In vitro atlas of dorsal spinal interneurons reveals Wnt signaling as a critical regulator of progenitor expansion

Graphical abstract

Highlights

- Creation of a full complement of spinal sensory interneurons from mouse ESCs
- ESC-derived and endogenous dls are transcriptionally indistinguishable
- BMP signaling regulates interneuron patterning and induces Wnt activation
- Wnt signaling regulates progenitor proliferation and expansion in vitro

Authors

Sandeep Gupta, Riki Kawaguchi, Eric Heinrichs, ..., Igor Mandric, Bennett G. Novitch, Samantha J. Butler

Correspondence
butlersj@ucla.edu (S.J.B.)

In brief

Gupta et al. develop protocols to derive a full complement of bona fide spinal sensory interneurons from mouse embryonic stem cells (mESCs). They further identify key cellular states that contribute to the interneuron diversity and identify Wnt signaling as a target to expand spinal progenitors in vitro for producing sensory interneurons in large quantities for therapeutic applications.
In vitro atlas of dorsal spinal interneurons reveals Wnt signaling as a critical regulator of progenitor expansion

Sandeep Gupta,1,2 Riki Kawaguchi,3 Eric Heinrichs,1,4 Salena Gallardo,1,5 Stephanie Castellanos,1,2,6 Igor Mandric,7 Bennett G. Novitch,1,2,8 and Samantha J. Butler1,2,8,9,*

1Department of Neurobiology, David Geffen School of Medicine, University of California, Los Angeles, Los Angeles, CA 90095, USA
2Eli and Edythe Broad Center of Regenerative Medicine and Stem Cell Research, University of California, Los Angeles, Los Angeles, CA 90095, USA
3Department of Psychiatry, David Geffen School of Medicine, University of California, Los Angeles, Los Angeles, CA 90095, USA
4Genetics and Genomics Graduate Program, University of California, Los Angeles, Los Angeles, CA 90095, USA
5Molecular Biology Interdepartmental Graduate Program, University of California, Los Angeles, Los Angeles, CA 90095, USA
6CIRM Bridges to Research Program, California State University, Northridge, Los Angeles, CA, USA
7Department of Computer Science, Samueli School of Engineering, University of California, Los Angeles, Los Angeles, CA 90095, USA
8Intellectual & Developmental Disabilities Research Center, University of California, Los Angeles, Los Angeles, CA 90095, USA
9Lead contact
*Correspondence: sandeepgupta@ucla.edu (S.G.), butlersj@ucla.edu (S.J.B.)

SUMMARY

Restoring sensation after injury or disease requires a reproducible method for generating large quantities of bona fide somatosensory interneurons. Toward this goal, we assess the mechanisms by which dorsal spinal interneurons (dIs; dI1–dI6) can be derived from mouse embryonic stem cells (mESCs). Using two developmentally relevant growth factors, retinoic acid (RA) and bone morphogenetic protein (BMP) 4, we recapitulate the complete in vivo program of dI differentiation through a neuromesodermal intermediate. Transcriptional profiling reveals that mESC-derived dIs strikingly resemble endogenous dIs, with the correct molecular and functional signatures. We further demonstrate that RA specifies dI4–dI6 fates through a default multipotent state, while the addition of BMP4 induces dI1–dI3 fates and activates Wnt signaling to enhance progenitor proliferation. Constitutively activating Wnt signaling permits the dramatic expansion of neural progenitor cultures. These cultures retain the capacity to differentiate into diverse populations of dIs, thereby providing a method of increasing neuronal yield.

INTRODUCTION

Spinal cord injuries (SCIs) can result in the loss of both coordinated movement and somatosensation (Ionta et al., 2016; Lenggenhager et al., 2012), as well as the ability to perceive the environment. Without somatosensation, patients can no longer communicate through touch, sense danger through pain, or control the functions of the gut and bladder (Furusawa et al., 2011). Although stem-cell-based cellular replacement therapies have shown promise for SCI (Antonic et al., 2013; Kong et al., 2021), the focus so far has been on the recovery of coordinated motor function (Faravelli et al., 2014; Trawczynski et al., 2019). Less attention has been paid to regaining sensory functions, despite their critical roles ensuring physical and emotional well-being, and providing modulatory feedback to the motor system through proprioceptive pathways. Somatosensory information is received peripherally and then relayed centrally by different populations of dorsal spinal interneurons (dIs) in the spinal cord. Each dI population is specialized for distinct somatosensory modalities and forms microcircuits that span different layers of the dorsal horn (Gupta and Butler, 2021; Lai et al., 2016). For example, dI1s (proprioceptors) (Lai et al., 2011), dI2s, dI3s (both mechanosensors) (Bui et al., 2013; Haimson et al., 2020), and dI6 (gait) (Perry et al., 2019) are located in the deep layers of the dorsal horn, while dI4s and dI5s regulate pain, heat, and itch in the superficial layers of the dorsal horn (Koch et al., 2018). To replace damaged spinal tissue, it will be critical to derive a complete complement of dorsal sensory interneurons that molecularly and functionally phenocopy endogenous dIs.

Toward this goal, we are working to identify the developmental mechanisms that specify sensory dIs in the spinal cord and then apply these insights to develop protocols to derive dIs from stem cells (Andrews et al., 2017; Gupta et al., 2018, 2021). Distinct classes of dIs arise during embryonic development, when growth factors, including retinoic acid (RA) and the bone morphogenetic proteins (BMPs), pattern six progenitor domains.
(dP1–dP6), which differentiate into d1–d6 (Andrews et al., 2019; Gupta and Butler, 2021). The addition of RA and BMP4 in embryoid body (EB) protocols (Wichterle et al., 2002) can induce some dIl fates in both mouse and human stem cell cultures (Andrews et al., 2017; Duval et al., 2019; Gupta et al., 2018). However, it has remained unclear how the addition of RA and BMP4 specifically induces dIl fates, and whether the dIs generated from these protocols mirror their endogenous counterparts. BMPs pattern multiple organ systems throughout development, including the non-neuronal cardiac mesoderm (Kattman et al., 2011; Ladd et al., 1998), as well as osteogenic tissues (Kawaguchi et al., 2005). The BMPs also have iterative activities both patterning dPs and controlling their proliferation to ensure that the precise number of specific dIs are generated (Andrews et al., 2012; Le Dreau et al., 2012), the downstream signals regulating the messenger complex mediates dI patterning (Hazen et al., 2011; Chi et al., 2005). The BMPs also have reiterative activities both patterning dPs and controlling their proliferation to ensure that the precise number of specific dIs are generated (Andrews et al., 2017; Ille et al., 2007). Although the Smad second messenger complex mediates dI patterning (Hazan et al., 2012; Le Dreau et al., 2012), the downstream signals regulating proliferation remain unresolved.

Here, we define RA ± BMP4 protocols that generate mouse embryonic stem cell (mESC)-derived dIs via a neuromesodermal progenitor (NMP) intermediate (Gouti et al., 2014) that faithfully mimic the normal developmental programs of the neural tube (Delile et al., 2019; Rodrigo Albors et al., 2018; Sagner et al., 2018). This approach has successfully generated a complete atlas of in-vitro-derived dIs. Transcriptional profiling of these dIs demonstrated that they possess sensory function-specific gene signatures that strikingly resemble endogenous dIs, indicating that our protocols generate bona fide dIs. Using these protocols as an in vitro model for dorsal spinal cord development, we identified the hierarchy of fate specification decisions mediated by RA and BMP signaling. We further identify Wnt/β-catenin signaling as an immediate downstream response to BMP signaling that maintains neural progenitors in a mitotic state. Remarkably, elevating Wnt signaling, using the CHIR99021 (CHIR) agonist, can dramatically extend the proliferative capacity of dPs, while preserving their ability to differentiate into specific sensory neurons on demand. Taken together, this study provides a mechanistic understanding of the developmental trajectories needed to generate the full complement of different dI identities. These insights were used to develop a protocol to expand the propagation of multipotential dPs, thereby paving the way for the production of in-vitro-derived dIs needed for both drug screening and cellular transplantation therapies.

## RESULTS

### Establishment of 2-d protocols to direct mESCs toward dorsal spinal interneuron fates

To direct dIl fates through an NMP intermediate, we adapted a mESC protocol previously used to derive ventral spinal neurons (Gouti et al., 2014; Sagner et al., 2018). In this protocol, treatment with basic fibroblast growth factor (bFGF) and the Wnt agonist CHIR (Figure 1B) results in ~90% of mESCs acquiring a bipotential NMP identity by day 3 in culture (Figure S1A) distinguished by the co-expression of both mesodermal (brachyury (T)) and neural progenitor cell (NPC; Sox2) markers (Figure S1A).

Using the developmentally relevant growth factors (Figure 1A) previously shown to direct ESCs toward dorsal spinal fates (Andrews et al., 2017; Duval et al., 2019; Gupta et al., 2018; Wilson et al., 2004), we then treated day 3 NMPs with 100 nM RA for 48 h either alone (protocol 1, Figure 1B) or in combination with 10 ng/ml BMP4 starting at either day 3 (protocol 2) or day 4 (protocol 3). Treatment with RA alone did not activate BMP signaling (distinguished by the absence of phosphorylated [p] Smad1/5/8 staining; Figure 1C) and resulted in dorsalized neural cultures, with Pax3+/Pax6+/Sox2+ neural rosettes extending Tuj1+ neurites (arrowheads, Figures 1F and 1I). When BMP4 was added in combination with RA, BMP signaling was now activated (arrowheads, Figures 1D and 1E). However, the timing of BMP4 addition critically affected NMP identity. When BMP4 was added concomitantly with RA at day 3, no Pax3+/Pax6+/Sox2+ (Figure 1G) or Sox2+ (Figure 1J) cells were observed. The aggregates exhibited Tuj1 staining but had no apparent neuronal morphology (Figure 1J). In contrast, when RA was added at day 3, followed by BMP4 at day 4, we again observed Pax3+/Pax6+/Sox2+ neural rosettes, which robustly extended Tuj1+ neurites (arrowheads, Figures 1H and 1K).

### Unbiased transcriptomic analyses identify the developmental trajectories leading to either spinal cord or cardiac fates

To assess the transcriptional changes resulting from the timed addition of RA ± BMP4, we conducted bulk RNA sequencing (RNA-seq) analyses along the timelines of the three protocols (Figure 1L). Principal-component (PC) analysis (PCA) (Son et al., 2018) revealed that PC1 and PC2 contribute 44% of the variation in the data and separate the least (D0, D3) from the most (D7, D9) differentiated samples (Figure S1B), while PC3...
contributes 15% variance and separates the samples from the different protocols (Figure S1C). A three-dimensional PCA plot combining PC1-PC3 reveals three differentiation trajectories (Figure 1M), connected by bifurcation points at days 3 and 4. The C-branch (RA + BMP4, protocol 2) follows a distinct trajectory from the R-branch (RA, protocol 1) after bifurcating at day 3, while the B-branch (RA + BMP4, protocol 3) bifurcates from the R-branch at day 4 (Figure 1M). The R- and B-branches follow similar trajectories and terminate in adjacent PCA spaces, distinct from the C-branch, suggesting that their differentiation program is largely equivalent. Supporting this conclusion, a Pearson correlation demonstrated R- and B-branch transcriptional profiles were the most similar to each other (Figure 1N).

We next examined whether the three protocols direct distinct differentiation outcomes, by assessing the groups of genes upregulated over time using weighted gene co-expression and network analysis (WGCNA) (Langfelder and Horvath, 2008). Distinct eigengene modules (Farhadian et al., 2021; Panahi et al., 2020), i.e., a collection of genes with similar expression trends, were identified for each branch (Figure S1E). Of these, the yellow-green, dark gray, and green modules show increasing expression from day 0 to 9 for protocols 1, 3, and 2, respectively (Figure 1O). A functional enrichment analysis of these modules using Enrichr (Chen et al., 2013) revealed that both the yellow-green and dark-gray gene modules were highly enriched for nervous-system-specific Gene Ontology (GO) terms (Z score > 30) (Figure 1O), supporting the hypothesis that protocols 1 and 3 promote neural identities. In contrast, the top GO term in the green module is “outflow tract septum” (Figure 1O), a structure that regulates the blood flow in the heart (Webb et al., 2003). Other cardiac-related terms, such as actomyosin, contractile actin filament, and stress fibers terms, are represented in the cellular component categories (Figure S1D). We additionally observed that multiple cardiac-specific genes are upregulated by day 9 in cells derived from protocol 2 (Figure S1F). Taken together, these analyses reveal the timeline over which RA and BMPs act sequentially to direct NPCs toward the neural lineages, rather than endocardial fates.

**Single-cell RNA-seq (scRNA-seq) of mESC-derived neurons identifies the full complement of spinal dIs**

To assess whether protocols 1 and 3 direct distinct classes of dls, we obtained single-cell transcriptomes from 9,704 cells using protocol 1 and 13,121 cells using protocol 3, taken at day 9 of the differentiation procedure (Figure 2A). We performed Louvain clustering followed by Uniform Manifold Approximation and Projection (UMAP) plots to visualize groups of transcriptionally distinct cell types (Becht et al., 2019). This pipeline identified 21 clusters of cells in both protocol 1 (Figure 2C) and protocol 3 (Figure 2D). In both conditions, ~30% of cells identify as being stressed; they cluster separately and display ~2,000 RNA counts (Figures 2C, 2D, S2A, and S2B). Approximately 2%–4% of cells are pluripotent stem cells, marked by Sox2, Pou5f1 (Oct4), and Nanog expression, while 5%–8% of cells are non-neural, either of mesodermal or cardiac neural crest identity, expressing Twist1, Runx2, and Hand2 (Han et al., 2021; Soldatov et al., 2019; Vincentz et al., 2013) (Figures 2C, 2D, S2A, and S2B). The remaining ~60% of cells have a neural identity, either Sox2+ NPCs or Tubb3+ (class III β-tubulin, Tuj1) differentiated neurons (feature plots, Figures 2C and 2D).

To distinguish the neuronal subtypes, we subclustered the Tubb3+ cells, resulting in 14 (protocol 1) and 13 (protocol 2) transcriptionally distinct clusters (Figures 2E and 2F). We then used a well-characterized panel of transcription factors (Figure 2B) to assign dl identities to these clusters (Gupta and Butler, 2021; Lai et al., 2016). This analysis revealed that protocol 1 primarily generates dl4, dl5, and dl6 that mediate pain, itch, and heat perception, while protocol 3 most notably generates dl1, dl2, and dl3 that regulate proprioception, gait, and mechanosensation. Specifically, in protocol 1, the majority of subclustered Tubb3+ cells divide into two major groups of closely aligned clusters (Figures 2B, 2E, and S2C): one group (clusters 7, 2, 0, and 1) expresses Lmx1b, Prx1, and Tbx1/3, which define dl5 identity (Lai et al., 2016); the second group (clusters 6, 12, 13, 4, 5, and 3) co-expresses Pax2 and Lhx1, which denote dl4/dl6 identity. Of these clusters, only cluster 6 also expresses Dmrt3, an established marker of dl6s (Andersson et al., 2012). The neurotransmitter profile supports these fate designations: phenocopying endogenous neurons (Figure 2B), the dl4/dl6 clusters express Gad2, a GABA-synthesizing enzyme (Pillai et al., 2007), while the dl5 clusters express S100a6 (Vglut2), a glutamate transporter specific to excitatory neurons (Figure 2E) (Cheng et al., 2004). Of the remaining four clusters, clusters 8 and 9 map to dl1 and dl2 identities (Figure S2C), while cluster 10 is enriched for neural- and neuroendocrine-specific genes, and cluster 11 is enriched for ribosomal genes (Table S1).

In protocol 3, the subclustered Tubb3+ cells divide into three groups of closely aligned clusters (Figures 2B, 2F, and
S2D): clusters 7, 1, 0, and 8 express Lhx2/9 and Barhl1/2, denoting dI1 identity; clusters 11, 5, and 6 express the dI2 markers Foxd3, Lhx1, and Lhx5, while clusters 12 and 10 express Isl1 and Tbx3, indicating dI3s. As in vivo, all three classes of in vitro-derived neurons are excitatory, expressing either Slc17a6 or Grin2b (Figure 2F). Of the remaining four clusters, clusters 2 and 9 express Atoh1 and Neurog1, respectively, which mark the dp1 and dp2 state and suggest these cells are immature dI1s and dI2s (Figure S2D); cluster 4 expresses dI4-specific markers (Figure S2D), and cluster 3 expresses multiple cytoskeletal and mitochondrial genes indicative of dying neurons (Illicic et al., 2016) (Table S1).

In summary, protocol 1 (RA) generates ~33% dI4s, 40.5% dI5s, 8% dI6s, 5% dI1s, 5% dI2s, and 8.5% unknown neural cell types (pie chart, Figure 2E). In contrast, protocol 3 (RA + BMP4) generates ~45.5% dI1s, 29% dI2s, 8% dI3s, 8% dI4s, and 9.5% unknown cell types (pie chart, Figure 2F). Both immunohistochemical and qRT-PCR analyses further support these designations (Figures S2E–S2H).

**GO analyses of ESC-derived dIs identify sensory modality-specific ontology modules**

We next assessed whether the functional identities of these in vitro-derived neurons could be predicted from the transcriptomic data. In vivo, the dI relay diverse somatosensory modalities either locally within the spinal cord (Fernandes et al., 2016; Todd, 2017) or by long-range afferent connections (Wercberger and Basbaum, 2019). Generally, the most dorsal populations regulate proprioception (dI1/dI2) (Sakai et al., 2012; Yuengert et al., 2015) and touch-induced mechanosensation (dI3s) (Bui et al., 2019), while the more intermediate classes form relay circuits for pain, itch, heat, and touch (dI4/dI5) (Koch et al., 2018; Todd, 2010). The dI6s regulate coordinated movement by forming inhibitory synapses with spinal motor neurons, while themselves receiving cholinergic and glutamatergic inputs (Perry et al., 2019).

To identify functionally relevant GOs, we selected the cluster for each dI population that highly expressed the differentiation markers (>0.2 log fold change [FC]) and subjected them to the DAVID pipeline (Huang et al., 2009a, 2009b), which identifies enriched biological processes and signaling pathways related to diseases or drugs (Figure 2G). Remarkably, each class of ESC-derived dIs displayed the relevant modality-specific signatures. For the dorsal-most neurons, we analyzed clusters 0 (dI1), 5 (dI2), and 10 (dI3) and specifically found enriched GO categories related to balance and walking behaviors, as well as excytatory synaptic signaling. Both dI1 and dI2 clusters additionally include proprioception-related terms (blue terms, Figure 2G; Table S2). For the more intermediate neurons, we analyzed clusters 4 (dI4), 1 (dI5), and 6 (dI6). The dI4/dI5 clusters include GO terms associated with pain and itch perception, specifically implicating both oxytocin, a neuropeptide that modulates pain processing (Boll et al., 2018), and β-alanine, an amino acid that induces itch (Liu et al., 2012). As in vivo, the dI4 cluster contains terms related to inhibitory synaptic signaling, while the dI5 cluster is enriched for excitatory synaptic terms (red terms, Figure 2G; Table S2). In contrast, the dI6 cluster contains terms associated with coordinated movement and synaptic signaling pathways, accurately reflecting their functional identity. Finally, we also found enriched terms in the dI clusters related to different drug addiction pathways, including amphetamine and endocannabinoids for dI1/dI2s, and amphetamine, morphine, and cocaine for dI4/dI5s (Figures 2G and S3A; Table S2). Such signatures suggest the dIs mediate the ability of psychoactive drugs to modulate pain and itch perception (Lipman and Yosipovitch, 2021) or proprioception (Downey et al., 2017). Other dl-subtype markers identified in this analysis include receptors, ion channels, and adhesion molecules (Figures S3A and S3B; Table S4).

**ESC-derived dIs are transcriptionally indistinguishable from endogenous spinal interneurons**

To assess whether the ESC-derived dIs are also transcriptionally similar to their in vivo counterparts, we used reciprocal PCA-based data integration and label transfer (Hao et al., 2021) to compare a single-cell dataset taken from embryonic stage (E) 9.5–E13.5 mouse spinal cords (Delile et al., 2019) with the in vitro-derived cell types in protocols 1 and 3. Projecting the combined in vivo and in vitro datasets into UMAP plots revealed a remarkable degree of overlap (Figures 3A–3C). This overlap was observed only for the neural progenitor and neural identities; the blood, mesoderm, neural crest, and skin lineages present in the in vivo dataset were notably absent from the in vitro datasets (Figure 3A). Although cells from protocols 1 and 3 mapped directly on top of the spinal cord cells (Figure 3B), these three datasets were distinct from trachea, lung, and kidney cells (Tabula Muris Consortium et al., 2018) (Figure 3E).

To compare the distribution of cell types in the in vivo and in vitro datasets, we performed unsupervised clustering on an
integrated dataset, yielding a total of 29 clusters (Figure 3C); 24 clusters (0–10 and 12–25) were common to all datasets, while 6 clusters were unique to the spinal cord dataset. A dot plot analysis of dI-specific markers within the shared 24 clusters showed striking similarities between the in-vivo- and in-vitro-derived datasets (Figure 3D). Although there are modest differences, there is overall concordance between the number of cells expressing a given dI marker and the strength of marker expression in the in vivo and in vitro cell clusters. This concordance continues as the neurons mature, demonstrated by the expression of neuropeptides, which modulate the intensity of sensory inputs in the adult spinal cord (Lai et al., 2016; Russ et al., 2021). Specific neuropeptides, including prepronociceptin (pnoc; dI2, dI4), neuromedin S (nms; dI4), neuropeptide Y (npy; dI4), and gastrin-releasing peptide (grp; dI1, dI5), are enriched in the same populations of dIs in both the in vitro and in vivo datasets (Figures S3C and S3D). Thus, the ESC-derived dIs appear to proceed along the same maturation process as endogenous spinal neurons.

Finally, to assess whether the in-vitro-derived dI populations specifically mirror their endogenous counterparts, we compared the protocol 1 versus protocol 3 datasets with an atlas of the neuronal classes of the spinal cord (Figure 3F), generated using the annotation provided by Delile et al. (2019). The in vivo dI1–dI3s and dI4–dI6s map to identical positions in protocol 3 (RA + BMP4) and protocol 1 (RA), respectively (Figure 3F). In contrast, there was no overlap between the ventral spinal neurons, such as the motor neurons (MNs), with the in vitro datasets (Figure 3F and data not shown). We also computationally extracted the cells expressing specific dI subtype markers from the in vitro and in vivo datasets and projected them in the same UMAP space (Figure 3G). This analysis further demonstrated that all six classes of in-vitro-derived dIs are transcriptionally indistinguishable from their spinal cord-derived counterparts.

Taken together, these studies suggest that our in vitro protocols are accurately replicating the in vivo developmental program of the spinal cord permitting the generation of bona fide sensory interneurons.

**RA activates a transcriptional network that specifies a dI4–dI6 default state**

Using the directed differentiation protocols, we investigated the unresolved mechanisms by which RA and BMP direct dI fate specification. We first assessed the genetic basis of RA-mediated fate decisions. During the 24-h pulse of RA at day 3, Pax3 levels gradually increase, whereas T expression concomitantly declines (Figure S5A). By day 4, ~90%–95% of cells are Pax3+ (Figure S5B), indicating NMPs have transitioned to a dorsal neural progenitor state (dP state1) (Figure S4A). To identify the transcriptional changes occurring during this first commitment step, we performed bulk RNA-seq at day 3 (NMP state), day 3.25 (early transcriptional response to RA), and day 4 (dP state1) (Figure S4B).

The differential expression analysis identified 138 upregulated and 19 downregulated genes after 6 h of RA treatment (Figure S4C). Upregulated genes include known RA-regulated factors, such as Meis1, Meis2, Stra8, Cyp26a1, and Rarb, confirming that cells are directly responding to RA. By 24 h of RA exposure, 389 genes were significantly upregulated, including genes present in intermediate dPs and the dI4/dI5s (Figure S4D), while 590 genes were significantly downregulated, which include mesodermal-specific genes (Figures S4E–S4G). These findings support the hypothesis that RA induces NMPs toward intermediate spinal identities, while also directing them away from mesodermal fates, thereby ensuring the transition from an NMP to Pax3+ dP state 1.

We next assessed whether the dP state 1 can be defined as a function-specific interaction network, by subjecting the upregulated genes to a Metascape protein-protein interaction (PPI) analysis (Zhou et al., 2019). At 6 h, a nascent PPI network exists, with one functional module for embryonic patterning containing the RA-target genes Meis1 and Meis2 (Figure S4H, inset). However, by 24 h, the PPI network has expanded into 13 functional modules (Figures S4I and S4C). Four modules, insulin-like signaling, pattern specification, sensory organ morphogenesis, and peptide ligand receptor, contained genes expressed in the intermediate spinal cord (insets, Figures S4I and S4C). Notably, Meis1 and Meis2 interact with Pax6, Lmx1b, Msx1, and Mbx2 in the pattern specification module. Using both qRT-PCR (Figure S5D) and transcriptional profiling of untreated day 4 cells (no RA control), or after a 24-h pulse of RA ± AGN193109, a pan-RAR inhibitor (Figure S4J), we independently validated that Meis1 and Meis2, but not Meis3, are upregulated by RA, including many patterning genes. Thus, Meis1/2 are good candidates for the key transcription factors that link RA signaling to the initiation of spinal cord patterning, i.e., dP state 1.

**Wnt signaling is induced as an immediate consequence of BMP4 treatment**

We hypothesized that the addition of BMP4 to NPCs in dP state 1 directs them to a more dorsal dP state, i.e., dP state 2 (Figure S4A). To define the transcriptional changes occurring after BMP4 addition, we performed bulk RNA-seq at day 4 (dP state 1), day 4.25 (early transcriptional response to RA + BMP4), and


day 5 (dP state 2) (Figure 4A). We identified 106 genes upregulated by 6 h and 305 genes upregulated after 24 h of BMP4 exposure (Figure S6A). Seventy-eight genes, out of 411, were common between both time points (Figures 4B and S6B). The GO analysis of these 78 common genes included the BMP signaling pathway (Figure 4C; Table S3), confirming that the analysis identified BMP-responsive genes.

In addition, we also found enriched GO terms related to Wnt signaling across three functional categories (Figure 4C). Both Wnt ligands, including Wnt1, Wnt2, Wnt3, Wnt3a, and Wnt4, and Wnt receptors, frizzled (Fzd) 8, Fzd3, Fzd10, and Lrp6 (Figure S6C), were upregulated in response to BMP4 treatment, along with Wnt signaling regulators, such as Dkk, Lmx1a (Hoekstra et al., 2013), and Bambi (Zhao et al., 2020) (Figures 4D, 4E, and S6E). Selected Wnt-signaling genes were further assessed in a qPCR analysis of day 5 NMPs to validate whether they are BMP4 responsive. We found that expression of all tested genes significantly increased after the addition of BMP4, compared with the RA alone condition (Figure 4F). The upregulation of Wnt expression was suppressed if Noggin, a BMP inhibitor, was concomitantly added with BMP4 on day 4 (Figure 4F).

Many of the upregulated Wnts, Wnt1, Wnt3a, and Wnt4, are expressed in the developing dorsal spinal cord in vivo (Agalliu et al., 2009; Daneman et al., 2009) and signal through the canonical Wnt signaling pathway (MacDonald et al., 2009), suggesting that canonical Wnt signaling mediates the ability of BMP4 to direct the dP state 1-to-dP state 2 transition.

Inhibiting canonical Wnt signaling blocks the induction of the dorsal-most fates

To assess the role of canonical Wnt signaling in dI fate specification, we added endo-IWR1 (IWR1e), a small-molecule inhibitor that blocks β-catenin function (Chen et al., 2009) to dPs from day 4 to 6 (Figure 5A). By day 9, the RA + BMP4 condition robustly induces dl1s–dl3s (Figures 5F–5I and 5N–5P). In contrast, the addition of IWR1e resulted in the loss of dl1s and dl2s (Figures 5J, 5K, 5N, and 5O), while dl3 differentiation was less affected (Figures 5L and 5P).

BMP4 directs the dorsal-most fates (dl1–dl3), while concomitantly suppressing RA-dependent intermediate dorsal fates (dl4–dl6; Figures 5E, 5I, S6D, and S6E). Supporting a role for the Wnts mediating this latter activity, there is no suppression of Pax2* dl4/dl6 and Lmx1b* dl5 fates when IWR1e is added to the dP cultures, along with RA + BMP4 (Figures 5M and 5Q). The addition of IWR1e did not erode Pax3* pan-dorsal identity or Pax7 expression, but did suppress Olig3 (Figure S6F), which may contribute to the loss of dl1/dl2 and perdurance of the dl4–dl6 fates.

Together, these results demonstrate that the BMP4-dependent dl fates require canonical Wnt/β-catenin signaling. In its absence, BMP4-mediated suppression is lost, and dPs retain their intermediate spinal cord identity.

Wnt signaling promotes BMP4-dependent fates by regulating proliferation, not patterning

Previous studies have suggested that Wnts act either as mitogens (Dickinson et al., 1994; Megasan and McMahon, 2002) or patterning factors (Alvarez-Medina et al., 2008; Muroyama et al., 2002) in the specification of the dorsal spinal cord. We assessed these models, first by determining the effect of canonical Wnts (Wnt1, Wnt2, Wnt3a, and Wnt4), a non-canonical Wnt (Wnt9b), and a small-molecule Wnt agonist (CHIR99021) on dl identity in the RA ± BMP4 protocols. When Wnts were added together with RA, during the period when NPCs are competent to respond to patterning factors, they were unable to induce the dorsal-most dl fates (Figure S7A). Similarly, when the Wnts were added concomitantly with BMP4, there was no synergistic effect on the specification of dorsal spinal identity (Figures 6A and 6B). Thus, Wnts are not sufficient to direct dl fate identity.

We next asked whether activating Wnt signaling affected the mitotic index of mESC-derived NPCs after the patterning period. To better visualize proliferation, we allowed mESCs to form EBs (Andrews et al., 2017; Duval et al., 2019; Gupta et al., 2018) and then treated them with RA ± BMP4 over the same timeline as protocols 1 and 3 (Figure 1B) to induce dorsal spinal cord patterning. EBs were then cultured with either DMSO or CHIR from day 5 to 9 (Figure 6C) and finally pulsed with 5-ethylmyl-2′-deoxyuridine (EdU) to label S-phase cells. Strikingly, there was an ∼3-fold increase in EdU+ cells in CHIR-treated EBs (Figures 6E, 6G, and 6H) compared with DMSO control (Figures 6D, 6F, and 6H) in both protocols. Similarly, the CHIR-treated EBs had an ∼2- to 4-fold increase in the number of phi-stone H3 (pH3)+ cells in M-phase, compared with control (Figures 6D–6G and 6I). Many of the dividing cells in the CHIR-treated cultures were Sox2+ NPCs (Figures 6E and 6G), while NPCs are largely depleted by day 9 in control cultures (Figures 6D and 6F). Thus, extending the window of Wnt signaling prolongs NPC proliferation without affecting their original identity, i.e., NPCs from protocol 1 expressed Pax3 and Pax7, while NPCs from protocol 3 expressed Pax3 and Olig2 (Figures 6J–6P).

Together, these studies support the hypothesis that Wnt/β-catenin signaling functions as a mitogen for dorsal spinal NPCs, rather than specifying distinct dl identities.

Wnt/β-catenin signaling can be modulated to extend the timeline of dP patterning

Finally, we assessed whether the mitogenic properties of Wnt signaling can be leveraged to expand the pool of mESC-derived NPCs. RA ± BMP4-patterned EBs were serially passaged at 1:3 dilution every 7 days in the presence of DMSO (control) or CHIR (Figure 7A). Although the number of EBs was comparable in the DMSO and CHIR conditions on the first passage, they declined sharply by the second passage in the controls (Figures 7B, 7C, 7E, 7F, 7G, 7I, 7B, and 7C). The ability of CHIR to expand the number of EBs was observed in both RA ± BMP4 conditions but was most robust for the RA + BMP4-patterned EBs (Figures 7E and 7I). CHIR-treated EBs remained Sox2+ after multiple passages, suggesting that they maintained their NPC identity (Figures 7D, 7H, 7D, and 7E).

To examine whether expanded NPCs can differentiate, we dissociated RA ± BMP4 CHIR-treated EBs at passages 6 (Figure 7J) or 7 (data not shown) onto Matrigel-coated plates and...
cultured them in the presence of CHIR, DMSO (i.e., CHIR withdrawal), or DAPT, a pro-differentiation agent (Crawford and Roelink, 2007) (Figure 7J). By day 10, while the CHIR-treated cultures remained Sox2+ (Figures 7K, 7N, and 7Q), the DMSO-treated cultures show robust induction of post-mitotic neural markers, neuronal nuclei (NeuN/Rbfox3), and Tubb3 (Figure 7Q) and contained many Tuj1+ neurites (Figures 7L and 7O). Thus, the removal of CHIR triggers spontaneous neural differentiation. The addition of DAPT did not significantly augment NeuN and Tubb3 expression compared with DMSO (Figure 7Q), although there was increased sprouting of Tuj1+ neurites (Figures 7M and 7P).
Figure 6. Wnt/β-catenin signaling mediates proliferation, not patterning, of mESC-derived spinal NPCs
(A) Timeline to assess whether Wnt signaling modulates the BMP-mediated dl fates through patterning activities.
(B) qRT-PCR analyses for dl1–dl3 marker genes show that neither Wnts (100 ng/mL) nor CHIR (5 μM) affect (one-way ANOVA) the BMP4-mediated dl fates. Expression levels were normalized to the day 0 and RA condition (n = 2).

(legend continued on next page)
We next asked whether specific dI fates are preserved in the expanded NPC pool. By day 10 (timeline in Figure 7J), passage 6/7 cultures derived from RA-patterned EBs contained many Lmx1b+ dI5s, but only a few Pax2+ dI4/dI6s, while cultures from the RA + BMP4-patterned EBs contained both Isl1+ dI3s and Lhx1b+ dI5s, but almost no Lhx2+ dI1s or Foxd3+dI2s (Figures S7I, S7J, S7K, S7L, 7M, and 7N). Thus, although some patterning information is preserved over many passages, prolonged CHIR treatment erodes its fidelity, with RA + BMP4 dPs now expressing high levels of Pax7 and low levels of Olig3, as if they have been returned to an intermediate spinal identity (Figures S7G and S7H). However, dorsal patterning information can be restored by including a pulse of RA ± BMP4 between days 3 and 5 during the differentiation of the expanded EBs on Matrigel (Figure 7J). The addition of RA resulted in an ~6-fold increase in the number of dI4/dI6s (Figures 7S and 7T), while the addition of RA + BMP4 induced ~6-fold more dI1s and ~2-fold dI2s (Figures 7V and 7X), compared with control (Figures 7R, 7T, and 7X). Unexpectedly, the pulse of RA + BMP4 suppressed the dI3 fate (Figures 7U, 7W, and 7X). This strategy was successful at restoring almost all dI identities in cultures taken from EBs both at the earliest passage and at the latest passage assessed (passage 7; Figure 7X).

Thus, these data show that the chemical activation of the canonical Wnt signaling can be successfully used to expand the size of the mESC-derived NPC population, and thereby to derive large numbers of specific dI populations.

**DISCUSSION**

Stem-cell-based directed differentiation protocols result in both a source of in-vitro-derived cell types and a model system to dissect developmental mechanisms. We have built on previous studies that derive spinal motor neurons in vitro from an NMP intermediate (Gouti et al., 2014; Sagner et al., 2018), to develop protocols that produce a complete in vitro atlas of dorsal spinal interneurons. Our transcriptomic analyses suggest that these neurons are indistinguishable, both molecularly and functionally, from their endogenous counterparts. We then used these protocols to investigate the mechanisms that regulate dI fate choices. These studies identified the hierarchy of dI fate specification and resolved the role of Wnt signaling as a mitogen in this process. We leveraged this mechanistic understanding to expand the numbers of dPs as a means of increasing dI yield.

**Hierarchy of dI fate specification**

Previous studies have shown that NMPs contribute to spinal cord and paraxial mesodermal lineages both in vivo (Attardi et al., 2018; Tzouanacou et al., 2009) and in vitro (Gouti et al., 2014). Our transcriptomic analyses revealed that the dI differentiation program conforms to the canonical Waddington model (Rajagopal and Stanger, 2016; Waddington, 1957), where NMPs are at the top of a “hill” of possible fates, and the time and sequence in which they receive RA and BMP4 signals control their differentiation into dIs versus mesodermal fates. Thus, the addition of RA at day 3 channels NMPs toward the intermediate dI fates. However, if BMP4 is added concomitantly with RA, the differentiation trajectory alters to specify endocardial lineages. BMP4 can induce only the dorsal-most dI fates after 24 h of RA treatment, suggesting that NMPs must move through an RA-induced early-competency state (dP state 1) before BMP4 can relay dorsal spinal patterning information. These studies thereby inadvertently developed a directed differentiation protocol (protocol 2) that is a starting point to derive the cardiac cell types needed to regenerate the heart (Duelen and Sampaolesi, 2017). BMP4 may direct NMPs to form lateral plate mesoderm (Row et al., 2018), which contributes multiple cell types to the embryonic heart. This finding also suggests that NMPs may have more pluripotency for mesodermal fates than previously realized. In this case, the use of alternative growth factors in protocol 2 will direct NMPs toward additional mesodermal derivatives.

Our studies suggest that RA is required to establish dP state 1, before specifying the dI4–dI6 fates. The Meis1/2 genes, which are dynamically expressed in the early spinal cord neuroepithelium (Sánchez-Guardado et al., 2011), may be central to this competency state. Meis1/2 are among the earliest genes to be induced by RA (Gouti et al., 2017; Oulad-Abdelghani et al., 1997), and they become central to a gene interaction network that contains the genes that specify intermediate spinal identity. Thus, the upregulation of Meis1/2 may drive the multipotential progenitor state (dP state 1) that differentiates into dI4–dI6. The addition of BMP4 both suppresses dI4–dI6 fates and induces another multipotent state, dP state 2, that differentiates into dI1–dI3. Interestingly, dP state 1 may be a default state, which was first suggested by earlier studies in the intermediate spinal cord (Diez Del Corral et al., 2003; Novitch et al., 2003). When RA + BMP4-patterned progenitors are expanded, they appear to revert to the default dP state 1 identity. This reversion can in turn be reversed: a pulse of RA + BMP4 remains sufficient to convert dP state 1 back to dP state 2 and elicit formation of the dorsal-most dIs.

**Extant to which in-vitro-derived dIs phenocopy endogenous sensory interneurons**

Our studies significantly extend previous studies seeking to generate dIs (Andrews et al., 2017; Duval et al., 2019; Gupta et al., 2018) by directly assessing the heterogeneity of the cultures and documenting the extent to which stem-cell-derived dIs resemble their endogenous counterparts. Using scRNA-seq analyses, we identified that our protocols generate the full...
complement of dls, that there is no heterogeneity beyond dorsal spinal differentiations. All CHIR-treated EBs maintain a Sox2+ NPC identity after passages 1 and 2 (D and H), but the expansion of RA-patterned EBs was less robust than that of RA + BMP4-patterned EBs (E and I). Bright-field images (B, C, F, and G) show neural differentiation in CHIR-expanded EBs following their dissociation. The right timeline determines the extent to which dI patterning can be restored in CHIR-treated EBs. The left timeline shows conditions to induce (J) Schematic of the timeline and experimental approaches to assay the differentiation potential of CHIR-treated EBs. The left timeline shows conditions to induce neural differentiation in CHIR-treated EBs following their dissociation. The right timeline determines the extent to which dI patterning can be restored in CHIR-treated EBs, with a 2-day pulse of RA ± BMP4. (K-Q) IHC analysis for Sox2+ NPCs and TuJ1+ neurites on passage 6 EBs (48 days) revealed that removal of CHIR induces spontaneous neural differentiation in both RA-treated (L) and RA + BMP4-treated (O) EBs. The addition of DAPT had a modest effect on the number of TuJ1+ neurites (M and P); qRT-PCR analyses confirm the upregulation of Tubb3 and NeuN in CHIR-depleted conditions. (Q) However, the number of Sox2+ cells remains unchanged. (R-X) Assessing the patterning profile of expanded EBs showed they can differentiate into dI5s (R and X; RA control) or dI3s (U; RA + BMP4 control), but not dI1s, dI2s, and dI4s (R, T, and X). A 2-day pulse of RA ± BMP4 from day 3 to 5 restored dI1 differentiation (V and X) with an ~5-fold increase in the number of Pax2+ dI4/dI6 neurons (S and X). The differentiation potential of passage (p) 1 and p6 (X) EBs did not show significant change over time, except for dI2s, where differentiation potential modestly improves over time (n = 6–10 EBs, Mann-Whitney test). Significance values: *p < 0.05, **p < 0.005, ***p < 0.0005. Scale bars: 2 mm (B, C, F, and G); 100 μm (Q-V); 2,000 μm (J-O).
cord. At present, it remains unresolved whether the pure neuronal populations (Fortin et al., 2018) will be more functionally beneficial than replacing populations of multipotential progenitors or mixtures of neurons (Kawai et al., 2021). The identification of dl-specific cell surface markers in this study (Table S4) enables the sorting of specific neural populations, which will make it possible to distinguish between these possibilities.

In summary, these studies uncover further mechanistic insights into the process of dl differentiation in vitro to derive bona fide spinal sensory neurons in the large numbers needed for clinical and drug discovery applications. They represent a critical step toward using ESC-derived dls for studying the effects of various addictive compounds and in vitro disease modeling for pain disorders.

Limitations of the study
Although this study shows that directed differentiation protocols can generate bona fide sensory modality-specific dorsal spinal interneurons, some hurdles remain. In each protocol, the dls arise as heterogeneous populations, and the yield of dl3 and dl6, the populations that regulate mechanosensation and gait, remain modest. Further transcriptomic analyses are needed to assess the mechanisms by which dp state1 resolves into dl4–dl6 and dp state2 resolves into dl1–dl3. The identification of any dl-specific regulators might permit the generation of large numbers of a specific dl population using the Chir-mediated expansion protocol. Finally, in vitro-derived dls need to be electrophysiologically assessed to determine their maturity; such an analysis requires an understanding of the electrophysiological signatures of endogenous spinal sensory neurons, which are not yet well defined.

STAR METHODS

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SUPPLEMENTAL INFORMATION

Supplemental information can be found online at https://doi.org/10.1016/j.celrep.2022.111119.

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AUTHOR CONTRIBUTIONS

S. Gupta and S.J.B. conceived the project. S. Gupta, S. Gallardo, and S.C. performed the differentiations. S. Gupta conducted bulk and single-cell RNA-seq. S. Gupta and R.K. analyzed the bulk RNA-seq data. S. Gupta, R.K., and E.H. analyzed the scRNA-seq data. This paper was written by S. Gupta and S.J.B. and edited by S. Gupta, B.G.N., and S.J.B.

DECLARATION OF INTERESTS

The authors declare no competing interests.

INCLUSION AND DIVERSITY

One or more of the authors of this paper self-identifies as underrepresented ethnic minority in science. One or more of the authors of this paper self-identifies as a member of the LGBTQ+ community. While citing references scientifically relevant for this work, we also actively worked to promote gender balance in our reference list.

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### STAR METHODS

#### KEY RESOURCES TABLE

| REAGENT or RESOURCE | SOURCE | IDENTIFIER |
|---------------------|--------|------------|
| **Antibodies**      |        |            |
| Goat polyclonal anti-Lhx2 | Santa Cruz | Sc-19344; RRID:AB_2135660 |
| Goat polyclonal anti-Isl1 | R&D Systems | AF1837, RRID:AB_2126324 |
| Rabbit polyclonal anti-Pax2 | Invitrogen | 71-6000, RRID:AB_2336046 |
| Mouse monoclonal anti-Lhx1/5 | Developmental Studies | 4F2, RRID:AB_531784 |
| Goat polyclonal anti-Pax3 | R&D Systems | AF2457, RRID:AB_416599 |
| Mouse monoclonal anti-Pax7 | Developmental Studies | AB_528428, RRID:AB_528428 |
| Rabbit monoclonal anti-phosphoSMAD1/5 | Cell Signaling Technology | 11971, RRID:AB_2797785 |
| Rabbit polyclonal anti-phospho Histone H3 | Cell signaling Technology | 9701, RRID:AB_331535 |
| Mouse monoclonal Tubulin III beta (clone-) | Covance | MMS-435P, RRID:AB_2315514 |
| Goat polyclonal anti-Sox2 | Santa Cruz | sc-17320, RRID:AB_2286684 |
| Rabbit polyclonal anti-Pax6 | MBL life science | PD-022, RRID:AB_1520876 |
| Goat polyclonal anti-T (Brachyury) | R&D Systems | AF-2085, RRID:AB_2200235 |
| Guinea pig polyclonal anti- FoxD3 sera | gift from Thomas Müller; (Müller et al., 2005) | N/A |
| Guinea pig polyclonal anti- Lmx1b sera | gift from Thomas Müller; (Müller et al., 2005) | N/A |
| Guinea pig polyclonal anti- Olig3 sera | gift from Thomas Müller; (Müller et al., 2005) | N/A |
| **Chemicals, peptides, and recombinant proteins** |        |            |
| CHIR99021 | Tocris Bioscience | 4423 |
| IWR1e | Millipore | 681669 |
| XAV939 | Stem Cell Technologies | 72674 |
| Matrigel membrane matrix | Fisher Scientific, Corning | CB-40234 |
| DAPT | Sigma Aldrich | 565784 |
| Human recombinant BMP4 | Thermo Fisher Scientific | PHC9534 |
| Retinoic acid | Sigma Aldrich | R2625 |
| Human recombinant Wnt1 | Thermo Fisher Scientific | PHC1804 |
| Human recombinant Wnt2 | Millipore | SRP-6560 |
| Human recombinant Wnt3a | R&D Systems | 5036-WN-010 |
| Mouse recombinant Wnt4 | R&D Systems | 475WN005 |
| Human recombinant Wnt5b | R&D Systems | 7347-WN |
| Mouse recombinant Wnt9b | R&D Systems | 3669-WN-025 |
| Human basic FGF (bFGF) | Thermo Fisher Scientific | PHG0023 |
| DMEM:F12 | HyClone | SH30023 |
| Neurobasal medium | Thermo Fisher Scientific | 21103049 |
| B27 supplement | Fisher Scientific | 17-504-044 |
| N2 supplement | Fisher Scientific | 17502048 |
| Mouse LIF | Millipore | ESG1107 |
| Trypsin Inhibitor 1x solution | Sigma-Aldrich | T7659 |
| Parafomaldehyde 16% | Sigma-Aldrich | 28906 |
| Phosphate buffer saline (PBS) 1X | HyClone | SH30028.02 |
| Prolong Gold | Thermo Fisher Scientific | P36930 |

(Continued on next page)
Continued

| REAGENT or RESOURCE SOURCE | IDENTIFIER |
|---------------------------|------------|
| RNEasy plus mini kit Qiagen | 74106 |
| Superscript IV First-Strand synthesis kit Thermo Fisher Scientific | 18091050 |
| Dead cell removal kit Miltenyi Biotec Inc. | 130-090-101 |
| EdU labelling assay Invitrogen | C10338 |
| Qia shredder Qiagen | 79654 |

Software and algorithms

| Corel DRAW version 20 Corel DRAW | www.coreldraw.com |
|----------------------------------|-------------------|
| Zeiss Blue lite Zeiss | https://www.zeiss.com/microscopy/us/products/microscope-software/zen-lite.html |
| Prism 9 Graphpad | https://www.graphpad.com/scientific-software/prism/ |
| Image J National Institute of Health | https://imagej.nih.gov/ij/ |
| BioRender | https://biorender.com |
| R-Package V.3.2.5 The R project for Statistical computing | https://www.r-project.org/ |
| Cell Ranger v 3.0.2 10x Genomics | https://support.10xgenomics.com/single-cell-gene-expression/software/ |
| Seurat v. 3.1.1 Satija Lab | https://satijalab.org/seurat/ |
| EnrichR Ma’ayan Lab | https://maayanlab.cloud/Enrichr/ |

Experimental model: Cell lines

| MM13 mouse embryonic stem cell line Tom Jessell Lab, Columbia University (Wichterle et al., 2002) | N/A |

Deposited data

| Bulk and single-cell RNA-Seq Gene Expression Omnibus (GEO) GEO: GSE185891 |

Oligonucleotides

| Oligonucleotides Mouse primers for qPCR analysis |
|-----------------------------------------------|
| GAPDH | Forward primer TGGTATACCTTGCCAGGTT |
| Lhx2 | Reverse primer TGTATACCTTGCCAGGTT |
| Foxd3 | CATCGTATGGACAGGACT |
| Isl1 | TTAAGGTTGGAAGGAGG |
| Lhx5 | CGGAGAGGCTGTTCAACTAC |
| Lbx1 | TGGTATACCTTGCCAGGTT |
| Lmx1b | CAGCTGAGTATGGACAGGACT |
| Meis1-spliced variant A | TTTGGTGTTCACT |
| Meis1-spliced variant B | TTCACTGGTACAGGACT |
| Meis2 | TGTGTATACCTTGCCAGGTT |
| Meis3 | CAGCTGAGTATGGACAGGACT |
| Irx1 | TTTGGTGTTCACT |
| Inr3 | CAGCTGAGTATGGACAGGACT |
| NeuN | TGTGTATACCTTGCCAGGTT |
| Tubb3 | CAGCTGAGTATGGACAGGACT |
| Sox2 | TGTGTATACCTTGCCAGGTT |
| Pax3 | CAGCTGAGTATGGACAGGACT |
| Pax7 | TGTGTATACCTTGCCAGGTT |
| Olig3 | CAGCTGAGTATGGACAGGACT |
| Wnt1 | TGTGTATACCTTGCCAGGTT |
| Wnt2 | (Continued on next page) |
RESOURCE AVAILABILITY

Lead contact
Samantha J. Butler, PhD. Email: butlersj@ucla.edu.

Materials availability
Further information regarding the resources and reagents should be directed to the lead contact- Samantha Butler, PhD (butlersj@ucla.edu).

Data and code availability
- Bulk and Single-cell RNA-seq data have been deposited at GEO and are publicly available. Accession numbers are listed in the key resources table. Microscopy data reported in this paper will be shared by the lead contact upon request.
- All original code is available from the lead contact upon request.
- Any additional information required to reanalyze the data reported in this paper is available from the lead contact upon request.

EXPERIMENTAL MODEL AND SUBJECT DETAILS

ESC maintenance and two-dimensional (2D) differentiation
The mouse ESC line, MM13, was routinely maintained in ES cell media (DMEM+20% FBS) with 100u/ml of mouse leukocyte inducing factor (LIF). Cells were maintained on mitotically inactive irradiated mouse embryonic fibroblasts (MEFs) and were passaged at least twice before starting the differentiation. To prepare cells for the differentiation, ESC colonies were dissociated with 0.25% trypsin and plated on gelatin coated plates at 1:10 dilution to reduce MEF transfer. Cells cultured on gelatin coated plates and allowed to proliferate for 1-2 days. To initiate differentiation, cells were plated on 0.1% gelatin coated 24-well CellBIND dishes (Corning) at 20,000 cells/ml density in 0.5mL/well N2/B27 medium that contains 10ng/ml basic fibroblast growth factor (bFGF). The N2/B27 medium contains 1:1 portions of Advanced Dulbecco's modified Eagle Media (DMEM) F12 (Hyclone) and Neurobasal media (Thermo Fisher Scientific) supplemented with 0.5x N2, 1x B27 supplement (with Vitamin A), 2mM L-glutamine, 0.1mM 2-mercaptoethanol (ME), and 1% BSA. On day 1, small colonies of cells can be observed attached to the bottom of the wells. On day 2, cells were supplemented with 10ng/ml bFGF and 5µM CHIR99021 for 24 h to induce a neuromesodermal identity (Gouti et al., 2014). On day 3, media was changed as follows: for protocol 1, cells were exposed to 100nM RA for 48hrs. For protocol 2, 10ng/ml human recombinant BMP4 (Thermo Fisher Scientific) was added concomitantly with 100nM RA (RA + BMP4) for 48hrs. For protocol 3, cells were exposed to 100nM RA for 24hrs, followed by a pulse of RA + BMP4 for another 24 h. To induce terminal differentiation in all three conditions, the medium containing growth factors was replaced with basic N2/B27 medium at day 5, and cell allowed to differentiate to day 9. At the end of the differentiation, the cultures were fixed for immunohistochemical analysis or lysed to obtain RNA for RNA-Seq or quantitative reverse transcriptase PCR analysis.

METHOD DETAILS

Bulk-RNA sequencing and data processing
Total cell lysate was obtained in buffer RLT (Qiagen) at different time points from three independent differentiations (biological replicates) conducted parallely. RNA extraction was performed using RNeasy mini kit (Qiagen) and the quality was determined using Agilent Technologies 2100 Bioanalyzer and only samples with a RIN score >8.0 were sequenced. Stranded libraries were constructed using Universal plus mRNA sequencing kit (NuGEN) and sequenced onto 2 lanes of NovaseqS1 to generate 30 million reads/sample. Obtained reads were aligned to the latest mouse reference genome (mm10) using the STAR spliced read aligner. Pearson correlation and principal component analysis (PCA) was performed to determine the similarities among the samples and replicates. Differentially expressed genes were identified using DESeq and EdgeR package on R platform. The candidate genes were selected using cutoff false discovery rate (FDR <0.05) and log fold change (>2). Differentially expressed gene lists were
WGCNA analysis
We used WGCNA package (Langfelder and Horvath, 2008) to identify the expression dynamics of gene modules associated with differentiations for protocol 1, 2, and 3. WGCNA was performed separately on RNA-Seq samples from protocol 1 (R-branch), 2 (C-branch), and 3 (B-branch) with day 0 and day 3 as common samples. First, differentially expressed genes were extracted for each group to determine the most variable genes. The Pearson correlation matrices were calculated for all RNA pairs in each group and were transformed into adjacency matrices using the power function. A dynamic tree-cut algorithm was used to identify gene co-expression modules where modules were defined as branches cut off from the tree and labeled in unique colors. The module eigengene (ME) represents the first principal component for each module. Gene ontologies associated with the modules were determined by supplying top 100 genes to the Enrichr platform.

Protein-protein interaction (PPI) network analysis
We used the metascape algorithm (www.metascape.org) to identify PPI networks induced by 6hrs and 24hrs of RA exposure. First, significant differentially expressed (DE) genes were extracted by pairwise comparison with the day3 dataset (day3 v/s 6hrs RA and day3 v/s 24hr RA) using log fold change > 2 and false discovery rate (FDR) < 0.01 as cut off for the upregulated genes. The entire gene list obtained from the DE analysis was then submitted to Metascape platform with Mus musculus as both input and analysis species. Both gene ontologies (GO) and PPI network were then retrieved for each gene list using the default Metascape parameters applied under the express analysis tool (minimum network size = 3, maximum network size 500).

Single-cell RNA sequencing and analysis
Preparation of single cell suspension
Day 9 cultures from both protocol1 (RA) and protocol3 (RA + BMP4) were dissociated with 0.25% cold trypsin for 5 min. Trypsin activity was stopped with the addition of trypsin inhibitor (Sigma-Aldrich) in 1:1 ratio. Cells were then pelleted by centrifugation at 1000RPM and resuspended in 1x PBS. The ratio of viable cells was determined using Trypan blue staining. Dead cells were removed using MACS dead cell removal kit (Miltenyi Biotec Inc.) by following the manufacturer’s protocol. Eluted live cells were then suspended in 1X PBS containing 0.04% BSA solution (400 μg/ml) for the library preparation.

Library preparation, and sequencing
~10,000 live cells/conditions were used to construct single-cell specific cDNA libraries using protocol described in 10x Genomics chromium single cell 3’ reagent kit (v3.1 Chemistry). Briefly, cells were partitioned into nanoliter-scale Gel-Beads-in-emulsion (GEM) using the 10x Chromium controller. Each GEM contains a unique barcode which is shared among the cDNA generated from a single cell. Cells were then lysed, and cDNA synthesis and feature barcoding were performed in the GEMs. The sequencing libraries were recovered using Magnetic separation and the quantity and quality of cDNA were assessed by Agilent 2100 expert High Sensitivity DNA Assay. cDNA samples were sequenced on 1 lane of NovaSeq 6000 S2 flow cell and reads were mapped to mouse mm10 genome using Cell Ranger v.3.0.2 to generate fastq files.

Filtering genes and cells
Seurat package v3.1.1 was used for the analysis. For each condition, genes expressing in less than 3 cells were removed and cells with >250 features were retained. PCA was used for dimension reduction (npcs = 50) with top 3000 most variable genes. Raw count data were normalized using regularized negative binomial regression with SCT transformation. Cell clustering is done using Shared Nearest Neighbor (SNN) Graph method and cluster specific markers were identified by Wilcoxon Rank-Sum test. Each cell cluster was annotated by a combination of the following methods 1) the expression of known canonical cell-type markers 2) Gene Set Enrichment Analysis (GSEA) of cluster specific markers determined by FindMarkers function in Seurat.

Reclustering of neuronal clusters
From the entire single-cell dataset, neuronal cells were first identified by the expression of pan-neuronal markers-Tubb3 and Elavl1/2. Roughly 30% cells express these markers in both protocol 1 and protocol 3 and were thus annotated as the mature neurons. The neuronal populations were then extracted using the Seurat package in R, by clustering the cells expressing Tubb3 >2 (median normalized log expression levels) using the Shared Nearest Neighbor (SNN) Graph method. This manipulation resulted in 12 clusters in protocol 1 (RA) and 13 clusters in protocol3 (RA + BMP4). Clusters were visualized with Uniform Manifold Approximation Projection (UMAP) method and the unique cluster-specific markers were identified using FindMarker function in Seurat.

GO analysis for neuronal clusters
We used DAVID functional annotation tool (Huang et al., 2009b) (https://david.ncifcrf.gov/tools.jsp) for identifying enriched biological processes and pathways associated with different neuronal clusters. For this, cluster-specific gene lists were sorted according to log fold change (FC) values and all genes with log FC > 0.2 were selected and submitted to the DAVID platform with default settings and Mus musculus as the input species. Ontological terms were then surveyed to determine if terms reflect the known functions of the dls that the given cluster belongs to. Selected GO terms were arranged in the decreasing order of -log10 (p value) and represented as bar plots.
Integration of single-cell datasets from different origins

The in vivo spinal cord single-cell dataset (Delile et al., 2019) was downloaded from https://www.ebi.ac.uk/arrayexpress/experiments/E-MTAB-7320/ and loaded into R. Gene names were standardized to match our dataset using the biomaRt function (Durinck et al., 2009). This data was then loaded into Seurat (version 4.0) and processed using the SCTransform pipeline (Hafemeister and Satija, 2019). In vivo cell type labels were simplified and transferred to the in vitro datasets. The in vivo dataset was then integrated with the in vitro protocol datasets using Seurat’s reciprocal PCA integration pipeline, and projected into a three-dimensional UMAP space. Tabula Muris reference datasets of kidney (2781 cells), lung (5449 cells) and trachea (11,269 cells) were downloaded from https://figshare.com/articles/dataset/Robject_files_for_tissues_processed_by_Seurat/5821263, loaded into R, and then processed through Seurat, as above. The three Tabula Muris datasets were then integrated with the in vitro and in vivo spinal cord datasets using Seurat’s reciprocal PCA integration pipeline and embedded into a three-dimensional UMAP space. Code for this analysis is available upon request.

**dl cell type extraction based on the marker gene expression**

Cutoffs for gene expression levels when extracting cells were initially determined through the cluster-specific violin plots from the integrated in vivo and in vitro spinal cord datasets for the SCT assay. Cutoffs were then refined by plotting the extracted cells on a UMAP plot adjusting the values to exclude non-specific cells with background levels of the genes. The cutoffs values for each gene are as follows: dI1 = Barhl1 > 1 (in vivo-417 cells, in vitro-1120 cells), dI2 = Foxd3 > 1 (in vivo-552 cells, in vitro-395 cells), dI3 = Isl1 > 1 Tbx3 >0.5 (in vivo-1669 cells, in vitro-89 cells), dI4 = Pax2 > 1 and Lhx1 > 1 (in vivo-1943 cells, in vitro-810 cells), dI5 = Lmx1b > 1 and Pou4f1 > 1 (in vivo-433 cells, in vitro-216 cells), dI6 = Dmrt3 > 1 (in vivo-73 cells, in vitro-60 cells). To extract Tubb3+ cells in the in vivo and in vitro combined dataset, expression by cluster was plotted, and the clusters expressing high levels of Tubb3 were extracted as a whole. Code is available upon request.

**Embryoid body (EB) culture, and expansion**

To initiate EB formation, dissociated embryonic stem cells were seeded in low attachment plates (Corning) at 10⁵ cells/ml density in N2/B27 medium supplemented with 10ng/ml bFGF. To induce posterior spinal cord and dorsal spinal patterning, RA and RA + BMP4 were added using the same timeline as described for the 2D differentiation, and the media was changed at every other day after day 5.

**For expansion cultures**

EBs were dissociated in single cell suspension at day 6 by treating with 0.25% Trypsin, followed by trituration to break up the larger cell clumps. Cells were pelleted and passaged in N2/B27 medium containing either DMSO or 5μM CHIR at 1:3 dilution. Media was changed at every third day during the expansion culture.

**Embryoid body (EB) differentiation**

To induce differentiation, EBs were dissociated into smaller clumps by a 5–7-min treatment with cold 0.25% Trypsin (Gibco), followed by trituration. Trypsin was neutralized with 1:1 addition of Trypsin inhibitor (Sigma-Aldrich) and cells were pelleted by centrifugation at 1000RPM. Cell pellets were resuspended in N2/B27 medium (without growth factors) and plated onto a Matrigel precoated plate. In our experience, the single cell suspension of 100-150 EBs provided enough cells to seed one 24-well plate. Spontaneous neural differentiation is induced when dissociated EBs are cultured in N2/B27 medium for 10 days. At the end of 10 days, multiple neural processes can be visualized by phase contrast microscopy and cultures were either fixed with 4% PFA for immunohistochemistry or lysed to extract RNA in RLT buffer (Qiagen).

**Immunohistochemistry**

For 2d differentiations, adherent cultures were first washed with 1xPBS and fixed with fresh cold 4% PFA for 10 min in the well. Following fixation, cultures were washed twice with 1xPBS to remove any remaining PFA. Cells were blocked with 1xPBS with 1% heat inactivated horse serum for 1 h and primary antibodies are added in the blocking solution for an overnight incubation at 4°C. Following washes, species appropriate secondary antibodies (Jackson Immuno Research Laboratories) were added in 1x PBT (PBS +0.1% Triton 20) for 1hr. The cultures were then washed with 1x PBT to remove traces of secondary antibodies and counterstained with DAPI to stain nuclei. Plates were then imaged on Zeiss LSM800 inverted confocal microscope at 10x magnification.

For EBs, EBs were collected and fixed with fresh 4% PFA for 20 min at 4°C. EBs were then equilibrated with 30% sucrose for overnight at 4C and embedded into the OCT compound for cryo-sectioning. 12um sections were collected onto positive charged slides (VWR), incubated with the antibody blocking solution for 1hr, and then stained as above. Sections were counterstained with DAPI, mounted in Prolong Gold and imaged using a Zeiss LSM800 inverted confocal microscope.

**Quantitative (q) RT-PCR analysis**

RNA was isolated with RNeasy Mini Kit (Qiagen, cat no. 74104), and converted to cDNA using Superscript IV First Strand Synthesis kit (Thermo Fisher Scientific). qRT–PCR was performed using SYBR Green PCR Master Mix (Applied Biosystems) on Roche 480 Lightcycler (Roche). The relative fold expression was calculated using 2−ΔΔCt method of comparing the expression of the target gene with that of the housekeeping gene glyceraldehyde 3-phosphate dehydrogenase (Gapdh). All experiments were repeated at least 2-3 times (technical replicates) from at least two independent differentiations (biological replicates), and data is represented as mean ± SEM. The primer sequences for all target genes are listed in the Table S5.
QUANTIFICATION AND STATISTICAL ANALYSIS

Image quantification
To count the number of nuclei, images were converted to 8bit image in ImageJ, their threshold intensity was adjusted to capture only the fluorescent nuclei, and the number of nuclei counted using the analyze particle tool in ImageJ. For each image, the area under the DAPI staining was determined using ImageJ and target cells were represented as % of cells in 100μm² DAPI+ area.

Statistics
Data are represented as mean ± SEM (standard error of the mean). Tests for statistical significance were performed using Prism software (version 9). Values of p < 0.05 were considered significant in all cases.