectively.

(legend continued on next page)

Since spinal Phox2a neurons develop from dl5 embryonic neurons, and since dl5 neuron identity is specified by the transcription factor *Ascl1* while *Ptf1a* suppresses dl5 identity and induces the neighboring dl4 identity (Glasgow et al., 2005; Helms et al., 2005), we assessed whether Phox2a expression was altered in *Ascl1* null (*Ascl1*^{GFP/GFP}) and *Ptf1a* null (*Ptf1a*^{CRE/CRE}) spinal cords. Compared to littermate controls, virtually no Phox2a neurons were found in E11.5 *Ascl1*^{GFP/GFP} spinal cords, while additional Phox2a neurons were found in E11.5 and E14.5 *Ptf1a*^{CRE/CRE} embryos (Figures 5C–5F, S5E, and S5F). To determine whether *Ascl1* and *Ptf1a* transcription factors control Phox2a expression directly or indirectly, we analyzed chromatin immunoprecipitation sequencing (ChIP-seq) data (Borromeo et al., 2014) for *Ascl1* and *Ptf1a* binding to the *Phox2a* locus. A genomic region (*ePhox2a*) located >30 kb downstream of the *Phox2a* transcription start site was bound by *Ascl1* and *Ptf1a*, but not the *Ptf1a* co-factor *Rbpj* or *Prdm13*, both of which act to repress dl5 and promote dl4 identity (Figure S5G) (Chang et al., 2013; Hori et al., 2008). To test the ability of *Ascl1*, *Ptf1a*, and *Prdm13* to regulate Phox2a through *ePhox2a*, we co-electroporated plasmids encoding these proteins together with a plasmid containing an *ePhox2a* activity reporter (*ePhox2a:GFP*; Figure S5H) into chick spinal neuron progenitors and monitored GFP expression. *ePhox2a:GFP* alone directed GFP expression in a small number of neurons located within the dl5 domain (Figures 5G and S5I). Ectopic *Ascl1* (but not ectopic *Ptf1a* or *Prdm13*) dramatically increased the number of GFP+ cells (Figures 5H–5J). Furthermore, Phox2a expression was entirely abolished in *Lmx1b*^{-/-} E11.5 mouse spinal cords (Figure S5J). Together, these data suggest that *Ascl1* and *Lmx1b* are required for Phox2a expression, with *Ascl1* acting directly through a 3' enhancer, while *Ptf1a* represses Phox2a transcription (Figure 5K).

Given that Phox2a labels a set of AS neurons, we sought to identify other genes expressed within AS neurons using available single-cell RNA-sequencing (RNA-seq) data from E9.5–E13.5 mouse spinal cords (Delille et al., 2019). Since Phox2a neurons are a subset of *Lmx1b*-expressing dl5 neurons, we performed uniform manifold approximation and projection (UMAP) dimensionality reduction analyses on two cohorts of dl5 neurons: (1) those found at all time points (E9.5–E13.5; 2,614 neurons; Figure 5L) and (2) an earlier subset of *Lmx1b* neurons (E9.5–E11.5; 186 neurons; Figure 5N) in order to attempt to separate

the early dl5 neurons (pre-lamina I neurons) into subsets. From these, we were able to isolate clusters of dl5 neurons enriched for Phox2a+ neurons (Figures 5M, 5P, S5K, and S5L), as well as an early dl5 cluster enriched for *Tac1* (Figures 5O and S5L). Top enriched transcripts for each cluster are listed in Table S1. Select transcripts were then validated using immunohistochemistry and RNAscope *in situ* hybridization in E11.5 and E16.5 spinal cords. At E11.5, Phox2a neurons were enriched for the expression of *Nms*, *Tm4sf4*, *Scn9a*, and *Zim1* mRNAs (Figures 5Q and 5R), which remained expressed in E16.5 Phox2a^{Lami} neurons (Figures 5T and 5U), providing further support that the E11.5 Phox2a cells populate lamina I. Other dl5-enriched transcripts and proteins, *Syt4*, *Pdzm3*, *Shox2*, and *Pou6F2*, were also highly co-expressed with Phox2a but were less specific to Phox2a neurons (Figures S5M–S5P). *Tac1*, however, was expressed in a complementary set of dl5 and lamina I neurons (Figures 5Q, 5R, 5T, and 5U) in line with the separation of dl5 neurons into two types suggested by our UMAP analyses, potentially describing a molecular division of superficial dorsal horn AS neurons (Figures 5S and 5V). Together, these experiments reveal the cellular and molecular mechanisms of AS neuron specification and unravel an array of AS-enriched mRNAs.

Phox2a Is Required for AS Neuron Development

Given the requirement of Phox2a for normal LC development (Morin et al., 1997), we hypothesized that its loss may also impact the development of spinal Phox2a neurons. As *Phox2a* null mice do not survive beyond birth, we used the *Hoxb8*^{Cre} mouse line to ablate *Phox2a* selectively in the caudal nervous system and a Cre reporter (*R26*^{LSL-tdT}) to visualize AS axons (Figure S6A) (Bourojeni et al., 2020; Witschi et al., 2010), producing Phox2a^{CKO} (*Hoxb8*^{Cre}; *Phox2a*^{fl/fl}; *R26*^{LSL-tdT}) and control (*Phox2a*^{fl/fl} or *Hoxb8*^{Cre}; *Phox2a*^{+/+}; *R26*^{LSL-tdT}) adult mice. Examination of tdT axons in adult Phox2a^{CKO} and control mice revealed that, while most spinofugal targets were normally innervated in Phox2a^{CKO} mice (Figure S6C), a dramatic loss of tdT axons was observed in the pBil (Figures 6A and 6B; additional examples are given in Figure S6B). Using retrograde tracing from the pBil of adult mice, we demonstrated that control and Phox2a^{CKO} mice had comparable numbers of cervical lamina I and lamina V/LSN neurons projecting to the pB (Figures S6D–S6F). In the lumbar spinal cord (within the *HoxB8*^{Cre} expression domain), however, while the tracer-labeled lamina I neuron

(G–J) Representative images from transverse sections of E4 chick neural tubes co-electroporated with the *ePhox2a:GFP* reporter and expression plasmids: control (Myc-tag only) (G), *Myc*^{Ascl1} (H), *Myc*^{Ascl1} and *Prdm13* (I), or *Myc*^{Ptf1a} (J). Insets show control Myc-tag, *Prdm13*, or *Ptf1a* expression.

(K) Diagram of *Phox2a* expression regulation showing direct transcriptional upregulation by *Ascl1*, upregulation by *Lmx1b* (Figure S5), and repression by *Ptf1a*.

(L–P) Single-cell RNA-seq data analysis of dl5 (*Lmx1b*+) neurons (Delille et al., 2019).

(L and M) UMAP analysis (L) of *Lmx1b*+ neurons from E9.5–E13.5 compared between each other, with volcano plot of cluster-6-enriched mRNAs (M) compared to all other neurons.

(N–P) UMAP analysis (N) of *Lmx1b*+ neurons from E9.5–E11.5 compared between each other, with volcano plots of cluster-2-enriched mRNAs (O) and cluster-3-enriched mRNAs (P) compared to all other neurons.

(Q–V) *In situ* hybridization of select mRNAs enriched in dl5 (Q; E11.5) and lamina I neurons (T; E16.5) based on UMAP analyses. The percentage of Phox2a+ neurons co-expressing selected mRNAs are quantified at both embryonic time points in (R) and (U). *Tac1* is present in dl5 *Lmx1b*+ neurons and lamina I neurons that do not express Phox2a, as depicted in diagrams in (S) and (V).

Phox2a^{Cre}; *R26*^{LSL-tdT} embryos: (A and B) n = 3 E10.5; (Q and R) n = 3 E11.5; (T and U) n = 3–4 E16.5 embryos. (C–F) n = 6 control, n = 3 *Ascl1*^{GFP/GFP}, and n = 4 *Ptf1a*^{Cre/Cre} embryos. (G–J) n = 6 E4 chicken embryos for each condition. Student's t test in (F); **p < 0.01; ***p < 0.001. Data are represented as mean ± SEM. Data in (L)–(P) are derived from Delille et al. (2019); data processing and statistics are described in STAR Methods. Scale bars: 50 μm for all, except 10 μm for insets in (A).

(A). DRG, dorsal root ganglion; FC, fold change; MN, motor neuron.

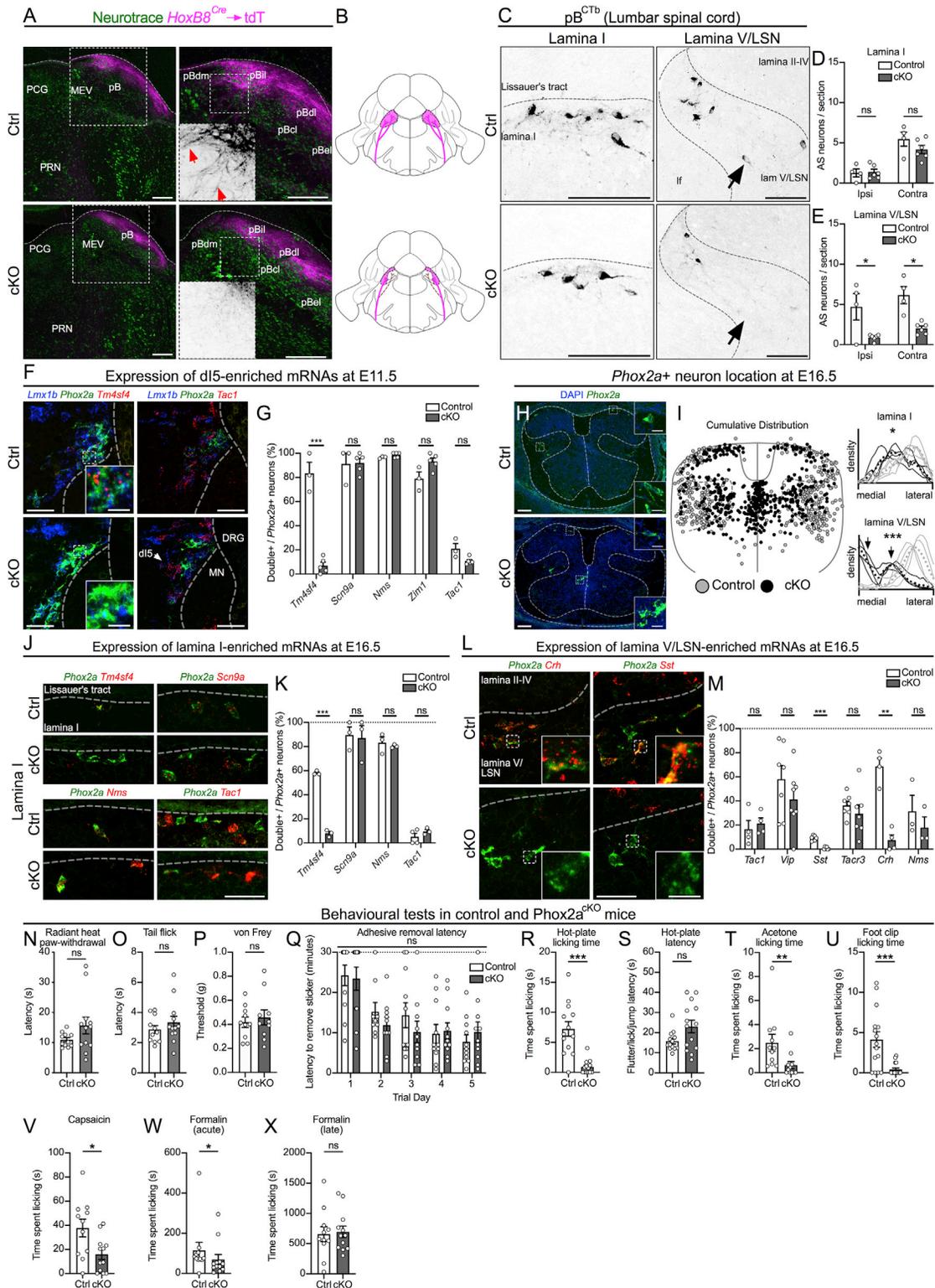


Figure 6. *Phox2a* Is Required for the Normal Development and Function of a Subset of AS Neurons

(A) The parabrachial nucleus (pB) of control (*Hoxb8^{Cre}; Phox2a^{+/+}; R26^{LSL-tdT/+}*; Ctrl, top row) and *Phox2a^{cKO}* (*Hoxb8^{Cre}; Phox2a^{fl/fl}; R26^{LSL-tdT/+}*; cKO, bottom row) adult mice, depicting spino-parabrachial axons labeled via *Hoxb8^{Cre}*-driven axonal tdT expression. Right panels show magnified views of the left panels, with insets depicting tdT axons (red arrows).

(B) Diagram depicting the loss of spinal afferents (magenta) to the pB of *Phox2a^{cKO}* mice.

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number was unchanged in *Phox2a*^{CKO} mice (Figures 6C and 6D), the number of Tracer+ ipsilateral and contralateral lamina V/LSN neurons was decreased by approximately 75% (Figures 6C and 6E). To investigate cellular changes leading to these connectivity phenotypes, we localized *Phox2a* mRNA in E16.5 control and *Phox2a*^{CKO} embryos, likely made possible by the persistence of the truncated *Phox2a* transcript. This analysis revealed similar numbers of *Phox2a* neurons in control and *Phox2a*^{CKO} E16.5 mice (12.6 ± 3.7 cells per section, n = 4; and 16.6 ± 1.5 cells per section, n = 4, respectively; p = 0.089, unpaired t test), although *Phox2a*^{Deep} neurons were displaced medially (Figures 6H and 6I), arguing that these cells are present in *Phox2a*^{CKO} mice but may be dysfunctional. *Phox2a* mRNA expression in *Phox2a*^{CKO} mice appeared elevated compared to that in controls, suggesting that *Phox2a* may negatively regulate its own expression.

To understand the molecular underpinnings of these phenotypes, we compared the expression of *Phox2a* AS neuron-enriched mRNAs (Figure 5) in control and *Phox2a*^{CKO} mice. Only the expression of *Tm4sf4*, a gene encoding a protein implicated in cellular differentiation, was affected by *Phox2a*^{CKO} mutation in E11.5 dl5 neurons and E16.5 lamina I neurons (Figures 6F, 6G, 6J, 6K, and S6G). Given the peptidergic heterogeneity of lamina V/LSN neurons (Leah et al., 1988), we also monitored the expression of neuromodulatory peptides and receptors in presumptive *Phox2a*^{Deep} neurons in E16.5 *Phox2a*^{CKO} and control spinal cords. Expression of genes encoding lamina V/LSN-enriched peptides *Sst* (somatostatin) and *Crh* (corticotrophin-releasing hormone) were reduced in *Phox2a*^{CKO} mice, while the expression of other *Phox2a* neuron-enriched transcripts remained unaffected (Figures 6L, 6M, S6K, and S6L). We also monitored the expression of selected neuromodulatory genes in lamina I neurons and found

elevated expression of *Vip* in *Phox2a*^{CKO} mice (Figure S6J); however, glutamatergic identity of *Phox2a* neurons was not changed (Figures S6H and S6I). Consistent with this, mRNAs encoding neuropeptides associated with inhibitory neurons were expressed sparsely among *Phox2a*^{Deep} neurons (Figure S6M). Together, these results demonstrate that *Phox2a* is essential for the normal axonal connectivity and migration of many AS neurons, as well as for their neuromodulatory identity.

Spinal *Phox2a* Loss Impairs Supraspinal Nocifensive Behaviors

Given the central role of the AS in supraspinal nociceptive signal relay, we reasoned that defects in spinoparabrachial connectivity and *Phox2a*^{Deep} neuromodulatory peptide expression in *Phox2a*^{CKO} mice might result in impaired nocifensive behaviors that are evoked by supraspinal circuits, with minimal effect on spinally mediated behaviors. Indeed, thermal and mechanical nociceptive reflex assays did not reveal any differences between control and *Phox2a*^{CKO} mice (Figures 6N–6P). Thermal preference to innocuous and noxious temperatures (Figures S6T–S6V) and behaviors evoked by light touch in the adhesive removal test (Figures 6Q and S6N) were also not affected by the *Phox2a*^{CKO} mutation. However, using a battery of behavioral assays requiring supraspinal transmission of noxious information, *Phox2a*^{CKO} mice showed deficits in hindpaw licking evoked by noxious stimuli—a nocifensive behavior requiring ascending spinal projections. Notable deficits emerged in the 53°C hot-plate assay, which monitors noxious heat sensitivity (Figures 6R, S6O, and S6Q); in the acetone assay, which monitors responses to cooling (Figures 6T and S6S); and following noxious mechanical stimulation (Figure 6U). Reflex-mediated, hot-plate-evoked behaviors remained unchanged (Figures 6S and S6P).

(C–E) Spinal neurons labeled by CTb injections in the pB, in lamina I (left column of C, quantified in D) and lamina V/LSN (right column of C, quantified in E), in control (top row) and *Phox2a*^{CKO} (bottom row) adult mice.
(F and G) Co-expression of *Phox2a* and *Lmx1b* with dl5-enriched mRNAs in control (top row) and *Phox2a*^{CKO} (bottom row) E11.5 spinal cords (F), quantified as a percentage of *Phox2a*+ neurons (G). See Figure S6 for additional images.
(H and I) Distribution of *Phox2a*+ neurons in E16.5 control and *Phox2a*^{CKO} mouse embryos.
(H) Insets show individual *Phox2a*+ cells magnified.
(I) Individual *Phox2a* neuron locations (left) and density plots of *Phox2a*+ neurons (right) derived from 3–5 sections (10 μm) of 4 control (gray) and 4 *Phox2a*^{CKO} (black) E16.5 spinal cords. Coordinates and mediolateral distribution are normalized to the width and height of an idealized spinal cord. Individual lines on density plots represent single animals, dotted lines represent mean distribution of 4 animals, gray lines represent control embryos, and black lines represent *Phox2a*^{CKO} *Phox2a*+ cells. Black arrows point to the bimodal distribution of *Phox2a*^{CKO} *Phox2a*+ cells in deep laminae. Average mediolateral positions of lamina I and lamina V/LSN neurons are statistically compared.
(J and K) Co-expression of *Phox2a* with dl5-enriched mRNAs in lamina I of control (top row) and *Phox2a*^{CKO} (bottom row) E16.5 spinal cords (J), quantified as a percentage of *Phox2a*+ neurons (K).
(L and M) Co-expression of *Phox2a* and candidate lamina V/LSN neuron marker mRNAs in lamina V/LSN neurons of control (top row) and *Phox2a*^{CKO} (bottom row) E16.5 spinal cords (L), quantified as a percentage of *Phox2a*+ neurons (M). See Figure S6 for additional images.
(N–X) Behavioral tests in control and *Phox2a*^{CKO} mice. (N) Radiant heat paw-withdrawal assay; n = 11 control, and n = 12 *Phox2a*^{CKO}. (O) Hot-water tail flick assay; n = 11 control, and n = 12 *Phox2a*^{CKO}. (P) von Frey test; n = 10 control, and n = 10 *Phox2a*^{CKO}. (Q) Adhesive removal latency; n = 11 control, and n = 10 *Phox2a*^{CKO}. (R and S) Time spent licking (R) and latency to any response (S) during the hot-plate test; n = 13 control, and n = 14 *Phox2a*^{CKO}. (T) Acetone test; n = 11 control, and n = 11 *Phox2a*^{CKO}. (U) Foot clip test; n = 15 control, and n = 16 *Phox2a*^{CKO}. (V) Capsaicin test; n = 11 control, and n = 12 *Phox2a*^{CKO}. (W and X) Formalin test (acute phase in W; late phase in X); n = 11 control, and n = 12 *Phox2a*^{CKO}.
In (A), n = 4 control, and n = 4 *Phox2a*^{CKO} adult mice. (C–E) n = 4 control, and n = 6 *Phox2a*^{CKO} adult mice. (F and G) n = 3 control, and n = 5 *Phox2a*^{CKO} E11.5 mice. (H and I) n = 4 control, and n = 4 *Phox2a*^{CKO} E16.5 mice. (J and K) n = 3–4 control, and n = 3 *Phox2a*^{CKO} E16.5 mice. (L and M) n = 3–8 control, and n = 3–8 *Phox2a*^{CKO} E16.5 mice. (N–X) The ns are given above.
In (D) and (E), two-way ANOVA with Tukey's multiple comparisons test; in (I), unpaired t test; in (G), (K), and (M), multiple t tests using the Holm-Sidak method; in (Q), mixed-effects analysis with Sidak's multiple comparisons test; and Mann-Whitney test in (N)–(P) and (R)–(X). ns, non-significant; *p < 0.05; **p < 0.01; ***p < 0.001. Data are represented as mean ± SEM. Scale bars: 250 μm in (A); 50 μm in (C); (H) 100 μm in (H); 20 μm in insets for (H); 50 μm in (F), (J), and (L); and 10 μm in insets for (F), (J), and (L). Images in Figure 5Q have been re-used in Figure 6F, and images in Figure 5T have been re-used in Figure 6J. Data from Figures 5R and 5U representing the above images are re-used in Figures 6G and 6K, respectively. These data were collected and analyzed as a single experiment.

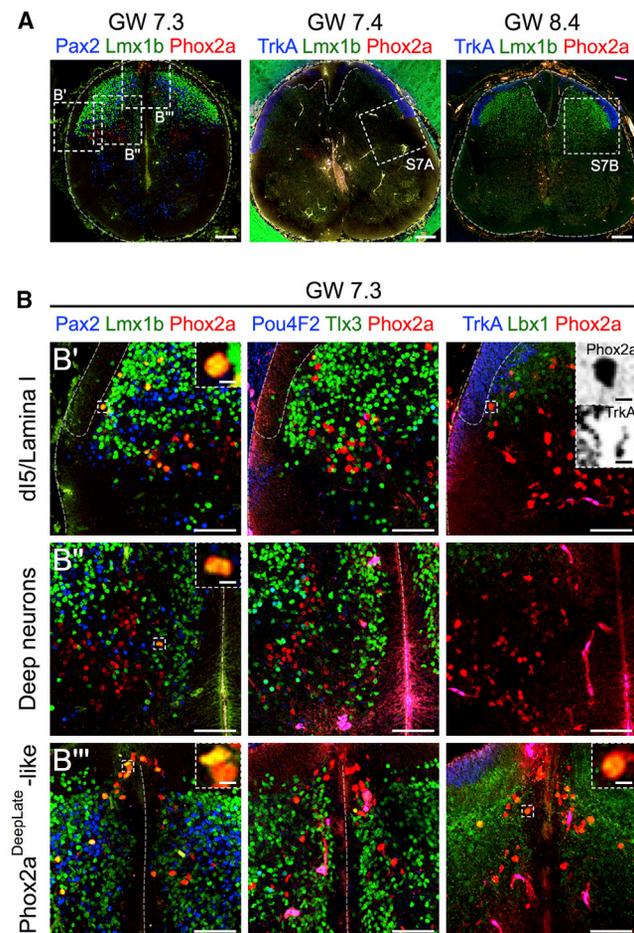


Figure 7. Phox2a Neuron Molecular Identity Is Conserved in the Developing Human Spinal Cord

(A) Sections of GW7.3–GW8.4 human spinal cords showing Phox2a, Lmx1b, TrkA, and Pax2 expression. Location of higher magnification panels is shown in boxed regions.

(B–B''') In (B): Phox2a, Lmx1b, Pax2, Pou4F2, Tlx3, Lbx1, and TrkA expression in the GW7.3 spinal cord, demonstrating co-labeling of Phox2a neurons with dorsal horn markers Lmx1b and Lbx1, but not with Pax2, Pou4F2, or Tlx3, in a Phox2a^{Lami}/Phox2a^{DeepEarly}-like cluster (B'), the deep dorsal horn (B''), and a Phox2a^{DeepLate}-like cluster near the roof plate (B'''). Top right panel shows apposition of Phox2a cells with TrkA⁺ sensory afferents, similar to that in mouse (Figure 4). Insets show higher magnification of boxed regions (TrkA and Phox2a channels split and inverted).

In (A), three human embryonic spinal cords (GW7.3, GW7.4, and GW8.4) are represented. In (B), one GW7.3 human embryonic spinal cord from (A) is represented. Scale bars: 200 μ m in (A), 100 μ m in (B), and 10 μ m in insets in (B')–(B''')

We attempted to measure nocifensive behaviors in the cold-plate assay, but neither control nor Phox2a^{ckO} mice responded (Figure S6R). In addition, Phox2a^{ckO} mice also exhibited less licking of hindpaws injected with the TRPV1 and TRPA1 agonists capsaicin and formalin, respectively, although the late/tonic phase of post-formalin injection licking was unaffected (Figures 6V–6X). Together, these results show that a loss of Phox2a during development disrupts AS neuron innervation of the pB, disrupts their molecular differentiation and concomitantly affects

the supraspinal aspects of a variety of nocifensive behaviors associated with AS function.

Phox2a Neuron Molecular Identity Is Conserved in the Developing Human Spinal Cord

Given that little is known about the molecular identity of human spinal neurons, we also asked whether Phox2a expression in the developing human spinal cord might allow insights into human AS development. We thus examined the expression of Phox2a protein in human spinal cords at developmental ages comparable to mouse mid-gestation: two at gestational week (GW)7.3 and one each at GW7.4, GW8.0, and GW8.4, three of which are shown in Figures 7A and S7 (Altman and Bayer, 2001). At GW7.3, Phox2a neurons (identified using a commercial Phox2a antibody; Figure S7C) were found in the superficial dorsal horn adjacent to TrkA⁺ fibers (Figures 7B', S7A, and S7B), in deeper laminae (Figure 7B''), and near the roof plate (Figure 7B'''), resembling the location of mouse Phox2a^{Lami} and Phox2a^{Deep} neurons. Human spinal Phox2a neurons co-expressed Lmx1b and Lbx1, but not Pax2 or Tlx3 (Figures 7B, S7A, and S7B). A previous study that examined Tlx3 expression in the human spinal cord did show its expression in the putative dl5 domain at GW5, although it was not compared to Phox2a expression (Marklund et al., 2014). As in mice, human Phox2a expression appeared weaker in older spinal cords (GW8.4; Figures 7A and S7B). Together, these data suggest that the spinal Phox2a neuron developmental program is evolutionarily conserved and that Phox2a expression is a molecular feature of developing human AS neurons.

DISCUSSION

Spinal neurons that express Phox2a^{Cre} during their development constitute a major tributary of the AS. Our studies reveal their developmental heterogeneity and requirement for normal nociception, as well as provide insights into a molecular logic that underlies their functions.

Diversity of AS Neuron Development

Nearly all spinal Phox2a^{Cre} neurons can be retrogradely labeled from the VPL thalamus and the pB, indicating that Phox2a is a genetic marker of a subpopulation of AS neurons. This has allowed us to investigate the development of AS neurons, revealing three neuronal classes that arise from embryonic dl5 neurons: Phox2a^{Lami}, Phox2a^{DeepEarly}, and Phox2a^{DeepLate}. A recent study argues that some AS neurons express *Tac1* (Huang et al., 2019); our molecular profiling suggests that the early population of dl5 Lmx1b⁺ Tac1⁺ neurons likely gives rise to Tac1⁺ AS neurons, that this population does not overlap with early-born Phox2a-expressing dl5 neurons, and that there at least two molecularly distinct Lamina I AS neuron subpopulations: Phox2a⁺ and Tac1⁺. Contrary to the notion that spinal neurons are born in a ventral-to-dorsal order (Nornes and Carry, 1978), superficial dorsal horn Phox2a neurons are born concurrently with motor neurons, as suggested recently for spinofugal neurons (Nishida and Ito, 2017). *Ascl1* (expressed in dl5 progenitors) and *Ptf1a* (expressed in dl4 progenitors) were previously shown to promote and inhibit dl5 neuron fates, respectively (Glasgow et al., 2005; Helms et al., 2005). Our data demonstrate that this may occur

via action at a 3' *Phox2a* enhancer defined in this study. The stereotyped birth order of *Phox2a* AS neurons raises the possibility that it is orchestrated by transcription factors involved in the temporal competence of *Ascl1*-expressing progenitors, as in the cerebral cortex and retina (Kohwi and Doe, 2013).

Following birth and early specification, *Phox2a*^{Lami}, *Phox2a*^{DeepEarly}, and *Phox2a*^{DeepLate} AS neurons migrate along distinct tangential and radial trajectories. The contacts between afferent axons and *Phox2a*^{Lami} neurons may be important for the settling of lamina I neurons in a somatotopic order corresponding to their dermatome-specific sensory afferents (Willis et al., 1974). At the molecular level, the neuronal migration cue *Reelin* likely mediates the radial migration of *Phox2a*^{Deep} neurons, since its intracellular signaling effector *Dab1* is required for normal positioning of lamina V/LSN neurons (Yvone et al., 2017). Conversely, *Netrin1* in the nascent dorsal horn prevents the premature ingrowth of primary afferents (Watanabe et al., 2006), and the *netrin-1* receptor *DCC* is required for the normal entry of *Phox2a*^{Lami} neurons into the dorsal horn (Ding et al., 2005). Thus, *netrin* signaling likely contributes to the interplay between sensory afferents and *Phox2a*^{Lami} neurons, while *Phox2a*^{Deep} neuron migration might rely on *reelin* signaling.

Phox2a Is Required for the Terminal Differentiation of AS Neurons

Phox2a-expressing neurons are present in normal numbers in embryonic *Phox2a*^{CKO} spinal cords, suggesting that *Phox2a* is not required for their early specification or survival. However, *Phox2a* mutation results in the loss of *Tm4sf4* and gain of *Vip* expression in *Phox2a*^{Lami} neurons, indicating that *Phox2a* is required for their molecular differentiation and, thus, possibly their function, despite apparent normal target connectivity. In contrast, lamina V/LSN *Phox2a* neurons are more dramatically affected, as nearly 75% of lamina V/LSN AS neurons fail to innervate the pB in *Phox2a*^{CKO} mice. This indicates that *Phox2a* is expressed in the vast majority of lamina V/LSN neurons, in agreement with our observation that *Phox2a*^{Cre} under-reports *Phox2a* expression in many of these neurons. Aberrant *Phox2a*^{Deep} neuron position in *Phox2a*^{CKO} mice, similar to that observed for lamina V and LSN neurons in *Reelin* signaling-deficient mice (Wang et al., 2012; Yvone et al., 2017), suggests that *Phox2a* may be required for expression of *Reelin* signaling genes. Together with the observation that *Phox2a* mutation also results in the loss of neuropeptide expression in *Phox2a*^{Deep} neurons, these defects argue that *Phox2a* specifies the terminal differentiation of a subset of AS neurons, and its absence likely impairs their function.

Phox2a AS Neuron Function in Supraspinal Nociception

The transmission of noxious and thermal information is a major function of the AS, and adult spinal *Phox2a* neuron morphologies, laminar organization, and their brain targets are consistent with such a function. The normal hindpaw adhesive-tape-evoked behaviors in *Phox2a*^{CKO} mice are consistent with the notion that light touch sensation is not a function of the AS (Hyndman and Wolkin, 1943). *Phox2a*^{CKO} mice also have normal spinal nociceptive reflexes, indicating that neither local reflex circuitry nor the descending pathways that modulate these behaviors (Ren and Dubner, 2009) depend on normal AS func-

tion. The apparently normal temperature preference of *Phox2a*^{CKO} mice may result from normal thermosensation in the forepaws and head, which is processed in the upper cervical spinal cord rostral to *Hoxb8*^{Cre} expression (Bourojeni et al., 2020; Witschi et al., 2010).

In comparison, *Phox2a*^{CKO} mice exhibit a reduction in the frequency and duration of behaviors evoked by the transmission of noxious information from the spinal cord to the brain, a function likely carried out by *Phox2a* AS neurons. While lamina I AS neurons have been proposed to transmit sensory-discriminative information and are likely to be impaired in *Phox2a*^{CKO} mice, a lack of motivation to respond to noxious stimuli—likely due to defects in *Phox2a* lamina V/LSN neurons—prevents us from testing this hypothesis. Our data, together with earlier studies, implicate lamina V/LSN neurons in directing motivated nociceptive behaviors via the spino-pBil-medial thalamus pathway (Bourgeois et al., 2001; Deng et al., 2020). Our observations are also in line with those made in mice with a loss of *Tac1* spinal neurons (Huang et al., 2019), in that approximately 20% of *Phox2a*^{Deep} neurons express *Tac1*, and so their loss may affect the transmission of motivational nociceptive information from the spinal cord to the brain. However, a loss of *Tac1* interneurons that control the function of *Phox2a*^{Deep} neurons may also contribute to the observed phenotypes. At the molecular level, *Phox2a*^{CKO} mice show decreased expression of the neuropeptides *Sst* (Leah et al., 1988) and *Crh*, normally enriched in *Phox2a*^{Deep} neurons of lamina V/LSN. Given the role of CRH in stress responses, *Crh*-expressing *Phox2a*^{Deep} neurons may convey motivational information linked to noxious stimuli.

The Molecular Logic of the AS

Although *Phox2a*^{Cre} labels a subpopulation of *Phox2a* neurons, we are compelled to draw some general inferences about the significance of *Phox2a* expression in AS neurons. Supraspinal *Phox2a* lineage-derived neurons exist in a variety of autonomic circuits, raising the question of whether these may be functionally intertwined with *Phox2a* AS neurons. Two lines of thought shed some light on this. First, *Phox2a* and its closely related transcription factor *Phox2b* specify the development of neurons afferent to medullary visceral reflex circuits that control many autonomic functions implicated in homeostasis (Brunet and Pattny, 2002). Our genetic tracing experiments reveal that *Phox2a* AS neurons participate in this connectivity logic by innervating brain stem autonomic regions such as the NTS and CVLM, as well as higher autonomic regulatory regions such as the pB. Second, because pain motivates behaviors that correct homeostatic changes, it has been proposed as a “homeostatic emotion” (Craig, 2003a). In light of this, the AS can be viewed as a pathway signaling deviations from homeostasis, such as changes in skin temperature, or the presence of noxious or pruritogenic stimuli, to brain regions that trigger compensatory autonomic responses (e.g., CVLM) or drive compensatory behavioral responses such as licking or scratching (e.g., pB). Given this, *Phox2a* AS neurons may specialize in transmitting somatic sensations with a motivational character such as cutaneous and deep pain, thermosensation, itch, visceral pain, nausea, and sexual arousal, all of which are abolished by anterolateral cordotomy in humans (Hyndman and Jarvis, 1940; Hyndman and Wolkin, 1943).

Our results suggest that the molecular identity of mouse Phox2a AS neurons is conserved in the developing human spinal cord, pointing to a conserved molecular logic of somatosensory circuit development, supported, in part, by the expression of *PHOX2A* in the human LC (Fan et al., 2018). A genetic proof of this idea remains out of reach because of the lack of obvious nociceptive or autonomic deficits in humans with *PHOX2A* mutations, which may be due to hypomorphic alleles (Nakano et al., 2001). Nevertheless, *PHOX2A* is a compelling molecular marker of human AS neurons and, given the effectiveness of cordotomy as a crude treatment of intractable chronic pain, a molecularly defined inactivation of a Phox2a AS neuron subpopulation could be its more refined iteration.

STAR★METHODS

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SUPPLEMENTAL INFORMATION

Supplemental Information can be found online at <https://doi.org/10.1016/j.celrep.2020.108425>.

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The authors declare no competing interests.

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STAR★METHODS

KEY RESOURCES TABLE

REAGENT or RESOURCE	SOURCE	IDENTIFIER
Antibodies		
Rabbit anti-Phox2a (1:10,000 from frozen stock; Figures 1, 2, 3, 4, 5, and 6)	Jean-François Brunet (École normale supérieure, Paris, France)	(Tiveron et al., 1996) RRID: AB_2315159
Rabbit anti-Phox2a (Figure 7 ; and 1:1000 for mouse, 1:500 for human embryo)	Abcam	Cat#: Ab155084 Lot#: GR117345-3 RRID: N/A
Rabbit anti-Phox2b (1:10,000 from frozen stock)	Jean-François Brunet (École normale supérieure, Paris, France)	(Pattyn et al., 1997) RRID: AB_2315160
Rabbit anti-RFP (red fluorescent protein) (1:1000)	Rockland	Cat#: 600-401-379 RRID: AB_2209751
Mouse anti-NeuN (1:1000)	Millipore	Cat#: MAB377 RRID: AB_2298772
Guinea Pig anti-vGlut2 (1:1000)	Synaptic Systems	Cat#: 135-404 RRID: AB_887884
Rat anti-Bromodeoxyuridine (BrdU) (1:10,000)	Abcam	Cat# ab6326: RRID: AB_305426
Goat anti-rTrkA (1:1000 for mouse, 1:500 for human embryo)	R&D Systems	Cat#: Af1056 RRID: AB_2283049
Mouse anti-Islet1 (1:100)	Developmental Studies Hybridoma Bank (DSHB)	Cat#: 39.3F7 RRID: AB_1157901
Mouse anti-Pax7 (1:100)	DSHB	Cat#: pax7 RRID: AB_528428
Mouse anti-Lhx2 (1:100)	DSHB	Cat#: Lhx2-1C11 RRID: AB_2618817
Mouse anti-Nkx6.1 (1:100)	DSHB	Cat#: F55A10 RRID: AB_532378
Mouse anti-Neurofilament-L (NF-L)	DSHB	Cat#: 2H3 RRID: AB_531793
Goat anti-hPax2 (1:1000 for mouse, 1:500 for human embryo)	R&D Systems	Cat#: AF3364 RRID: AB_10889828
Guinea Pig anti-Lbx1 (1:10000 for mouse, 1:5000 for human embryo)	Carmen Birchmeier (Max Delbruck Center, Berlin, Germany)	RRID: AB_2532144
Guinea Pig anti-Lmx1b (1:10000 for mouse, 1:5000 for human embryo)	Carmen Birchmeier (Max Delbruck Center, Berlin, Germany)	RRID: AB_2314752
Guinea Pig anti-Tlx3 (1:10000 in mouse, 1:10000 for chick, 1:5000 for human embryo)	Carmen Birchmeier (Max Delbruck Center, Berlin, Germany)	RRID: AB_2532145
Guinea pig anti-PRDM13 (1:1000)	Takahisa Furukawa (Osaka University, Osaka, Japan)	(Watanabe et al., 2015) RRID: N/A
Guinea pig anti-PTF1A (1:10000)	Jane Johnson (University of Texas Southwestern, Dallas, United States)	TX507 RRID: N/A
Mouse anti-MYC (1:1000)	Abcam	Cat# ab32 RRID: AB_303599
Goat anti-Brn3b (Pou4F2), (1:1000 for mouse, 1:500 for human embryo)	Santa Cruz Biotechnology	Cat#: sc-6026 RRID: AB_673441
Mouse anti-Ascl1 (1:100)	Santa Cruz Biotechnology	Cat#: sc-390794 RRID: N/A
Mouse anti-Ascl1 (1:100)	Santa Cruz Biotechnology	Cat#: sc-374550 RRID: AB_10985986
Mouse anti-Ascl1 (1:100)	Santa Cruz Biotechnology	Cat# sc-374104 RRID: AB_10918561
Goat anti-hRobo3	R&D Systems	Cat# AF3076 RRID: AB_2181865
Goat anti-mDcc	R&D Systems	Cat# AF844 RRID: AB_2089765
Rat anti-Pou6F2 (1:2000)	Jay Bikoff (Thomas Jessell Laboratory, HHMI Columbia University, New York, United States)	Cat# CU1796 RRID: AB_2665427
Rabbit anti-Shox2 (1:200)	Laskaro Zagoraiou (Thomas Jessell Laboratory, HHMI Columbia University, New York, United States)	(Dougherty et al., 2013) RRID: N/A
Sheep anti-FoxP2 (1:2000)	R&D Systems	Cat#: AF5647 RRID: AB_2107133

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REAGENT or RESOURCE	SOURCE	IDENTIFIER
Alexa 488 Donkey anti-Rabbit (1:500)	Jackson Immunoresearch Laboratories	Cat#: 711-545-152 Lot#: 141848 RRID: AB_2313584
Alexa 488 Donkey anti-Guinea Pig (1:500)	Jackson Immunoresearch Laboratories	Cat#: 706-545-148 Lot#: 138058 RRID: AB_2340472
Alexa 488 Donkey anti-Mouse (1:500)	Jackson Immunoresearch Laboratories	Cat#: 715-545-150 Lot#: 136831 RRID: AB_2340846
Alexa 488 Donkey anti-Goat (1:500)	Jackson Immunoresearch Laboratories	Cat#: 705-545-147 Lot#: 136089 RRID: AB_2336933
Alexa 488 Donkey anti-Rat (1:500)	Jackson Immunoresearch Laboratories	Cat#: 712-545-153 Lot#: 138117 RRID: AB_2340684
Alexa 488 Donkey anti-Sheep (1:500)	Jackson Immunoresearch Laboratories	Cat#: 713-545-003 Lot#: N/A RRID: AB_2340744
Cy3 Donkey anti-Rat (1:500)	Jackson Immunoresearch Laboratories	Cat#: 712-165-153 Lot#: 139289 RRID: AB_2340667
Cy3 Donkey anti-Rabbit (1:500)	Jackson Immunoresearch Laboratories	Cat#: 711-165-152 Lot#: 138270 RRID: AB_2307443
Cy3 Donkey anti-Mouse (1:500)	Jackson Immunoresearch Laboratories	Cat#: 715-165-150 Lot#: N/A RRID: AB_2340813
Cy3 Donkey anti-Goat (1:500)	Jackson Immunoresearch Laboratories	Cat#: 705-165-147 Lot#: 134527 RRID: AB_2307351
Cy5 Donkey anti-Rabbit (1:500)	Jackson Immunoresearch Laboratories	Cat#: 711-175-152 Lot#: 138336 RRID: AB_2340607
Cy5 Donkey anti-Mouse (1:500)	Jackson Immunoresearch Laboratories	Cat#: 715-175-150 Lot#: 135323 RRID: AB_2340819
Cy5 Donkey anti-Goat (1:500)	Jackson Immunoresearch Laboratories	Cat#: 705-175-147 Lot#: 134531 RRID: AB_2340415
Cy5 Donkey anti-Guinea Pig (1:500)	Jackson Immunoresearch Laboratories	Cat#: 706-175-148 Lot#: 136607 RRID: AB_2340462
Chemicals, Peptides, and Recombinant Proteins		
NeuroTrace 435/455 Blue Fluorescent Nissl Stain	ThermoFisher Scientific	Cat#: N21479 RRID: N/A
NeuroTrace 500/525 Green Fluorescent Nissl Stain	ThermoFisher Scientific	Cat#: N21480 RRID: N/A
2-Hydroxystilbene-4,4'-dicarboximidine (fluorogold)	ThermoFisher Scientific	Cat#: H22845 RRID: N/A
Alexa 488-conjugated Cholera toxin B	ThermoFisher Scientific	Cat#: C22841 Lot#: 2038245 RRID: N/A
5-bromo-2'-deoxyuridine (BrdU)	ThermoFisher Scientific	Cat#: B23151 Lot#: 1916418 RRID: N/A
Paraformaldehyde	Millipore Sigma	Cat#: P6148 RRID: N/A
(E)-Capsaicin	Tocris	Cat#: 0462 Lot#: 7A/218361 RRID: N/A
4',6-diamidino-2-phenylindole (DAPI)	Thermo Fisher Scientific	Cat# D1306
Mowiol (Polyvinyl alcohol)	Millipore Sigma	Cat#: 81381 RRID: N/A
Experimental Models: Organisms/Strains		
Mice - Phox2a ^{Cre}	This manuscript	RRID: N/A
Mice - HoxB8 ^{Cre} (Tg(Hoxb8-cre)1403Uze)	Hanns Ulrich Zeilhofer, ETH Zürich, Zürich, Switzerland)	Cat#: MGI: 4881836 RRID: N/A
Mice - Cdx2 ^{FipO} (Tg(CDX2-flpo)#Gld)	Martyn Goulding (Salk Institute, San Diego, United States)	Cat#: MGI: 5911680 RRID: N/A
Mice - Ai14 (B6;129S6-Gt(ROSA)26Sortm14(CAG-tdTomato)Hze/J)	The Jackson Laboratory	Cat#: JAX:007908 RRID: IMSR_JAX:007908
Mice - Ai65 (B6;129S-Gt(ROSA)26Sortm65.1(CAG-tdTomato)Hze/J)	The Jackson Laboratory	Cat#: JAX:021875 RRID: IMSR_JAX:021875
Mice - Phox2alox (B6D2.129S2-Phox2atm2Jbr/Orl)	European Mutant Mouse Archive (EMMA)	Cat#: EM:04758 RRID:IMSR_EM:04758
Mice - TrkA ^{-/-} (Ntrk1 ^{tm1Par})	Lino Tessarollo (National Cancer Institute, Frederick, MD, United States)	(Liebl et al., 2000) Cat#: MGI: 1933963 RRID: N/A

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REAGENT or RESOURCE	SOURCE	IDENTIFIER
Mice - <i>Ptf1atm1</i> (cre)Wri	Christopher Wright (Vanderbilt University, Nashville, United States)	(Kawaguchi et al., 2002) Cat#: MGI: 2387812 RRID: N/A
Mice - <i>Ascl1^{tm1Reed}/J</i>	The Jackson Laboratory	Cat# JAX:012881 RRID: IMSR_JAX:012881
Mice - C57BL/6J	The Jackson Laboratory	Cat# JAX:000664 RRID: IMSR_JAX:000664
Mice - 129S1/SvImJ	The Jackson Laboratory	Cat# JAX:002448 RRID: IMSR_JAX:002448
Mice - B6C3F1/J	The Jackson Laboratory	Cat# JAX: 100010 RRID: IMSR_JAX: 100010
Oligonucleotides		
RNA Scope Probe – Mm-Tac1 C1	Advanced Cell Diagnostics	Cat#: 410351 Lot#: 18354A RRID: N/A
RNA Scope Probe – Mm-Phox2a C2	Advanced Cell Diagnostics	Cat#: 520371-C2 Lot#: N/A RRID: N/A
RNA Scope Probe – Mm-Lmx1b C3	Advanced Cell Diagnostics	Cat#: 412931-C3 Lot#: N/A RRID: N/A
RNA Scope Probe – Mm-Nms C1	Advanced Cell Diagnostics	Cat#: 472331 Lot#: TBD RRID: N/A
RNA Scope Probe – Mm-Tm4sf4 C1	Advanced Cell Diagnostics	Cat#: 819831 Lot#: N/A RRID: N/A
RNA Scope Probe – Mm-Zim1 C1	Advanced Cell Diagnostics	Cat#: 819821 Lot#: N/A RRID: N/A
RNA Scope Probe – Mm-Scn9a C1	Advanced Cell Diagnostics	Cat#: 313341 Lot#: N/A RRID: N/A
RNA Scope Probe – Mm-Pdzn3 C1	Advanced Cell Diagnostics	Cat#: 517061 Lot#: 17269A RRID: N/A
RNA Scope Probe – Mm-Syt4 C1	Advanced Cell Diagnostics	Cat#: 574731 Lot#: N/A RRID: N/A
RNA Scope Probe – Mm-VIP C1 (Vasoactive Intestinal Polypeptide)	Advanced Cell Diagnostics	Cat#: 415961 Lot#: 19045A RRID: N/A
RNA Scope Probe – Mm-Sst C1	Advanced Cell Diagnostics	Cat#: 404631 Lot#: N/A RRID: N/A
RNA Scope Probe – Mm-TacR3 C1	Advanced Cell Diagnostics	Cat#: 481671 Lot#: 18254A RRID: N/A
RNA Scope Probe – Mm-Crh C1	Advanced Cell Diagnostics	Cat#: 316091 Lot#: N/A RRID: N/A
RNA Scope Probe – Mm-Slc17A6 C1 (vGlut2)	Advanced Cell Diagnostics	Cat#: 319171 Lot#: 19052B RRID: N/A
RNA Scope Probe – Mm-Slc32A1 C1 (vGAT)	Advanced Cell Diagnostics	Cat#: 319191 Lot#: 19057A RRID: N/A
RNA Scope Probe – Mm-TacR1 C1	Advanced Cell Diagnostics	Cat#: 428781 Lot#: 19057A RRID: N/A
RNA Scope Probe – Mm-Cck C1 (Cholecystokinin)	Advanced Cell Diagnostics	Cat#: 402271 Lot#: 19057A RRID: N/A
RNA Scope Probe – Mm-Lypd1 C1	Advanced Cell Diagnostics	Cat#: 318361 Lot#: 18353B RRID: N/A
RNA Scope Probe – Mm-Gal C1 (Galanin)	Advanced Cell Diagnostics	Cat#: 400961 Lot#: 18277C RRID: N/A
RNA Scope Probe – Mm-pDyn C1 (preproDynorphin)	Advanced Cell Diagnostics	Cat#: 318771 Lot#: 18303A RRID: N/A
RNA Scope Probe – Mm-pNoc C1 (prepronociceptin)	Advanced Cell Diagnostics	Cat#: 437881 Lot#: 19016B RRID: N/A
Genotyping primers		
Cre-1: 5'-AGG TGT AGA GAA GGC ACT TAG C –3' Expected band size: 412 bp (only one band)	This manuscript	N/A
Cre-2: 5'-CTA ATC GCC ATC TTC CAG CAG G-3'	This manuscript	N/A
FLPo-1: 5'-TGA GCT TCG ACA TCG TGA AC-3' Expected band size: 350 bp (only one band)	Martyn Goulding (Salk Institute, San Diego, United States)	N/A
FLPo-2: 5'-ACA GGG TCT TGG TCT TGG TG –3'	Martyn Goulding (Salk Institute, San Diego, United States)	N/A
Ai14-1: 5'-TCA ATG GGC GGG GGT CGT T-3' Expected band sizes: WT: 350 bp, Mutant: 250 bp	This manuscript	N/A
Ai14-2: 5'-CTC TGC TGC CTC CTG GCT TCT-3'	This manuscript	N/A
Ai14-3: 5'-CGA GGC GGA TCA CAA GCA ATA-3'	This manuscript	N/A
Ai65 WT1: "oIMR9020" 5'-AAG GGA GCT GCA GTG GAG TA-3' Expected band size: 315 bp	The Jackson Laboratory	N/A

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REAGENT or RESOURCE	SOURCE	IDENTIFIER
Ai65 WT2: "oIMR9021" 5'-CCG AAA ATC TGT GGG AAG TC-3'	The Jackson Laboratory	N/A
Ai65 Mutant1: "oIMR9103" 5'-GGC ATT AAA GCA GCG TAT CC-3' Expected band size: 297 bp	The Jackson Laboratory	N/A
Ai65 Mutant2: "oIMR9105" 5'-CTG TTC CTG TAC GGC ATG G-3'	The Jackson Laboratory	N/A
Phox2a ^{fl-1} : 5'-GCC TCC AAC TCC ATA TTC C-3' Expected band sizes: WT: 150bp, Flox: 200bp	Jean-François Brunet (École normale supérieure, Paris, France)	N/A
Phox2a ^{fl-2} : 5'-ATC AGG AGT CAG TCG TCT G -3'	Jean-François Brunet (École normale supérieure, Paris, France)	N/A
TrkA-WT-5': 5'-TGT ACG GCC ATA GAT AAG CAT-3' Expected WT band size: 160 bp	Lino Tessarollo (National Cancer Institute, Frederick, MD, United States)	N/A
TrkA-WT-3': 5'-TTG CAT AAC TGT GTA TTT CAC-3'	Lino Tessarollo (National Cancer Institute, Frederick, MD, United States)	N/A
TrkA-mutant (pGKneopolyA) forward primer: 5'-CGC CTT CTT GAC GAG TTC TTC TG-3' Expected mutant band size: 550 bp	Lino Tessarollo (National Cancer Institute, Frederick, MD, United States)	N/A
Recombinant DNA		
Plasmid: pMiVIII-Myc- ASCL1	Jane Johnson (University of Texas Southwestern, Dallas, United States)	N/A
Plasmid: pMiVIII-Myc- PTF1A	Jane Johnson (University of Texas Southwestern, Dallas, United States)	N/A
Plasmid: pMiVIII-Prdm13	Jane Johnson (University of Texas Southwestern, Dallas, United States)	N/A
Plasmid: pMiVIII-Myc-tag	Jane Johnson (University of Texas Southwestern, Dallas, United States)	N/A
Plasmid: pMCSIII-ePhox2a	This manuscript	N/A
Software and Algorithms		
Graphpad Prism 8 (macOS) – Version 8.3.0	Graphpad Software	https://www.graphpad.com/scientific-software/prism/ RRID: SCR_002798
Bioseb T2CT – Version 2.2.4	Bioseb	https://www.bioseb.com/en/pain-thermal-allodynia-hyperalgesia/897-thermal-place-preference-2-temperatures-choice-nociception-test.html RRID: N/A
ImageJ – Version 2.0.0	National Institutes of Health	https://imagej.nih.gov/ij/ RRID: SCR_003070
Photoshop and Illustrator	Adobe	N/A
Excel for Mac 2011 v14.7.7	Microsoft	N/A
Aegisub v3.2.2	Aegisub	http://www.aegisub.org/
Other		
Stereotaxic frame and digital display	David Kopf Instruments	Cat#: 940, 960, 1770, 900C, 922, 933-B RRID: N/A
Stereotaxic syringe pump with Micro4 controller	David Kopf Instruments	Cat#: UMP3-1, 1770-C RRID: N/A
Touch Test Sensory Evaluators (0.008, 0.02, 0.04, 0.07, 0.16, 0.4, 0.6, 1, 1.4, 2g)	North Coast Medical	Cat#: NC12775-01 - NC12775-10 RRID: N/A
IITC Hot Cold-Plate Analgesia Meter for Mice and Rats	IITC	Cat#: PE34 RRID: N/A
Thermal Place Preference, 2 Temperatures Choice Nociception Test	Bioseb	Cat#: BIO-T2CT RRID: N/A
iPhone SE – Version 12.3.1	Apple	Cat#: iPhone SE. RRID: N/A

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REAGENT or RESOURCE	SOURCE	IDENTIFIER
Micro Toothless Alligator Test Clip (Copper Plated)	https://www.amazon.com/Toothless-Alligator-Copper-Plated-Microscopic/dp/B0187MIUU4	Cat#: CECOMINOD005515 RRID: N/A
Confocal Microscope	Zeiss	Cat#: LSM-710
Confocal Microscope	Leica	Cat#: SP8
Epifluorescence Upright Microscope	Leica	Cat#: DM6000
Epifluorescence Upright Microscope	Leica	Cat#: DM6
Epifluorescence Dissecting Stereomicroscope	Leica	Cat#: MZ16FA

RESOURCE AVAILABILITY

Lead Contact

Further information and requests for resources and reagents should be directed to and will be fulfilled by the Lead Contact, Artur Kania (artur.kania@ircm.qc.ca).

Materials Availability

Phox2a^{Cre} mice are available from the Lead Contact upon request.

Data and Code Availability

The published article includes all datasets generated in this study. Single cell RNA-Seq data analyzed here was generated by [Delile et al. \(2019\)](#) and was obtained per their instructions from Array Express (<https://www.ebi.ac.uk/arrayexpress/>) with accession number “E-MTAB-7320.”

EXPERIMENTAL MODEL AND SUBJECT DETAILS

Mouse lines and Phox2a^{Cre} mouse line generation

Adult male and female mice, between 6–19 weeks of age, were used in this study. Sex ratios were kept as close to 1:1 as possible in all experiments, though not all experiments had the power to distinguish sex differences. Mice were kept on a 12 hour light: 12 hour dark cycle (light 6:00–18:00) with food and water provided *ad-libitum*. All procedures (except those involving *TrkA*^{-/-}, *Ptf1a*^{CRE} and *Ascl1*^{GFP} mice) were approved by the IRCM Animal Care Committee, using regulations and guidelines provided by the Canadian Council for Animal Care (CCAC). *TrkA*^{-/-} mouse use was approved by the Committee of Animal Care and Use of the National Cancer Institute, while the use of *Ptf1a*^{CRE} and *Ascl1*^{GFP} mouse lines (maintained on a mixed background of ICR and C57BL/6), was approved by the Institutional Animal Care and Use Committee at University of Texas Southwestern. Phox2a^{Cre} mice were generated at the IRCM where *Phox2a*-containing BAC RP23-333J21 ([GENSAT, 2008](#)) was modified by insertion of a Cre-PolyA sequence into the ATG site of *Phox2a* using GalK recombineering strategies ([Warming et al., 2005](#)). The Cre-containing BAC was injected into fertilized ova, and the resulting offspring were screened for genomic insertion of the BAC using Cre PCR. In total, we screened 230 pups, and were able to produce one founder from which all mouse lines containing *Phox2a*^{Cre} were derived. Genotyping was done by PCR for *Cre*, *FlpO*, *R26*^{LSL-tdT/+} (Ai14), *R26*^{FSF-LSL-tdT/+} (Ai65), *Phox2a*^{fl/fl} and *TrkA*^{-/-} as previously described ([Glasgow et al., 2005](#); [Kim et al., 2008](#)). The *Ptf1a*^{CRE} mouse line replaces the coding sequence for *Ptf1a* with that for *Cre* recombinase ([Kawaguchi et al., 2002](#)) and the *Ascl1*^{GFP} (*Ascl1*^{tm1Reed/J}) mouse strain replaces the coding sequence of *Ascl1* with that for *GFP* ([Leung et al., 2007](#)).

Generation of mice and mouse embryos

Mice containing the following transgenes or alleles were generated: Phox2a^{Cre}, R26^{LSL-tdT/+}, Phox2a^{Cre}, Cdx2^{FlpO}, R26^{FSF-LSL-tdT/+}, Hoxb8^{Cre}, Phox2a^{fl/fl}, R26^{LSL-tdT/+}, *Ascl1*^{GFP/GFP}, *Ptf1a*^{Cre/Cre}, and *TrkA*^{-/-} by breeding parents bearing one or more of the necessary alleles/transgenes. Vaginal plugs were checked daily at 6:00am, and the day of plug detection was noted as embryonic day 0.5 (e0.5). Mothers were anesthetised with a 0.3 mL intra-peritoneal injection of Ketamine/Xylazine solution. Embryos were dissected in ice-cold 1x phosphate-buffered saline (1x PBS), transferred to 4% paraformaldehyde in 1x PBS (4°C) and left to fix for two hours on a moving shaker (except *Ptf1a*^{Cre/Cre} and *Ascl1*^{GFP/GFP} embryos which were fixed for one hour). After fixation, embryos were washed briefly in 1x PBS, then cryoprotected in 30% sucrose for 1–2 days or until sunk. Embryos were harvested and fixed on the following embryonic days: *TrkA*^{-/-} on E14.5, *Ptf1a*^{Cre/Cre} and *Ascl1*^{GFP/GFP} both on E11.5 and E14.5, Hoxb8^{Cre}, Phox2a^{fl/fl}, R26^{LSL-tdT/+} on E11.5 and E16.5, and Phox2a^{Cre}, R26^{LSL-tdT/+} on E9.5, E10.5, E11.5, E12.5, E13.0, E13.5, E14.5, E15.5, E16.5 and E18.5.

Acquisition of human embryonic spinal cords

Human embryos were obtained with the parent's written informed consent (Gynaecology Hospital Jeanne de Flandres, Lille, France) with approval of the local ethic committee. Tissues were made available via the INSERM-funded Human Developmental Cell Atlas resource (HuDeCA) in accordance with the French bylaw (Good practice concerning the conservation, transformation and transportation of human tissue to be used therapeutically, published on December 29, 1998). Permission to use human tissues was obtained from the French agency for biomedical research (Agence de la Biomédecine, Saint-Denis La Plaine, France). Human embryo spinal cords were fixed by immersion for 12–24 hours in 4% paraformaldehyde in 0.12 M phosphate buffer, pH 7.4 (PFA) over night at 4°C. Samples were cryoprotected in a solution of 10% sucrose in 0.12 M phosphate buffer (pH 7.2), frozen in isopentane at 50°C and then cut at 20 μm with a cryostat (NX70 Thermo Fisher). Spinal cords from five separate embryos were used in this study: two from G.W. 7.3, and one each from G.W. 7.4, 8.0 and 8.4.

METHOD DETAILS

Neuronal birthdating

Pregnant female mice were given an i.p. injection of BrdU on E9.5, E10.5, E11.5 or E12.5 and embryos were harvested and fixed at E11.5, E12.5, E13.5 or E16.5. The BrdU dose was 50 mg/kg for all time points except E9.5, where this dose produced ubiquitous BrdU+ immunoreactivity in the spinal cord and thus was reduced to 25 mg/kg.

Stereotaxic surgery

Prior to surgery mice were given 1 mg/kg buprenorphine for analgesia, then anesthetised using a mixture of 5% isoflurane in oxygen and maintained using 2% isoflurane in oxygen. Eyes were coated in eye ointment to prevent drying during anesthesia. Prior to incision, the top of the head was shaved and decontaminated using an iodine solution. Mice were fitted into a stereotaxic frame with digital coordinate display and an incision was made longitudinally along the scalp to bare skull sutures. Injections were made via a hole drilled in the skull, which was made using medial-lateral and anterior-posterior coordinates for underlying brain regions as defined by the coronal Allen Brain reference atlas (Dong, 2008). Retrograde tracers (fluorogold or CTb-488) were injected using a 5 μL Hamilton syringe fitted with a pulled glass needle backfilled with mineral oil, which were injected in the VPL thalamus (coordinates AP –1.7, ML –2.0, DV –3.2), the MD thalamus (AP –1.25, ML –0.4, DV –3.2), the cerebellar vermis (AP –6.2, ML –0.8, DV –2.0), or the parabrachial nucleus (AP –5.35, ML –1.4, DV –3.05), identified using the coronal Allen Brain reference atlas (Allen Institute for Brain Science, 2004). Injection volumes of 500 nL (fluorogold, 2%) were injected into the VPL and 300 nL (CTb-488, 1%) into the MD thalamus, cerebellum or parabrachial nucleus. The needle was left in place for 5 minutes before slowly withdrawing to prevent reflux. The incision was then stitched together using silk sutures and mice were allowed to recover under a heating lamp before being returned to their home cage. Mice were perfused at 7 days post-injection and spinal cords were dissected.

Mouse behavioral assays

R. B. R. performed all behavioral assays, and was blinded to genotypes. R. B. R. and M. B. analyzed video-recorded mouse behavior, though each experimenter analyzed equal numbers of mice from each sex and genotype per assay. Mice of both sexes were used in each behavioral assay. Mice from control and Phox2a^{CKO} groups were always littermates and the same sex, to prevent confounding effects of litter versus sex. Control and Phox2a^{CKO} groups thus always contained an equal proportion of mice from each sex, and the proportion of male to female mice within groups was kept as close to 50% as possible, constrained only by the number of Phox2a^{CKO} mice generated (at an expected rate of 12.5% in a given litter). Mice were habituated in a dedicated mouse behavior room for at least 30 minutes prior to onset of tests. Mice received no other treatments other than the test itself. Mice were habituated in a small plexiglass chamber measuring 4 cm long, 2.2 cm wide and 2.5 cm high for von Frey, radiant heat paw-withdrawal, acetone and adhesive removal tests. For the von Frey and acetone tests, the chambers were placed atop a perforated stainless-steel floor due to the need for physical hind paw manipulations. For the radiant heat paw-withdrawal and adhesive removal test, the chambers were placed atop a transparent glass sheet. For all other assays mice were habituated in their home cages. When necessary, all behavioral tests were filmed using an iPhone SE except for the temperature preference assay, where the video camera included in the apparatus was used.

The **von Frey test** involved using a set of nylon filaments (0.008, 0.02, 0.04, 0.07, 0.16, 0.4, 0.6, 1.0, 1.4 g) to stimulate the hind paw plantar surface of each mouse in order to determine the median force which produces a withdrawal reflex. Mice were tested with a series of filaments using the “up-down” method of Dixon, as described previously (Chaplan et al., 1994; Mogil et al., 1999), with an inter-trial interval of at least 5 minutes.

The radiant heat paw-withdrawal (**Hargreaves**) test involved stimulating the hind paw plantar surface from below with a focused beam of light (set to 10% maximum intensity of the machine) and verifying latency to withdraw either hind paw. Each hind paw was stimulated eight times (16 total stimulations), and data was represented as the average of 16 withdrawal latencies, with an inter-trial interval of at least 2 minutes, performed as previously described (Hargreaves et al., 1988; Mogil et al., 1999).

The **hot water tail-withdrawal** test was performed as described previously (Mogil et al., 1999). Mice were placed in a small cloth pouch into which they entered voluntarily, the distal portion of the tail was dipped into a hot water bath maintained at 49 ± 1°C and the latency to withdraw the tail was recorded. Mice were tested three times with an inter-trial interval of at least 2 minutes, and data was represented as the average of 3 withdrawal latencies.

The **adhesive removal test** was performed as described previously (Bouet et al., 2009). Mice were tested on five consecutive days for the ability/motivation to remove an adhesive placed on the plantar surface of the hind paw. The adhesive was half of a 1.5 mL Eppendorf tube cap label, cut into a semicircle, and placed on the plantar surface. The latency to remove the label was recorded to the nearest minute, and these data were reported exactly as recorded (with only one test per day and no averaging between trials). If mice did not remove the adhesive within 30 minutes of the start of the test, latency was recorded as “30 minutes” for the purpose of data analysis, and mice were then returned to their home cage.

The **two-plate temperature preference assay** was performed as described previously (Minett et al., 2012). Two temperature-controlled metal plates were abutted together within a plexiglass enclosure. Mice were given the choice to travel between a probe temperature plate and a control temperature (always 30°C) plate for 10 minutes and the time spent per plate, distance traveled per plate and transitions between plates were recorded via a video camera above the enclosure (included with apparatus) and analyzed automatically via the accompanying software. Mice were tested twice for each probe temperature, and data for time/distance/transitions were represented as the average of both trials. In order to prevent mice from associating one plate as the control plate, the control plate was switched for each trial. Moreover, between testing for different probe temperatures, the initial position of the control plate was switched with the probe plate to prevent mice from associating the order of trials with the location of the control plate. As well, to encourage mice to sample both plates, mice were placed randomly on either the control plate or the probe plate for the first trial, and this order was then switched for the second trial.

The **hot-plate test** was performed as described previously (Mogil et al., 1999), and the **cold-plate test** was performed using similar methods. Mice were placed within the hot-cold plate apparatus (IITC PE34) on a stainless-steel metal plate heated to $53 \pm 0.1^\circ\text{C}$ or cooled to $0 \pm 0.1^\circ\text{C}$ and were video-recorded from the side (with a mirror opposite the test chamber to view each side of the mouse) for 60 s at which point they were returned to their home cage. The latency to either lick the hind paw, flutter of the hind paw or to attempt to escape via jumping was recorded. Additional behaviors were recorded: total time spent licking either hind paw, total hind paw licking episodes, total jumps and total hind paw flutters. Mice were tested once, and data were represented directly based on behaviors recorded in one 60 s trial. Entirely different cohorts of mice were used for the hot and cold-plate tests respectively, to prevent behavioral adaptation to the test.

The **acetone test** was performed as described previously (Colburn et al., 2007). Briefly, the mouse’s hind paw was stimulated with a drop of acetone extruded from the blunt end of a 1ml syringe. Mice were recorded for 60 s following the application, and total time spent licking was recorded as well as the magnitude of behavior on a 0–2 scale as reported previously (Colburn et al., 2007). Mice were stimulated 5 times, with an inter-trial interval of at least 5 minutes. Total licking time was reported as a sum of 5 trials, and the behavioral score (0–2) was reported as an average of 5 trials.

The **foot clip test** was performed as described previously (Pan et al., 2019). Briefly, a toothless mechanical clip was used to pinch skin on the plantar surface of the hind paw, and mice were placed in a plexiglass cylinder (dimensions) on the glass sheet used previously and video recorded from below for 60 s (this recording setup is identical to the following formalin and capsaicin tests). The total amount of time licking the clipped hind paw was recorded, and data is presented as the total time licking during the one trial.

The **capsaicin and formalin tests** (Mogil et al., 1999; Sakurada et al., 1992) were performed similarly – mice were injected with approximately 20 μL of capsaicin solution (1.5 $\mu\text{g}/20 \mu\text{L}$ in 1x PBS) or formalin solution (2% in 1x PBS) in the plantar surface of the right hind paw using a standard 28G insulin syringe (BD) and video recorded from below for either 15 or 60 minutes respectively. Mice were tested only once on each test, with different cohorts of mice used for each respective test. Data were represented as time spent licking the injected hind paw. For formalin-injected animals, these data were analyzed separately acutely after injection (0–10 minutes) or chronically after injection (11–60 minutes).

Tissue fixation, freezing and sectioning

Adult mice were first anesthetised with a 0.3 mL i.p injection of Ketamine/Xylazine solution (10 mg/ml Ketamine, 1 mg/ml Xylazine, in 0.9% saline). Transcardial perfusion was done with a peristaltic pump (Gilson miniPuls2). Mice were perfused with 10 mL of ice cold 1x PBS followed by 20 mL of ice cold 4% PFA in 1x PBS. Brains and spinal cords were dissected and post-fixed in 4% PFA in 1x PBS at 4°C for two hours, washed briefly in 1x PBS, and acclimated to 30% sucrose for 1–2 days or until sunk. After cryoprotection, tissue was frozen in OCT Compound and cryosectioned at -22°C . Tissue was cut into 25 μm sections for all experiments other than RNA Scope, in which case 10 μm sections were used, and those involving *Ptf1a*^{CRE} and *Ascl1*^{GFP} lines where 30 μm sections were used.

Immunohistochemistry

For mouse tissue, sections were heated at 37°C for 15 minutes prior to immunohistochemistry. Following this, sections were washed three times in 1x PBS for 10 minutes, blocked using a solution of 5% heat-inactivated horse serum (HIHS) and 0.1% Triton X-100 in 1x PBS (0.1% tPBS) for 30 minutes, and incubated with a primary antibody solution (in 1% HIHS, 0.1% tPBS) overnight at 4°C. The following day, sections were again washed three times in 1x PBS for 10 minutes, and incubated with a secondary antibody solution (in 1% HIHS, 0.1% tPBS) at room temperature for 1 hour. Following this, sections were washed three more times in 1x PBS for 10 minutes and coverslipped using a Mowiol solution (10% Mowiol - Sigma, 25% glycerol). Slides were allowed to dry in the dark at room temperature and subsequently imaged using fluorescent microscopy. For immunohistochemistry involving the anti-BrdU antibody, two rounds of immunohistochemistry were done: the first round involved staining for RFP or Phox2a and the second round for BrdU with some modifications. Prior to the anti-BrdU primary antibody incubation, slides were treated in a 2 N hydrochloric acid solution at

37°C for 30 minutes. Subsequently, slides were neutralized by washing in a Tris-buffered saline solution (pH 8.5, 50 mM Tris, 150 mM NaCl) for 10 minutes at room temperature, after which primary antibody incubation was done. BrdU immunohistochemistry proceeded in two steps, as acid denaturation of DNA reveals anti-BrdU epitopes but destroys RFP/Phox2a epitopes; however, acid denaturation does not destroy secondary antibody-conjugated fluorophores from the first round of immunohistochemistry. Immunohistochemistry on human tissue was performed on cryostat sections after blocking in 0.2% gelatin in PBS containing 0.25% Triton X-100 (Sigma). Sections were then incubated overnight with respective primary antibodies, all used at 1:500 dilutions, followed by 2 hours incubation in appropriate secondary antibodies. In figures demonstrating tdT signal, amplification was done using an anti-RFP antibody (see key resources table).

In situ-hybridization (ISH)

ISH was done using RNA Scope® Multiplex Fluorescent v2 kits, according to manufacturer's instructions. All experiments used Mm-Phox2a-C2 coupled to Opal™ 520, Mm-Lmx1b-C3 coupled to Opal™ 690, and all other candidate probes being compared to Phox2a (all Mm C1 probes) were coupled to Opal™ 570.

In-ovo chicken electroporation and tissue processing

Fertilized White Leghorn eggs were obtained from the Texas A&M Poultry Department (College Station, TX, USA) and incubated for 48 hours at 39°C. The supercoiled reporter plasmid *ePhox2a-GFP* was diluted to 1.5 mg/mL in H₂O/1X loading dye and injected into the lumen of the closed neural tube at stages Hamburger-Hamilton (HH) stages 13-15 (~E2) along with either a *pMiWIII-Myc* epitope tagged plasmid serving as an electroporation control or the same plasmid containing the coding region of *Ascl1*, *Ptf1a*, or *Prdm13* (Hamburger and Hamilton, 1951). The injected embryos were then electroporated with 5 pulses of 25 mV each for 50 msec with intervals of 100 msec. Embryos were harvested 48 hours later at HH stages 22-23 (~E4), fixed with 4% paraformaldehyde for 45 minutes, and processed for cryosectioning and immunofluorescence.

Generation of reporter constructs and expression vectors

Previously published ChIP-seq data for *Ascl1*, *Ptf1a*, *Rbpj*, and *Prdm13* (Borromeo et al., 2014; Meredith et al., 2013; Mona et al., 2017) (GSE55840; GSE90938) were used to identify a putative enhancer for *Phox2a* in the chicken dorsal neural tube. An 851 bp region (chr7:101834344-101835194 from mm10) encompassing two peaks bound by *Ascl1* and *Ptf1a* was cloned into the MCSIII GFP reporter cassette (*ePhox2a-GFP*). This reporter cassette contains the β -globin minimal promoter, a nuclear localized fluorescence reporter, and the 3' cassette from the human growth hormone. The *ePhox2a* sequence was PCR amplified from ICR mouse DNA. *Prdm13*, *mycPtf1a*, and *mycAscl1* were expressed in the pMiWIII expression vector (Chang et al., 2013; Gowan et al., 2001; Matsunaga et al., 2001). All constructs were sequence-verified and expression of the transcription factor confirmed by immunohistochemistry with antibodies to the myc tag or with factor-specific antibodies.

Epifluorescence and Confocal Microscopy

Micrographs of tissue sections were taken either with epifluorescence microscopes (Leica DM6, DM6000) or confocal microscopes (Leica SP8 or Zeiss LSM710). All RNA Scope images, for quantification and for analysis, were taken using confocal microscopes on a 40x objective in order to resolve single puncta.

Human sections were imaged with a laser scanning confocal microscope (FV1000, Olympus) and processed using ImageJ (NIH) and Adobe Photoshop.

QUANTIFICATION AND STATISTICAL ANALYSIS

Bioinformatics

scRNA-seq data used in this study were previously published (Delile et al., 2019), using spinal cord cells from E9.5, E10.5, E11.5, E12.5 and E13.5 mouse embryos and processed via the 10x Genomics Chromium Single Cell 3' v2 protocol. Raw data were extracted from ArrayExpress E-MTAB-7320. Cell Ranger v 2.1.1 (Zheng et al., 2017) was used to align reads to the 10X mm10 mouse reference genome v2.1.0, filter barcodes and quantify genes. Biological replicates from the same time points were then merged using Cell Ranger's *aggregate* function. All downstream analyses were performed on these 41 009 cells, using the Seurat v.3.1.1. R package.

Using Delile et al. (2019) annotation, only cells classified as neurons were kept for further analysis (18 048 cells). From these, data for each time point was normalized and highly variable genes identified using SCTransform's normalization and variance stabilizing methods (Hafemeister and Satija, 2019). Different time points were then integrated using CCA (Stuart et al., 2019). From the integrated dataset, 2 subsets were created: 1) all cells expressing *Lmx1b* (*Lmx1b+*) were isolated (2614 cells) and 2) all early (E9.5, E10.5 and E11.5) *Lmx1b+* cells (186 cells). For each of these subsets, dimensionality reduction (PCA and UMAP) was applied and clusters identified using Seurat's SNN modularity optimization-based clustering Louvain algorithm. Differential expression analysis was performed on the non-integrated assay to identify markers for each cluster (Wilcoxon Rank Sum test).

A differential expression analysis was then performed on clusters of interest in each of these 3 subsets (cluster 6 from *Lmx1b+* cells and clusters 2 and 3 in early *Lmx1b+* cells). Each of these clusters was compared to all other neuron cells in order to identify specific

markers within these clusters. This analysis was limited to genes which had on average, at least 0.1 log-Fold difference between the two groups compared and present in at least 10% of cells of either group. Markers were then considered significantly differentially expressed if adjusted p values < 0.05.

Cell counts

All cell counts were done with ImageJ v. 2.0.0 software, using the cell counter plugin. Data was recorded and sorted in Microsoft Excel.

Animal Behavior

Video-recordings of mice were quantified by R.B.R and M.B using Aegisub (free subtitling software), which includes video annotation functions allowing precise start and stop times for specified behaviors to be recorded. Data was exported and then sorted in Microsoft Excel.

Data representation

Graphs display data points from individual animals (hollow circles), mean data from all animals (bars), and \pm standard error of the mean (error bars).

Numbers

All numbers are noted in figure legends. In all experiments using adult mice, n represents unique individuals. In all experiments using embryonic mice, n represents unique embryos. No data represented as single ns have been pooled from multiple individual animals.

Statistics

All statistical analyses were performed using GraphPad Prism v8.3.0 software, except those involving single cell RNA-Seq data processing, which are described in Bioinformatics (above). Statistical tests used are described in figure legends. Significance is represented as ns: non significant, *p < 0.05, **p < 0.01 or ***p < 0.001.

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Supplemental Information

Phox2a Defines a Developmental Origin of the Anterolateral System in Mice and Humans

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SUPPLEMENTAL INFORMATION:

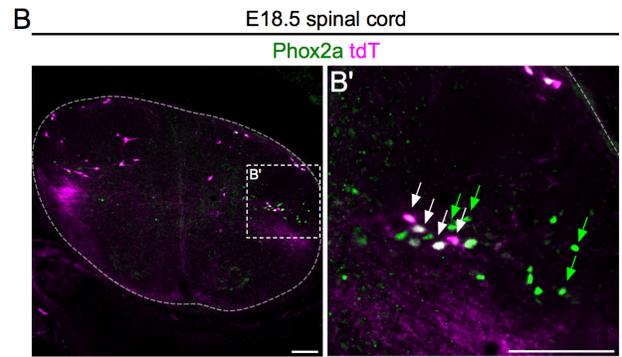
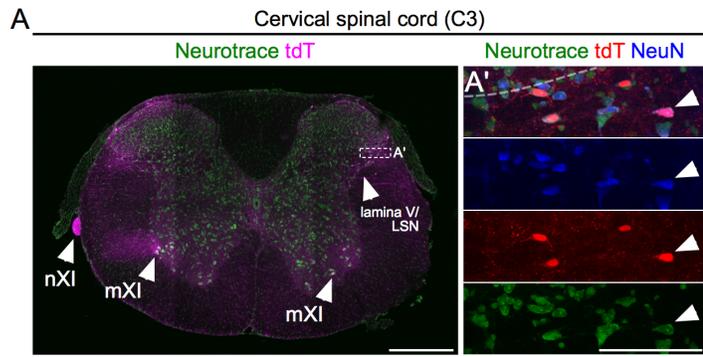
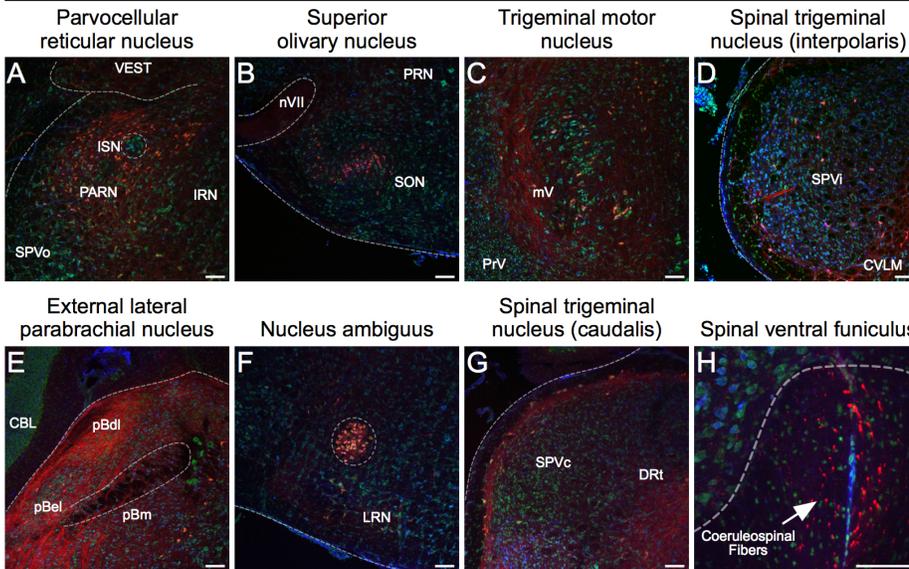


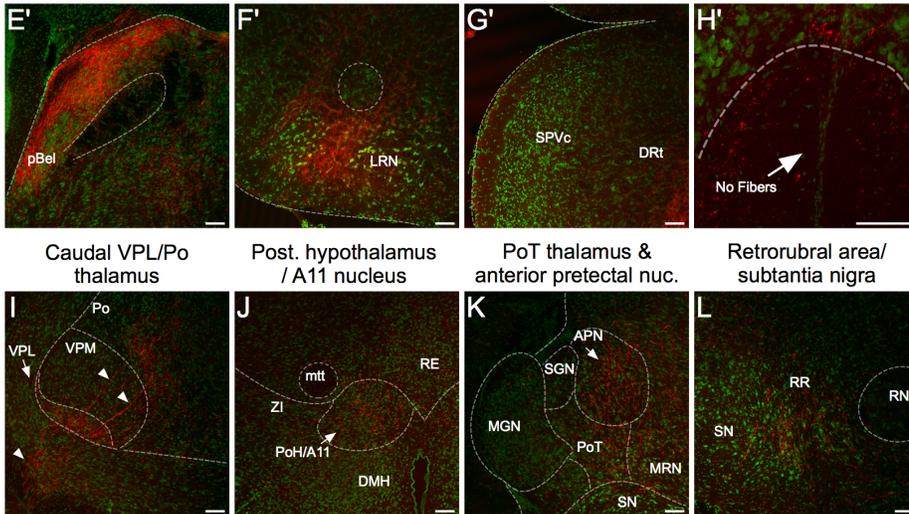
Figure S1: Spinal *Phox2a*^{Cre} neurons reside in lamina I, V and LSN. Related to Figure 1.

All images are of *Phox2a*^{Cre}; *R26*^{LSL-tdT/+} mice. (A) TdT+ neurons in the cervical spinal cord of adult mice. (A') Magnified box from (A) showing lamina V/LSN Neurotrace, tdT and NeuN co-labelling. (B) Expression of tdT in *Phox2a*+ neurons in an E18.5 mouse embryo. Numbers: (A) n=3 adult mice, (B) n=3 E18.5 embryos. Scale bars: (A) 500 μ m, (A', B) 100 μ m. Abbreviations: nXI (accessory motor nerve).

NeuN Neurotrace tdT (*Phox2a^{Cre}; R26^{LSL-tdT}*)



Neurotrace tdT (*Phox2a^{Cre}; Cdx2^{FipO}; R26^{FSF-LSL-tdT}*)



FoxP2 tdT (*Phox2a^{Cre}; Cdx2^{FipO}; R26^{FSF-LSL-tdT}*)

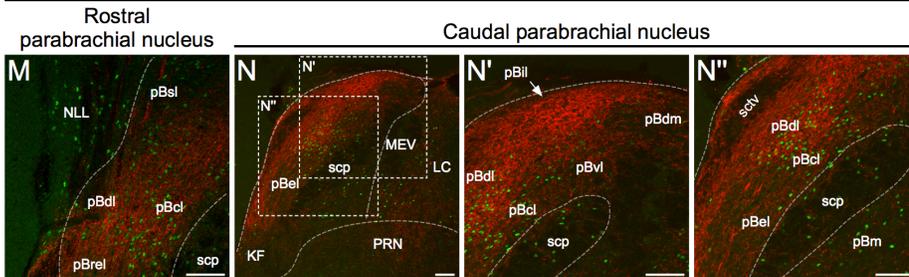


Figure S2: Spinal *Phox2a*^{Cre} neurons innervate AS targets. Related to Figure 2.

(A–G) NeuN, Neurotrace and tdT staining in brain regions expressing *Phox2a*^{Cre}; *R26*^{LSL-tdT/+} not depicted in Fig. 2. (E'–G') Lack of cellular tdT expression in *Phox2a*^{Cre}; *Cdx2*^{FLPo}; *R26*^{FSF-LSL-tdT/+} mice compared to the same brain regions of *Phox2a*^{Cre}; *R26*^{LSL-tdT/+} mice (E–G). (H, H') Large axons in the spinal ventral funiculus of *Phox2a*^{Cre}; *R26*^{LSL-tdT/+} mice (H), absent in *Phox2a*^{Cre}; *Cdx2*^{FLPo}; *R26*^{FSF-LSL-tdT/+} mice (H'), are likely coeruleospinal. (I–L) Additional brain regions receiving *Phox2a*^{Cre+} spinofugal axons not depicted in Fig. 2. (M–N'') Spinal *Phox2a*^{Cre+} axons in parabrachial subnuclei identified by FoxP2 expression (present in pBdl, pBcl but absent from pBel), in *Phox2a*^{Cre}; *Cdx2*^{FLPo}; *R26*^{FSF-LSL-tdT/+} mice. N' and N'': higher magnification of boxes in N.

Numbers: (A–H) n=3 *Phox2a*^{Cre}; *R26*^{LSL-tdT/+} adult mice, (E'–H', I–N, N', N'') n=3 *Phox2a*^{Cre}; *Cdx2*^{FLPo}; *R26*^{FSF-LSL-tdT/+} adult mice.

Scale bars: 100 μ m.

Abbreviations: APN (anterior pretecal nucleus), CBL (cerebellum), DMH (dorsomedial hypothalamus), DRt (dorsal reticular nucleus), IRN (intermediate reticular nucleus), ISN (inferior salivatory nucleus), KF (Kölliker-Fuse nucleus), LC (locus coeruleus), MGN (medial geniculate nucleus), MRN (midbrain reticular nucleus), mtt (mamillothalamic tract), mV (trigeminal motor nucleus), nVII (facial motor nerve), pBrel (rostral external-lateral parabrachial nucleus), PoH (posterior hypothalamus), PoT (posterior triangular thalamus), RE (Reuniens nucleus), RN (red nucleus), RR (retrotrubral nucleus), sctv (ventral spinocerebellar tract), SGN (suprageniculate nucleus), SN (substantia nigra), SON (superior olivary nucleus), SPVc (spinal trigeminal nucleus, caudalis), SPVi (spinal trigeminal nucleus, interpolaris), SPVo (spinal trigeminal nucleus, oralis), VEST (vestibular nuclei), VPM (ventral posteromedial thalamus).

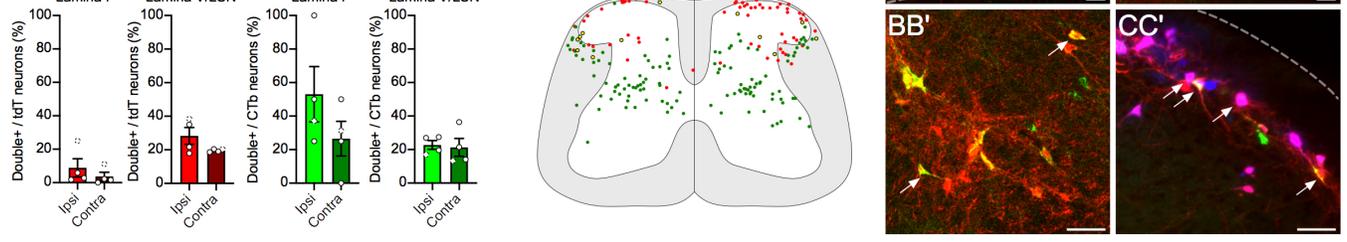
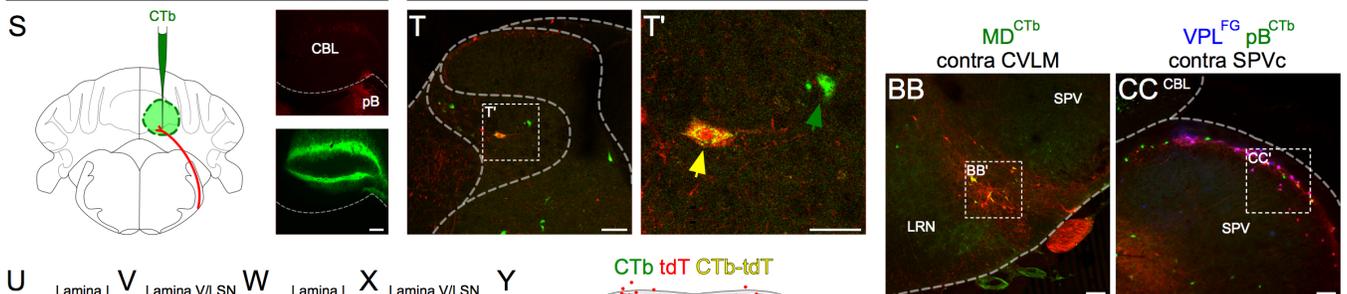
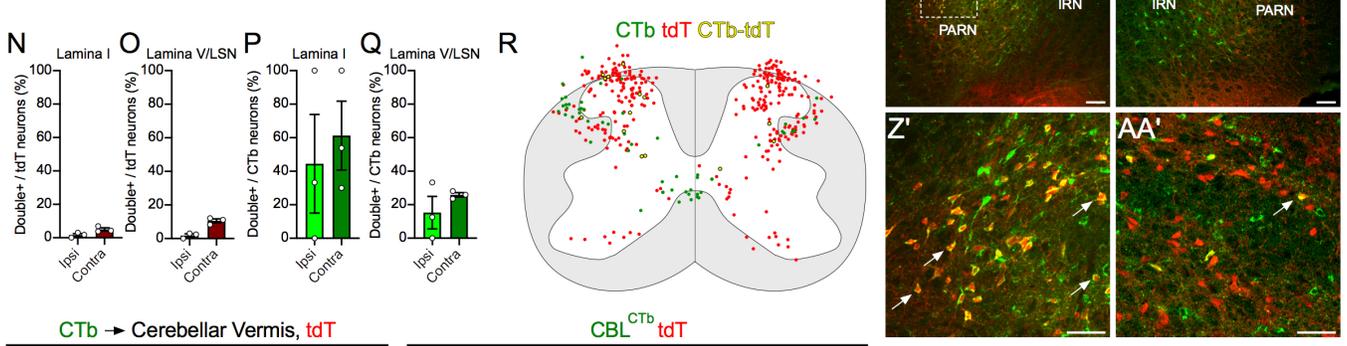
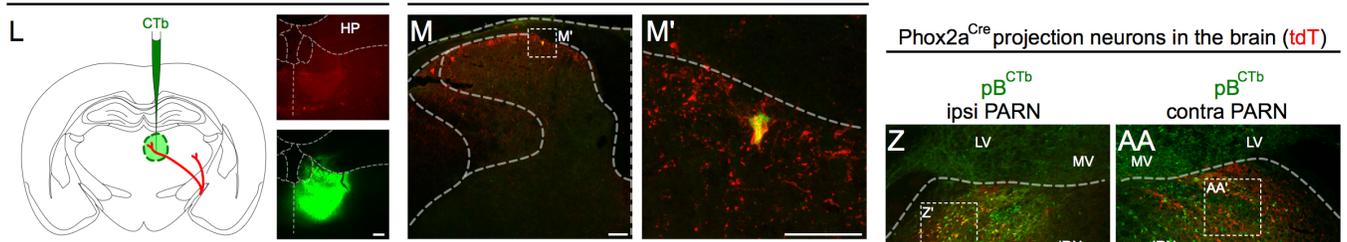
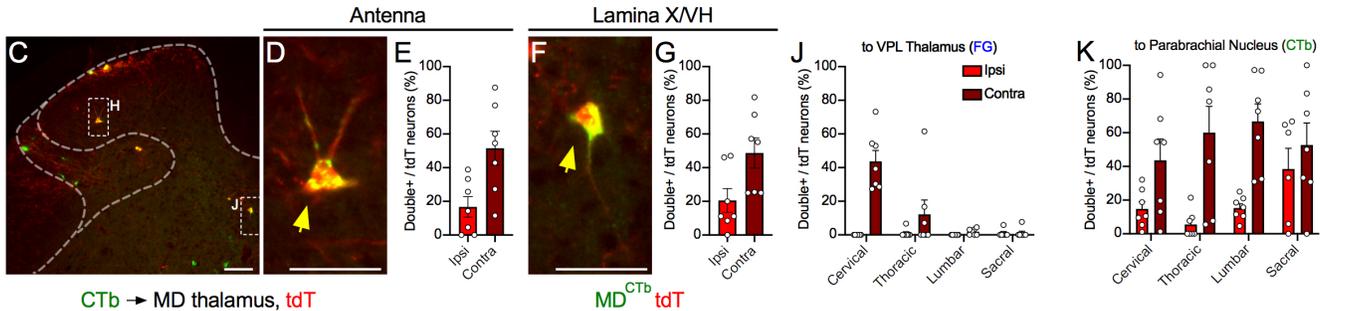
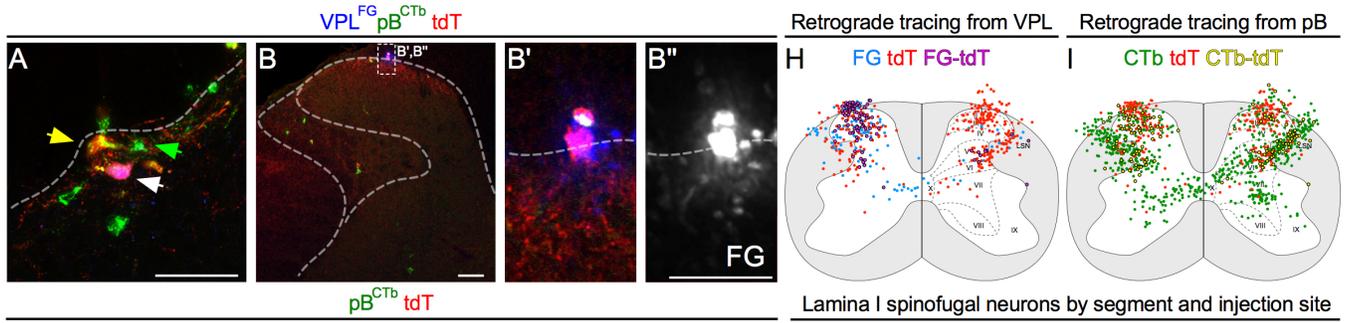


Figure S3: Spinal *Phox2a*^{Cre} neurons are predominantly AS neurons. Related to Figure 3.

(A) Inset of Fig. 3G, demonstrating tdT and retrograde tracer co-localization in lamina I of *Phox2a*^{Cre}; *R26*^{LSL-tdT/+} mice with VPL-injected FG and pB-injected CTb. White arrow: tdT+ cell labelled with both FG and CTb; green arrow: CTb-only cell; yellow arrow: tdT+ cell labelled with CTb. (B–B'') Examples of FG-labelled neurons. As FG+ cells were difficult to detect with the confocal microscope, an epifluorescence microscope was used to photograph them prior to cover-slipping (B'') after which, CTb and tdT were imaged using a confocal microscope. FG and CTb/tdT images were then overlaid in register and merged; (B', B'') are higher magnifications of boxed lamina I area in B. White arrow: tdT+ cell doubly labelled with FG and CTb; magenta arrow: tdT+ cell labelled with FG; yellow arrow: tdT+ cell labelled with CTb. (C–G) Rare *Phox2a*^{Cre} neuron types such as antenna neurons (D) and lamina X/VH neurons (F). Percent of tdT+ neurons labeled with either or both tracers in Antenna (E) or lamina X/VH (G) neurons. (H, I) Diagram of the location of tdT+ neurons (red) and VPL-injected FG-labelled neurons (H, FG+ neurons in blue, and tdT+/FG+ co-labelled neurons in magenta) or pB-injected CTb-labelled neurons (I, CTb+ neurons in green and tdT+/CTb+ neurons in yellow) in five, non-sequential 25 μm sections of the cervical spinal cord of one representative *Phox2a*^{Cre}; *R26*^{LSL-tdT/+} mouse. (J, K) Percent of lamina I tdT+ neurons labelled with VPL-injected FG (J) or pB-injected CTb (K) in cervical, thoracic, lumbar and sacral spinal segments. (C–G) (L–R) Analysis of *Phox2a*^{Cre}; *R26*^{LSL-tdT/+} mice injected in the MD thalamus with CTb. (L) Representative image of a CTb injection site. HP: Hippocampus (M, M') Representative cervical spinal cord section with a lamina I tdT+ neuron labelled with CTb (M'). (N–O) Percent of tdT neurons labelled with CTb in lamina I (N) or lamina V/LSN (O) of all spinal segments. (P–Q) Percent of CTb-labelled neurons expressing tdT in lamina I (P) or lamina V/LSN (Q) of all spinal segments. (R) Diagram of the location of tdT+ only (red), CTb+ only (green) or tdT+/CTb+ co-labelled (yellow) neurons, in 5 non-sequential 25 μm sections of the cervical spinal cord compiled from all 3 animals analysed. (S–Y) Analysis of *Phox2a*^{Cre}; *R26*^{LSL-tdT/+} mice injected in the cerebellar (CBL) vermis with CTb (green). (S) Representative image of a CTb injection site. (T, T') Representative lumbar spinal cord section where a lamina V/LSN tdT+ neuron is labelled with CTb and magnified in (T'; yellow arrow). Green arrow shows a CTb-only neuron. (U–V) Percent of tdT neurons labelled with CTb in lamina I (U) or lamina V/LSN (V) at different spinal levels. (W–X) Percent of CTb-labelled neurons expressing tdT in lamina I (W) or lamina V/LSN (X) of all spinal segments. (Y) Diagram of the location of tdT+ only (red), CTb+ only (green) or tdT+/CTb+ co-labelled (yellow) neurons, in 10 non-sequential 25 μm sections of the lumbar spinal cord compiled from one representative animal. A CTb injection in one of the four mice spread into the dorsal pB – data points from this animal are represented by circles with dashed borders; increased incidence of lamina I tdT+ neurons labeling with

CTb is notable. (Z–CC) Examples of tdT+ neurons in various brain regions (arrows), labelled by retrograde tracer injections in the pB, VPL and MD thalamus. (Z, Z') tdT+ parvocellular reticular nucleus (PARN) neurons project to the ipsilateral parabrachial nucleus (AA, AA') but rarely to the contralateral parabrachial nucleus, (BB, BB') tdT+ caudal ventrolateral medulla (CVLM) neurons project to the MD Thalamus, and (CC, CC') lamina I/paratrigeminal neurons of the spinal trigeminal nucleus (SPVc) project to the VPL thalamus and, much more sparsely, to the parabrachial nucleus.

Numbers: (S3A–K, Z–CC) n=7 *Phox2a*^{Cre}; *R26*^{LSL-tdT/+} adult mice (4 male, 3 female), (S3L–R) n=3 *Phox2a*^{Cre}; *R26*^{LSL-tdT/+} adult mice (1 male, 2 female), (S3S–Y) n=4 *Phox2a*^{Cre}; *R26*^{LSL-tdT/+} adult mice (2 male, 2 female).

Data are represented as mean ± SEM.

Scale bars: 100 μm, except (L, S) 250 μm, and (A, B', B'', D, F, M', T', Z'–CC') 50 μm.

Abbreviations: HP (Hippocampus).

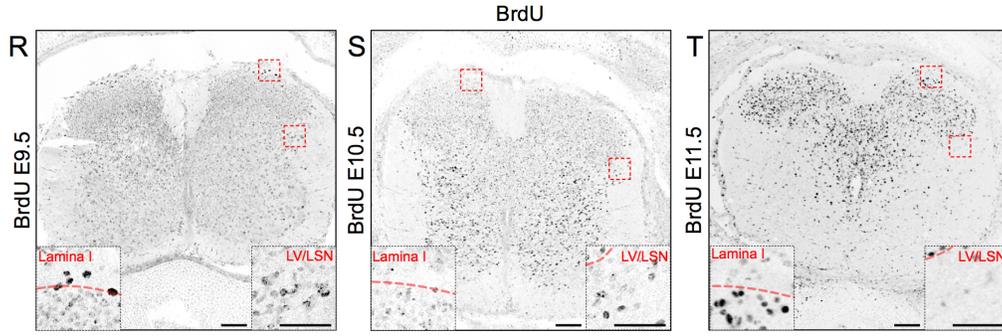
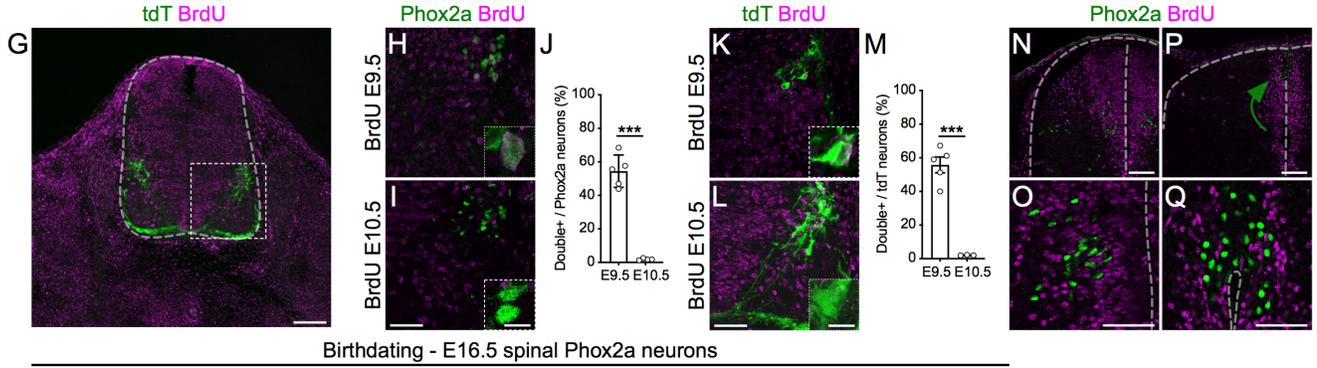
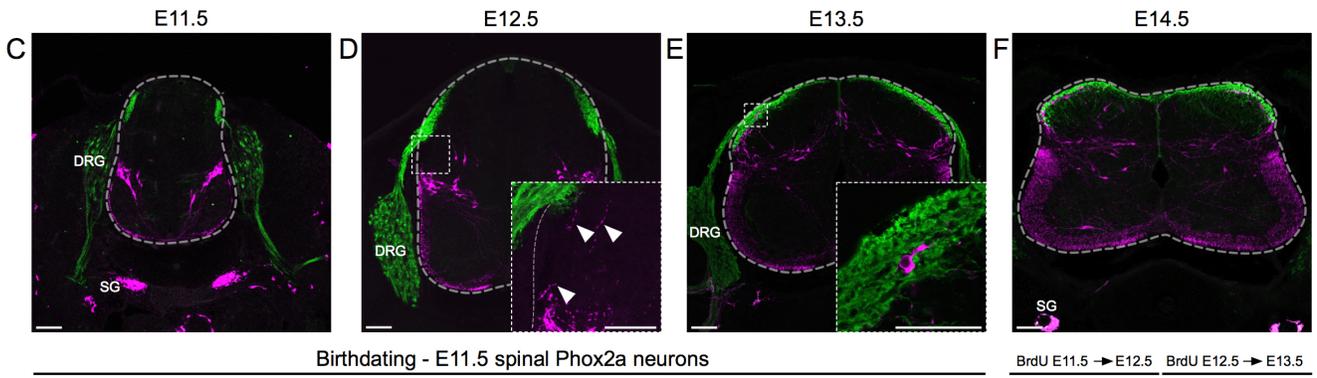
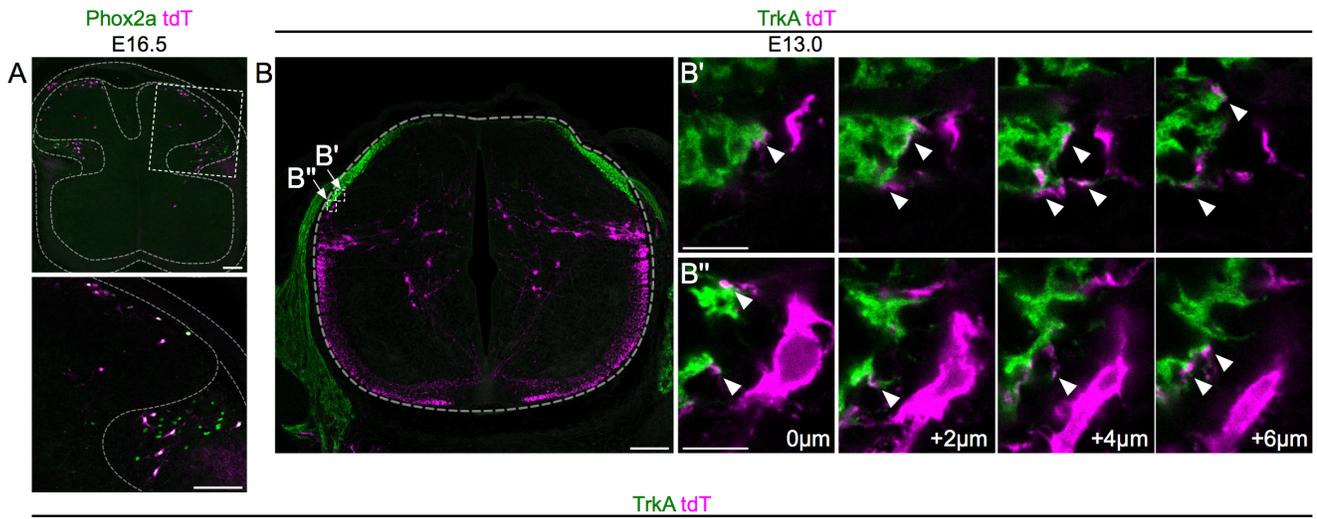


Figure S4: Heterogeneity of spinal Phox2a neuron migration, sensory afferent interaction and birth time. Related to Figure 4.

(A) Position of Phox2a⁺, tdT⁺ and Phox2a⁺ tdT⁺ (white) neurons in E16.5 embryonic spinal cords of *Phox2a*^{Cre}; *R26*^{LSL-tdT/+} mice. (B–F) Development of contacts between lamina I Phox2a neurons and TrkA⁺ primary afferents in E13.0 (B), E11.5 (C), E12.5 (D), E13.5 (E) and E14.5 (F) *Phox2a*^{Cre}; *R26*^{LSL-tdT/+} mouse spinal cords. (B', B'') Consecutive confocal microscopy z-stack images of lamina I neuron (magenta) contacts with TrkA primary afferents in E13.0 *Phox2a*^{Cre}; *R26*^{LSL-tdT/+} mouse embryos, magnified boxed areas from (B). (D) inset: Phox2a neuron fibres near the dorsal root entry zone, prior to Phox2a cell soma entry into the dorsal horn. (E) inset: a lamina I Phox2a cell surrounded by TrkA afferents. (G–M) Birthdating of spinal Phox2a neurons in E11.5 *Phox2a*^{Cre}; *R26*^{LSL-tdT/+} mouse embryos, labelled by BrdU exposure at E9.5 (H, K) or E10.5 (I, L). (G) The spinal cord of an E11.5 embryo exposed to BrdU at E10.5, showing BrdU and tdT labelling. (J, M) Percent of all tdT cells labelled with BrdU (J) and the percent of all Phox2a⁺ cells labelled with BrdU (M), in E11.5 *Phox2a*^{Cre}; *R26*^{LSL-tdT/+} embryos exposed to BrdU at E9.5 or E10.5 (n=4-5). (N, O) E12.5 spinal cord of *Phox2a*^{Cre}; *R26*^{LSL-tdT/+} embryos showing Phox2a^{DeepLate} neurons (Phox2a⁺) intermingled with newly born (BrdU⁺) neurons exposed to BrdU at E11.5. (P, Q) E13.5 spinal cord of *Phox2a*^{Cre}; *R26*^{LSL-tdT/+} mouse embryos with Phox2a^{DeepLate} neurons clustered near the roof of the central canal, separate from other newborn (BrdU⁺) neurons, labelled with BrdU at E12.5. (R–T) E16.5 spinal cord of *Phox2a*^{Cre}; *R26*^{LSL-tdT/+} mouse embryos pulsed with BrdU at E9.5 (R), E10.5 (S) or E11.5 (T), showing BrdU⁺ cells in lamina I and lamina V/LSN (insets). Lamina I cells are labelled by a BrdU pulse at E9.5 but E10.5 and E11.5 BrdU pulses result in sparser labelling. Lamina V/LSN cells show labelling by E9.5 and E10.5 BrdU pulses but minimal labelling by a E11.5 BrdU pulse.

Numbers: *Phox2a*^{Cre}; *R26*^{LSL-tdT/+} embryos: (A) n=3 E16.5, (B) n=3 E13.0, (C) n=3 E11.5, (D) n=3 E12.5, (E) n=3 E13.5, (F) n=3 E14.5, (G–M) n=3-5 E11.5 per condition, (N, O) n=3 E12.5, (P, Q) n=3 E13.5, (R) n=3 E16.5, (S) n=3 E16.5, (T) n=3 E16.5.

Statistics: (J, M) Unpaired t-test, ***: p<0.0001. Data are represented as mean ± SEM.

Scale bars: (A–G, N, P, R–T) 100 μm. (H, I, K, L, O, Q) and insets of (D, E, R, S, T) 50 μm, (B', B'') and insets of (H, I, K, L) 10 μm.

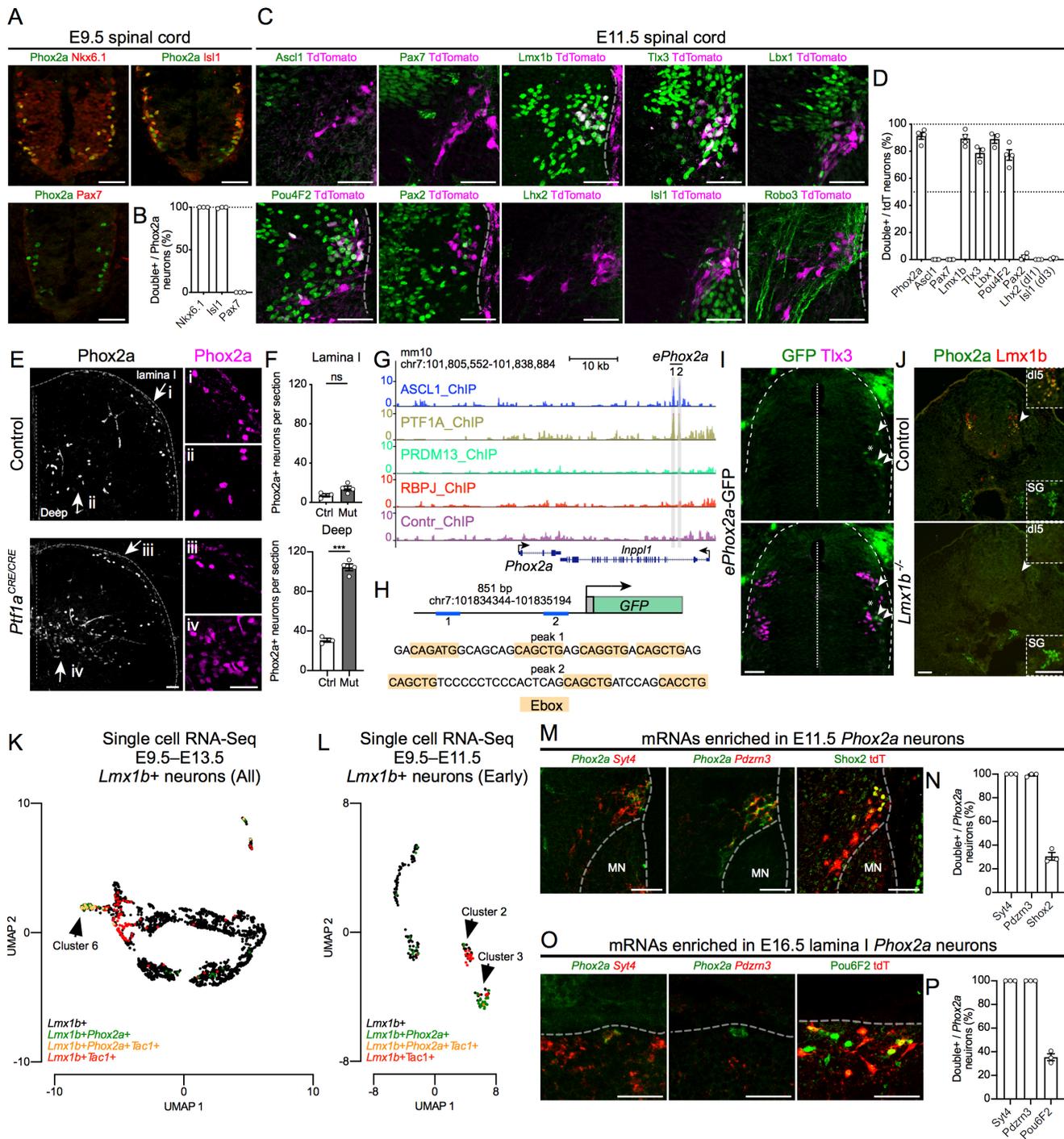


Figure S5: The molecular identity and specification of spinal Phox2a neurons. Related to Figure 5.

(A, B) Molecular characterisation of spinal Phox2a neurons in the E9.5 *Phox2a^{Cre}; R26^{LSL-tdT/+}* spinal cord. Phox2a neurons co-express ventral motor neuron markers Nkx6.1, Isl1, but not the progenitor marker Pax7, and are most likely accessory motor neurons. The percent of Phox2a+ cells co-expressing Nkx6.1, Isl1 and Pax7 is quantified. (C, D) Co-expression of tdT with embryonic spinal neuron markers in the in the E11.5 *Phox2a^{Cre}; R26^{LSL-tdT/+}* spinal cord using immunohistochemistry, with (D) quantification of each marker done as a percent of tdT+ neurons. (E) Detection of Phox2a cells in spinal cords of control and *Ptf1a^{Cre/Cre}* E14.5 mice. Panels with Roman numerals are magnifications of regions marked by arrows. (F) Number of Phox2a+ cells per average section in lamina I and deep laminae of E14.5 control and *Ptf1a^{Cre/Cre}* (mut) mice. (G, H) Genome sequences ChIP-Seq hits detected using antibodies against Ascl1, Ptf1a, Prdm13, and Rbpj at the *Phox2a* locus (G). The two sites bound by both Ascl1 and Ptf1a are highlighted in gray. (H) Diagram of the *ePhox2a* GFP reporter construct with the cis-regulatory sequence containing Ascl1 and Ptf1a sites. The sequence at the center of each ChIP-seq peak is shown with the E-box basic helix-loop-helix transcription factors binding motif is highlighted in orange. The active Ptf1a containing transcription complex requires a TC-Rbpj binding sequence that is not present in this enhancer. (I) Transverse sections of embryonic day 4 (E4) chicken neural tubes electroporated with the *ePhox2a-GFP* reporter at embryonic day 2. *ePhox2a* directs dl5-specific GFP expression in Tlx3+ cells. (J) E11.5 *Lmx1b^{-/-}* spinal cords lack Phox2a expression (dl5 inset), but sympathetic ganglia maintain it (SG inset). (K, L) UMAP plots of E9.5-E13.5 *Lmx1b⁺* neurons (K) and E9.5-E11.5 *Lmx1b⁺* neurons (L) showing *Lmx1b⁺/Phox2a⁺*, *Lmx1b⁺/Tac1⁺*, and *Lmx1b⁺/Phox2a⁺/Tac1⁺* cell clusters. (M, N) Validation of selected dl5-enriched mRNAs at E11.5 by RNA-Scope in situ detection of *Synaptotagmin 4*, *Pdzrn3* mRNAs and Shox2 protein immunofluorescence (Z). (N) Percent of E11.5 Phox2a+ or *Phox2a* mRNA+ neurons co-expressing the selected mRNAs or proteins. (O, P) Validation of selected dl5-enriched mRNAs at E16.5 in lamina I, by RNA-Scope in situ detection of *Synaptotagmin 4* and *Pdzrn3* mRNAs and Pou6F2 protein immunofluorescence. (P) Percent of E16.5 Phox2a+ or *Phox2a* mRNA+ neurons co-expressing the selected mRNAs or proteins. Numbers: *Phox2a^{Cre}; R26^{LSL-tdT/+}* embryos (A, B) n=3 E9.5, (C, D) n=3-4 E11.5, (M, N) n=3 E11.5, (O, P) n=3 E16.5. (E, F) n=3 control, n=4 *Ptf1a^{Cre/Cre}* E14.5 embryos, (I) n=6 E4 chicken embryos, (J) n=3 control, n=3 *Lmx1b^{-/-}* E11.5 embryos. Statistics: (F) Groups compared with Student's t-test, ***: p<0.001. (K, L) Data derived from Delile et al., (2019); data processing and statistics described in STAR Methods. Data are represented as mean ± SEM. Scale bars: All 50 µm, except (J) 100 µm.

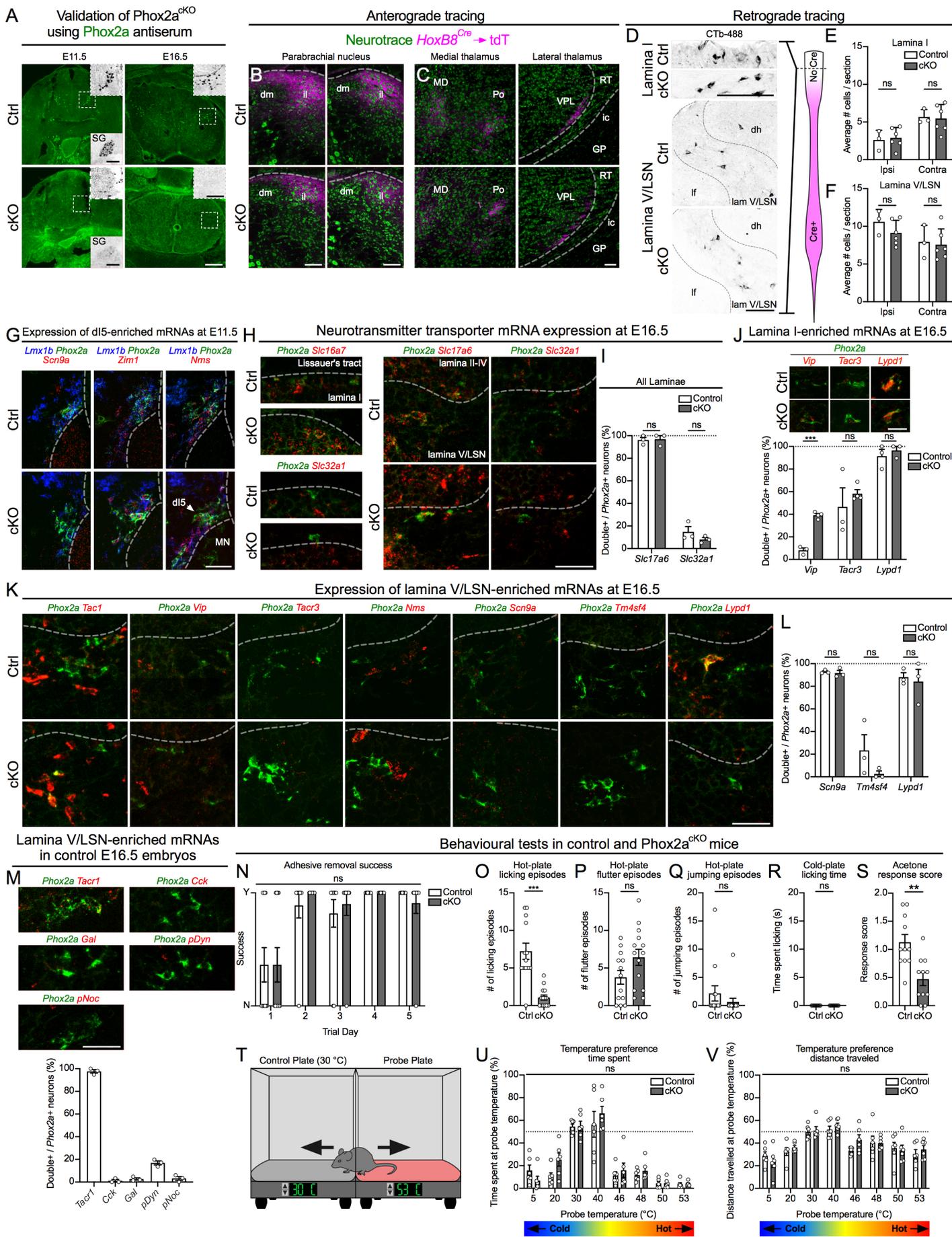


Figure S6: Phox2a is required for AS neuron development and function. Related to Figure 6.

(A) Phox2a expression in E11.5 (left) and E16.5 (right), control (top row, Ctrl) and Phox2a^{ckO} (bottom row; ckO) spinal cords. Insets at the top right are magnified boxed areas, depicting the loss of Phox2a expression (black) in ckO. Bottom right inset shows loss of Phox2a expression in sympathetic ganglia. (B, C) Brains of *HoxB8^{Cre}; Phox2a^{+/+}; R26^{LSL-tdT/+}* (Ctrl, top row) and *HoxB8^{Cre}; Phox2a^{ff}; R26^{LSL-tdT/+}* (Phox2a^{ckO} or ckO, bottom row) mice, in which spinofugal axons are labelled via *HoxB8^{Cre}*-driven axonal tdT (magenta). (B) Brain sections from two different animals per genotype show decreased Pb innervation in the parabrachial nucleus of ckO mice (as in Fig. 6). (C) Normal innervation of the medial (left) and lateral (right) thalamus was noted in both control and ckO mice. (D–F) CTb-labeled neurons in lamina I (upper panels, quantified in E) and lamina V/LSN (lower panels, quantified in F), in the cervical spinal cord of control and Phox2a^{ckO} mice, rostral to *HoxB8^{Cre}* expression domain, diagrammed in magenta. (G) Expression of dl5-enriched mRNAs in E11.5 control (top row) and Phox2a^{ckO} (bottom row) spinal cords not shown in Fig. 6 but quantified in Fig. 6G. (H) Expression of *Slc17a6* and *Slc32a1* mRNAs encoding, respectively, the neurotransmitter transporters vGlut2 and vGAT, in lamina I and lamina V/LSN of E16.5 control and Phox2a^{ckO} embryos. (I) Percent of control and Phox2a^{ckO} *Phox2a*⁺ neurons expressing *Slc17a6* and *Slc32a1* mRNAs (n=3 control, n=3 Phox2a^{ckO}). (J) Expression and quantification of candidate AS-enriched mRNAs in lamina I *Phox2a*⁺ neurons of E16.5 control and Phox2a^{ckO} embryos (n=3 control, n=3 Phox2a^{ckO}). (K) Expression of mRNAs encoding neuropeptides and neuropeptide receptors in lamina V/LSN *Phox2a*⁺ neurons of E16.5 control (top row) and Phox2a^{ckO} (bottom row) spinal cords not shown in Fig. 6 but quantified in Fig. 6M. (L) Expression and quantification of mRNAs encoding neuropeptides expressed in lamina V/LSN *Phox2a*⁺ neurons, not depicted in Fig. 6L. (M) Expression and quantification of mRNAs encoding neuropeptides and neuropeptide receptors in *Phox2a*⁺ neurons of control E16.5 embryos – high co-expression of *Phox2a* and *Tacr1* (a known AS marker) is noted. (N) Adhesive removal success ratio between control and Phox2a^{ckO} mice over 5 days. (O–S) Responses to non-injected noxious stimuli: episodes of (O) licking, (P) hind paw flutter, and (Q) jumping during the hot-plate test (n=13 control, n=14 Phox2a^{ckO}) – though jumping frequency was not significantly different between groups, 4/13 control mice exhibited jumping versus 1/14 ckO mice. (R) Time spent licking during the cold-plate test (n=9 control, n=10 Phox2a^{ckO}). (S) The acetone response score after hind paw application of acetone (n=6 control, n=6 Phox2a^{ckO}). (T–V) The two-choice temperature preference assay (T), demonstrating time spent at the probe temperature versus 30 °C (U), and the distance traveled in the probe temperature compartment versus 30 °C (V) (n=6 control, n=6 Phox2a^{ckO}).

Numbers: (A) n=3 control, n=3 Phox2a^{ckO} E11.5 and E16.5 mice. (B, C) n=4 control, n=4 Phox2a^{ckO} adult mice; (D–F) n=3 control, n=6 Phox2a^{ckO} adult mice; (G) n=3 control, n=5 Phox2a^{ckO} E11.5 mice; (H, I) n=3 control, n=3 Phox2a^{ckO} E16.5 mice; (J) n=3 control, n=4 Phox2a^{ckO} E16.5 mice; (K) n=3 control, n=3 Phox2a^{ckO} E16.5 mice; (L) n=3 control E16.5 mice; (N–V) Numbers described above.

Statistics: (E, F) Two-way ANOVA with Tukey's multiple comparisons test, (I, J, K) multiple t-tests using Holm-Sidak method, (M) mixed-effects analysis with Sidak's multiple comparisons test, (O, P, Q, R, S) Mann–Whitney test, and (U, V) two-way ANOVA with Sidak's multiple comparisons test. ns: non-significant, **: p<0.01, ***: p<0.001. Data are represented as mean ± SEM.

Scale bars: (A) 200 μm, insets in (A) and (B–D) 100 μm, (G, H, K, M) 50 μm and (J) 25 μm.

Images in Fig. 5Q have been re-used in Fig. S6G.

Abbreviations: dh (dorsal horn), lf (lateral funiculus).

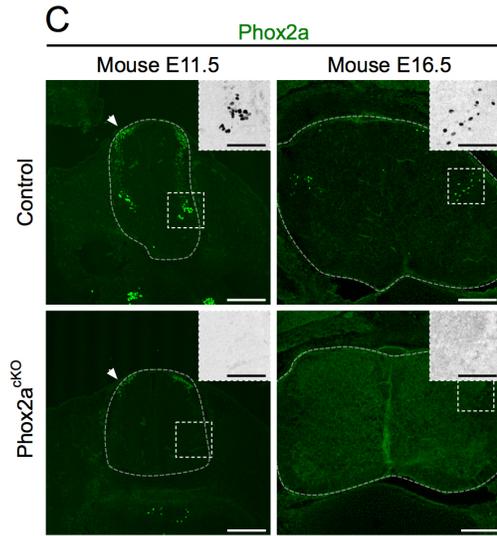
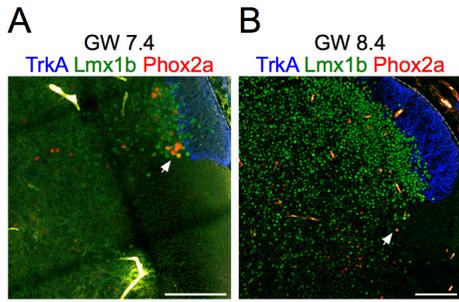


Figure S7: Phox2a neuron molecular identity is conserved in the developing human spinal cord. Related to Figure 7.

(A, B) Insets from G.W 7.4 (A) and G.W 8.4 (B) human spinal cords in Fig. 7, demonstrating Phox2a and Lmx1b co-expression, and apposition of TrkA fibres with Phox2a neurons. (C) Validation of the Abcam Phox2a antibody in mouse Phox2a^{ckO} tissue at both E11.5 and E16.5. Arrows show immunostaining in dorsal dl neurons (likely dl1), which was not observed with the Phox2a antibody used in previous experiments (from JF Brunet) or by RNA in situ detection of *Phox2a* mRNA, and it is not eliminated in the Phox2a^{ckO}. These neurons may contribute to some of the Phox2a+/Lmx1b-negative neurons observed in Fig. 7.

Numbers: (A) Single G.W 7.4 human spinal cord, (B) single G.W 8.4 human spinal cord, (C) n=3 E11.5 control, n=3 E11.5 Phox2a^{ckO}, n=3 E16.5 control, n=3 E16.5 Phox2a^{ckO} mouse embryos.

Scale bars: (A, B) and insets in (C) 100 μ m, (C) 200 μ m.

Tables:

Table S1: mRNAs enriched in embryonic AS neurons. Related to Figure 5.

Top 25 enriched mRNAs per three clusters identified in Fig. 5L and 5N, when compared to all other spinal neurons in the dataset at the same embryonic ages. For each cluster, the transcript name, log₁₀fold change (log₁₀FC) and -log₁₀p values are represented. P-values rounded to 0 (<2.22x10⁻³⁰⁸) were recorded as having a -log₁₀P value of “>307.65”. Transcripts with high -log₁₀P values that were not expressed in non-Lmx1b+ neurons were considered to be useful markers for Phox2a neurons.

E9.5-E13.5 <i>Lmx1b</i> Cluster 6			E9.5-E11.5 <i>Lmx1b</i> Cluster 2			E9.5-E11.5 <i>Lmx1b</i> Cluster 3		
Name	log ₁₀ FC	-log ₁₀ p	Name	log ₁₀ FC	-log ₁₀ p	Name	log ₁₀ FC	-log ₁₀ p
<i>Phox2a</i>	0.97	>307.65	<i>Lmx1b</i>	0.75	218.80	<i>Lmx1b</i>	0.74	270.53
<i>Nms</i>	0.44	>307.65	<i>Tac1</i>	1.16	94.60	<i>Nms</i>	0.65	177.44
<i>Syt13</i>	0.66	271.21	<i>C13002</i>	0.53	48.54	<i>Phox2a</i>	1.11	158.15
<i>Zim1</i>	0.20	176.74	<i>Pou4f2</i>	0.45	37.76	<i>C13002</i>	0.67	99.06
<i>633040</i>	0.28	146.78	<i>Gabra2</i>	0.50	29.19	<i>Car10</i>	0.64	88.30
<i>Arhgdig</i>	0.66	138.31	<i>Tlx3</i>	0.60	24.04	<i>Tm4sf4</i>	0.14	86.40
<i>Snca</i>	1.01	123.62	<i>Syt13</i>	0.42	18.00	<i>Syt13</i>	0.66	63.69
<i>Syt4</i>	2.18	120.03	<i>Chl1</i>	0.86	17.43	<i>Mar1</i>	0.49	50.42
<i>Lmx1b</i>	0.56	119.42	<i>Tcerg1l</i>	0.51	16.68	<i>Rit2</i>	0.31	48.88
<i>Scn9a</i>	0.50	112.81	<i>Mab21l2</i>	1.07	16.43	<i>Pde8b</i>	0.14	46.44
<i>Sema3</i>	0.44	107.43	<i>Pou4f1</i>	1.26	15.95	<i>Fam19a</i>	0.47	46.09
<i>Kctd4</i>	0.33	103.94	<i>Tmeff2</i>	0.57	14.85	<i>Tcerg1l</i>	0.74	41.81
<i>Sncg</i>	1.77	103.89	<i>Kif26b</i>	0.36	13.68	<i>Mdga1</i>	0.41	40.92
<i>Tac1</i>	0.48	101.61	<i>Cadm2</i>	0.40	13.52	<i>Pdzm3</i>	0.91	39.54
<i>Rit2</i>	0.45	101.60	<i>Slc17a6</i>	0.54	13.39	<i>Nrxn1</i>	1.26	38.57
<i>Car10</i>	0.50	99.29	<i>Tshz2</i>	1.03	12.96	<i>Tmeff2</i>	0.67	37.60
<i>Pcp4</i>	1.18	94.51	<i>Mab21l1</i>	0.42	12.76	<i>Negr1</i>	0.58	36.64
<i>Tcerg1l</i>	0.62	92.38	<i>Ntrk3</i>	0.50	12.00	<i>Scn3b</i>	0.65	31.99
<i>Pde8b</i>	0.12	91.23	<i>Ppp3ca</i>	0.78	11.09	<i>Syt4</i>	1.91	30.60
<i>Tspan1</i>	0.65	88.46	<i>Nrxn1</i>	0.77	10.56	<i>Stk39</i>	0.86	27.30
<i>Scn3b</i>	0.63	83.81	<i>Nms</i>	0.16	10.19	<i>Shox2</i>	0.85	26.70
<i>Adap1</i>	0.27	81.32	<i>Mtus2</i>	0.70	10.11	<i>Stk32b</i>	0.55	25.69
<i>Fam19a</i>	0.48	80.39	<i>Scn3b</i>	0.46	9.95	<i>Chl1</i>	0.88	25.24
<i>Nptxr</i>	0.28	76.58	<i>Stk39</i>	0.69	9.84	<i>Sema3a</i>	0.45	25.20
<i>Gap43</i>	1.91	76.21	<i>Hs3st2</i>	0.10	9.56	<i>Necab2</i>	0.50	24.31