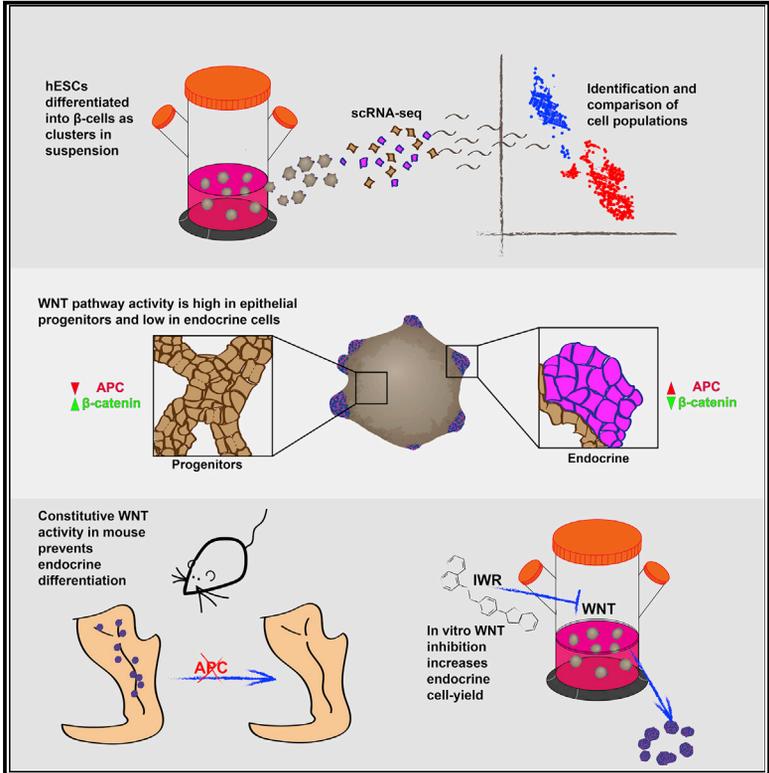


Wnt Signaling Separates the Progenitor and Endocrine Compartments during Pancreas Development

Graphical Abstract



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In Brief

In vitro differentiation of pluripotent cells into β cells is a promising alternative to cadaveric islet transplantation as a cure for type 1 diabetes. Sharon et al. use scRNA-seq to identify the cell populations that form during the process and uncover a role for WNT pathway inhibition during endocrine differentiation.

Highlights

- scRNA-seq of hESC differentiation into β cells *in vitro*
- Progenitor and endocrine compartments express different WNT pathway components
- APC deletion in mouse endocrine precursors prevents endocrine differentiation
- Small molecule inhibition of the WNT pathway *in vitro* increases endocrine yield



Wnt Signaling Separates the Progenitor and Endocrine Compartments during Pancreas Development

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SUMMARY

In vitro differentiation of pluripotent cells into β cells is a promising alternative to cadaveric-islet transplantation as a cure for type 1 diabetes (T1D). During the directed differentiation of human embryonic stem cells (hESCs) by exogenous factors, numerous genes that affect the differentiation process are turned on and off autonomously. Manipulating these reactions could increase the efficiency of differentiation and provide a more complete control over the final composition of cell populations. To uncover *in vitro* autonomous responses, we performed single-cell RNA sequencing on hESCs as they differentiate in spherical clusters. We observed that endocrine cells and their progenitors exist beside one another in separate compartments that activate distinct genetic pathways. WNT pathway inhibition in the endocrine domain of the differentiating clusters reveals a necessary role for the WNT inhibitor APC during islet formation *in vivo*. Accordingly, WNT inhibition *in vitro* causes an increase in the proportion of differentiated endocrine cells.

INTRODUCTION

Type 1 diabetes (T1D) is caused by autoimmune destruction of the insulin-producing β cells in the pancreatic islets. Transplantation of cadaveric islets can cure the disease (Shapiro et al., 2000), but donor scarcity and high cost limit its feasibility. In an attempt to develop a ready supply of β cells for transplantation, several protocols for the *in vitro* differentiation of pluripotent cells into β cells were developed in recent years (Pagliuca et al., 2014; Reznia et al., 2014; Russ et al., 2015). Our protocol directs differentiation of human embryonic stem cells (hESCs) into β cells that resemble cadaveric β cells in both gene expression and

function, including the ability to secrete insulin in response to changing glucose levels (Pagliuca et al., 2014). Still, under these *in vitro* conditions, only about 30% of the generated cells are, in fact, β cells, and finding ways to increase the efficiency of the differentiation will be valuable.

An obstacle to protocol improvement is our incomplete understanding of the complex process of β cell differentiation. During normal embryonic development, the nascent pancreas contains a network of monolayered tubules composed of epithelial progenitors, called epithelial cords (Pan and Wright, 2011). As cells in the cords divide, some turn on NEUROG3 and form “peninsulas”—bud-like structures that grow and develop to become the islets (Sharon et al., 2019). Current protocols aim to recapitulate embryonic islet development by stepwise application of defined factors. Here, we use single-cell RNA sequencing (scRNA-seq) to characterize the cell populations that appear during the *in vitro* differentiation process and identify pathways that affect β cell yield.

RESULTS

Single-Cell RNA Sequencing of Differentiating β Cells

hESCs were differentiated into stem-cell-derived β cells as clusters in suspension using a six-stage protocol (Pagliuca et al., 2014) (Figure 1A). scRNA-seq was performed on undifferentiated cells and on 10 consecutive time points, representing the end of each of the differentiation stages and select intermediate points (Figures S1A and S1B). To analyze the relationships between the cells, we combined SIMLR analysis (single-cell interpretation via multikernel learning) with topic modeling (TM). SIMLR is a method that groups cells based on cell-to-cell similarity and then displays them in lower dimensional space (Wang et al., 2017) (Figure 1B). TM is a probabilistic unsupervised learning algorithm that, in the context of gene expression analysis, identifies groups of genes that are frequently expressed together in the same cell and gathers them into “expression profiles” (EPs) (Blei, 2012; Gerber et al., 2007; Teh et al., 2006). For each EP, every gene receives a “relevance value,” which



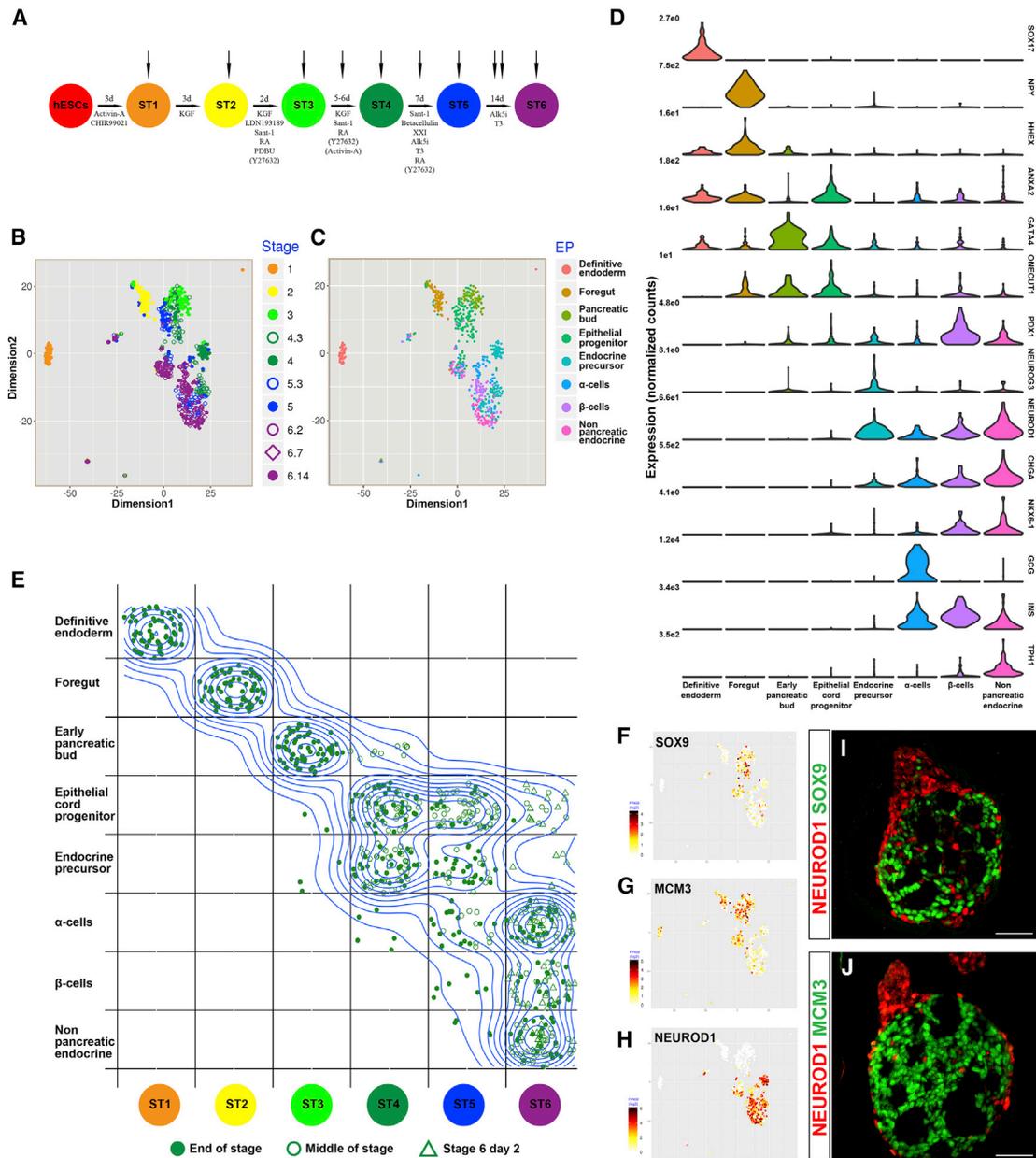


Figure 1. scRNA-Seq Analysis of the Directed Differentiation of β Cells *In Vitro*

(A) Overview of the protocol for the directed differentiation of hESCs into β cells (Pagliuca et al., 2014). Cells were collected from undifferentiated hESCs at the end of stages 1 through 5, on the 3rd day of stages 4 and 5, and on days 2, 7, and 14 of stage 6.

(B) SIMLR visualization of the 720 differentiating cells analyzed (excluding hESCs). Colors represent time of collection.

(C) SIMLR plot, with colors representing the most highly used cell-type EP.

(D) Violin plots represent gene expression distributions within the cell groups defined by cell-type EPs.

(E) A density-based topographic map is overlaid on top of a bins table to emphasize the changes that occur in the identity of the cells during *in vitro* differentiation. Cells are binned based on stage of collection (columns) and developmental identity (rows) and are randomly dispersed within each bin. For a numerical summary of the bins table, see Figure S1C.

(F–H) SIMLR map overlaid with the relative expression level of the indicated gene in each cell. (F) SOX9; (G) MCM3; and (H) NEUROD1.

(I and J) Immunofluorescent staining of clusters at the end of stage 4: (I) NEUROD1 and SOX9; (J) NEUROD1 and MCM3. Scale bars: 50 μ m.

describes the gene’s weight in the identification of this particular EP. While establishing which genes constitute an EP, the TM algorithm simultaneously quantifies how active each EP is within

a particular cell via a “usage value.” Cells that tend to use genes from the same EPs (have high usage values for similar EPs) can be grouped together. Whereas customarily used clustering

methods, such as hierarchical clustering, assume that the relationships between genes are strict (e.g., Euclidean distance, correlation), TM analyzes these relationships as probability distributions. This allows the clustering of cells and genes in a flexible arrangement. Rather than forcing each gene to one expression module, with TM, a gene can be relevant to several EPs, reflecting its possible expression in the context of different biological processes. Similarly, since each cell uses several biological processes, a single cell may use several EPs, to varying extents. Furthermore, since conventional clustering methods allow a gene to belong only to a single expression module, many genes can be lost to artificial modules caused by technical noise. However, the inherent flexibility of TM allows these genes to appear in biologically meaningful EPs as well. Altogether, the advantages of TM analysis over conventional clustering methods are especially relevant for discovering hidden structures in highly complex datasets, including scRNA-seq of heterogeneous populations.

TM identified 36 EPs in the data (Figure S2). Of these, 8 EPs contain markers related to cell-type identity and, combined, are used by all differentiating cells (excluding undifferentiated hESCs). The identity of each cell could now be determined according to the EP it uses most (Figures 1C and 1D; Figures S1C and S1F). Cells collected in the first three stages correspond with the first three cell types, whereas cells collected at the 4th stage and beyond are heterogeneous and can be divided into five cell types (Figure 1E). An interactive database that combines SIMLR with TM and describes the major cell populations and their relative similarity to each other can be found at <https://ifx.rc.fas.harvard.edu/invitrobetacells/>.

Early *In Vitro* Differentiation Is Linear and Uniform, whereas Late Differentiation Gives Rise to Heterogeneous Populations

At the end of the first stage, marked by SOX17 expression (Pagliuca et al., 2014), all cells analyzed primarily use an EP comprising other definitive endoderm genes, including CER1 and LEFTY2. By the end of the second stage, cells primarily use an EP corresponding to foregut endoderm (HHEX, TTR) (Figures 1D and 1E; Figures S1C and S1F; Table S1) (Bort et al., 2004; Thomas et al., 1998). Almost all cells analyzed at the end of the third stage, when PDX1 is activated (Pagliuca et al., 2014), use an EP that contains additional pancreatic-bud markers such as GATA4, ONECUT1, and ONECUT2 (Figures 1D and 1E; Figure S1F; Table S1) (Decker et al., 2006; Jacquemin et al., 2003). The absence of NKX6.1 and expression of the foregut markers HOXA1, SHISA2, and HHEX suggest that these cells represent an early stage in pancreatic-bud development (Figure 1D; Table S1) (Filipe et al., 2006; Godwin et al., 1998; Jennings et al., 2013; Rodríguez-Seguel et al., 2013). Unlike the first two EPs, the early pancreatic-bud EP is also used by cells from later stages, albeit with low usage values and not as the primary EP (Figure 1E; Figures S3A and S3B). This suggests that many of the genes expressed at this point remain activated in later stages of development.

From stage 4 onward, the clusters contain heterogeneous populations that can be divided into two major cell groups (Figure 1E). The cells in one group express genes from an

EP containing epithelial-cord markers, including SOX9 and ONECUT1 (HNF6), together with cell-cycle-related genes such as MCM3 and CCND2 (Figures 1E–1G; Table S1). A second group expresses genes from several EPs containing endocrine markers including CHGA, ISL1, and NEUROD1 (Figures 1D, 1E, and 1H; Table S1). The latter group also expresses lower levels of genes associated with cell proliferation (Figures 1G and 1H).

It was previously shown that treatment with Activin A and the ROCK inhibitor Y-27632 during stage 4 induces the appearance of buds on the clusters' surface (Sharon et al., 2019). Immunostaining shows that NEUROD1⁺ non-proliferating endocrine cells reside primarily in peninsula-like buds that formed in stage 4, whereas SOX9⁺ proliferating epithelial progenitors are found in the cluster core (Figures 1I and 1J). Altogether, we conclude that the uniform population that formed in the first three stages of the protocol differentiates and splits into epithelial-cord-like progenitors and endocrine cells. These two cell types are physically separated within differentiating clusters.

The Endocrine Population Includes Four Distinct Cell Types

The endocrine EPs demarcate four major endocrine cell types (Figures 1C–1E). One group represents endocrine precursors and includes cells that express the endocrine initiating gene NEUROG3 (Figure 1D), with some cells co-expressing both progenitor markers (e.g., SOX11) and early-onset endocrine genes (e.g., RUNX1T1) (Osipovich et al., 2014) (Figure S1F). The hormones INS and GCG are expressed at low levels, and some cells express GHRL or SST. These endocrine precursors can be further classified into an early sub-group, collected mostly in the middle of stage 4, and a late sub-group, collected at the end of stages 4 and 5 (EPs 28 and 22, respectively, in Figure S2 and Table S1).

The remaining endocrine cells form three major types of hormonal cells. The first group expresses α cell markers including GCG, ETV1, and ARX, together with the non- α marker INS, and occasionally SST and PPY (Figures 1D and S1F; Table S1). Flow cytometry analysis indicates that, as cells progress into stage 6, these polyhormonal cells resolve into GCG⁺INS⁻ α cells (Figures S1D and S1E). A second group is that of β cells, identified by high expression of INS, NKX6.1, PDX1, and LMO1 and scarcity of GCG transcripts (Figures 1D and S1F; Table S1). Finally, a group of CHGA⁺ endocrine cells express low levels of the pan-islet marker ISL1, serotonin-related genes such as TPH1, and non-islet genes (Segerstolpe et al., 2016; Uhlén et al., 2015) such as ADH6 and LMX1A (Figures 1D, 1E, and S1F; Table S1). These non-pancreatic endocrine cells may represent byproducts of the *in vitro* differentiation process that resemble enteroendocrine populations (Gunawardene et al., 2011; Veres et al., 2019).

Two Waves of Endocrine Differentiation

The presence of epithelial-progenitors side by side with endocrine cells observed from stage 4 onward is reminiscent of normal embryonic pancreas formation, wherein new endocrine cells emerge from epithelial-cord progenitors in a continuous, asynchronous flux (Johansson et al., 2007; Miyatsuka et al.,

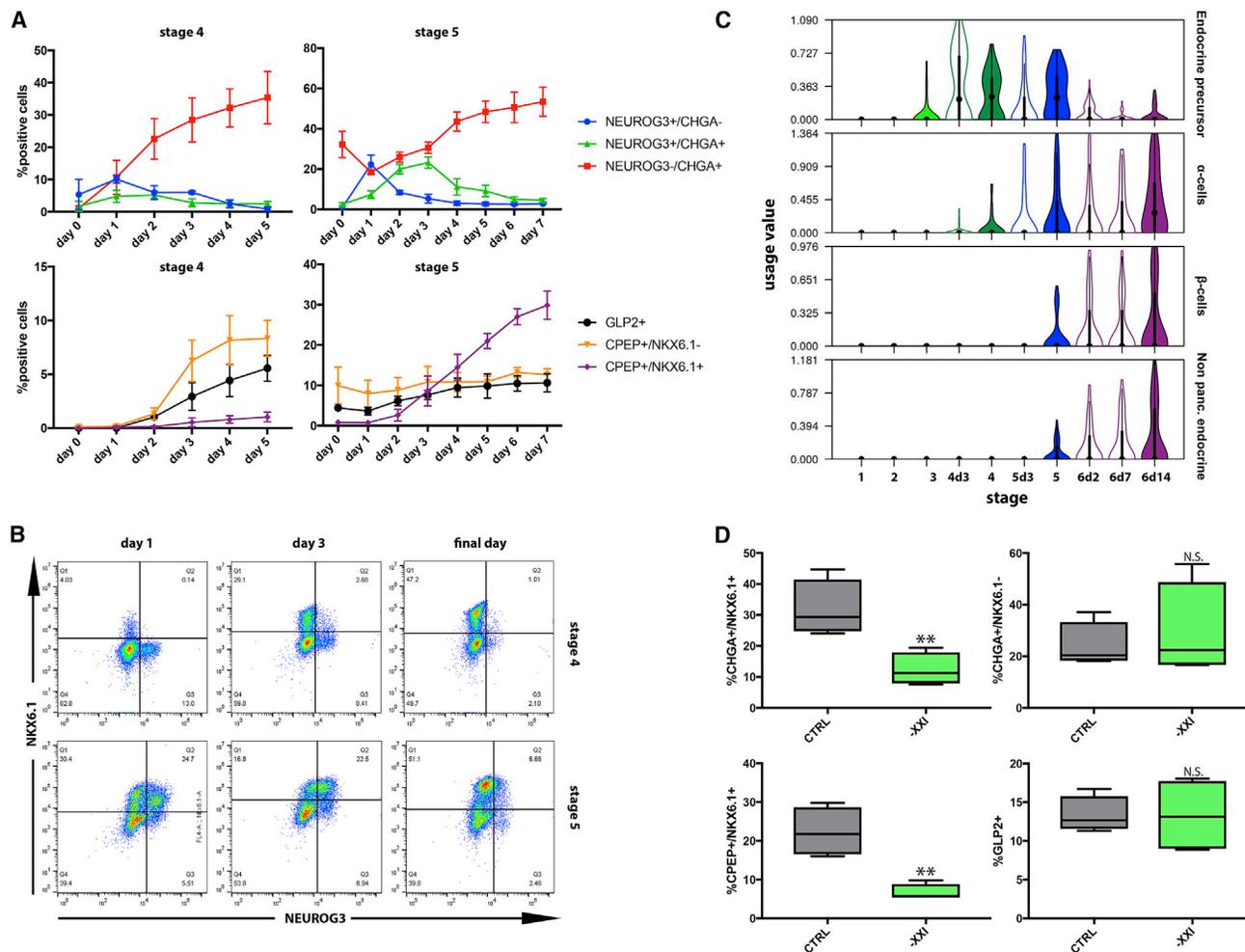


Figure 2. Stages 4 and 5 Produce Different Endocrine Cell Types

(A) Daily flow cytometry analysis of endocrine induction during stages 4 and 5. $n = 3$; data points represent mean \pm SD.

(B) Representative flow cytometry plots show the relationship between NKX6.1 and NEUROG3⁺ expression during stages 4 and 5.

(C) Violin plots present the usage value distributions of endocrine EPs in cells collected in different stages.

(D) Flow cytometry analysis on cells at the end of stage 5, with (control) or without XXI. $n = 4$; p values, clockwise from top left: 0.0022, 0.3933, 0.1848, and 0.0066, Student's t test.

2009). To see whether such a flux occurs *in vitro*, we investigated the dynamics of endocrine differentiation during stages 4 and 5.

Flow cytometry reveals a wave of newly formed NEUROG3⁺ endocrine precursors during the first days of stage 4. A few precursors appear at the last day of stage 3, and within the first day of stage 4 their numbers peak to about 10% of the entire population (Figure 2A). In the following days, fewer NEUROG3⁺ cells appear, and as they differentiate, NEUROG3 is gradually replaced by the pan-endocrine gene CHGA. Less than 2% of cells are NEUROG3⁺CHGA⁻ by the end of stage 4 (Figure 2A). During stage 5, we observed a second wave of endocrine precursor formation. Here, too, the wave peaks on the first day of the stage, when ~20% of the cells are NEUROG3⁺. Similar to stage 4, the initial rise is followed by a daily decline in the number of precursors, as NEUROG3⁺ is replaced with CHGA, until less than 3% of the cells are NEUROG3⁺ by the last day of stage 5 (Figure 2A).

These data show that, unlike embryonic development, *in-vitro*-generated endocrine cells form in two waves, one in stage 4 and the other in stage 5 of the protocol.

It should be noticed that, during stage 6, the proportion of endocrine cells continues to increase, but scRNA-seq shows that this is accompanied with a decline in the number of endocrine precursors detected (Figures 1E and S1C). Thus, this is most likely not the result of a third wave of endocrine differentiation but, rather, the result of existing endocrine precursors completing their differentiation, accompanied by a decline in the numbers of epithelial progenitors.

α Cells Start Forming in the First Wave of Endocrine Differentiation, whereas β Cells Form in the Second Wave

Several findings suggest that the two waves of endocrine formation differ in the populations generated: (1) The α cell EP is

utilized by cells already in the middle of stage 4, whereas the β cell and the non-pancreatic-endocrine programs are only used by cells from the end of stage 5 onward (Figure 2C). (2) NKX6.1, which is expressed together with INS in β cells (Figure 1D), is rarely detected together with either NEUROG3 or CHGA during stage 4. However, at the first day of stage 5, most of the newly emerging NEUROG3⁺ cells express NKX6.1 at medium levels and then gradually turn into CHGA⁺NKX6.1^{high} endocrine cells (Figures 2B and S3C). (3) The GCG⁺ INS⁺ polyhormonal cells that appear at stage 4 express ARX (Table S1), indicative of an α cell state. Conversely, bona fide CPEP⁺NKX6.1⁺ β cells appear almost exclusively during stage 5 (Figure 2A).

To test this hypothesis, we removed XXI from the media. XXI serves to inhibit the Notch pathway during stage 5, and induce general endocrine differentiation (Apelqvist et al., 1999; Behr et al., 2001; Murtaugh et al., 2003). Its removal is, therefore, expected to hamper endocrine differentiation during stage 5, without affecting cells that had differentiated earlier. Indeed, XXI removal during stage 5 dramatically reduced the number CPEP⁺NKX6.1⁺ β cells but did not significantly affect the number of GLP2⁺ α cells (Figure 2D). Thus, interference with endocrine differentiation at stage 5 limits β cell differentiation but does not affect the earlier formation of α cells. This confirms that the first wave of endocrine differentiation, during stage 4, produces primarily α cells, whereas the second wave, at stage 5, gives rise to β cells. This is reminiscent of normal mouse embryonic development, where α cell differentiation precedes that of β cells (Johansson et al., 2007; Sharon et al., 2019).

Identifying Pathways that Affect Pancreas Compartmentalization

The distinct fates of endocrine cells differentiating in the two waves suggests that controlling the timing of differentiation could affect the composition of the clusters. To control the timing of differentiation, we set out to find molecular mechanisms that act in the clusters autonomously rather than through the direct effect of factors already included in the protocol. Ligand-receptor interaction analysis suggested multiple potential pairs that could act either in an autoregulatory process or between different cell types (Figures S3D and S3E; Table S3).

To focus on genetic pathways that control the overall balance between undifferentiated progenitors and endocrine cells, we compared cells with high usage values of the epithelial progenitors EP with cells that use the general endocrine EP (EPs 19 and 6, respectively) (Figure 3A). The expression of more than 4,800 genes is significantly different between the two groups (Figure S3F). As expected, the endocrine population is enriched for transcripts related to islet function and with maturity onset diabetes of the young (MODY), including PAX4, PDX1, PAX6, and NEUROD1; and it expresses higher levels of transcripts related to insulin or glucagon signaling (INS, GCG, ABCC8, GCK, FOXO1, and G6PC2) (Figure 3C; Figures S3F and S4A). Additional genes enriched in these cells are related with general mechanisms of secretion, including the endoplasmic reticulum, endocytosis, and calcium reabsorption (Table S2). The mechanisms shared by endocrine cells and

neurons are also evident with the enrichment of transcripts related with synaptic transport (GABRB3 and CAMK2B). Lastly, *in-vitro*-generated endocrine cells are enriched for transcripts related to several molecular pathways that have a role in endocrine function, including cAMP (RYR2 and AKT3), cGMP/PKG (MAPK1 and MAPK3), AMPK (RAB2A and CREB3), mTOR (PTEN and PRKCA), VEGF (VEGFA and SHC2), and PPAR (PPARA) signaling (Table S2).

Conversely, the epithelial progenitors are enriched for transcripts related to cell adhesion (CDH1, CDC42, and ERBB2) and cell division (MCM7, CCND2, CHEK1, and PCNA). They also express genes that indicate a high activity of several signaling pathways known to have a role in progenitor-to-endocrine differentiation, including Notch (Apelqvist et al., 1999) (NOTCH2, NOTCH3, JAG1, and HES1), hippo-Yap (Gao et al., 2013; Rosado-Olivieri et al., 2019; Mamidi et al., 2018) (YAP1 and AMOT), activin (SMAD2/3, TGFBR1, and ACVR2A), and BMP (BMP2, BMP7, and BMPR1B) (Chung et al., 2010) (Figures S4C–S4F). The progenitor population also expressed higher levels of transcripts related with the PI3K (phosphatidylinositol 3-kinase)-AKT pathway (PIK3R1/3), possibly in context with fibroblast growth factor (FGF) activity and the enrichment of FGF receptors 1 and 2 (FGFR1 and FGFR2) in these cells (Figure S4G).

Unexpectedly, in spite of its known role in pancreas development, we did not observe enrichment of hedgehog pathway activity in either cell type (Kawahira et al., 2003, 2005). Upon examination of specific components of the pathway, we noticed that progenitors express significantly higher levels of both SHH and its signal transducer SMO (Alcedo et al., 2000), but neither of the pathway's inhibitors PTCH1 or PTCH2 show a significant difference in RNA levels (Figure S4H). Accordingly, *in situ* hybridization showed that transcripts of PTCH1 are spread uniformly across the cluster, whereas SMO mRNA is dense in the non-endocrine region of the differentiating cluster and sparse in the endocrine bud (Figure S4I). Surprisingly, unlike its mRNA transcripts, high levels of PTCH1 protein were confined to the endocrine region both in clusters (Figure S4J) and in developing mouse islets (Figure S4K). This suggests that a post-translational mechanism acts to inhibit hedgehog signaling, specifically in the endocrine compartment.

Finally, transcripts upregulated in progenitors are enriched for factors related to the active form of WNT, including the ligand WNT3, the receptors FZD5 and -7, and the major pathway transducer β -catenin (CTNNB1), which shows a similar pattern on the protein level (Figures 3B, 3D, and 3E). Conversely, APC, which inhibits WNT activity, is upregulated in endocrine cells (Figure 3D). In mouse embryos, Apc protein is detected exclusively in the endocrine peninsula area, whereas β -catenin levels are high in the epithelial cords (Figure 3F).

WNT Pathway Inhibition Is Necessary for Endocrine Differentiation in Mice

High WNT pathway activity in progenitors and the opposing expression of APC, the WNT inhibitor, in the endocrine compartment suggest that WNT inhibition plays a role in endocrine differentiation. The Wnt pathway has a role in early murine pancreas formation and in the later expansion of the acinar compartment,

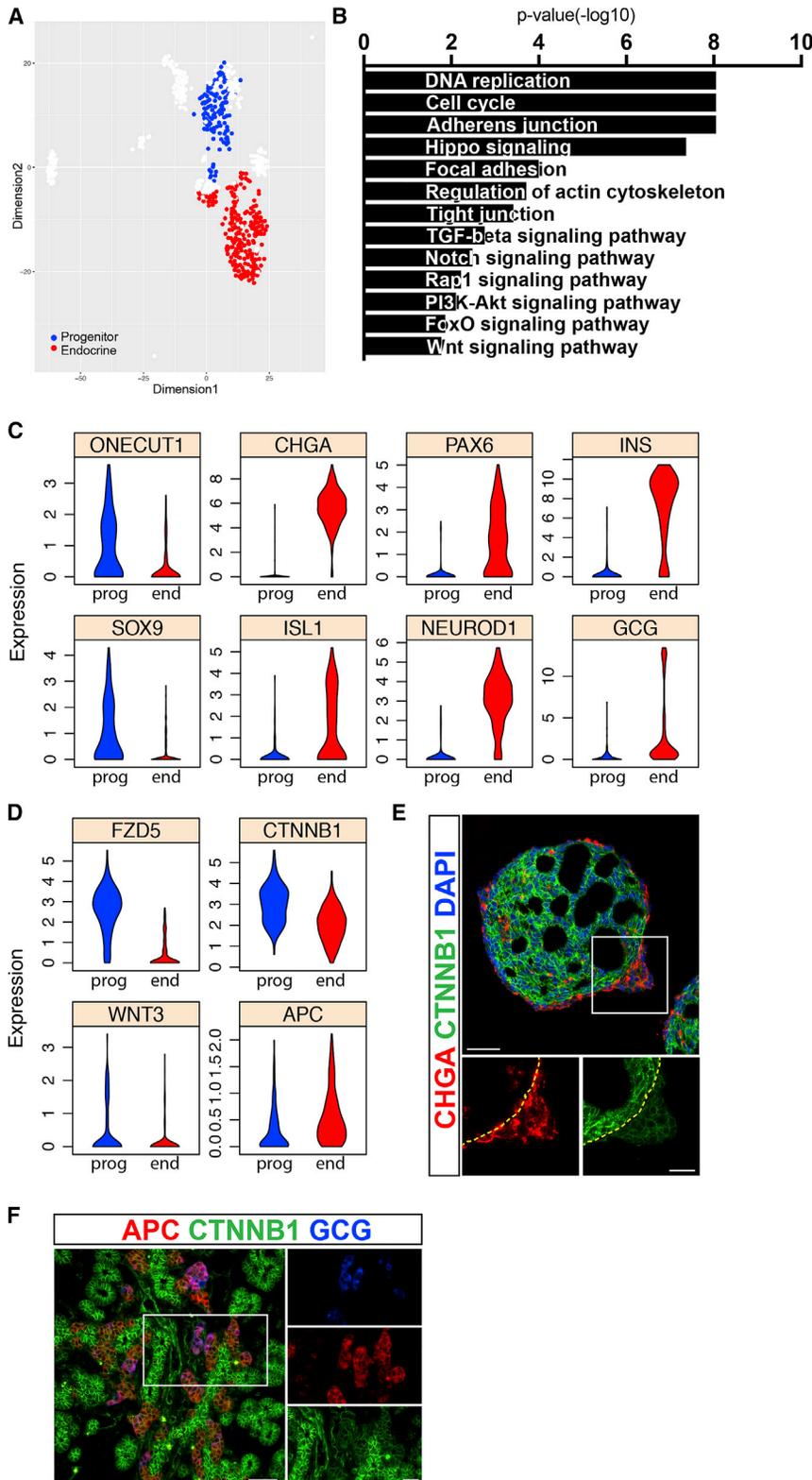


Figure 3. Identification of Genetic Pathways that Separate Progenitors and Endocrine Cells

(A) SIMLR map presents the cells used for differential expression analysis. Cells with transitory intermediate characteristics were omitted to obtain clearer distinction between the populations.

(B) Representative gene categories enriched in the progenitor compartment.

(C) Violin plots show the expression distributions of various genes in progenitor (prog) and in endocrine (end) cells.

(D) Violin plots show the expression distributions of WNT-pathway related genes in progenitor (prog) and in endocrine (end) cells.

(E) Immunofluorescent staining of clusters at the end of stage 4. Dashed line marks the base of the endocrine bud. Scale bars: 50 μ M.

(F) Immunofluorescent staining of a mouse embryonic pancreas at E15.5. Scale bars: 25 μ m.

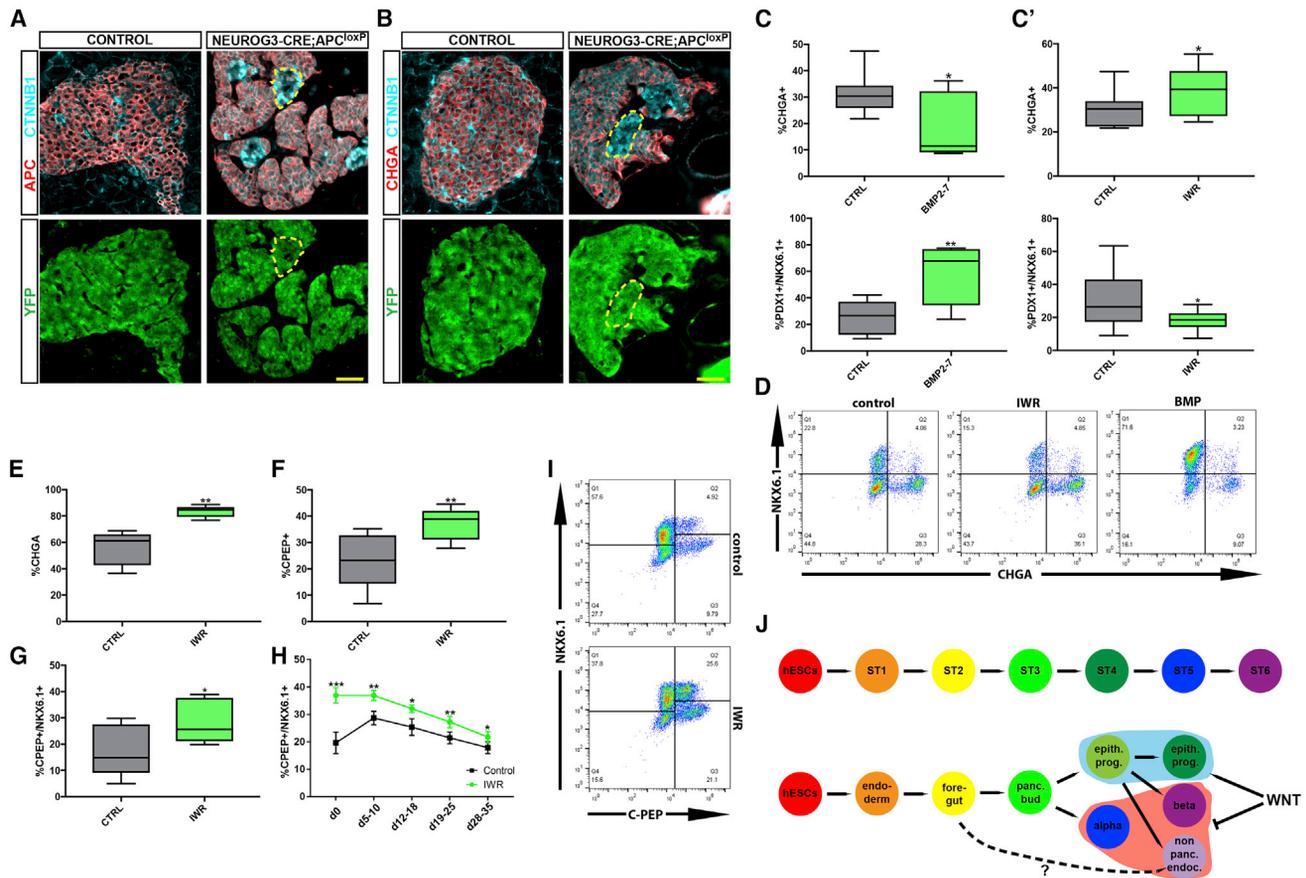


Figure 4. WNT Pathway Inhibition in Endocrine Differentiation

(A and B) Immunofluorescent staining of pancreata from a 1-month-old Neurog3-cre female mouse (control) and a 2-month-old Neurog3-CRE;APC^{loxP/loxP} (APC-endocrine-knockout [KO]) female mouse. Both mice bear a loxP-stop-loxP-YFP sequence in their ROSA26 locus to trace cells that activated CRE. Notice that APC was not deleted in all YFP⁺ cells. Scale bars: 25 μ m. (A) APC⁻ cells express high levels of β -catenin. (B) β -catenin^{high} cells do not express CHGA.

(C) Cell-population proportions calculated using flow cytometry analysis at the end of stage 4, following treatment with BMP2-7. n = 7. Top: p = 0.0235. Bottom: p = 0.0098, Student's t test.

(C') Flow cytometry analysis at the end of stage 4 following treatment with IWR1-endo. n = 9. Top: p = 0.0078. Bottom: p = 0.0314, Student's t test.

(D) Representative flow cytometry plots at the end of stage 4.

(E-G) Flow cytometry analysis at the end of stage 5 following treatment with IWR1-endo. (E) n = 5; p = 0.0045. (F) n = 5; p = 0.0037. (G) n = 5; p = 0.0138. Student's t test.

(H) Flow cytometry analysis of stage 6 cells following treatment with IWR1-endo during stage 5. Values represent mean \pm SEM. d, days into stage 6; d0, last day of stage 5. 2-way ANOVA: p < 10⁻⁴ for change in β cell proportion over days and for overall difference in β cell proportion between conditions. Time-point-specific differences: d0, n = 8, p = 1.6 \times 10⁻⁴; d5-10, n = 7, p = 0.002; d12-18, n = 7, p = 0.03; d19-25, n = 7, p = 0.001. d28-35, n = 7, p = 0.03, Student's t test.

(I) Representative flow cytometry plots at the end of stage 5.

(J) An overview of the protocol (top) and the actual dynamics of cell differentiation *in vitro* (bottom).

but previous reports are inconclusive about a possible role for the WNT pathway in endocrine differentiation (Dessimoz et al., 2005; Heiser et al., 2006; Murtaugh, 2008; Murtaugh et al., 2005; Strom et al., 2007). Since our protocol does not involve manipulation of the pathway at this advanced stage, we decided to examine WNT pathway's role in endocrine differentiation.

First, we examined the effect of Apc deletion on islet formation *in vivo*. Apc is a central component of the destruction mechanism that degrades Ctnnb1, and, in its absence, excess Ctnnb1 molecules accumulate in the nucleus and constitutively activate WNT pathway targets (MacDonald et al., 2009). In agreement with the importance of WNT inactivation during early

pancreas development (Muñoz-Bravo et al., 2016), induced deletion of Apc in the pancreatic bud of mouse embryos resulted in the formation of a rudimentary pancreas (Figures S4L–S4N). To examine WNT's later role in the formation of the endocrine compartment, we crossed mice carrying a conditional knockout of Apc (APC^{MIN}, Apc^{loxP/loxP}) (Kuraguchi et al., 2006; Moser et al., 1990) with mice expressing CRE recombinase specifically in cells expressing Neurog3 (Schönhoff et al., 2004). Cells that efficiently deleted Apc presented nuclear Ctnnb1 accumulation (Figure 4A) and failed to express either the pan-endocrine marker Chga (Figure 4B) or the hormones insulin and glucagon (Figure S4O). Since Neurog3 is

also active during enteroendocrine cell differentiation (Jenny et al., 2002), APC was deleted in gut cells as well, leading to hyperplastic lesions (Figure S4P). Altogether, this shows that WNT pathway inhibition is necessary for proper endocrine differentiation in the mouse pancreas.

WNT Inhibition and BMP Activation Modulate the Ratio between Progenitors and Endocrine Cells

Finally, we examined the potential of compartmentalizing pathways to regulate the ratio between progenitors and endocrine cells *in vitro*. We focused on WNT and BMP pathways, since they are not directly manipulated in the critical stages of our current protocol.

BMP is active in the pool of epithelial progenitors (Figures 3B and S4F), and it was previously shown to attenuate endocrine differentiation (Nostro et al., 2011; Russ et al., 2015). Accordingly, the addition of recombinant BMP2-7 at stage 4 resulted in the downregulation of endocrine genes (Figure S5D), and a nearly 2-fold decrease in the number of CHGA⁺ endocrine cells (Figures 4C and 4D). Concomitantly, the number of PDX1⁺/NKX6.1⁺ progenitors increased by more than 2-fold on average (Figures 4C and 4D). We confirmed this in an independent cell line (Figure S5C) but did not observe a similar effect when BMP2-7 was added during stage 5.

Conversely, the role we found for WNT inhibition in islet formation suggests that it could be used to increase the number of endocrine cells generated *in vitro*. Indeed, treatment with the WNT-tankyrase inhibitor IWR1-endo (Chen et al., 2009) during both stages 4 and 5 increased the expression of endocrine markers and downregulated progenitor markers (Figure S5D). Similar results were obtained using XAV939, another WNT-tankyrase inhibitor (Figure S5D) (Huang et al., 2009). To estimate the effect of WNT pathway activity on the actual yield of the protocol, we measured the effect of its manipulation on cell proportions *in vitro*. Activation of the WNT pathway using the GSK3 inhibitor CHIR99021 (Ring et al., 2003) decreased the overall number of endocrine cells at stage 4 (Figure S5A). In contrast, inhibition of the pathway with IWR1-endo during stage 4 significantly raised the proportion of CHGA⁺ endocrine cells and reduced the proportion of PDX⁺NKX6.1⁺ progenitors (Figures 4C' and 4D). IWR treatment during stage 5 caused an even higher relative increase in the proportion of CHGA⁺ endocrine cells (Figures 4E and S5C'), and, specifically, the proportion of C-PEP⁺/NKX6.1⁺ β cells increased by more than 50% on average, relative to controls (Figures 4F and 4G). The higher proportion of β cells upon IWR addition in stage 5 was maintained during stage 6 of the protocol (Figure 4H), and overall cell numbers were slightly higher following treatment (Figure S5E), indicating that IWR increases the overall yield of *in-vitro*-generated β cells. IWR-treated cells maintain their ability to secrete insulin in response to high glucose levels (Figure S5F), but upon consecutive stimulations with high glucose, IWR-treated cells release less insulin in the second challenge, compared to controls (Figure S5G).

These experiments show that the ratio between the progenitor pool and the differentiated endocrine cells can be regulated by the activity of either the WNT pathway or the BMP pathway.

The exogenous manipulation of the ratio between the two compartments at the right stage in the differentiation protocol affects β cell yield. Specifically, expanding the endocrine compartment through inhibition of the WNT pathway during β cell differentiation at stage 5 improves the yield of *in vitro*-generated β cells.

DISCUSSION

Protocols for *in vitro*-directed differentiation of hESCs into β cells are commonly described as linear processes that homogeneously drive cells from one step to the next (Pagliuca et al., 2014; Petersen et al., 2017; Reznia et al., 2014; Russ et al., 2015). Using scRNA-seq, we found that differentiation is homogeneous only at the first three stages of the protocol, after which epithelial progenitors and endocrine cells reside side by side. A wave of endocrine differentiation at stage 4 tends to produce α cells, and a second wave, at stage 5, gives rise to β cells and non-pancreatic endocrine cells (Figure 4J).

These findings indicate that the *in vitro* differentiation protocol recapitulates multiple aspects of embryonic pancreas development. In both cases, the epithelial cord progenitors and the endocrine peninsulas that emerge from them exist side by side, and in both cases, α cell differentiation precedes that of β cells (Johansson et al., 2007; Pan and Wright, 2011; Sharon et al., 2019). However, we also noticed differences between the protocol and embryonic development, including the absence of early Cpa1⁺ multipotent progenitors that give rise to acinar, ductal, and endocrine cells (Zhou et al., 2007), and the formation of non-pancreatic enteroendocrine-like cells. It should be noted that the differentiating clusters may contain additional rare cell types, which have not been detected by our analysis.

Cell heterogeneity may be attributed to physical constraints such as the effect of cluster thickness on diffusion, but even planar differentiation protocols are often not homogeneous (Kelly et al., 2011). Here, we show that cells initiate autonomous molecular programs, and in the case of WNT and hedgehog pathways, for example, we observe opposing activity levels in the progenitor and endocrine compartments, suggesting a regulatory cross-talk between the various cell types. By intervening with potential regulatory mechanisms, we can manipulate the outcome of the differentiation process. Exogenous BMP activation increases the progenitor compartment in stage 4, whereas small-molecule WNT inhibitors increase the opposite, endocrine population.

The ultimate goal of *in vitro* differentiation protocols is to provide diabetic patients with β cells derived from pluripotent stem cells instead of cadaveric islets. However, it should be noticed that the currently practiced medical protocol involves whole islet transplantation, rather than β cells alone. In fact, there is no certainty that transplanting a single cell type will be able to regain the function of the entire islet. It is, therefore, sensible to aim efforts at generating whole islets *in vitro*, rather than pure populations of β cells. This work joins others (Petersen et al., 2017; Veres et al., 2019) in the attempt to characterize protocols for *in vitro* production of pancreatic endocrine cells at the single-cell level. Our use of scRNA-seq generates a large, unbiased

dataset, which enabled the thorough characterization of the various cell types generated, the timing of their differentiation, and the identification of molecular pathways that affect cell fate. Looking ahead, the ability to finely tune the timing and balance of progenitor versus endocrine fates can be used to control the ratio between α and β cells and promote the *in vitro* formation of islets with proper cell composition.

STAR★METHODS

Detailed methods are provided in the online version of this paper and include the following:

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SUPPLEMENTAL INFORMATION

Supplemental Information can be found online at <https://doi.org/10.1016/j.celrep.2019.04.083>.

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AUTHOR CONTRIBUTIONS

Conceptualization, N.S. and D.A.M.; Methodology, N.S., J.S., D.G., and D.A.M.; Software, J.S., J.M., R.C., and C.T.; Validation, N.S., J.V., E.N.E., and Q.Z.; Formal Analysis, N.S., J.S., J.M., Q.Z., and E.N.E.; Investigation, N.S., J.V., Q.Z., and E.N.E.; Resources, C.T., D.K.G., and D.A.M.; Data Curation, J.S., R.C., J.M., and C.T.; Writing – Original Draft, N.S.; Writing – Review

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DECLARATION OF INTERESTS

D.A.M. is a founder and SAB member for Semma Therapeutics.

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STAR★METHODS

KEY RESOURCES TABLE

REAGENT or RESOURCE	SOURCE	IDENTIFIER
Antibodies		
Rat anti C-peptide	DSHB	Cat#GN-ID4; RRID: AB_2255626
Rabbit anti Chaga	Abcam	Cat#Ab15160; RRID: AB_301704
Chicken anti GFP	Aves Labs inc.	Cat#GFP-1020; RRID: AB_10000240
Goat anti glp2	Santa Cruz Biotechnology	Cat#sc-7781; RRID: AB_2107346
Mouse anti Glucagon	Santa Cruz Biotechnology	Cat#sc-514592; RRID: AB_2629431
Goat anti Neurod1	R&D systems	Cat#AF2746; RRID: AB_2149217
Sheep anti Neurogenin3	R&D systems	Cat#AF3444; RRID: AB_2149527
Mouse anti Nkx6.1	DSHB	Cat#F55A12; RRID: AB_532379
Rabbit anti Sox9	Abcam	Cat#Ab-5535; RRID: AB_2239761
Goat anti PDX1	R&D systems	Cat#AF2419; RRID: AB_355257
Mouse anti somatostatin	Santa Cruz Biotechnology	Cat#sc-55565; RRID: AB_831726
Rabbit anti MCM3	Cell Signaling Technology	Cat#4012s; RRID: AB_2235150
Mouse anti Ctnnb1	ThermoFisher Scientific	Cat#MA1-2001; RRID: AB_326078
Goat anti Patched1	Santa Cruz Biotechnology	Cat#sc-6147; RRID: AB_2253263
Rat anti PTCH	R&D systems	Cat#MAB41051; RRID: AB_2174045
Rabbit anti APC	Abcam	Cat#Ab-15270; RRID: AB_301806
Mouse anti E-cadherin	BD Transduction Laboratories	Cat#610181; RRID: AB_397580
Chemicals, Peptides, and Recombinant Proteins		
ActivinA	R&D Systems	Cat#338-AC
Rock Inhibitor Y-27632	DNSK	Cat#DNSK-KI15-02
Chir99021	Stemgent	Cat#04-0004-10
KGF	Peptotech	Cat#100-19
Sant1	Sigma-Aldrich	Cat#S4572
PDBu	EMD Millipore	Cat#524390
XXI	EMD Millipore	Cat#565790
Alk5i II	Axxora	Cat#ALX-270-445
T3	EMD Millipore	Cat#642511
Betacellulin	ThermoFisher Scientific	
IWR1-endo	Sigma-Aldrich	Cat#I0161
XAV939	Santa Cruz Biotechnology	Cat#sc-296704A
Recombinant Human BMP2/BMP-7 Heterodimer	ThermoFisher Scientific	Cat#3229-BM
Critical Commercial Assays		
ViewRNA ISH Tissue Assay Kit (2-plex)	ThermoFisher	Cat#QVT0012
Anti human SMO ViewRNA probeset	ThermoFisher	Cat#VA1-14719
Anti human PTCH1 ViewRNA probeset	ThermoFisher	Cat#VA6-20367
SMARTer® Ultra Low Kit for the Fluidigm® C1 System	Clontech	Cat#634833
C1 Single-Cell Auto Prep Array for mRNA Seq (17-25 μm)	Fluidigm	Cat#100-5761
Deposited Data		
Raw and analyzed single cell sequencing data	This paper	SRA BioProject PRJNA532884
Experimental Models: Cell Lines		
HUES8 hESCS	HSCI	hES Cell line: HUES-8
hiPSC 1016	HSCI	hiPSC 1016

(Continued on next page)

Continued

REAGENT or RESOURCE	SOURCE	IDENTIFIER
Experimental Models: Organisms/Strains		
Mouse: C57BL/6j-Apc ^{Min} /J	Jackson Laboratories	RRID:IMSR_JAX:002020
Mouse: B6.Cg-Apc ^{tm2Rak} /Nci	NCI Mouse Repository	RRID:IMSR_NCIMR:01XAA
Mouse: Tg(Pdx1-cre) ^{89.1Dam}	In house	RRID:MGI:5902759
Mouse: Tg(Neurog3-cre) ^{C1Able} /J	Jackson Laboratories	RRID:IMSR_JAX:005667
Mouse: B6.129X1-Gt(ROSA)26Sor ^{tm1(EYFP)Cos} /J	Jackson Laboratories	RRID:IMSR_JAX:006148
Software and Algorithms		
Topic modeling	This paper	http://groups.csail.mit.edu/cgs/geneprogram.html
Interactive database	This paper	https://ifx.rc.fas.harvard.edu/invitrobetacells/

CONTACT FOR REAGENT AND RESOURCE SHARING

Further information and requests for resources and reagents should be directed to and will be fulfilled by the Lead Contact, Douglas A. Melton (dmelton@harvard.edu).

EXPERIMENTAL MODEL AND SUBJECT DETAILS**Animals and animal handling**

To generate mice lacking Apc in the pancreatic bud, we crossed heterozygote C57BL/6j-Apc^{Min}/J (Moser et al., 1990) (Jackson laboratory, Bar Harbor, ME) mice, carrying a mutation in the gene (APC^{min/+}), with B6.Cg-Apc^{tm2Rak}/Nci (NCI/NIH) mice (Kuraguchi et al., 2006) (APC^{loxP}) carrying a conditional knock out of Apc, and with Tg(Pdx1-cre)^{89.1Dam} (Gu et al., 2002) mice that express CRE recombinase under a Pdx1 regulatory element. To delete Apc in endocrine cells, we used Tg(Neurog3-cre)^{C1Able}/J (Jackson laboratory, Bar Harbor, ME) (Schonhoff et al., 2004) mice expressing CRE recombinase under a Neurog3 regulatory element with the previous Apc mice, to generate either heterozygotes that carry one mutated allele and one conditional allele, or mice that are homozygotes for the conditional allele. To validate CRE activity, these mice were also crossed with B6.129X1-Gt(ROSA)26Sor^{tm1(EYFP)Cos}/J (Jackson laboratory, Bar Harbor, ME) mice, that express YFP form the ROSA26 locus upon CRE mediated recombination. Both male and female mice were used for breeding all strains, except for C57BL/6j-Apc^{Min}/J, where only heterozygote males were used for breeding and crossed with females from other strains. Sections of adult mice are from 1 and 2 month old females. Sex of the newborns analyzed was not determined. Mice were euthanized through CO₂ inhalation prior to tissue isolation.

Animal studies were performed in strict accordance with the recommendations in the Guide for the Care and Use of Laboratory Animals of the National Institutes of Health. All of the animals were handled according to approved institutional animal care and use committee (IACUC) protocol number [16-05-269]. The protocol was approved by the Committee on the Use of Animals in Research and Teaching of Harvard University Faculty of Arts & Sciences (HU/FAS). The HU/FAS animal care and use program is AAALAC International accredited, has a PHS Assurance (A3593-01) on file with NIH's Office of Laboratory Animal Welfare, and is registered with the USDA (14-R-0128).

HESC culture and differentiation

HUES8 (male) human embryonic stem cells and human induced pluripotent stem cells (hiPSCs) from line 1016 (male) were obtained from the Human Embryonic Stem Cell Facility and iPS Core Facility of the Harvard Stem Cell Institute. Cells were differentiated into beta like cells in suspension as described previously (Pagliuca et al., 2014). Briefly, 150Xe⁶ cells were suspended in 300ml of supplemented mTeSR1 and grown in spinning flasks (70rpm, 37°C, 5%CO₂) for 48 hours prior to the beginning of a stepwise differentiation protocol:

Stage 1 – 24 hours in S1 medium supplemented with ActivinA (100ng/ml) and CHIR99021 (1.4μg/ml), followed by 48 hours without CHIR99021.

Stage 2 – 72 hours in S2 medium supplemented with KGF (50ng/ml).

Stage 3 – 48 hours in S3 medium supplemented with KGF (50ng/ml), LDN193189 (200nM), Sant1 (0.25μM), retinoic acid (2μM), and PDBU (500nM).

Stage 4 – 5 days in S3 medium supplemented with KGF (50ng/ml), Sant1 (0.25μM) and retinoic acid (0.1μM).

Stage 5 – 7 days in BE5 medium supplemented with Beta Cellulin (20ng/ml), XXI (1μM), Alk5i11(10 μM) and T3 (1μM). Sant1 (0.25μM) was added in the first three days, and retinoic acid was added at 0.1μM in the first three days, then at 0.025μM.

Stage 6 – CMRLS medium supplemented with Alk5i11 (10μM) and T3 (1μM).

In many of the experiments following cell collection for scRNA-seq, the protocol was slightly modified to add Y-27632 (10 μ M) at stage 3-5, and ActivinA (5ng/ml) in stage 4. In some cases, an extra day at stage 4 was also added. For the presented functional and flow cytometry analysis in stage 6, CMRLS based media was replaced with S3 medium, without supplements.

Factors were tested in either 6-well plates placed on a rocker, or in Biott ABLE Bioreactors (Stemgent). IWR1-endo (Sigma-Aldrich) was used at 5 μ M. XAV939 (Santa Cruz, SC-296704A) was used at 5 μ M. Recombinant human BMP2/BMP7 (Thermo-Fisher) was used at 100 μ g/ml. when added during stage 4, CHIR99021 was used at a concentration of 1.4-1.55 μ g/ml.

METHOD DETAILS

Preparation of single-cell suspension

Clusters were collected prior to commencement of differentiation, at the last day of stages 1-5, on the 3rd day of stages 4 and 5, and on the 2nd, 7th and 14th days of stage 6. The clusters were allowed to settle, washed in PBS and dissociated by 5min incubation in TrypLE Express (GIBCO) at 37°C, followed by mechanical agitation through pipetting. TrypLE activity was terminated by washing with PBS+1% FCS.

Single cell library preparation and sequencing

Single-cell capture, cell lysis, reverse transcription of full-length mRNA and cDNA amplification were performed using the Fluidigm C1 system and the Smart-seq protocol (Ramsköld et al., 2012). Sequencing libraries were prepared using the Illumina Nextera XT DNA library preparation kit and samples were sequenced on an Illumina HiSeq.

DATA ANALYSIS

Alignment and processing of sequencing reads, data conversion and cell filtering

Reads were aligned to the human reference genome build hg19 with TopHat (v2.0.13), which was provided with gene annotations from GENCODE (v17). Gene expression profiles for each cell were computed using Cufflinks (2.2.0) as previously described in Trapnell et al., 2014. Per-cell fragments per kilobase million (FPKM) measures were converted into relative transcript counts using the Census algorithm from the Monocle package (Qiu et al., 2017a, 2017b; Trapnell et al., 2014). Cells with < 1k detected genes or with > 45k total mRNA were removed. Additional potential outliers were removed by defining lower and upper thresholds based on the appearance of the per-cell total mRNA distribution curve, separately for each sample. This approach automatically accounted for all cells lying outside of 2 standard deviations of each sample's mean log-transformed per-cell total mRNA count and resulted in final number of 773 single-cell expression profiles (949 pre-filtering), with a median of 898,228 (mean: 945,907) usable paired end reads per cell (Figure S1B).

Inference of cellular gene expression programs via topic modeling

The topic model (TM) is a probabilistic unsupervised learning algorithm which is designed to discover latent topics present in a set of text documents, based solely on the observed usage of each word in each document (Blei, 2012). For text data, a topic is defined as set of (*a priori* unspecified) related words which probabilistically tend to co-occur. For example, statistical co-occurrence of words such as “banks,” “market,” and “GDP” within the same document provides strong evidence that a topic such as finance is being discussed. Once the topics have been inferred, a document can be more interpretably represented in terms of the relative presence of each topic rather than a massive collection of word frequencies. In our analysis, we utilize the topic model and inference algorithm used in the GeneProgram software tool (Gerber et al., 2007), which employs a Hierarchical Dirichlet Process (HDP) prior. This enables automatic inference of the number of latent topics (along with the topics themselves) from the data (Teh et al., 2006).

TM is adapted to scRNA-seq data by treating each cell as a document and each gene as a word. Within the cell, the unit amount of mRNA from the gene is therefore the genomic equivalent of a word's frequency within a document. In this context, we refer to a latent topic as an Expression Program (EP), which represents a set of genes that behave coordinately in particular cells. Unlike traditional clustering techniques, our TM analysis allows many EPs to be active within a given cell, and a single gene may also be used by different EPs (one gene can regulate multiple different biological processes depending on the context in which it is expressed). For each EP, a gene receives a relevance value (quantifying how active the gene is within the EP) and a cell is assigned a usage value (quantifying how active the EP is within the cell). After inferring the EPs underlying our scRNA-seq measurements via Markov-Chain Monte Carlo (Gerber et al., 2007), we can interpretably represent and classify each cell in terms of its relative usage of different EPs, rather than the raw expression values of many thousand genes. Similarly, each EP is itself represented as a collection of genes along with their “relevance” in defining the EP. The biological significance of an EP can thus be elucidated by investigating which genes are the most actively used (Gerber et al., 2007).

The TM software used to infer the EPs in our analysis is available at <http://groups.csail.mit.edu/cgs/geneprogram.html>, and was run with the following settings: 4 levels of discretization for the expression values, 100 posterior samples (taken every 1,000 steps of the MCMC-chain with an initial burn-in of 30,000 steps), minimum of 2 genes per EP with a minimum usage value of 0.1, with the default prior HDP concentration and minimum-similarity/merging parameters used in (Gerber et al., 2007). See (Gerber et al., 2007) for detailed descriptions of these parameters.

SIMLR analysis

HESCs were removed from the 773 cleaned cells, retaining 720 cells for all subsequent analyses. A polynomial trend curve was fitted to the mean-variance plot of gene FPKM counts, and used as an estimate for technical variability. Altogether 3453 genes showed higher variance, and were used to cluster the cells using the SIMLR method implemented in the SIMLR R library (Wang et al., 2017). SIMLR calculates a cell-cell similarity metric based on parametrized Gaussian Kernels and uses the similarity values as input to the stochastic neighbor embedding method to plot the cell data in reduced dimensions. K-means clustering was subsequently used to assign the cells to different numbers of clusters, eventually settling on 10 predetermined clusters.

Differential expression was determined using the ROTS algorithm (Suomi et al., 2017). P values were corrected for multiple testing using the false discovery rate (FDR), and 4863 genes with FDR of less or equal to 0.05 were further characterized using Enrichr pathway analysis (Chen et al., 2013). The KEGG database (Kanehisa et al., 2017) was used as a source of curated lists of pathway genes.

Ligand-receptor interaction analysis

Ligands and receptors from Ramilowski et al. (Ramilowski et al., 2015) were introduced into the stage-specific analysis if expressed in 20% or more of the cells analyzed from that stage. The expression of these genes was compared between the cells belonging to the EPs analyzed on a given stage, and those that were significantly differentially expressed between EPs were retained and assigned to the EP where their average expression level is highest. This analysis was performed separately for stages 4 and 5.

Flow cytometry analysis

Differentiated cell clusters were dispersed into single-cell suspension by incubation in TrypLE Express at 37°C for 10-30 minutes, quenched with PBS+2% FBS, fixed with 4% PFA for at least 30 min at 4°C, washed once in PBS, and resuspended in PBS+ 2% FBS. Staining was done in 96-well plates. Cells were incubated in blocking buffer (PBS+0.1% Triton X-100+5% donkey serum) at 4°C for 30-60 min. Cells were then re-suspended in blocking buffer with primary antibodies and incubated at room temperature for 1 hour or at 4°C overnight. Cells were washed twice in blocking buffer and incubated in blocking buffer with secondary antibodies at 4°C for 1 hour. Cells were then washed three times in PBS+5% Donkey serum and analyzed using either the BD Accuri c6 flow cytometer or the LSR-II flow cytometer (BD Biosciences). Analysis of the results was performed using FlowJo software.

Nano-string gene expression analysis

Cells were collected for RNA extraction in RLT Lysis Buffer (QIAGEN) and frozen at –80°C. RNA was extracted using the RNeasy Mini Kit (QIAGEN), according to the manufacturer's instructions - including DNA digestion, and 100ng of RNA was used for each reaction. Nanostring set up was performed according to the protocol for the nCounter XT Gene Expression Assay. Data was analyzed in the nSolver 2.5 Software. All genes were normalized to housekeeping genes (ITCH, RPL15, RPL19, TCEB1, UBE2D3). Analysis of changes in gene expression of markers for endocrine and progenitor cells was done using R.

Immunofluorescence staining and *In situ* hybridization

Isolated tissue was fixed overnight in 4% paraformaldehyde, washed in PBS, embedded in paraffin and sectioned on a microtome. Paraffin was removed using Histo-Clear (National Diagnostics) and serial washes in declining ETOH concentrations. After antigen retrieval, samples were blocked in PBS with 5% donkey serum and 0.1% Triton x-100 for 1 hour in room temperature. Primary antibody was diluted in the blocking solution and incubated in room temperature for 1hr, followed by washing. Secondary antibody was incubated for 1 hour in room temperature, washed, and DAPI staining was added.

In situ hybridization was performed using ViewRNA ISH tissue assay kit. For detection of human SMO and PTCH1 transcripts, VA1-14719 and VA6-20367 probes were used, respectively.

Glucose Stimulated Insulin Secretion (GSIS)

Differentiated cells were washed twice in Krebs buffer (Krb) containing 2.8mM glucose, and then distributed into 24-well plates containing Millicell Cell culture inserts for easy transfer between conditions. Each well received 0.5ml buffer with approximately 10⁶ cells. After fasting in 2.8mM glucose for 1 hour in 37°C, the analysis of the cells' response to changing glucose levels was performed by a series of 1 hour incubations in 37°C, in the following order: 1) 2.8mM (low) glucose. 2) 20mM (high) glucose. 3) brief wash in 2.8mM glucose. 4) 1 hour incubation in 2.8mM glucose. 5) 20mM glucose. 6) 2.8mM glucose with 30mM KCL. At the end of each step (excluding the wash), the incubation buffer was collected and the amount of insulin secreted during the hour's incubation was determined using human insulin ELISA kit (ALPCO diagnostics; 80-INSHUU-E01.1). Clusters were then dispersed into single cells using TrypLE Express (Life Technologies), and cell number was counted automatically by a Vi-Cell (Beckman Coulter).

QUANTIFICATION AND STATISTICAL ANALYSIS

All statistical methods used are described in the legend to the relevant figures. n marks the number of times an experiment was performed on independent differentiation batches, except for Figure S5B, which describes a triplicate of samples taken from the same differentiation batch. Data was analyzed and plotted using Prism software from GraphPad. When boxplots are presented, the bottom

and top box borders represent the 25th and 75th percentiles, respectively. The line inside the box represents the median, and the whiskers represent the minimum and maximum values observed.

DATA AND SOFTWARE AVAILABILITY

Single-cell RNA-sequencing data was deposited to the NCBI Sequence Read Archive (SRA). The accession number for the raw sequences is: PRJNA532884.

ADDITIONAL RESOURCES

An interactive website which presents gene expression and EP usage in the cells is accessible at <https://ifx.rc.fas.harvard.edu/invitrobotacells/>