Loss of Wt1 in the murine spinal cord alters interneuron composition and locomotion

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Running head: Wt1 in neurons of the spinal cord

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Abstract
Rhythmic and patterned locomotion is driven by spinal cord neurons that form neuronal circuits, referred to as central pattern generators (CPGs). Recently, dI6 neurons were suggested to participate in the control of locomotion. The dI6 neurons can be subdivided into three populations, one of which expresses the Wilms tumor suppressor gene Wt1. However, the role that Wt1 exerts on these cells is not understood. Here, we aimed to identify behavioral changes and cellular alterations in the spinal cord associated with Wt1 deletion. Locomotion analyses of mice with neuron-specific Wt1 deletion revealed that these mice ran slower than controls with a decreased stride frequency and an increased stride length. These mice showed changes in their fore-hindlimb coordination, which were accompanied by a loss of contralateral projections in the spinal cord. Neonates with Wt1 deletion displayed an increase in uncoordinated hindlimb movements and their motor neuron output was arrhythmic with a decreased frequency. The population size of dI6, V0 and V2a neurons in the developing spinal cord of conditional Wt1 mutants was significantly altered. These results show that the development of particular dI6 neurons depends on Wt1 expression and loss of Wt1 is associated with alterations in locomotion.

Introduction
In vertebrates, rhythmic activity is generated by a network of neurons, commonly referred to as central pattern generators (CPGs) (Grillner and Zangger 1979; Grillner 1985). CPGs do not require sensory input to produce rhythmical output; however, the latter is crucial for the refinement of CPG activity in response to external cues (Shik and Orlovsky 1976; Rossignol S 1988; Pearson 2003). The locomotor CPGs are located in the spinal cord and consist of distributed networks of interneurons and motor neurons (MN), which generate an organized motor rhythm during repetitive locomotor tasks like walking and swimming (Grillner 1985; McCrea and Rybak 2008).
The spinal cord develops from the caudal region of the neural tube. The interaction of secreted molecules including sonic hedgehog (Shh) and bone morphogenetic proteins (BMPs) provides instructive positional signals to the 12 progenitor cell domains that reside in the neuroepithelium (Alaynick et al., 2011). Each domain is characterized by the expression of specific transcription factor encoding genes that are used to selectively identify these populations. The dI1-dI5 interneurons are derived from dorsal progenitors and primarily contribute to sensory spinal pathways. The dI6, V0-V3 interneurons and MN arise from intermediate or ventral progenitors and are involved in the locomotor circuitry (Goulding 2009).

Whereas the involvement of V0 - V3 neurons in locomotion has been well documented, the role for dI6 neurons in locomotion has only recently been investigated (Andersson et al. 2012; Dyck et al. 2012). In particular, a part of the dI6 population shows rhythmically active neurons (Dyck et al. 2012), and a more defined subpopulation of dI6 neurons expressing the transcription factor Dmrt3, is critical for normal development of coordinated locomotion (Andersson et al. 2012). Another group of dI6 neurons is suggested to express the Wilms tumor suppressor gene Wt1 but has not yet been characterized (Goulding 2009; Andersson et al. 2012).

Wt1 encodes a zinc finger transcription factor that is inactivated in a subset of Wilms tumors, a pediatric kidney cancer (Call et al. 1990; Gessler et al. 1990). Wt1 fulfills a critical role in kidney development; however, the function of Wt1 is not limited to this organ. Phenotypic anomalies of Wt1 knockout mice can be found, among others, in the gonads, heart, spleen, retina and the olfactory system (Kreidberg et al. 1993; Herzer et al. 1999; Moore et al. 1999; Wagner et al. 2002; Wagner et al. 2005). In one of the first reports on Wt1 expression, the spinal cord was described as a prominent Wt1+ tissue (Armstrong et al. 1993; Rackley et al. 1993), however, until now there are no further reports regarding the function of Wt1 in the central nervous system (CNS).
Here, we have examined the role of Wt1 in the developing spinal cord. We performed locomotor analyses of conditional Wt1 knockout mice and used molecular biological and electrophysiological approaches to elucidate the function of Wt1 expressing neurons for locomotion. Our data suggest that Wt1 expressing dl6 neurons contribute to the coordination of locomotion and that Wt1 is needed for proper dl6 neuron specification during development.

Results

Wt1 expressing cells in the spinal cord are dl6 neurons

In order to determine the spatial and temporal pattern of Wt1 expressing cells in the spinal cord, we performed immunohistochemical analyses. Wt1+ cells were detected in the medioventral mantle zone of the developing spinal cord at embryonic day (E) 12.5 (Fig. 1A). Until E15.5, embryonic spinal cords showed a constant amount of Wt1+ cells; thereafter, their number gradually decreased until they could no longer be detected in adult mice (Fig. 1B).

We next wanted to determine the birthdate of Wt1+ cells, defined as the time point when progenitor cells cease to proliferate, leave the ventricular zone and start to differentiate. Using Bromodeoxyuridine (BrdU), the proliferative cells in the ventricular zone were labelled at different embryonic stages (E9.5, E10.5 and E11.5). Immunostaining of these cells for Wt1 at E12.5 revealed that prospective Wt1 expressing cells still proliferate at E9.5 and even at E10.5 (Fig. 1C). At E11.5 Wt1+ cells no longer showed incorporation of BrdU suggesting that they had left the ventricular zone and started their migration and differentiation in the mantel zone at this time-point.

Wt1 has been proposed to label dl6 neurons (Goulding 2009), however, the only available primary data has so far only suggested its presence in a subpopulation of dl6 neurons expressing Dmrt3 (Andersson et al., 2012). In order to closer examine the nature of Wt1+ cells, we performed immunostainings of embryonic spinal cords at
E12.5. Cells expressing Wt1 were positive for Pax2 and Lim1/2 labelling dl4, dl5, dl6, V0d and V1 neurons (Tanabe and Jessell 1996; Burrill et al. 1997) while being negative for the post-mitotic V0v marker Evx1 (Moran-Rivard et al. 2001) (Fig. 1D, E). Wt1 expression did not overlap with Lmx1b, a marker specific for dl5 neurons, but did coincide with Lbx1 (Gross et al. 2002) and Bhlhb5 (Skaggs et al. 2011), which commonly occur in the ventral most dl4-dl6 Lbx1+ domain giving rise to dl6 neurons. Thus, these data supports and extends on the previous observations that Wt1 is a marker for a subset of dl6 neurons.

**Deletion of Wt1 affects locomotor behavior**

To investigate the function of the Wt1+ neurons in the spinal cord, we made use of a Nes-Cre;Wt1fl/fl mouse line (Fig. 2A). At E12.5, no Wt1 mRNA or protein was detected in neurons from this mouse line (Fig. 2A, B). Given the location of the Wt1+ neurons within the ventral dl6 population that has been shown to be involved in regulating locomotion, we performed behavioral tests associated with locomotion to investigate potential phenotypic consequences of deleting Wt1 in spinal cord neurons. Footprints of adult mice walking on a transparent treadmill at fixed speeds (0.15, 0.25, 0.35 m/s) were recorded to analyze different gait parameters (Supplemental Fig. S1A). Nes-Cre;Wt1fl/fl mice revealed a significant reduction in stride frequency for both the fore- and hindlimbs relative to control (Wt1fl/fl) animals at all speeds measured. Heterozygous Wt1 knockout mice (Nes-Cre;Wt1fl/+ ) did not differ significantly from controls. Stride length, accordingly, was significantly longer in Nes-Cre;Wt1fl/fl animals compared to wild type mice and Nes-Cre;Wt1fl/fl. Thus, although Nes-Cre;Wt1fl/fl mice were slightly smaller compared to controls (body mass Wt1fl/fl vs Nes-Cre;Wt1fl/fl: males 33 +/- 3.9 vs 25 +/- 3.7 g; females 25 +/- 3.2 g vs 22 +/- 1.4 g; body length males 9.9 +/- 0.4 g vs 9.4 +/- 0.4 cm; females 9.9 +/- 0.4 cm vs 9.8 +/- 0.3 cm), they made longer strides with lower frequency.
To further explore gait alterations, we used X-ray fluoroscopy as a complementary method in a larger cohort of mice (Fig. 2C; Supplemental Fig. S1B; Supplemental Movie 1 and 2). When animals walked voluntarily at their preferred speed, deviations in stride frequency and stride length from the expected value (control baseline) for the given speed were again observed in Nes-Cre;Wt1fl/fl (Fig. 2D), but statistical significance is confirmed only for females. The changes were accompanied by a significant reduction of raw speed and size-corrected speed (= Froude number) in Nes-Cre;Wt1fl/fl mice of both sexes (Supplemental Fig. S1C).

While both the duration of stance and swing phase and the distance covered by the trunk and the limbs, respectively, differ between controls and Nes-Cre;Wt1fl/fl by more than 10 percent in males and more than 15 percent in females, the ratio between the two phases, expressed by the Duty factor, remains unaffected (Supplemental Fig. S1D). Thus, the temporal coordination between stance and swing phase in adult Nes-Cre;Wt1fl/fl mice is normal.

We tested whether changes in gait parameters are accompanied by changes in the phase relationships between the limbs (Fig. 2E, F). The footfall pattern of control and Nes-Cre;Wt1fl/fl females did not show significant differences at the same speed of 0.21 m/s (Supplemental Fig 1E). However, the different spread along the X-axis indicates the evenly elongated stance and swing phases.

The symmetry of left and right limb movements expressed as the time-lag between footfalls in percent stride duration of a reference limb was unaffected in the Nes-Cre;Wt1fl/fl mice (Fig. 2E,F – 1 and 2). Also, the timing of forelimb footfalls relative to the hindlimb cycles is very similar between Wt1fl/fl mice and Nes-Cre;Wt1fl/fl mice (Fig. 2E,F – 3 and 4). Significant differences between Wt1fl/fl mice and Nes-Cre;Wt1fl/fl mice were observed in the timing of the hindlimb footfalls relative to the forelimb cycles (Fig. 2E,F – 3 and 4). The touchdown of the ipsilateral and the contralateral hindlimb fall in a later fraction of the forelimb stride cycle in Nes-Cre;Wt1fl/fl mice compared to the Wt1fl/fl mice. The deviation cannot be explained by
the differences in animal speed, because the hind-to-forelimb coordination does not show speed-dependent variation.

So far, the limb kinematics of adult Nes-Cre;Wt1\textsuperscript{fl/fl} mice compared to the Wt1\textsuperscript{fl/fl} mice shows subtle differences in gait parameters and interlimb coordination with a high degree of variation. In sum, these differences result in a performance reduction indicated by the overall lower walking velocities.

**Deletion of Wt1 results in a disturbed and irregular postnatal locomotor pattern**

After having observed altered gait parameters in adult Nes-Cre;Wt1\textsuperscript{fl/fl} animals, we wondered whether gait also would be affected in younger mice. Indeed, Nes-Cre;Wt1\textsuperscript{fl/fl} pups had more difficulty coordinating their fore- and hindlimbs compared to controls when performing air stepping. Although there was no increase in hindlimb synchronous steps, left/right alternating steps were decreased and the number of uncoordinated steps were increased in Nes-Cre;Wt1\textsuperscript{fl/fl} animals (Supplemental Fig 2; Supplemental Movie 3 and 4). We next performed fictive locomotion experiments on isolated spinal cords from control and Nes-Cre;Wt1\textsuperscript{fl/fl} mice (P0-P3). Fictive locomotor drugs induced a markedly slower, disturbed, more variable pattern of locomotor-like activity in Nes-Cre;Wt1\textsuperscript{fl/fl} spinal cords (n = 6) compared to the stable, rhythmic pattern of locomotor-like activity in control mice (n = 5). Control spinal cords had recorded activity bursts that showed clear left/right (L2 vs L2) and flexor/extensor (L2 vs L5) alternation that persisted throughout activity periods, whereas activity bursts in Nes-Cre;Wt1\textsuperscript{fl/fl} spinal cords were uncoordinated and did not maintain strict left/right or flexor/extensor alternation (Fig 3A-B). The relationship between left/right and flexor/extensor alternation was examined and gave a strong phase preference for alternating bursts in control (Fig. 3C; l/r control, average phase preference: 183.4° R = 0.93; f/e control, 185.2°, R = 0.84). However, spinal cords from mice with a Wt1 deletion showed an irregular locomotor pattern with inconsistent alternation as indicated by the length and direction of the phase
vector (Fig. 3C; l/r average phase preference: 165.3°, R = 0.60; f/e 155.2°, R = 0.44). Additionally, the frequency of the ventral root output was decreased (Fig 3D: control; 0.30 ± 0.024 Hz: Nes-Cre;Wt1*/*; 0.18 ± 0.08 Hz ). This slower rhythm in Nes-Cre;Wt1*/* cords could be attributed to altered L2 and L5 activity burst parameters, as Nes-Cre;Wt1*/* mice had significantly longer burst, interburst and cycle periods compared to control (Fig. 3E, F). Thus, the deletion of Wt1 results in a disturbed and irregular locomotor pattern, which suggests that there are changes to the neuronal locomotor circuitry that occur following Wt1 deletion.

**Wt1+ neurons receive various synaptic inputs and can project commissurally**

In order to assess how Wt1+ dI6 neurons are connected within the CPG network, we focused on the innervation pattern of these cells. We used the Wt1-GFP reporter mouse line (Hosen et al. 2007) where Wt1+ neurons are labeled by GFP. In contrast to the restricted localization of Wt1 in the nucleus, GFP is distributed throughout the cytoplasm and labels the soma and major processes (Fig 4A). In combination with antibodies against particular vesicular synaptic transporters, we observed that excitatory (VGLUT2), inhibitory (VGAT) and modulatory (VMaT2) synapses contact the soma of Wt1+ dI6 neurons (Fig. 4B). This shows that Wt1+ dI6 neurons receive excitatory, inhibitory and modulatory inputs suggesting that Wt1+ neurons are positioned to receive a multitude of signals and could act during locomotion to integrate different CPG signals.

Using the Wt1-GFP reporter mouse, we found GFP+ fibers crossing the spinal cord midline beneath the central canal suggesting that Wt1+ neurons project commissural fibers (Fig. 4C). Fluorescent dextran amine retrograde tracing of contralateral projections confirmed that at least part of the Wt1+ dI6 neurons project commissurally (Supplemental Fig 3). We analyzed spinal cord commissural neurons in control (Nes-Cre;Wt1*/*) and homozygous (Nes-Cre;Wt1*/*) mice (P1-5) to determine if the deletion of Wt1 alters the total number of commissural neurons and
investigated ascending (aCIN), descending (dCIN) and bifurcating (adCIN) subpopulations (Fig. 4 D, E). All traced subpopulations were markedly reduced in *Nes-Cre;Wt1<sup>fl/fl</sup>* cords spinal cords compared to controls (Fig. 4F – H).

**Loss of Wt1 leads to altered interneuron composition**

To assess the possible impact of *Wt1* deletion for interneuron development, we analyzed dl6 and non-dl6 populations situated in the embryonic ventral spinal cord. The number of *Dmrt3* expressing cells, which constitutes a distinct but partly overlapping dl6 population (Andersson et al. 2012), was significantly decreased in the embryos harboring a loss of *Wt1* in the spinal cord already at E12.5 (Fig. 5A) persisting throughout development (E16.5 and P1). At any investigated time point, neurons co-expressing both *Wt1* and *Dmrt3* were not detected in *Nes-Cre;Wt1<sup>fl/fl</sup>* embryos and neonates.

Loss of the transcription factor *Dbx1* that is involved in differentiation of the V0 population results in a fate switch of some V0 neurons to become dl6 interneuron-like cells (Lanuza et al., 2004). Thus, we investigated whether populations flanking the dl6 population were affected in *Nes-Cre;Wt1<sup>fl/fl</sup>* mice. The Lmx1b<sup>+</sup> dl5 population was similar in number when comparing *Nes-Cre;Wt1<sup>fl/fl</sup>* with wild type embryos, whereas the number of Evx1<sup>+</sup> V0<sub>v</sub> neurons was significantly increased already at E12.5 (Fig. 5B). This increase was still detectable at E16.5. No differences could be seen in Foxp2<sup>+</sup> V1 neurons, Chx10 (V2a) and Gata3 (V2b) neurons and Islet 1/2<sup>+</sup> motor neurons between conditional *Wt1* knockout and control embryos at E12.5. However, at E16.5 Chx10<sup>+</sup> V2a neurons showed a significant decrease in cell number.

To verify the changes of interneuron composition found in the developing *Nes-Cre;Wt1<sup>fl/fl</sup>* mice, we made use of a second mouse line, namely *Lbx1-Cre;Wt1<sup>fl/fl</sup>* mice. At embryonic stage E16.5, we observed a decrease in the amount of dl6 neurons and an increase in the cell number of Evx1<sup>+</sup> neurons similar to *Nes-
Cre;Wt1<sup>fl/fl</sup> mice (Fig. 5C). This decline in the number of dI6 neurons and the concomitant increase in the amount of Evx1+ neurons might point to a change in the developmental fate from dI6 neurons into V0 neurons prompted by deletion of Wt1. To test this hypothesis, we ablated the cells destined to express Wt1. We used Lbx1-Cre;Wt1-GFP-DTA mice in which the Diphtheria toxin subunit A (DTA) is expressed from the endogenous Wt1 locus after Cre-mediated excision of a GFP cassette harboring a translational STOP-codon. Cre expression driven by the Lbx1 promoter targets the dI4 to dI6 interneuron populations (Müller et al. 2002). In Lbx1-Cre;Wt1-GFP-DTA embryos, nearly all Wt1+ neurons were ablated at E16.5 (Fig. 5D). The ablation of Wt1+ neurons coincided with a significantly decreased number of Dmrt3+ neurons in Lbx1-Cre;Wt1-GFP-DTA embryos, but did not affect the number of Evx1+ neurons (Fig. 5D). Taken together, the results from the Wt1 deletion and the ablation of the Wt1 neurons suggests that the fate switch from dI6 neurons into Evx1+ V0 neurons occurs due to the deletion of Wt1. A postnatal phenotypic behavioral analysis of these mice was not possible because neonates died immediately after birth due to serious respiratory deficits (data not shown).

The analyses of the interneuron composition in developing conditional Wt1 knockout mice and embryos with an ablation of Wt1+ neurons suggest a fate switch within a specific subset of dI6 and V0<sub>y</sub> neurons that depends on the presence of the cells destined to express Wt1.

**The transition of dI6 neurons into Evx1+ V0<sub>y</sub> neurons upon loss of Wt1 is not direct**

In order to further investigate the cellular fate change upon deletion of Wt1 we combined Wt1-GFP and Nes-Cre;Wt1<sup>fl/fl</sup> animals to generate Nes-Cre;Wt1<sup>fl/GFP</sup> mice. These mice harbor a constitutive knockout allele of Wt1 due to the insertion of a GFP coding sequence and another conditional Wt1 knockout allele. GFP and Wt1 were co-localized in the ventral spinal cord of Wt1<sup>fl/GFP</sup> control animals at E13.5, whereas
GFP, but not Wt1, was detected in spinal cords of Nes-Cre;Wt1^fl/GFP embryos of the same age (Fig 6A). Thus, Nes-Cre;Wt1^fl/GFP mice allowed us to inactivate Wt1 while the cells destined to express Wt1 are labelled by GFP.

To investigate whether Wt1 deletion leads to apoptosis in the respective cells, terminal deoxynucleotidyl transferase dUTP nick end labeling (TUNEL) was used. TUNEL+ cells were present in the ventrolateral spinal cords of Wt1^fl/GFP control and Nes-Cre;Wt1^fl/GFP embryos (Fig 6A). However, TUNEL signals never overlapped with GFP+ dl6 neurons destined to express Wt1, suggesting that Wt1 inactivation in dl6 neurons did not result in cell death.

In order to find out whether cells destined to express Wt1 would directly convert to V0_V neurons upon Wt1 inactivation, we performed immunohistochemical analyses. The presence of Dmrt3 and Evx1 in GFP+ dl6 neurons was analyzed in Wt1^fl/GFP control and Nes-Cre;Wt1^fl/GFP embryos at E12.5 (Fig 6B). The number of GFP+ cells per hemicord was determined and set to 100%. The proportion of Dmrt3+ cells was approximately 13% of all GFP+ cells in spinal cord of E12.5 control embryos. When Wt1 was absent, the amount of Dmrt3+ GFP cells significantly decreased to 4%. In contrast, the proportion of GFP+ dl6 neurons that also showed Evx1 staining was not changed between Wt1^fl/GFP control and Nes-Cre;Wt1^fl/GFP animals (below 1% for both). Thus, the increase in the amount of Evx1+ V0_V neurons observed in mice lacking Wt1, does not seem to result from a direct transition of future Wt1+ dl6 neurons into Evx1+ V0_V neurons.
Discussion

In this study, we have examined Wt1, which marks a subset of dl6 neurons. We found that Wt1 is required for proper differentiation of spinal cord neurons and that deletion of Wt1 results in locomotor aberrancies in neonate and adult mice.

Neonates lacking Wt1 in the spinal cord increased the number of uncoordinated steps, which was supported by a markedly slower and variable pattern of locomotor-like activity in isolated Nes-Cre;Wt1fl/fl spinal cords. Adult Nes-Cre;Wt1fl/fl animals showed an increased stride length and a decreased stride frequency resulting in slower absolute locomotor speed, however, these alterations were more subtle compared to locomotion abnormalities seen in neonates. Compensatory adaptation of functional properties during postnatal maturation of neuronal circuits could possibly act to reduce the severity of the deficits seen in neonates. For instance, the corticospinal tract, which allows control of the spinal circuits directly by the motor cortex and overrides the spinal circuits, does not reach the lumbar spinal cord until approximately P7-P9 (Kamiyama et al. 2015).

Wt1+ neurons are unlikely to participate in locomotor rhythm generation per se since a rhythm is established when Wt1 is deleted. We hypothesize that these neurons are involved in the maintenance or modulation of this rhythm. Adult Nes-Cre;Wt1fl/fl animals show a decreased stride frequency. The hindlimb-to-forelimb phase relationship is altered in animals with Wt1 deletion in the spinal cord. This supports a possible role of the Wt1+ dl6 neurons in both timing and limitation of the stride cycle. An involvement in timing of the stride cycle would require an integrative position in the locomotor CPGs, which is compatible with the observed multi-synaptic input to Wt1+ dl6 neurons.

The timing of hindlimb footfalls relative to forelimb footfalls differed between Wt1fl/fl and Nes-Cre;Wt1fl/fl mice, particularly at the contralateral limbs, suggesting Wt1 cells to play a role for long range coordination between various spinal cord segments. This phenotype is supported by the observation that at least a fraction of
Wt1+ dl6 neurons possess commissural projections and thus are involved in the contralateral communication between the spinal cord halves. If and how the loss of this communication affects the adult locomotor phenotype does not become apparent from our data set of footfall timing but we expect to detect more details by the analysis of intrinsic limb kinematics. Deletion of Wt1 leads to a decline in the number of commissural neurons suggesting an involvement of Wt1 in establishing proper projections of the Wt1+ dl6 neurons. Thus, it will be of future interest to investigate the transcriptome of Wt1+ dl6 neurons and screen for potential target genes involved in axon guidance.

Lack of Wt1 in the spinal cord causes alterations in the differentiation of dl6, V0 and V2a spinal cord neurons (Fig 6C). The inverse alterations in the dl6 and V0 populations suggest a fate change from dl6 to V0-like neurons when Wt1 is inactivated. The putative transition from dl6 to V0-like neurons occurs at the time-point when Wt1 expression would normally start. This instantaneous effect might be due to the derivation of both interneuron populations from neighboring progenitor domains sharing common transcription factors such as Dbx2 (Alaynick et al., 2011). Thus, loss of Wt1 might lead to a switch in developmental programs that are normally repressed; whether this repression occurs cell-autonomously or non-cell-autonomously still has to be determined. In any case, when future Wt1+ cells are ablated an increase of V0-like neurons is no longer observed, suggesting that the fate switch requires the cells about to express Wt1.

The fate change of prospective dl6 to V0-like neurons is complex since dl6 neurons can be subdivided into at least three subsets based on expression of the transcription factor encoding genes Wt1 and Dmrt3 (Fig 6C). Loss of Wt1 not only affects the small number of dl6 neurons that possess Wt1 and Dmrt3 but also the number of neurons that only express Dmrt3. This population is significantly decreased. The presence of Wt1+ neurons therefore is essential to maintain the character of a subset of Dmrt3 neurons. If Wt1 is inactivated, in addition to the cells
which are programmed to express \textit{Wt1}, possibly also Dmrt3+ neurons may differentiate into V0-like neurons.

Two main subpopulations exist within the V0 population (Alaynick \textit{et al.}, 2011): the Evx1+, more ventrally derived V0\textsubscript{v} and the Evx1 negative, more dorsally derived V0\textsubscript{d} population, for which no distinct marker has yet been described. The knockout of \textit{Dbx1} results in loss of the whole V0 population, whereby Evx1+ V0\textsubscript{v} neurons acquire a more ventral fate and become V1 neurons, whereas Evx1 negative V0\textsubscript{d} neurons acquire characteristics of dl6 neurons (Lanuza \textit{et al.}, 2004). This suggests that the V0\textsubscript{d}, rather than the V0\textsubscript{v}, neurons closer resemble the dl6 neurons and poses the question whether the fate change from Wt1+ dl6 neurons to Evx1+ V0\textsubscript{v}-like neurons represents a direct or an indirect transition. The investigations using the \textit{Nes-Cre;Wt1\textsuperscript{fl/GFP}} mice suggest that the \textit{Wt1}-deficient dl6 cells do not change their fate directly into Evx1+ V0\textsubscript{v}-like neurons suggesting an indirect transition. That points to the possibility that the fate change might be achieved by a transition of Wt1+ dl6 neurons into the more closely related Evx1 negative V0\textsubscript{d}-like neurons, which leads to a putative increase of the V0\textsubscript{d} population (Fig 6C). The Evx1+ V0\textsubscript{v} population might, in turn, increase its number to compensate for a higher proportion of V0\textsubscript{d}-like neurons.

In addition to the changes in the dl6 and V0 population that occur upon \textit{Wt1} deletion in the spinal cord, Chx10+ V2a neurons show a slight but significant decrease in their cell number at E16.5 (Fig 6C). This might represent a secondary effect of the alterations in the dl6 and V0 population, which occur already at E12.5. It was reported that V2a neurons directly innervate V0\textsubscript{v} neurons (Crone \textit{et al.}, 2008). This secondary effect might thus be due to a potential adaptation to the altered interneuron composition in the spinal cord and the necessity to form proper contacts with target cells to build up the neuronal circuits responsible for locomotion.

In sum, the results obtained in this study not only shed light on the so far undescribed necessity for Wt1 in the development of spinal cord neurons and their
functional implementation in circuits responsible for locomotion. The data also broadens our view on the complex interplay of the various neuron subpopulations within the spinal cord.
Materials and Methods

Mouse husbandry

All mice were bred and maintained in the Animal Facility of the Leibniz Institute on Aging – Fritz Lipmann Institute (FLI), Jena, Germany, according to the rules of the German Animal Welfare Law. Sex- and age-matched mice were used. Animals were housed under specific pathogen-free conditions (SPF), maintained on a 12 hour light/dark cycle, fed with mouse chow and tap water ad libitum. Mice used for analysis of fictive locomotion and projection tracing were kept according to the local guidelines of Swedish law. Wt1\(^{fl/fl}\) mice were maintained on a mixed C57B6/J x 129/Sv strain. Wt1-GFP mice (Hosen et al. 2007) were maintained on a C57B6/J strain. Conditional Wt1 knockout mice were generated by breeding Wt1\(^{fl/fl}\) females (Gebeshuber et al. 2013) to Nes-Cre;Wt1\(^{fl/fl}\) (Tronche et al. 1999) or Lbx1-Cre;Wt1\(^{fl/fl}\) mice (Sieber et al. 2007). To generate mice with Wt1 ablated cells, Wt1-GFP-DTA mice were bred with Lbx1-Cre mice. Control mice were sex- and age-matched littermates (wild type or Wt1\(^{fl/fl}\)). For plug mating analysis, females of specific genotypes were housed with males of specific genotypes and were checked every morning for the presence of a plug. For embryo analysis, pregnant mice were sacrificed by CO\(_2\) inhalation at specific time points during embryo development and embryos were dissected. Typically, female mice between 2 and 6 months were used.

Generation of Wt1-GFP-DTA mice

The Wt1-GFP-DTA mouse line bares an IRES-lox-GFP-lox-DTA cassette that was inserted into intron 3 of the Wt1 locus. This cassette consists of a GFP encoding sequence that ends in a translational STOP-codon and is flanked by loxP sites. Downstream of GFP, the coding sequence for the Diphtheria toxin subunit A (DTA) was incorporated. Before Cre-induction, the IRES cassette ensures the generation of a functional GFP protein. After Cre-mediated excision of the floxed GFP sequence, the DTA is expressed from the endogenous Wt1 promotor.
The Wt1-GFP-DTA model was generated by homologous recombination in embryonic stem (ES) cells. After ES cell screening using PCR and Southern Blot analyses, recombined ES cell clones were injected into C57BL/6J blastocysts. Injected blastocysts were re-implanted into OF1 pseudo-pregnant females and allowed to develop to term. The generation of F1 animals was performed by breeding of chimeras with wild type C57BL/6 mice to generate heterozygous mice carrying the Wt1 knockin allele.

Immunohistochemistry

Embryonic and postnatal spinal cords were dissected. They were either frozen unfixed after 15 min dehydration with 20% sucrose (in 50 % TissueTec/PBS) (post-fix) or fixed for 75 min in 4% paraformaldehyde in PBS (pre-fix). Pre-fixed tissue was cryo-protected in 10%, 20% and 30% sucrose (in PBS) before freezing in cryo-embedding medium (Neg-50 - Thermo Scientific, Kalamazoo, USA). Post- and pre-fix samples were sectioned (12 μm). Post-fixed samples were fixed for 10 min after sectioning and washed with 2% Tween in PBS (PBS-T). For pre-fixed samples, antigen retrieval was performed by incubation in sub boiling 10 mM sodium citrate buffer pH6.0 for 30 min. After blocking with 10% goat serum and 2% BSA in PBS-T (post-fix) or (prefix), sections were incubated with primary antibodies (in blocking solution) using the following dilutions: gBhlhb5 1:50 (Santa Cruz Biotechnology, Inc., Santa Cruz, California, USA), BrdU 1:100 (abcam, Cambridge, UK), shChx10 1:100 (abcam, Cambridge, UK), gpDmrt3 1:5000 (custom made (Andersson et al. 2012)), mEvx1 1:100 (1:3000 pre-fix) (Developmental Studies Hybridoma Bank, University of Iowa, Iowa City, IA), chGFP 1:1000 pre-fix (abcam, Cambridge, UK), mGFP 1:100 (Santa Cruz Biotechnology, Inc., Santa Cruz, California, USA), rFoxP2 1:800 (abcam, Cambridge, UK), mIslet1/2 1:50 (Developmental Studies Hybridoma Bank, University of Iowa, Iowa City, IA), gpLbx1 1:20,000 (gift from C. Birchmeier, MDC, Berlin, Germany), Lim1/2 1:50 (Developmental Studies Hybridoma Bank, University
NeuN, 1:500 (Merck, Darmstadt, Germany), rbPax2 1:50 (Thermo Fisher Scientific, Waltham, Massachusetts, USA), rbLmx1b 1:100 (gift from R. Witzgall, University, Regensburg, Germany), rbWt1 1:100 (Santa Cruz Biotechnology, Inc., Santa Cruz, California, USA). Secondary antibodies were applied according to species specificity of primary antibodies. Hoechst was used to stain nuclei. Quantitative analysis of the antibody staining was statistically analyzed using student t-test and two-way ANOVA followed by Tukey’s post hoc test.

Bromodeoxyuridine labeling

To label proliferating cells in the embryonic spinal cord, pregnant mice at E9.5, E10.5 and E11.5 were injected intraperitoneally with 100 µg/g of Bromodeoxyuridine (BrdU) dissolved in 0.9% sodium chloride solution. Embryos were harvested at E12.5 to isolate spinal cords and stain for BrdU and Wt1. Spinal cords were frozen unfixed after 15 min dehydration with 20% sucrose (in 50 % TissueTec/PBS) and sectioned (12 µm). After any of the following treatments, sections were washed with PBS. Antigen retrieval was performed by incubation in 98°C sub boiling 10 mM sodium citrate buffer pH6.0 for 30 min. After treatment with 2N HCl at 37°C for 30 min, sections were incubated with primary antibodies using dilutions mentioned above (Immunohistochemistry). Secondary antibodies were applied according to species specificity of primary antibodies.

RNA isolation and qRT-PCR analysis

Total RNA was isolated from E12.5 embryonic spinal cords using Trizol (Invitrogen) according to manufacturer’s protocol. Subsequently, 0.5 µg of RNA was reverse transcribed with iScript™ cDNA Synthesis Kit (Bio-Rad) and used for qRT-PCR. The primer sequences used for RT-PCR analyses are as follows: TGT TAC CAA CTG GGA CGA CA (Act_for); GGG GTG TGG AAG GTC TCA AA (Act_rev); AGT TCC CCA ACC ATT CCT TC (Wt1_qRT_for); TTC AAG CTG GGA GGT CAT
TT (WT1_qRT_rev). Real time PCR was carried out in triplicates for each sample using Syber®GreenER™ (Thermo Fisher Scientific, USA) and Bio-Rad iCycler™ (Bio-Rad). PCR efficiencies of primer pairs were calculated by linear regression method. Ct values were normalized to the mean of the reference gene, Actin. Relative expression was determined by comparing normalized Ct values of WT1 conditional knockout and control samples.

Analysis of locomotor behavior

In order to characterize gait parameters, 10 animals per sex and genotype were used. Body masses of the mice varied considerably within the groups and among the groups with significant differences between male WT1<sup>fl/fl</sup> and Nes-Cre;WT1<sup>fl/fl</sup> mice (WT1<sup>fl/fl</sup>: 28 g ± 3 g vs. Nes-Cre;WT1<sup>fl/fl</sup>: 23 g ± 3 g; F<sub>s</sub> = 31.98; t<sub>s</sub> = 3.28, P > 0.001) and moderate differences between the female WT1<sup>fl/fl</sup> and Nes-Cre;WT1<sup>fl/fl</sup> mice (WT1<sup>fl/fl</sup>: 25 g ± 5 g vs. Nes-Cre;WT1<sup>fl/fl</sup>: 22 g ± 4 g; F<sub>s</sub> = 3.80; t<sub>s</sub> = 1.62, n.s.). We recorded the voluntary walking performance of this larger cohort using high-resolution X-ray fluoroscopy (biplanar C-arm fluoroscope Neurostar, Siemens AG, Erlangen, Germany). Because of body size variation within and among groups, we adjusted treadmill speed dynamically to the individual preferences and abilities of the mice. This method of motion analysis has been described in detail in several recent publications (e.g. Böttger et al., 2011; Andrada et al., 2015; Niederschuh et al., 2015) and will be only briefly summarized here: The X-ray system operates with high-speed cameras and a maximum spatial resolution of 1536 dpi x 1024 dpi. A frame frequency of 500 Hz was used. A normal-light camera operating at the same frequency and synchronized to the X-ray fluoroscope was used to document the entire trial from the lateral perspective. Footfall sequences and spatio-temporal gait parameters were quantified by manual tracking of the paw toe tips and two landmarks on the trunk (occipital condyles, iliosacral joint) using SimiMotion 3D.
Speed, stride length, stride frequency, as well as the durations of stance and swing phases, as well as the distances that trunk or limb covered during these phases were computed from the landmark coordinates collected at touchdown and lift-off of each limb. The phase relationships between the strides of left and right limbs as well as fore- and hindlimbs were determined from footfall sequences as expression of temporal interlimb coordination. As the animals frequently accelerated or decelerated relative to the treadmill speed, the actual animal speed was obtained by offsetting trunk movement against foot movement during the stance phase of the limb. The resulting distance was divided by the duration of the stance phase. Animal speed as well as all temporal and spatial gait parameters were then scaled to body size following the formulas published by Hof (1996): non-dimensional speed = v/g \( l_0 \), where v is raw speed, g is gravitational acceleration and \( l_0 \) is the cube root of body mass as characteristic linear dimension, which scales isometrically to body mass; non-dimensional frequency = f/g \( l_0 \), where f is raw frequency; non-dimensional stride length = l/\( l_0 \), where l is raw stride length. The scaled spatio-temporal gait parameters change as a function of non-dimensional speed. Therefore, linear regression analyses were computed for each parameter in the male and the female \textit{Wt1\( ^{fl/fl} \)} group. The power formulas obtained from regression computation (Y = a + bX) were then used to calculate the expected value for a given non-dimensional speed for each gait parameter (baseline) in each animal of all four groups. The coefficient of determination \( r^2 \) was computed. The deviations of the measured values of Y from the expected values, the residuals, were determined and are given in percent of deviation. Using these residuals, one-way ANOVA were computed in order to establish the significance of the differences between the means of \textit{Wt1\( ^{fl/fl} \)} and \textit{Nes-Cre; Wt1\( ^{fl/fl} \)} in males and females. Group means were calculated from the means of 10 animals. Sample size per mouse and limb ranged between 5 and 41 stride cycles with an average sample size of 22 ± 9.
**Fictive locomotion**

Animals (P0-P3) were euthanized and the spinal cords eviscerated in ice-cold cutting solution containing (in mM): 130 K-Gluconate, 15 KCl, 0.05 EGTA, 20 HEPES, 25 Glucose, pH adjusted to 7.4 by 1M KOH) and then equilibrated in artificial cerebrospinal solution (aCSF) (Perry et al. 2015) for at least 30 minutes before the beginning of experimental procedures. Suction electrodes were attached to left and right lumbar (L) ventral roots 2 and 5 (L2 and L5). A combination of NMDA (5 µM) + 5-HT (10 µM) + dopamine (DA) (50 µM) were added to the perfusing aCSF to induce stable locomotor-like output. All chemicals were obtained from Sigma. Recorded signals containing compound action potentials were amplified 10,000 times, and band-pass filtered (100-10 kHz) before being digitized (Digidata 1322A, Axon instruments) and recorded using Axoscope 10.2 (Axon Instruments Inc.) for later off-line analysis. The data was rectified and low-pass filtered using a third-order Butterworth filter with a 5 Hz cut-off frequency before further analysis. Coherence plots between L2 and L2/L5 traces were analysed using a morlet wavelet transform in SpinalCore (Version 1.1, (Mor and Lev-Tov, 2007)). Preferential phase alignment across channels are shown in the circular plots and burst parameters were analysed for at least 20 sequential bursts, as previously described (Kiehn and Kjaerulff 1996) using an in-house designed program in Matlab (Mathworks R2014b). Ventral root recording preferential phase alignment was assessed by means of circular statistics (Rayleight test) for 20 consecutive cycles as described (Kiehn and Kjaerulff 1996). Burst parameters, including frequency, are presented as the mean ± standard deviation (SD). Burst parameters were compared using the two-tailed Mann-Whitney test or the Kruskal-Wallis analysis of variance test followed by a Dunns post-test comparing all groups.
Tracing of commissural neurons

To examine whether the loss of Wt1 affects spinal cord populations, tracing experiments were conducted as previously described (Rabe et al. 2009; Andersson et al. 2012). Nes-Cre;Wt1fl/fl and Nes-Cre;Wt1+/- littermate control mice P0-P5, were prepared as described above (Fictive locomotion). Two horizontal cuts (intersegmental tracing targeting commissural ascending/descending/bifurcating neurons) were made in the ventral spinal cord at lumbar (L) level 1 and between L4 and 5. Fluorescent dextran-amine (FDA, 3000 MW, Invitrogen) was applied at L1 and rhodamine-dextran amine (RDA, 3000 MW, Invitrogen) was applied between the L4/5 ventral roots. Spinal cords were incubated overnight at room temperature, subsequently fixed in 4% formaldehyde (FA) and stored in the dark at 4°C until transverse sectioning (60µm) on a vibratome (Leica, Germany).

Fluorescent images were acquired on a fluorescence microscope (Olympus BX61W1). For quantitative analyses of traced cords, consecutive images were taken between the two tracer application sites using Volocity software (Improvision, Lexington, USA). Captured images were auto-levelled using Adobe Photoshop software. Only cords with an intact midline, as assessed during imaging, were used for analysis.

Traced neurons in Wt1fl/fl control, Nes-Cre;Wt1fl/+ and Nes-Cre;Wt1fl/fl cords were examined for significance using the Kruskal-Wallis analysis of variance test followed by a Dunns post-test comparing all groups. Tracing data are presented as the mean ± standard error of the mean (SEM).

TUNEL-Assay

To detect apoptosis in situ, the TUNEL assay was performed prior to antibody binding. Slides were incubated with TUNEL reaction solution (1x Reaction Buffer TdT and 15 U TdT in ddH2O from Thermo Scientific; 1 mM dUTP-biotin from Roche) at 37 °C for 1 h and washed with PBS afterwards.
**Imaging and picture processing**

Fluorescent images were viewed in a Zeiss Axio Imager and a Zeiss Observer Z1 equipped with an ApoTome slider for optical sectioning (Zeiss, Germany). Images were analyzed using the ZEISS ZEN2 image analysis software. For quantitative analyses of traced spinal cords, the application sites were identified and consecutive photographs were taken between the two application sites using the OptiGrid Grid Scan Confocal Unit (Qioptiq, Rochester, USA) and Volocity software (Improvision, Lexington, USA). Confocal images were captured on a ZEISS LSM 710 ConfoCor 3 confocal microscope and analyzed using the ZEISS ZEN2 image analysis software. Captured images were adjusted for brightness and contrast using ZEN2 image analysis software and Adobe Photoshop software.

**Statistical Analyses**

Data are expressed as mean ± SD or as indicated. Groups were compared using two-way ANOVA or two-tailed two-sample equal variance student t-test as determined by group and sample size. If normal distribution of a sample was not confirmed, sample means are compared by using non-parametric Mann-Whitney U test. All statistical analyses were done using GraphPad Prism Software (GraphPad Software inc., San Diego, USA), IBM SPSS Statistics 24 (IBM Corporation, New York, USA), Microsoft Excel (Microsoft Corporation, Redmond, USA) or Matlab (Mathworks, R2014b). Normal distribution was assessed using the D’Agostino-Pearson normality test or Kolmogorov-Smirnov test. Significance was determined as * = P <0.05, ** = P<0.01, *** = P<0.001.
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Figure Legends

Figure 1. Characterization of Wt1+ neurons in the developing spinal cord. (A) Schematic illustration and Wt1 immunolabeling analysis of a transverse section (12 µm) from E12.5 spinal cord showing the position of Wt1+ neurons (red) in the mantle zone of the developing spinal cord. Stippled line represents the border between the ventricular and mantle zones. Scale bar: 50 µm. (B) Plot showing the average cell number of Wt1+ neurons per 12 µm spinal cord section from different embryonic and postnatal stages. Wt1+ neurons are first found at E12.5 and decrease in cell number postnatally. (C) Determination of the birthdate of Wt1+ neurons by Bromodeoxyuridine (BrdU) proliferation assay. Proliferating cells situated in the ventricular zone were labelled by BrdU incorporation at different embryonic stages (E9.5, E10.5 and E11.5). Additional immunolabeling of these cells for Wt1 and BrdU at E12.5 revealed that prospective Wt1 expressing cells still proliferate at E9.5 and at E10.5 but not at E11.5. Scale bar: 10 µm. Insets show higher magnifications of respective areas. Scale bar: 5 µm. (D) Schematic illustration of an E12.5 spinal cord section with markers and their occurrence in different neuron populations. These markers were used to establish the origin of Wt1+ neurons as dl6 neurons (red). (E) Immunolabeling of Wt1+ neurons with markers present in dl6 and adjacent interneuron populations. The partly overlapping location of Wt1 with Pax2, Lim1/2, Lbx1 and Bhlhb5 supports a dl6 character. Scale bar 10 µm. Insets show higher magnifications of respective areas. Scale bar: 5 µm.

Figure 2. Mice with Wt1 inactivation display altered locomotion. (A) Schematic illustration of the Wt1fl allele. loxP sites flanking exon 2 and 3 of the Wt1 coding sequence allow Cre- mediated excision and conditional knockout of Wt1. Confirmation of a functional conditional Wt1 knockout in Nes-Cre;Wt1fl/fl at E12.5 using qRT-PCR (quantification to the right). (B) Loss of Wt1 immunopositive signals in Nes-Cre;Wt1fl/fl embryos at E12.5 corroborates the loss of Wt1 protein. Schematic
illustration shows the position where pictures were taken. Stippled line represents the border between ventricular and mantle zone. Scale bar: 40 µm. (C) X-ray radiograph of a walking mouse in lateral perspective. (D) Graphs displaying stride parameters collected in the X-ray radiograph. Stride frequency is significantly lower in female *Nes-Cre;Wt1*⁰⁰⁰ mice in both forelimbs and hindlimbs. The stride length in female *Nes-Cre;Wt1*⁰⁰⁰ mice is increased compared to female *Wt1*⁰⁰⁰ mice, whereas smaller differences are found in male mice. Box plots indicate the median (bold white or black line), the 25th and the 75th percentile (box), and the data range (whiskers). Significance level of *F*: *** P < 0.001; ** P < 0.01; * P < 0.05. (E,F) Interlimb coordination expressed as the time-lag between footfall events in percent stride duration. Left limbs are reference limbs. The scheme (E) illustrates, which phase relationships are shown by which graph. Phase relationships (F) between forelimbs (1) and between hindlimbs (2) illustrate overall symmetry of the walk. Timing of forelimb touchdown relative to hindlimb touchdown for ipsilateral (3) and contralateral (4) limbs show only minor differences between *Wt1*⁰⁰⁰ and *Nes-Cre;Wt1*⁰⁰⁰ mice. The timing of hindlimb footfalls relative to forelimb footfalls (5,6) differ between *Wt1*⁰⁰⁰ and *Nes-Cre;Wt1*⁰⁰⁰ mice, particularly at the contralateral limbs. Box plots indicate the median (bold white or black line), the 25th and the 75th percentile (box), and the data range (whiskers). Significance level: *** P < 0.001; ** P < 0.01; * P < 0.05.

**Figure 3.** Locomotor activity is variable and uncoordinated in *Nes-Cre; Wt1*⁰⁰⁰ pups. (A) Representative traces showing locomotor-like activity during fictive locomotion from left and right lumbar (L) 2 and right L5 ventral roots from control (*Nes-Cre; Wt1⁺/⁺*) and *Wt1* conditional knockout (*Nes-Cre; Wt1*⁰⁰⁰) mice. Rhythmic activity was induced by application of NMDA, serotonin and dopamine. Raw traces in black; rectified, low-pass filtered signal of IL2 trace in blue; activity burst shown in green. Spinal cord schematic depicts the attached suction electrodes to the right (r) and left (l) L2 and rL5 ventral roots. Scale bar: 5 seconds. (B) Phase analysis and associated
coherence power spectra of left/right (L2/L2) and flexor/extensor (L2/L5) recordings. Regions of persistent coherence emerge for control mice at 0.30 Hz, whereas spinal cords from Nes-Cre; Wt1^{fl/fl} mice show a reduced coherency region at 0.18 Hz. Color graded scale indicates normalized coherence. Scale bar: 125 seconds. (C) Locomotor patterns, analyzed from 20 consecutive bursts, reveal impaired and variable left/right and flexor/extensor alternation in Nes-Cre; Wt1^{fl/fl} mice (black dots). Normal left/right and flexor/extensor alternation is maintained in control (white dots) mice. Each dot represents one cord; arrows represent the mean phase. The length of the vector is a measure of the statistical significance of the preferred phase; dashed grey line indicates region of high significance at 0.8 (Rayleigh test). Control, n = 5, Nes-Cre;Wt1^{fl/fl}, n=7. (D) Nes-Cre;Wt1^{fl/fl} mice have a slower locomotor frequency than control mice, p = 0.0060. (E - F) The slower locomotor frequency in Nes-Cre; Wt1^{fl/fl} mice is mirrored by an increased cycle period, burst and interburst duration in both L2 (E) and L5 (F) roots. Data expressed as mean ± SD. Significance level: ** p < 0.01, *** p < 0.001.

**Figure 4.** Innervation of Wt1+ neurons and number of commissural neurons in neonatal mice. (A) Spinal cords of Wt1-GFP embryos (stage E16.5) show a Wt1+ interneuron immunopositive for GFP. Wt1 is localized in the nucleus and GFP throughout the cell. Scale bar: 2 µm. (B) Wt1+ neurons (green) receive excitatory, inhibitory and monoaminergic synaptic contacts. Synaptic terminals are identified with synaptophysin (blue). Glutamatergic terminals were immunolabeled for VGLUT2; inhibitory synapses immunolabeled for VGAT, monoaminergic terminals immunolabeled for VMAT2. Arrows point to individual synaptic terminals (magenta) present on Wt1+ neurons (green). Boxed areas show higher magnification panels of separated channels. Scale bar: 2 µm. (C) GFP immunolabeled dl6 neurons in the spinal cord of E16.5 Wt1-GFP embryos. GFP antibody staining (green; left panel) and merge with Hoechst (blue; right panel) is shown. Boxed areas represent location
of higher magnification panels shown on the right of each panel. Contralateral projections crossing the midline (dashed line) of the spinal cord are visible (arrow heads in magnified images). Scale bar: 20 μm for overview images and 50 μm for magnified images. (D) Photomicrographs of transverse, 60μm, lumbar, spinal cord sections with applied fluorescein-dextran amine (FDA, green) and Rhodamine-dextran amine (RDA, red) tracers. Higher magnification images of wild type (Nes-Cre;Wt1<sup>+/+</sup>) and homozygous (Nes-Cre; Wt1<sup>fl/fl</sup>) segments, showing intersegmental retrograde FDA (white arrow), RDA (open arrow) and double labelled (triangle arrow) neurons. Scale bar: 200μm. (E) Schematic illustration of FDA (lumbar (L)1) and RDA (L4/5) application sites tracing descending (green), ascending (red) and bifurcating (yellow) neurons. The area of analysis (L2/3) is indicated by black dashed line. (F - H) Quantification of descending FDA labelled neurons (F), ascending RDA labelled neurons (G), and bifurcating, double labelled neurons per section (H). Descending, ascending and bifurcating CINs are significantly fewer in homozygous spinal cords compared to control cords. Data expressed as mean ± SEM. Significance level:*P < 0.05, **P < 0.01, ***P < 0.001.

Figure 5. Alterations in the composition of ventral neurons upon Wt1 knockout. (A) Average cell number of Wt1<sup>-</sup>, Dmrt3<sup>-</sup> and Wt1/Dmrt3 neurons per 12 μm spinal cord section from different embryonic and postnatal stages of control (Wt1<sup>fl/fl</sup>) and Wt1 conditional knockout (Nes-Cre;Wt1<sup>fl/fl</sup>) mice. Number of Dmrt3 neurons significantly decrease in Nes-Cre;Wt1<sup>fl/fl</sup>. No Wt1/Dmrt3 neurons are detected in Nes-Cre;Wt1<sup>fl/fl</sup> animals. (B) Average cell number of Lmx1b, Evx1, Foxp2, Chx10, Gata3 and ventral Islet1/2 neurons per 12 μm spinal cord section from control (Wt1<sup>fl/fl</sup>) and homozygous (Nes-Cre;Wt1<sup>fl/fl</sup>) mice at E12.5 and E16.5. Number of Evx1<sup>+</sup> V0 neurons is significantly increased. (C) Average cell number of Wt1<sup>-</sup>, Dmrt3<sup>-</sup> and Wt1/Dmrt3 dl6 neurons and Evx1 V0 neurons per 12 μm spinal cord section from E16.5 control (Wt1<sup>fl/fl</sup>) and Wt1 conditional knockouts (Lbx1-Cre;Wt1<sup>fl/fl</sup>) mice. Lbx1-Cre based
conditional *Wt1* knockouts show a decrease in the amount of dl6 neurons and an increase in the cell number of Evx1 neurons as for *Nes-Cre; Wt1*<sup>fl/fl</sup> animals. (D) Schematic illustration of *Wt1-GFP-DTA* allele. Cassette consisting of *loxP* sites flanking *GFP* coding sequence upstream of *Diphtheria toxin subunit A* (*DTA*) was inserted into the *Wt1* locus. This cassette allows Cre-mediated ablation of *Wt1*+ neurons. Graph shows average cell number of *Wt1-*-, *Dmrt3-* and *Wt1/Dmrt3* dl6 neurons and *Evx1+ V0* neurons per 12 µm spinal cord section from E16.5 wild type control and *Lbx1-Cre; Wt1-GFP-DTA* mice. Nearly all *Wt1*+ neurons are absent. The number of *Dmrt3* neurons is significantly decreased. Population size of *Evx1* neurons is not altered after ablation of *Wt1*+ neurons. Data expressed as mean ± SD. Significance level:*P < 0.05, **P < 0.01, ***P < 0.001.

**Figure 6.** (A) Immunofluorescence staining of spinal cord sections from E13.5 *Wt1*<sup>+/GFP</sup> (control) and *Nes-cre;Wt1*<sup>fl/GFP</sup> embryos. GFP is depicted in green, *Wt1* in red, TUNEL+ cells in white and Hoechst in blue. Orientation: dorsal at the top and ventral at the bottom. Scale bar: 50 µm. (B) Quantification of GFP+ cells harbouring the interneuron markers *Dmrt3* and *Evx1*. Analysis was performed using E12.5 *Wt1*<sup>+/GFP</sup> (control; n=3) and *Nes-cre;Wt1*<sup>fl/GFP</sup> (n=3) embryos. The number of cells showing co-localization of GFP and the respective markers was determined and normalized to the total number of GFP+ cells, which was set to 100%. Upon *Wt1* knockout the amount of GFP+ cells possessing *Dmrt3* is significantly decreased. The amount of GFP+ cells possessing *Evx1* is not altered, suggesting an indirect fate change of dl6 neurons into Evx1+ V0<sub>V</sub>-like neurons upon *Wt1* knockout. Data expressed as mean ± SD. Significance level: **P < 0.01. (C) Scheme represents various progenitor cell domains (pd5, pd6, pV0, pV1 and pV2) that give rise to different populations of spinal cord neurons (shown as circles) in wild type and tissue specific *Wt1* knockout. In wild type animals, progenitor cells leave these domains, become postmitotic and differentiate into distinct interneuron populations that further
subdivide. The dl6 interneuron population consists of neurons either positive for 
Dmrt3 (dl6_D), Wt1 (dl6_W) or both (dl6_DW). Due to the knockout of Wt1, no dl6_W and 
dl6_DW are detectable and the number of dl6_D neurons is decreased. In contrast, the 
number of Evx1+ V0_V neurons increases, which is an indirect effect as potential dl6_W 
cells that lack Wt1 did not show a Evx1 signal. This effect might be explained by a 
hypothetical fate change of dl6 neurons into V0_D like neurons (dashed light grey 
circle). The increased number of V0_D neurons would thus prompt the pV0 progenitor 
cells to differentiate preferentially into V0_V neurons, which would compensate the 
excess amount of V0_D neurons and lead to an increase in the population size of 
Evx1+ V0_V neurons. As a secondary effect, the number of V2a neurons, which 
nervate the V0_V neurons, declines at later developmental stages when neurons 
start to connect to each other potentially compensating the increased number of V0_V 
neurons. Only the subsets of interneuron populations are shown that are affected by 
the tissue specific Wt1 knockout. Red indicates decrease in population size, green 
indicates increase in population size.
A

B

C