A human ESC-based screen identifies a role for the translated IncRNA LINC00261 in pancreatic endocrine differentiation

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Abstract Long noncoding RNAs (lncRNAs) are a heterogenous group of RNAs, which can encode small proteins. The extent to which developmentally regulated lncRNAs are translated and whether the produced microproteins are relevant for human development is unknown. Using a human embryonic stem cell (hESC)-based pancreatic differentiation system, we show that many lncRNAs in direct vicinity of lineage-determining transcription factors (TFs) are dynamically regulated, predominantly cytosolic, and highly translated. We genetically ablated ten such lncRNAs, most of them translated, and found that nine are dispensable for pancreatic endocrine cell development. However, deletion of LINC00261 diminishes insulin+ cells, in a manner independent of the nearby TF FOXA2. One-by-one disruption of each of LINC00261’s open reading frames suggests that the RNA, rather than the produced microproteins, is required for endocrine development. Our work highlights extensive translation of lncRNAs during hESC pancreatic differentiation and provides a blueprint for dissection of their coding and noncoding roles.

Introduction Defects in pancreatic endocrine cell development confer increased diabetes risk later in life (Bakhti et al., 2019). Therefore, a detailed understanding of the factors that orchestrate endocrine cell differentiation is highly relevant to human disease. Many of the molecular mechanisms that underlie the formation of pancreatic endocrine cells have been defined (Romer and Sussel, 2015; Schiesser and Wells, 2014). However, despite some evidence that long noncoding RNAs (lncRNAs) are important for proper development and function of pancreatic beta cells (Arnes et al., 2016; Morán et al., 2012; Wong et al., 2019), a systematic functional assessment of the noncoding transcriptome during pancreas development is lacking.

Most lncRNAs with to date demonstrated roles in the regulation of fundamental developmental processes are active in the cell’s nucleus (Daneshvar et al., 2016; Jiang et al., 2015; Klattenhoff et al., 2013; Kurian et al., 2015; Lin et al., 2014; Luo et al., 2016; Ramos et al., 2015). However, a large proportion of lncRNAs is predominantly cytosolic (Cabilly et al., 2015; van Heesch et al., 2014), and the functional relevance of these lncRNAs has remained unexplored in the context of human development. It is now widely accepted that many cytosolic lncRNAs possess short, ‘non-canonical’ open reading frames (sORFs) that are translated (Bazzini et al., 2014;
Makarewich and Olson, 2017; Ruiz-Orera et al., 2014). What fraction of these non-canonical ORFs is functional, and whether sORF translation serves a pure regulatory purpose or results in the production of stable microproteins, remains an active topic of debate (Levy, 2019; Ruiz-Orera et al., 2018). Since high rates of conservation have historically been employed for the identification and annotation of canonical protein coding sequences (Lin et al., 2011; Mudge et al., 2019), a primary reason for doubting the protein-coding capacity of sORFs in presumed lncRNAs is their generally poor sequence conservation across species. To address these questions, several recent studies have systematically assessed the biological activity of newly discovered sORFs, revealing that many produce evolutionary young microproteins with roles across cellular organelles and processes, and a subset being essential for cell survival (Chen et al., 2020; Martinez et al., 2020; Prensner et al., 2018). This previously unrecognized coding capacity of supposedly non-coding RNAs illustrates their functional diversity and has called into question the noncoding classification of some lncRNAs. Thus, there is a need for careful investigation and dissection of any gene’s coding and noncoding functions.

LncRNAs, translated or fully noncoding, are not randomly distributed in the genome but are frequently located close to, and coregulated with, canonical protein-coding genes in cis (Luo et al., 2016; Neumann et al., 2018; van Heesch et al., 2019). For example, the lncRNAs DIGIT (also known as GSC-DT) and Gata6as (also known as IncGata6 or GATA6-AS1) have been reported to enhance expression of divergently expressed endoderm regulators Goosecoid (GSC) and Gata6, respectively (Daneshvar et al., 2016; Luo et al., 2016; Neumann et al., 2018). Similarly, the Pax6- associated lncRNA Paupar promotes pancreatic islet alpha cell formation through the alternative splicing of Pax6 transcripts in mice (Singer et al., 2019). Furthermore, LINC00261 (also known as DEANR1) and its neighboring TF FOXA2 are both induced in endoderm formation, during which LINC00261 has been proposed to positively regulate FOXA2 expression (Jiang et al., 2015). However, whether such cis-acting lncRNAs are translated and may exert cytosolic functions through trans-acting, microprotein-dependent mechanisms relevant for endoderm and pancreas development is not known.

In this study, we classified lncRNAs based on their dynamic regulation, subcellular localization, and translation in a hESC differentiation system that recapitulates in vivo pancreas development. Next, we used this classification to prioritize select dynamically regulated and highly translated lncRNAs for deletion in hESCs, followed by extensive phenotypic characterization across multiple intermediate states of pancreas development. Nine out of the ten selected lncRNAs were not essential for pancreatic development and, despite their vicinity to lineage-determining TFs, none of these lncRNAs regulated the expression of these TFs in cis.

The deletion of one lncRNA, LINC00261, impaired human endocrine cell development and led to a significant reduction in the number of insulin-producing cells. Contrary to previous studies of LINC00261 knockdown hESCs (Jiang et al., 2015), deletion of LINC00261 had no effect on the expression of nearby TF FOXA2 or other proximal genes, suggesting control of endocrine cell formation through a trans- rather than cis-regulatory mechanism. LINC00261 was among the most highly translated lncRNAs based on ribosome profiling (Ribo-seq) and produced multiple microproteins with distinct subcellular localizations upon overexpression in vitro. To systematically assess LINC00261’s coding and noncoding functions, we separately introduced frameshift mutations into each of seven identified LINC00261 sORFs. However, rigorous phenotypic characterization revealed no apparent consequences of loss of each of the seven LINC00261-sORF-encoded microproteins on endocrine cell development. Our comprehensive assessment of functional lncRNA translation identified a likely trans-regulatory role for LINC00261 in endocrine cell differentiation that appears to be independent of the seven microproteins that were individually deleted. With this detailed investigation we provide a blueprint for the proper dissection of a gene’s coding and noncoding roles in a human disease-relevant system.
Results

LncRNAs and nearby lineage-determining transcription factors exhibit dynamic coregulation during pancreas development

To identify lncRNAs involved in the regulation of pancreas development, we profiled RNA expression at five defined stages of hESC differentiation toward the pancreatic lineage: hESCs (ES), definitive endoderm (DE), primitive gut tube (GT), early pancreatic progenitor (PP1), and late pancreatic progenitor (PP2) (Figure 1A). While some lncRNAs were constitutively expressed (n = 592; 25.3%), the majority showed dynamic expression patterns (n = 1745; 74.7%), being either strongly enriched in (n = 874; 37.4%) or specific to (n = 871; 37.3%) a single developmental intermediate of pancreatic lineage progression (Figure 1B and Figure 1—source data 1A). The expression of many of these
dynamically regulated IncRNAs correlated with that of proximal coding genes (Figure 1—figure supplement 1A–D and Figure 1—source data 1B,C), further exemplified by a subset of IncRNAs that was specifically coregulated with the key endodermal and pancreatic TFs GATA6, FOXA2, PDX1, and SOX9 (Figure 1C,D). The expression coregulation of these IncRNA-TF pairs is likely explained by a shared chromatin environment (Figure 1—figure supplement 1E–H), which raises the possibility that like the TFs, the function of the IncRNAs is also required for endoderm and pancreas development.

Many pancreatic progenitor-expressed IncRNAs are cytoplasmically enriched and translated

Although most functional roles described for IncRNAs to date have been predominantly nuclear (Marchese et al., 2017), multiple recent studies have shown that many IncRNAs are cytosolic and translated into sometimes biologically active microproteins (reviewed in Makarewich and Olson, 2017). To further characterize the above-identified dynamically regulated IncRNAs, we analyzed their subcellular localization and translation potential using fractionation RNA-seq and Ribo-seq across multiple hESC clones independently differentiated into PP2 stage pancreatic progenitors (Figure 2A). Of all IncRNAs expressed in two replicate differentiations into PP2 cells, we classified 21% (n = 347) as localized to the nucleus, whereas a larger number (n = 563; 34%) primarily resided in the cytosol (Figure 2—figure supplement 1A and Figure 2—source data 1A). This subcellular distribution of pancreatic IncRNAs is in agreement with previous IncRNA localization studies by us and others (Cabili et al., 2015; Clark et al., 2012; Sun et al., 2015; van Heesch et al., 2014). IncRNAs enriched in the cytosol were expressed at higher levels than nucleus-localized IncRNAs, with expression levels similar to canonical protein-coding mRNAs (Figure 2—figure supplement 1B). Intriguingly, almost half (49.4%) of all cytosol-enriched IncRNAs (278 out of 563) displayed dynamic expression regulation during the differentiation of hESCs to pancreatic progenitors, raising the possibility that many IncRNAs with putative developmental functions do not act in the nucleus, but instead in the cytosol where they may be translated.

To investigate the translation potential of these cytosolic IncRNAs, we used Ribo-seq, through which we obtained exceptionally deep and high quality translome coverage across six replicate differentiations (Figure 2—figure supplement 1C and Figure 2—source data 1B). As nearly 90% of the sequenced ribosomal footprints exhibited clear 3-nucleotide codon movement characteristic of translation (Figure 2—figure supplement 1D–F), these data have strong predictive value for the computational detection of non-canonical ORFs, such as upstream ORFs (uORFs) in the 5’ leader sequences of mRNAs and sORFs in genes annotated as IncRNAs (Figure 2—source data 1C). Requiring stringent reproducibility criteria (the exact ORF needed to be detected by RiboTaper (Calviello et al., 2016) in at least four out of six replicates), we identified a total of 625 new sORFs in IncRNAs with a median length of 47 amino acids (aa) (Figure 2—source data 1D). The majority of detected sORFs (76%; n = 477/625) is currently not present in the sORFs.org database (Olexiouk et al., 2016). The translated sORFs are located within 285 cytosolically localized IncRNAs (25.3% of all expressed IncRNAs) (Figure 2—figure supplement 1B), which are expressed at higher levels than untranslated IncRNAs (Figure 2—figure supplement 1G) and exhibit translational efficiencies similar to mRNAs (Figure 2—figure supplement 1H and Figure 2—source data 1E). Of note, almost none of the newly identified sORFs are highly conserved across species, as judged by their low PhyloCSF scores (Lin et al., 2011; Figure 2—source data 1D).

Using approaches similar to ours, non-canonical sORFs have previously been characterized in multiple immortalized human cell lines (Bazzini et al., 2014; Calviello et al., 2016; Chen et al., 2020; Ji et al., 2015; Martinez et al., 2020; Prensner et al., 2020; Raj et al., 2016) and human tissues (van Heesch et al., 2019). However, to our knowledge, our data constitute the first comprehensive set of non-canonical human ORFs generated from a non-transformed human cell model of development, providing a valuable resource for future functional studies.

Transcribed IncRNAs in pancreatic progenitors produce microproteins with distinct subcellular localizations

Having established that many stage-specific pancreatic IncRNAs are translated, we next sought to validate their translation potential through independent experimental approaches, additionally
Figure 2. Cytosolic lncRNAs contain translated small open reading frames. (A) Overview of experimental strategy for subcellular fractionation and Ribo-seq-based identification of translated small open reading frames (sORFs) from lncRNAs expressed in PP2 cells. Replicates from six independent differentiations to PP2 stage each for total (polyA) RNA-seq and Ribo-seq experiments, and two biological replicates for the subcellular fractionation were analyzed. The histogram on the far right depicts the size distribution of the sORF-encoded small peptides as number of amino acids (aa). The pie
Figure 2 continued on next page
investigating the production of the predicted microproteins at the protein level. To this end, we first performed coupled in vitro transcription-translation assays on endogenous and complete transcript isoforms of four of the most highly translated lncRNAs (LINC00261, RP11-834C11.4, LHFPL3-AS2, and MIR7-3HG; Figure 2-figure supplement 1; expression and ORF information in Figure 2B–E).

Second, we generated a series of in vivo translation reporter constructs to assess the subcellular localization of microproteins translated from each of ten sORFs derived from the same four lncRNAs. Transient expression of individual constructs carrying in-frame GFP fusions in HEK293T cells produced GFP signal for all ten assayed microproteins, which was abolished upon introduction of a frameshift within the sORF or a stop codon following the sORF sequence (Figure 2F and Figure 2-figure supplement 1J–L). To rule out a possible localization bias induced by the GFP fusion, we also expressed a FLAG-tag fusion peptide (RP11-834C11.4 sORF-1xFLAG), which revealed a cytoplasmic localization identical to the one observed for the GFP construct (Figure 2-figure supplement 1J).

While most sORF-GFP fusion products were ubiquitously distributed throughout transfected cells, LINC00261 sORF4-GFP specifically localized to mitochondria (Figure 2-figure supplement 1K), and LINC00261 sORF7-GFP exhibited a perinuclear accumulation pattern reminiscent of aggresomes (Figure 2-figure supplement 1L). Taken together, our results validate the translation potential of sORFs encoded by pancreatic progenitor-expressed lncRNAs and show that, upon ectopic expression, these translation events result in the production of microproteins with different subcellular localizations.

**Deletion phenotypes of translated lncRNAs during hESC pancreatic differentiation**

To identify potential functional roles of translated lncRNAs during pancreas development, we selected ten candidates for CRISPR/Cas9-based genome editing in hESCs through excision of the lncRNA promoter or entire lncRNA locus (Figure 3A,B). These ten lncRNAs were prioritized based on (i) high expression and endodermal tissue-specificity, (ii) dynamic regulation during pancreas development, (iii) abundant translation of sORFs, and (iv) proximity to TFs with known roles in endoderm and pancreas development. For seven of the selected lncRNAs, translation was highly abundant and reproducibly detected across Ribo-seq replicates: LINC00617 (also known as TUNAR; Lin et al., 2014), GATA6-AS1 (also known as GATA6-AS; Neumann et al., 2018), LINC00261, RP11-834C11.4, SOX9-AS1, MIR7-3HG, and LHFPL3-AS2. Although for two additional lncRNAs the translation potential could not be determined, they were nonetheless included because of a previously reported requirement for definitive endoderm formation (DIGIT, also known as GSC-DT) (Daneshvar et al., 2016) and genomic localization adjacent to the definitive endoderm TF LHX1 (RP11-44SF12.1, also known as LHX1-DT). Lastly, LINC00479 was chosen as a non-translated control with expression dynamics and a subcellular localization similar to LINC00261. Of note, for each of the ten selected lncRNAs, we generated at least two independent hESC knockout (KO) clones and used different combinations of single guide RNAs where possible (Figure 3-source data 1A).

We next differentiated each of the IncRNA KO hESC lines stepwise toward the pancreatic endocrine cell stage, conducting up to 16 replicate differentiations for each KO clone. Because
Figure 3. A small-scale CRISPR loss-of-function screen for dynamically expressed and translated lncRNAs during pancreatic differentiation. (A) qRT-PCR analysis of candidate lncRNAs during pancreatic differentiation of H1 hESCs relative to the ES stage. Data are shown as mean ± S.E.M. (mean of n = 2–6 independent differentiations per stage; from H1 hESCs). Individual data points are represented by dots. See also Figure 3—source data 2. (B) CRISPR-based lncRNA knockout (KO) strategy in H1 hESCs and subsequent phenotypic characterization. (C) Immunofluorescence staining for OCT4

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and SOX17 in DE from control (ctrl) and KO cells for the indicated lncRNAs (representative images, n ≥ 3 independent differentiations; at least two KO clones were analyzed). (D) qRT-PCR analysis of DE lineage markers in DE from control and lncRNA KO (-/-) cells. TF genes in cis to the lncRNA locus are highlighted in red. Data are shown as mean ± S.E.M. (n = 3–16 replicates from independent differentiations and different KO clones). Individual data points are represented by dots. NS, p-value>0.05; t-test. See also Figure 3—source data 3. (E) Flow cytometry analysis at DE stage for SOX17 in control and KO (-/-) cells for indicated lncRNAs. The line demarks isotype control. Percentage of cells expressing SOX17 is indicated (representative experiment, n

3 independent differentiations). (F) Immunofluorescence staining for FOXA2 or GATA6 in DE from control and LINC00261, GATA6-AS1, and DIGIT KO cells. (G) Immunofluorescence staining for insulin (INS) in endocrine cell stage (EC) from control and KO hESCs for the indicated lncRNAs (representative images, n ≥ 3 independent differentiations from at least two KO clones). Boxed areas (dashed boxes) are shown in higher magnification. (H) qRT-PCR analysis of INS in EC stage cultures from control and lncRNA KO (-/-) hESCs. Data are shown as mean ± S.E.M. (n ≥ 4 replicates from independent differentiations of at least two KO clones). Individual data points are represented by dots. NS, p-value>0.05; t-test. See also Figure 3—source data 4 (I) Flow cytometry analysis at EC stage for INS in control and KO (-/-) cells for indicated lncRNAs. The line demarks isotype control. Percentage of cells expressing insulin is indicated (representative experiment, n ≥ 3 independent differentiations each from at least two KO clones). Scale bars = 100 μm. See also Figure 3—figure supplement 1 and Figure 3—source data 1–4. The online version of this article includes the following source data and figure supplement(s) for figure 3:

Source data 1. Differentially expressed genes after lncRNA deletion.
Source data 2. Source data used for the qRT-PCR quantification of gene expression presented in Figure 3A.
Source data 3. Source data used for the qRT-PCR quantification of gene expression presented in Figure 3D.
Source data 4. Source data used for the qRT-PCR quantification of INS expression presented in Figure 3H.
Figure supplement 1. Minor gene expression changes in definitive endoderm or pancreatic progenitor cells after lncRNA deletion.

LINC00617, RP11-445F12.1, DIGIT, GATA6-AS1, LINC00479, and LINC00261 were first expressed at, or before, the definitive endoderm stage (Figure 3A), we determined whether KO hESCs for these lncRNAs exhibited defects in definitive endoderm formation. Despite efficient lncRNA depletion (Figure 3—figure supplement 1A,B), neither quantification of definitive endoderm marker gene expression by qRT-PCR, nor immunofluorescence staining or flow cytometric analysis of the definitive endoderm marker SOX17 showed differences indicative of impaired endoderm formation in lncRNA KO lines (Figure 3C–E). Importantly, expression of TFs located in the direct vicinity of these lncRNAs, including GSC (DIGIT), LHX1 (RP11-445F12.1), GATA6 (GATA6-AS1), and FOXA2 (LINC00261), was unaffected by the lncRNA KO (Figure 3F, Figure 3—figure supplement 1C, Figure 3—source data 1B–D), arguing against cis-regulation by these lncRNAs. These findings are in contrast to prior reports that have shown a requirement for LINC00261 and DIGIT in definitive endoderm formation and the regulation of neighboring TFs FOXA2 and GSC, respectively (Amaral et al., 2018; Daneshvar et al., 2016; Jiang et al., 2015; Swarr et al., 2019).

Next, we further differentiated control and KO lines for eight out of ten lncRNAs toward the endocrine cell stage, excluding DIGIT and RP11-445F12.1 because they are not expressed after the definitive endoderm stage (Figure 3A). In KO hESC lines of seven out of these eight lncRNAs, we observed no effect on pancreatic progenitor cell formation or gene expression, with the exception of a handful of dysregulated genes in LHFPL3-AS2 and RP11-834C11.4 KO cells (Figure 3—figure supplement 1C and Figure 3—source data 1E–K). Furthermore, deletion of seven out of the eight lncRNAs did not impair endocrine cell formation, as determined by quantification of insulin+ cells and insulin mRNA levels (Figure 3G–I). Similar to the RNA expression results obtained at the definitive endoderm stage, deletion of none of the lncRNAs close to pancreatic TFs (e.g. GATA6-AS1 and SOX9-AS1) altered the expression of these TFs, once more arguing against cis-regulation of these TFs by the neighboring lncRNA (Figure 3—figure supplement 1C). Thus, nine out of ten endoderm- and pancreatic progenitor-enriched lncRNAs functionally investigated here appear to be nonessential for induction of the pancreatic fate and formation of insulin+ cells. Furthermore, these lncRNAs do not appear to control the transcript levels of proximal TFs.

**LINC00261 knockout impairs endocrine cell development**

The exception was the endoderm-specific lncRNA LINC00261, which is highly expressed and translated in pancreatic progenitors (Figure 4—figure supplement 1A and Figure 2C). While deletion of LINC00261 caused no discernable phenotype in definitive endoderm (Figure 3C–F and Figure 3—figure supplement 1C), we observed a significant 30–50% reduction in the number of insulin+ cells at the endocrine cell stage (Figure 4A,B). This reduction in insulin+ cell numbers was consistent.
Figure 4. LINC00261 deletion impedes pancreatic endocrine cell differentiation. (A) Flow cytometry analysis at endocrine cell stage (EC) for insulin (INS) in control (ctrl) and LINC00261-/- H1 hESCs. Top panel: Schematic of the LINC00261 locus. The dashed box represents the genomic deletion. Middle panel: The line demarks isotype control. Percentage of cells expressing INS is indicated (representative experiment, n = 4 deletion clones generated with independent sgRNAs). Bottom panel: Bar graph showing percentages of INS-positive cells. Data are shown as mean ± S.D. (n = 5 (clone

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across four separately derived LINC00261 KO hESCs lines, each independently differentiated to endocrine cell stage 5–8 times. In agreement with reduced insulin cell numbers, insulin content and insulin mRNA levels were also reduced in LINC00261 KO endocrine stage cultures (Figure 4C,D). Analysis of insulin median fluorescence intensities by flow cytometry further showed no reduction in insulin levels per cell in one LINC00261 KO clone and a mild reduction in the three other clones (Figure 4E), indicating that LINC00261 predominately regulates endocrine cell differentiation rather than maintenance of insulin production in beta cells.

To determine the molecular effects of LINC00261 deletion, we performed RNA-seq in pancreatic progenitors derived from LINC00261 KO and control hESCs. Similar to the absence of cis-regulatory functions observed in the other IncRNA KOs, we found no evidence for cis-regulation of FOXA2 by LINC00261 (Figure 4F and Figure 4—figure supplement 1C). However, we observed downregulation of the TFs MAFB and PAX4 (Figure 4F, Figure 4—figure supplement 1B, Figure 4—source data 1A), which are important regulators of beta cell differentiation (Artner et al., 2007; Sosa-Pineda et al., 1997). Of note, genes differentially expressed in LINC00261 KO cells mapped to all chromosomes and showed no enrichment for chromosome 20 where LINC00261 resides (Figure 4G). These results suggest a trans- rather than cis-regulatory function for LINC00261, consistent with its predominantly cytosolic localization, translation, and diffuse distribution within the nucleus (Figure 2C and Figure 4—figure supplement 1D). Trans-regulatory roles of LINC00261 have also been observed in previous studies (Aguet et al., 2019; Shi et al., 2019; Wang et al., 2019; Wang et al., 2017; Yan et al., 2019). This potential trans-functionality prompted us to further investigate whether LINC00261’s coding or noncoding features are essential for endocrine cell differentiation.
One-by-one disruption of LINC00261’s sORFs does not impact endocrine cell differentiation

We established that LINC00261 harbors multiple distinct and highly translated sORFs, which raises the possibility that the translation of these sORFs is functionally important for endocrine cell differentiation. To systematically discriminate LINC00261’s coding and noncoding roles, we individually mutated its seven most highly translated sORFs independently in hESCs, leaving the lncRNA sequence, and hence any noncoding function coupled to RNA sequence or structure, grossly intact.

Each of these hESC lines either carried a homozygous frameshift mutation near the microprotein’s N-terminus (for sORFs 1–6) or a full sORF deletion (sORF7; Figure 4—source data 1B). After verifying that CRISPR editing of the LINC00261 locus did not impact LINC00261 transcript levels (Figure 4—figure supplement 1E), we quantified (i) insulin mRNA levels, (ii) insulin+ cells, and (iii) total insulin content in endocrine cell stage cultures. We observed no difference between sORF loss-of-function and control hESC lines for most of these endpoints (Figure 4H,I and Figure 4—figure supplement 1E), although we noticed that the number of insulin+ cells, but not the amount of insulin produced, was reduced in one of the two sORF4 and sORF7 KO clones. Transcriptome analysis of pancreatic progenitors with frameshifts in sORF3 (the most highly translated LINC00261-sORF; Figure 2C and Figure 2—source data 1D) revealed no differentially expressed genes between LINC00261-sORF3 frameshift and control cells (Figure 4J and Figure 4—source data 1C), contrasting observations in LINC00261 RNA KO pancreatic progenitors (Figure 4F and Figure 4—source data 1A). These results indicate that there is not one dominant LINC00261 sORF that is required for endocrine cell formation, suggesting a functional role of the LINC00261 transcript and not the individual sORFs mutated here. However, it is possible that the different sORFs, or the microproteins translated from these sORFs, are functionally redundant and capable of phenotypic rescue.

It has been suggested that ribosome association can control lncRNA transcript levels by inducing nonsense-mediated decay (NMD) (Carlevaro-Fita et al., 2016; Tani et al., 2013). Therefore, we determined whether the presence of multiple sORFs could regulate LINC00261 stability. To this end, we simultaneously mutated start codons of all seven sORFs (ΔATG+sORF1-7 LINC00261) and expressed either wild type or ΔATG+sORF1-7 LINC00261 ectopically in HEK293T cells, where LINC00261 is normally not expressed. LINC00261 half-life measurements upon transcriptional inhibition with actinomycin D revealed no difference in LINC00261 levels between wild type and ΔATG+sORF1-7 LINC00261 (Figure 4K), suggesting that the translation of the seven sORFs does not reduce LINC00261 transcript stability.

In sum, through the systematic, one-by-one removal of sORFs within a highly translated lncRNA with functional importance for pancreatic endocrine cell formation, we found no evidence to implicate the individual sORFs, or the microproteins they produce, in endocrine cell development. Although LINC00261’s sORFs may share functional redundancy or have developmental roles that do not affect the production of insulin+ cells, our findings strongly suggest that by themselves, these sORFs are not functionally required for endocrine cell formation.

Discussion

Limited cis-regulatory consequences of lncRNA deletion

In this study we globally characterized molecular features of lncRNAs expressed during progression of hESCs toward the pancreatic lineage, including their subcellular localization and potential to be translated. We performed a phenotypic CRISPR loss-of-function screen, focusing on ten developmentally regulated, highly expressed, and highly translated lncRNAs proximal to TFs known to regulate pancreas development. The first important observation from this screen is that we found no evidence to implicate the lncRNAs LINC00261, DIGIT, GATA6-AS1, SOX9-AS1, and RP11-445F12.1 in the cis-regulation of their neighboring TFs FOXA2, GSC, GATA6, SOX9, and LHX, respectively, despite tight transcriptional coregulation of the lncRNA-TF pairs.

Contrasting our findings, a number of studies have reported cis-regulation of FOXA2 by LINC00261 (Amaral et al., 2018; Jiang et al., 2015; Swarr et al., 2019). However, several lines of evidence strongly support the conclusion that FOXA2 is not regulated by LINC00261 in our experimental system. First, we examined FOXA2 mRNA expression in LINC00261+/− cells at both the definitive endoderm and pancreatic progenitor cell stages. Second, we analyzed FOXA2 expression
using two independent methods, namely qRT-PCR and RNA-seq. Third, immunofluorescence staining in definitive endoderm revealed no difference in FOXA2 protein expression between control and LINC00261/− cells.

While different cellular contexts and species could explain the discrepancy between our findings and the ones by Amaral et al., 2018 and Swarr et al., 2019, Jiang et al., 2015 reported FOXA2 regulation by LINC00261 in hESC-derived definitive endoderm. One important difference between our study and the study by Jiang et al. is that we employed CRISPR-Cas9-mediated deletion, whereas Jiang et al. used shRNA-mediated knockdown to inactivate LINC00261. It is possible that lncRNA deletion triggers compensatory mechanisms that are not activated after shRNA-mediated knockdown. For coding genes, mutant mRNA degradation has been shown to trigger genetic compensation (El-Brolosy et al., 2019). Another difference between our study and the one by Jiang et al. is that our differentiation protocol was more efficient in generating definitive endoderm. It is conceivable that the stability of the cell fate and identity of neighboring cells could influence how LINC00261 loss-of-function affects gene regulation.

Translation of short, non-canonical ORFs in lncRNAs: regulatory, microprotein-producing, or just tolerated?

Although lncRNAs are now appreciated as a novel and abundant source of sORF-encoded biologically active microproteins (Makarewich and Olson, 2017), it remains largely unknown which translation events lead to the production of microproteins, which solely have regulatory potential, or which have no functional roles, but are not negatively selected against. The cytosolic localization and translation of many RNAs classified as lncRNAs provides a strong rationale for considering both, coding and noncoding functions.

In this study, we identified the translated lncRNA LINC00261 as a novel regulator of pancreatic endocrine cell differentiation, as evidenced by a severe reduction in insulin− cell numbers upon LINC00261 deletion. We show that LINC00261 transcripts are highly abundant in pancreatic progenitors and, albeit present in the nucleus, are predominantly localized to the cytoplasm. Here, they frequently associate with ribosomes which leads to the translation of multiple independent sORFs. We show that the sORFs are capable of producing microproteins with distinct subcellular localizations upon expression in vitro. In contrast to LINC00261 deletion, individual frameshift mutations in each of LINC00261’s sORFs did not impair endocrine cell development, suggesting that the requirement of LINC00261 for endocrine cell development can be uncoupled from the translation of its multiple sORFs. However, this does not exclude the possibility that these sORFs or the microproteins they produce could possess functions that become relevant under specific environmental, developmental, or disease conditions not examined in this study.

We found that mutating all translated LINC00261 sORFs simultaneously, thereby likely reducing LINC00261’s ability to bind ribosomes, did not affect LINC00261 transcript levels in HEK293T cells. This indicates that, in contrast to reports suggesting that translated sORFs can regulate RNA stability by promoting nonsense-mediated RNA decay (Carlevaro-Fita et al., 2016; Tani et al., 2013), the high translation levels and multiple sORFs of LINC00261 are unlikely to be part of a LINC00261 decay pathway. It would have been interesting to determine how concurrent mutation of all sORFs in LINC00261 affects pancreatic cell differentiation. However, given the size of the LINC00261 locus and the many sORFs, such an approach comes with technical challenges and significant caveats.

**LINC00261 - a potential trans regulator of endocrine cell differentiation?**

Several lines of evidence suggest that LINC00261 regulates endocrine cell differentiation in trans: (i) LINC00261 transcripts show a diffuse distribution in multiple subcellular compartments, (ii) genes differentially expressed in LINC00261 KO cells are randomly distributed throughout the genome, (iii) expression of the nearby TF FOXA2 is not affected by LINC00261 deletion. Such a trans regulatory mechanism for LINC00261 is supported by a recent study from the GTEx Consortium, where LINC00261 is highlighted as one of a few lncRNAs that forms a potential trans regulatory hotspot through genetic interactions that influence the expression of multiple distant genes (Aguet et al., 2019). Consistent with its preferential cytosolic localization, and further supporting the notion of a trans regulatory mechanism, LINC00261 has been suggested to regulate gene expression through
non-nuclear mechanisms, e.g. by preventing nuclear translocation of β-catenin (Wang et al., 2017) or by acting as a miRNA sponge (Shi et al., 2019; Wang et al., 2019; Yan et al., 2019). Although our observations and current literature strongly hint to a function in trans independent of the produced microproteins, the exact mechanism by which LINC00261 regulates gene expression in pancreatic progenitors remains to be determined.

Limitations and future directions
In this study, we have characterized the role of translated IncRNAs, and in particular LINC00261, in a hESC differentiation system that mimics pancreas development. However, there are several potential limitations that need to be considered when interpreting the results. First, a small subset of analyses in this study was based on low numbers of replicate differentiations, in particular the cytosolic versus nuclear fractionation RNA-seq experiments, where only two replicate differentiations into pancreatic progenitor cells were analyzed. Second, although we provide evidence that LINC00261 can produce microproteins using Ribo-seq, which is further supported by in vitro translation assays and overexpression of LINC00261 constructs with different in-frame tags, we provide no protein-level evidence for the endogenous production and stability of LINC00261’s microproteins in this differentiation system or in human pancreas development in vivo. Moreover, due to its highly specific expression pattern, LINC00261 has not been previously detected by sORF analyses in other cell types (Bazzini et al., 2014; Calviello et al., 2016; Chen et al., 2020; Ji et al., 2015; Martinez et al., 2020; Prenger et al., 2020; Raj et al., 2016; van Heesch et al., 2019). Even though we show microprotein production in vitro, it is possible that the act of translation has a key regulatory role rather than the protein products of LINC00261’s sORFs. Lastly, LINC00261’s microproteins and sORFs may have redundant functions, which could explain why deletion of individual sORFs produces no apparent phenotype. Thus, despite limited sequence similarity and stark differences in translation rates between the identified translated sORFs in LINC00261, we cannot rule out that different microproteins produced by LINC00261 compensate when one sORF is deleted. Future studies of LINC00261’s precise mechanisms of action could be aimed at further dissecting the potential regulatory features of sORF translation and possibility of redundancy between sORFs.

Conclusions
In summary, we here present a rigorous, in-depth characterization of dynamically regulated and translated IncRNAs in a disease-relevant cell model of human developmental progression. Our combination of ultra-high-coverage RNA- and Ribo-seq, in vitro protein-level validation of microprotein production and localization, and the systematic, one-by-one deletion of all individual microproteins encoded by a single translated IncRNA, not only provides a detailed resource of translated ‘non-canonical’ sORFs and their microproteins in pancreatic development, but also serves as a blueprint for the systematic functional interrogation of translated IncRNAs.

Materials and methods

| Key resources table |
|---------------------|
| **Reagent type**    | **Designation**     | **Source or reference** | **Identifiers** | **Additional information** |
| Gene (Homo sapiens) | LINC00617; TUNAR    | Ensembl 87               | ENSG00000250366 |                         |
| Gene (Homo sapiens) | RP11-445F12.1; LHX-DT | Ensembl 87               | ENSG00000250366 |                         |
| Gene (Homo sapiens) | DIGIT; GSC-DT       | HGNC and NCBI RefSeq    | HGNC:53074; NCBI RefSeq 108868751 |                     |
| Gene (Homo sapiens) | GATA6-AS1; GATA6-AS | Ensembl 87               | ENSG00000277268 |                         |
| Gene (Homo sapiens) | LINC00479         | Ensembl 87               | ENSG00000236384 |                         |

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| Reagent type (species) or resource | Designation | Source or reference | Identifiers | Additional information |
|-----------------------------------|-------------|--------------------|-------------|------------------------|
| Gene (Homo sapiens)               | LINC00261;  | Ensembl 87         | ENSG00000236384 |                        |
|                                   | DEANR1; ALIEN- |                   |             |                        |
|                                   | onco-lncRNA-17;  |                   |             |                        |
|                                   | Inc-FOXA2-2 |                   |             |                        |
| Gene (Homo sapiens)               | RP11-834C11.4 | Ensembl 87       | ENSG00000250742 |                        |
| Gene (Homo sapiens)               | SOX9-AS1     | Ensembl 87         | ENSG00000234899 |                        |
| Gene (Homo sapiens)               | MIR7-3HG     | Ensembl 87         | ENSG000000176840 |                        |
| Gene (Homo sapiens)               | LHFPL3-AS2   | Ensembl 87         | ENSG00000225329 |                        |
| Strain, strain background (Escherichia coli) | Stbl3 | ThermoFisher Scientific | Cat# C737303 | Chemically competent cells |
| Strain, strain background (Escherichia coli) | DH5α | New England Biolabs | Cat# C2987I | Chemically competent cells |
| Cell line (Homo sapiens)          | H1 (embryonic stem cells) | WiCell Research Institute | NIHhESC-10-0043, RRID:CVCL_9771 |                        |
| Cell line (Homo sapiens)          | HEK293T (embryonic kidney) | ATCC | Cat# CRL-3216, RRID:CVCL_0063 |                        |
| Antibody                          | anti-human OCT-4A (Rabbit monoclonal) | Cell Signaling Technology | Cat# 2890, RRID:AB_2167725 | IF (1:1000) |
| Antibody                          | anti-human SOX17 (Goat polyclonal) | R and D Systems | Cat# AF1924, RRID:AB_355060 | IF (1:250) |
| Antibody                          | anti-human FOXA2 (Goat polyclonal) | Santa Cruz Biotechnology | Cat# sc-6554, RRID:AB_2262810 | IF (1:250) |
| Antibody                          | anti-human GATA6 (Goat polyclonal) | Santa Cruz Biotechnology | Cat# sc-9055, RRID:AB_2108768 | IF (1:50) |
| Antibody                          | anti-human Insulin (Guinea pig polyclonal) | Dako | Cat# A0564, RRID:AB_10013624 | IF (1:1000) |
| Antibody                          | Alexa Fluor 488 AffiniPure anti-Goat IgG (Donkey polyclonal) | Jackson ImmunoResearch Labs | Cat# 706-545-148, RRID:AB_2340472 | IF (1:1000) |
| Antibody                          | Alexa Fluor 488 AffiniPure anti-Rabbit IgG (Donkey polyclonal) | Jackson ImmunoResearch Labs | Cat# 711-545-152, RRID:AB_2313584 | IF (1:1000) |
| Antibody                          | Cy3 AffiniPure anti-Goat IgG (Donkey polyclonal) | Jackson ImmunoResearch Labs | Cat# 705-165-147, RRID:AB_2307351 | IF (1:1000) |

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| Reagent type (species) or resource | Designation | Source or reference | Identifiers | Additional information |
|-----------------------------------|-------------|--------------------|-------------|------------------------|
| Antibody                          | anti-human Insulin-PE (Rabbit monoclonal) | Cell Signaling Technology | Cat# 8508S, RRID:AB_11179076 | Flow cytometry (1:50) |
| Antibody                          | anti-human SOX17-PE (Mouse monoclonal) | BD Biosciences | Cat# 561591, RRID:AB_10717121 | Flow cytometry (5 ul per test) |
| Antibody                          | IgG-PE (Rabbit monoclonal) | Cell Signaling Technology | Cat# 5742S, RRID:AB_10694219 | Flow cytometry isotype control antibody (1:50) |
| Antibody                          | IgG1, κ antibody (Mouse monoclonal) | BD Biosciences | Cat# 556650, RRID:AB_396514 | Flow cytometry isotype control antibody (1:50) |
| Recombinant DNA reagent           | pSpCas9(BB)–2A-Puro (PX459, V2.0) | Feng Zhang | RRID:Addgene_62988 | Cas9 from S. pyogenes with 2A-Puro, and cloning backbone for sgRNA |
| Recombinant DNA reagent           | pSpCas9(BB)–2A-GFP (PX458) | Feng Zhang | RRID:Addgene_48138 | Cas9 from S. pyogenes with 2A-EGFP, and cloning backbone for sgRNA |
| Recombinant DNA reagent           | pCMVR8.74 | Didier Trono | RRID:Addgene_22036 | 2nd generation lentiviral packaging plasmid |
| Recombinant DNA reagent           | pMD2.G | Didier Trono | RRID:Addgene_12259 | VSV-G envelope expressing plasmid |
| Recombinant DNA reagent           | pENTR-D-TOPO-LINC00261 | Kurian et al., 2015 (PMID:25739401) | Leo Kurian (University of Cologne) |
| Recombinant DNA reagent           | pRRLSIN.cPPT.PGK-GFP.WPRE | Didier Trono | RRID:Addgene_12252 | 3rd generation lentiviral backbone |
| Recombinant DNA reagent           | pRRLSIN.cPPT.PGK-LINC00261.WPRE | This paper | Transient (transfection) or stable (lentiviral integration) expression of wild type LINC00261 |
| Recombinant DNA reagent           | pRRLSIN.cPPT.PGK-ΔATG<sup>ORF1-7</sup>LINC00261.WPRE | This paper | Transient (transfection) or stable (lentiviral integration) expression of ΔATG<sup>ORF1-7</sup>LINC00261 |
| Recombinant DNA reagent           | pRRLSIN.cPPT.PGK-LINC00261-sORF1-GFP.WPRE | This paper | Transient (transfection) or stable (lentiviral integration) expression of LINC00261-sORF1-GFP fusion protein |
| Recombinant DNA reagent           | pRRLSIN.cPPT.PGK-LINC00261-sORF2-GFP.WPRE | This paper | Transient (transfection) or stable (lentiviral integration) expression of LINC00261-sORF2-GFP fusion protein |
| Recombinant DNA reagent           | pRRLSIN.cPPT.PGK-LINC00261-sORF3-GFP.WPRE | This paper | Transient (transfection) or stable (lentiviral integration) expression of LINC00261-sORF3-GFP fusion protein |

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| Reagent type                  | Designation                                                                 | Source or reference | Identifiers                                                                 | Additional information                                                                 |
|------------------------------|-----------------------------------------------------------------------------|--------------------|-----------------------------------------------------------------------------|----------------------------------------------------------------------------------------|
| Recombinant DNA reagent      | pRRLSIN.cPPT. PGK-LINC00261-sORF4-GFP.WPRE                                   | This paper         |                                                                             | Transient (transfection) or stable (lentiviral integration) expression of LINC00261-sORF4-GFP fusion protein |
| Recombinant DNA reagent      | pRRLSIN.cPPT. PGK-LINC00261-sORF5-GFP.WPRE                                   | This paper         |                                                                             | Transient (transfection) or stable (lentiviral integration) expression of LINC00261-sORF5-GFP fusion protein |
| Recombinant DNA reagent      | pRRLSIN.cPPT. PGK-LINC00261-sORF6-GFP.WPRE                                   | This paper         |                                                                             | Transient (transfection) or stable (lentiviral integration) expression of LINC00261-sORF6-GFP fusion protein |
| Recombinant DNA reagent      | pRRLSIN.cPPT. PGK-LINC00261-sORF7-GFP.WPRE                                   | This paper         | LINC00261-sORF3-frameshift-GFP control plasmid                              | Transient (transfection) or stable (lentiviral integration) expression of LINC00261-sORF7-GFP fusion protein |
| Recombinant DNA reagent      | pRRLSIN.cPPT. PGK-LINC00261-sORF2-STOP-GFP.WPRE                             | This paper         | LINC00261-sORF2-STOP-GFP control plasmid                                     | Transient (transfection) or stable (lentiviral integration) expression of RP11-834C11.4-sORF-GFP fusion protein |
| Recombinant DNA reagent      | pRRLSIN.cPPT. PGK-RP11-834C11.4-sORF-GFP.WPRE                               | This paper         |                                                                             | Transient (transfection) or stable (lentiviral integration) expression of RP11-834C11.4-sORF-GFP fusion protein |
| Recombinant DNA reagent      | pRRLSIN.cPPT. PGK-RP11-834C11.4-sORF-FLAG.WPRE                              | This paper         |                                                                             | Transient (transfection) or stable (lentiviral integration) expression of RP11-834C11.4-sORF-FLAG fusion protein |
| Recombinant DNA reagent      | pRRLSIN.cPPT. PGK-MIR7-3HG-sORF-GFP.WPRE                                    | This paper         |                                                                             | Transient (transfection) or stable (lentiviral integration) expression of MIR7-3HG-sORF-GFP fusion protein |
| Recombinant DNA reagent      | pRRLSIN.cPPT. PGK-RP11-LHFPL3-AS2-sORF-GFP.WPRE                             | This paper         |                                                                             | Transient (transfection) or stable (lentiviral integration) expression of LHFPL3-AS2-sORF-GFP fusion protein |
| Peptide, recombinant protein | Activin A                                                                   | R and D Systems    | Cat# 338-AC/CF                                                              |                                                                                         |
| Peptide, recombinant protein | Wnt3a                                                                       | R and D Systems    | Cat# 1324-WN-010                                                            |                                                                                         |
| Peptide, recombinant protein | KGF/FGF7                                                                     | R and D Systems    | Cat# 251 KG                                                                 |                                                                                         |

Continued on next page
| Reagent type (species) or resource | Designation | Source or reference | Identifiers | Additional information |
|-----------------------------------|-------------|-------------------|------------|------------------------|
| Peptide, recombinant protein      | Noggin      | R and D Systems   | Cat# 3344 NG-050 |                        |
| Commercial assay or kit           | RNase-Free DNase Set (50) | QIAGEN | Cat# 79254 |                        |
| Commercial assay or kit           | TruSeq Ribosome Profile (Mammalian) Library Prep Kit | illumina | Cat# RPYSC12116 |                        |
| Commercial assay or kit           | TruSeq Stranded mRNA Library Prep | illumina | Cat# 20020594 |                        |
| Commercial assay or kit           | TruSeq Stranded Total RNA Library Prep Gold | illumina | Cat# 20020599 |                        |
| Commercial assay or kit           | KAPA mRNA HyperPrep Kit | Roche | Cat# KK8581 |                        |
| Commercial assay or kit           | Qubit ssDNA assay kit | Thermo Fisher Scientific | Cat# Q10212 |                        |
| Commercial assay or kit           | KOD Xtreme DNA Hotstart Polymerase | Millipore | Cat# 71975 |                        |
| Commercial assay or kit           | GoTaq Green Master Mix | Promega | Cat# M7123 |                        |
| Commercial assay or kit           | TOPO TA Cloning Kit | Thermo Fisher Scientific | Cat# K452001 |                        |
| Commercial assay or kit           | Monarch Plasmid Miniprep Kit | NEB | Cat# T1010L |                        |
| Commercial assay or kit           | MinElute PCR Purification Kit | QIAGEN | Cat# 28006 |                        |
| Commercial assay or kit           | iScript cDNA Synthesis Kit | Bio-Rad | Cat# 1708890 |                        |
| Commercial assay or kit           | iQ SYBR Green Supermix | Bio-Rad | Cat# 1708880 |                        |

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| Reagent type (species) or resource | Designation | Source or reference | Identifiers | Additional information |
|-----------------------------------|-------------|--------------------|-------------|-------------------------|
| Commercial assay or kit           | Human Stem Cell Nucleofector Kit 2 | Lonza | Cat# VPH-5022 |                        |
| Commercial assay or kit           | XtremeGene 9 DNA Transfection Reagent | Sigma-Aldrich | Cat# 06365779001 |                        |
| Commercial assay or kit           | Cytofix/Cytoperm W/Golgi Stop Kit | BD Biosciences | Cat# 554715 |                        |
| Commercial assay or kit           | Insulin ELISA Jumbo | Alpco | Cat# 80-INSHU-E10.1 |            |
| Commercial assay or kit           | Pierce BCA Protein Assay Kit | Thermo Fisher Scientific | Cat# 23227 |                        |
| Commercial assay or kit           | TnT Coupled Wheat Germ Extract System | Promega | Cat# L4130 |                        |
| Chemical compound, drug           | Penicillin-Streptomycin | Thermo Fisher Scientific | Cat# 15140122 |                        |
| Chemical compound, drug           | Puromycin dihydrochloride | Sigma-Aldrich | Cat# P8833 |                        |
| Chemical compound, drug           | ALK5 Inhibitor II | Enzo Life Sciences | Cat# ALX-270–445 |                        |
| Chemical compound, drug           | Retinoic Acid | Sigma-Aldrich | Cat# R2625 |                        |
| Chemical compound, drug           | Ascorbic Acid | Sigma-Aldrich | Cat# A4403-100MG |                        |
| Chemical compound, drug           | LDN-193189 | Stemgent | Cat# 04–0074 |                        |
| Chemical compound, drug           | SANT-1 | Sigma-Aldrich | Cat# 54572 |                        |
| Chemical compound, drug           | TPB | Calbiochem | Cat# 565740 |                        |
| Chemical compound, drug           | TGFβ R1 kinase inhibitor IV | EMD Biosciences | Cat# 616454 |                        |
| Chemical compound, drug           | KAAD-Cyclopamine | Toronto Research Chemicals | Cat# K171000 |                        |
| Chemical compound, drug           | TTNPB | Sigma-Aldrich | Cat# T3757 |                        |
| Chemical compound, drug           | Cycloheximide | Sigma-Aldrich | Cat# C7698 |                        |
| Chemical compound, drug           | Actinomycin D | Sigma-Aldrich | Cat# A9415 |                        |

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| Reagent type (species) or resource | Designation | Source or reference | Identifiers | Additional information |
|----------------------------------|-------------|---------------------|-------------|------------------------|
| Chemical compound, drug          | Polyethylenimine (PEI) | Polysciences | Cat# 23966–1 |
| Chemical compound, drug          | Hoechst 33342, Trihydrochloride, Trihydrate | Thermo Fisher Scientific | Cat# H3570 |
| Chemical compound, drug          | MitoSOX Red | Thermo Fisher Scientific | Cat# M36008 |
| Chemical compound, drug          | D-(+)-Glucose Solution, 45% | Sigma-Aldrich | Cat# G8769 |
| Chemical compound, drug          | Sodium Bicarbonate | Sigma-Aldrich | Cat# NC0564699 |
| Chemical compound, drug          | ROCK Inhibitor Y-27632 | STEMCELL Technologies | Cat# 72305 |
| Software, algorithm              | Flowjo-v10 | FlowJo, LLC | RRID:SCR_008520 | http://www.flowjo.com/download-newest-version/ |
| Software, algorithm              | STAR 2.5.3a | Dobin et al., 2013 | RRID:SCR_015899 | https://github.com/alexdobin/STAR |
| Software, algorithm              | Bowtie 1.1.1 | Langmead et al., 2009 | RRID:SCR_005476 | http://bowtie-bio.sourceforge.net/index.shtml |
| Software, algorithm              | Cufflinks 2.2.1 | Trapnell et al., 2010 | RRID:SCR_014597 | https://github.com/cole-trapnell-lab/cufflinks |
| Software, algorithm              | HTSeq 0.6.1 | Anders et al., 2015 | RRID:SCR_005514 | https://htseq.readthedocs.io/en/master/install.html |
| Software, algorithm              | DEseq2 1.10.1 | Love et al., 2014 | RRID:SCR_015687 | https://www.bioconductor.org/packages/devel/bioc/html/DESeq2.html |
| Software, algorithm              | RiboTaper | Calviello et al., 2016 | RRID:SCR_018880 | https://ohlerlab.mdc-berlin.de/software/ |
| Software, algorithm              | R 3.5.0 | | RRID:SCR_001905 | https://cran.r-project.org/ |
| Software, algorithm              | SAMtools 1.3 | Li et al., 2009 | RRID:SCR_002105 | https://github.com/samtools/samtools |
| Software, algorithm              | BEDTools 2.17.0 | Quinlan and Hall, 2010 | RRID:SCR_006646 | https://bedtools.readthedocs.io/en/latest/content/installation.html |
| Software, algorithm              | HOMER 4.10 | Heinz et al., 2010 | RRID:SCR_010881 | http://homer.ucsd.edu/homer/download.html |
| Software, algorithm              | GREAT 3.0.0 | McLean et al., 2010 | RRID:SCR_005807 | http://great.stanford.edu/public/html/ |

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| Reagent type (species) or resource | Designation | Source or reference | Identifiers | Additional information |
|-----------------------------------|-------------|---------------------|-------------|------------------------|
| Software, algorithm               | Adobe Illustrator CS5 | Adobe | RRID:SCR_010279       |
| Software, algorithm               | Adobe Photoshop CS5  | Adobe | RRID:SCR_014199       |
| Software, algorithm               | GraphPad Prism v7.05 | GraphPad Software, LLC | RRID:SCR_002798 |
| Other                             | Novex 16% Tricine Protein Gel | Thermo Fisher Scientific | Cat# EC66955BOX |
| Other                             | Novex Tricine SDS Sample Buffer (2X) | Thermo Fisher Scientific | Cat# LC1676 |
| Other                             | Immobilon-PSQ PVDF Membrane | Merck Millipore | Cat# ISEQ00010 |
| Other                             | Stellaris RNA FISH Hybridisation Buffer | LGC Biosearch Technologies | Cat# SMF-HB1-10 |
| Other                             | Stellaris RNA FISH Wash Buffer A | LGC Biosearch Technologies | Cat# SMF-WA1-60 |
| Other                             | Stellaris RNA FISH Wash Buffer B | LGC Biosearch Technologies | Cat# SMF-WB1-20 |
| Other                             | QuickExtract DNA Extraction Solution | Lucigen | Cat# QE09050 |
| Other                             | Vectashield Antifade Mounting Medium | Vector Laboratories | Cat# H-1000 |
| Other                             | FastDigest BpiI | Thermo Fisher Scientific | Cat# FD1014 |
| Other                             | FastDigest BshTI | Thermo Fisher Scientific | Cat# FERFD1464 |
| Other                             | FastDigest SalI | Thermo Fisher Scientific | Cat# FD0644 |
| Other                             | TRIzol | Thermo Fisher Scientific | Cat# 15596018 |
| Other                             | Matrigel | Corning | Cat# 356231 |
| Other                             | GlutaMAX | Thermo Fisher Scientific | Cat# 35050061 |
| Other                             | DPBS (without calcium and magnesium) | Corning | Cat# 45000–434 |
| Other                             | mTeSR1 Complete Kit - GMP | STEMCELL Technologies | Cat# 85850 |
| Other                             | RPMI 1640 Medium, HEPES | Thermo Fisher Scientific | Cat# 22400–089 |
| Other                             | DMEM/F12 with L-Glutamine, HEPES | Corning | Cat# 45000–350 |

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HEK293T cell culture

HEK293T cells (female) were cultured in a humidified incubator at 37°C with 5% CO₂ using Dulbecco’s Modified Eagle Medium (Corning; 4.5 g/L glucose, [+L-glutamine, [-sodium pyruvate) supplemented with 10% fetal bovine serum (FBS; Corning, Cat# 35011CV). HEK293T cells were purchased from ATCC (Cat# CRL-3216, RRID:CVCL_0063) and tested for mycoplasma prior to the experiment.

hESC culture and maintenance

H1 hESCs (male) were obtained from WiCell (NIHhESC-10–0043, RRID:CVCL_9771) and tested for mycoplasma on a yearly basis. H1 hESCs were grown in feeder-independent conditions on Matrigel-coated dishes (Corning, Cat# 356231) with mTeSR1 media (STEMCELL Technologies, Cat# 85850). Propagation was carried out by passing the cells every 3 to 4 days using Accutase (eBioscience, Cat# 00-4555-56) for enzymatic cell dissociation. hESC research was approved by the University of California, San Diego, Institutional Review Board and Embryonic Stem Cell Research Oversight Committee.

Pancreatic differentiation

H1 hESCs were differentiated in a monolayer format as previously described (Rezania et al., 2012), with minor modifications. Undifferentiated hESCs were seeded into 24-wells at 0.4 x 10⁶ cells/well in 500 µl mTeSR1 medium. The next day the cells were washed in RPMI media (Thermo Fisher Scientific, Cat# 22400–089) and then differentiated with daily media changes. In addition to GlutaMAX (Thermo Fisher Scientific, Cat# 35050061), RPMI medium was supplemented with 0.12% (w/v) NaHCO₃ and 0.2% (Day 0) or 0.5% (Day 1–3) (v/v) FBS (Corning, Cat# 35011CV). DMEM/F12 medium (Corning, Cat# 45000–350) was supplemented with 2% (v/v) FBS and 0.2% (w/v) NaHCO₃, and DMEM High Glucose medium (HyClone, Thermo Fisher Scientific, Cat# SH30081.FS) was
supplemented with 0.5X B-27 supplement (Thermo Fisher Scientific, Cat# 17504044). Human Activin A, mouse Wnt3a, human KGF, and human Noggin were purchased from R and D Systems (Cat# 338-AC/CF, Cat# 1324-WN-010, Cat# 251 KG, Cat# 3344 NG-050). Other media components included TGFβ R1 kinase inhibitor IV (EMD Bioscience, Cat# 616454), KAAD-Cyclopamine (Toronto Research Chemicals, Cat# K171000), the retinoid analog TTNPB (Sigma Aldrich, Cat# T3757), the protein kinase C activator TPB (EMD Chemicals, Cat# 565740), the BMP type one receptor inhibitor LDN-193189 (Stemgent, Cat# 04-0074), and an inhibitor of the TGF-β type one activin like kinase receptor ALK5, ALK5 inhibitor II (Enzo Life Sciences, Cat# ALX-270–445).

Stage 1 (DE; collection on day 3):
Day 0: RPMI/FBS, 100 ng/mL Activin A, 25 ng/mL mouse Wnt3a
Day 1–2: RPMI/FBS, 100 ng/mL Activin A
Stage 2 (GT; collection on day 6):
Day 3: DMEM/F12/FBS, 2.5 μM TGFβ R1 kinase inhibitor IV, 50 ng/mL KGF
Day 4–5: DMEM/F12/FBS, 50 ng/mL KGF
Stage 3 (PP1, collection on day 10):
Day 6–9: DMEM/B27, 3 nM TTNPB, 0.25 mM KAAD-Cyclopamine, 50 ng/mL Noggin
Stage 4 (PP2; collection on day 13):
Day 10–12: DMEM/B27, 100 nM ALK5 inhibitor II, 100 nM LDN-193189, 500 nM TPB, 50 ng/mL Noggin
Stage 5 (endocrine cell stage; collection on day 16):
Day 13–15: DMEM/B27, 100 nM ALK5 inhibitor II, 100 nM LDN-193189, 500 nM TPB, 50 ng/mL Noggin

For ribosome profiling experiments, a scalable suspension culture protocol was employed for differentiation of H1 cells to the PP2 stage (Rezania et al., 2014). Undifferentiated hESCs were aggregated by preparing a single cell suspension in mTeSR1 media (STEMCELL Technologies; supplemented with 10 μM Y-27632) at $1 \times 10^6$ cells/mL and overnight culture in six-well ultra-low attachment plates (Costar) with 5.5 ml per well on an orbital rotator (Innova2000, New Brunswick Scientific) at 100 rpm. The following day, undifferentiated aggregates were washed in MCDB 131 media (Thermo Fisher Scientific, Cat# 10372–019) and then differentiated using a multistep protocol with daily media changes and continued orbital rotation at either 100 rpm or at 115 rpm from days 8 to 14. In addition to 1% GlutaMAX (Gibco, Thermo Fisher Scientific, Cat# 35050061) and 10 mM (days 0–10) or 20 mM (days 11–14) glucose, MCDB 131 media was supplemented with 0.5% (days 0–5) or 2% (days 6–14) fatty acid-free BSA (Proliant Biologicals, Cat# 68700), 1.5 g/L (days 0–5 and days 11–14) or 2.5 g/L (days 6–10) NaHCO$_3$ (Sigma-Aldrich), and 0.25 mM ascorbic acid (days 3–10).

Human Activin A, mouse Wnt3a, and human KGF were purchased from R and D Systems (Cat# 338-AC/CF, Cat# 1324-WN-010, Cat# 251 KG). Other media components included Insulin-Transferin-Selenium-Ethanolamine (ITS-X; Thermo Fisher Scientific, Cat# 51500–056; days 6–14), retinoic acid (RA) (Sigma-Aldrich, Cat# R2625), the sonic hedgehog pathway inhibitor SANT-1 (Sigma-Aldrich, Cat# S4572), the protein kinase C activator TPB (EMD Chemicals, Cat# 565740), the BMP type one receptor inhibitor LDN-193189 (Stemgent, Cat# 04–0074), and the TGFβ type one activin like kinase receptor ALK5, ALK5 inhibitor II (Enzo Life Sciences, Cat# ALX-270–445).

Stage 1 (DE; collection on day 3):
Day 0: MCDB 131, 100 ng/mL Activin A, 25 ng/mL mouse Wnt3a
Day 1–2: MCDB 131, 100 ng/mL Activin A
Stage 2 (GT; collection on day 6):
Day 3 – Day 5: MCDB 131, 50 ng/mL KGF
Stage 3 (PP1; collection on day 8):
Day 6 – Day 7: MCDB 131, 50 ng/mL KGF, 0.25 μM SANT-1, 1 μM RA 100 nM LDN-193189, 200 nM TPB
Stage 4 (PP2; collection on day 11):
Day 8 – Day 10: MCDB 131, 2 ng/mL KGF, 0.25 μM SANT-1, 0.1 μM RA, 200 nM LDN-193189, 100 nM TPB

**CRISPR/Cas9-mediated IncRNA knockout**

To generate clonal IncRNA knockout hESC lines, combinations of pSpCas9(BB)–2A-Puro plasmid pairs (Addgene plasmid #62988, RRID:Addgene_62988, gift from Feng Zhang) expressing Cas9 and
single sgRNAs targeting upstream and downstream regions of the IncRNA promoter/locus were co-transfected into 1.5 × 10^6 H1 hESCs using the Human Stem Cell Nucleofector Kit 2 (Lonza) and the Amazix Nucleofector II (Lonza). 24 hr after plating into Matrigel-coated six-well plates, nucleofected cells were selected with puromycin (1 μg/mL mTeSR1 media) for 2–3 consecutive days. Individual colonies that emerged within 7 days after transfection were subsequently transferred manually into 96-well plates for expansion. Genomic DNA for PCR genotyping with GoTaq Green Mastermix (Promega) and Sanger sequencing was then extracted using QuickExtract DNA Extraction Solution (Lucigen).

To generate sORF frameshift mutations, sgRNA sequences targeting the N-terminal region of the predicted small peptides were inserted into pSpCas9(BB)-2A-GFP (Addgene plasmid #48138, RRID: Addgene_48138, gift from Feng Zhang) via its BpiI cloning sites. 3 μg of the resulting plasmids were then transfected into 500,000 H1 cells plated into Matrigel-coated six-wells the day prior, using XtremeGene 9 Transfection Reagent (Sigma-Aldrich) according to the manufacturer’s instructions. 24 hr post-transfection, 10,000 GFP^+ cells were sorted on an Influx Cell Sorter (BD Biosciences) into Matrigel-coated six-wells containing 1 mL mTeSR1 media supplemented with 10 μM ROCK inhibitor and 1X penicillin/streptomycin. Seven days after sorting, emerging colonies were hand-picked and transferred into 96-well plates for genotyping. Frameshifts inside the targeted sORFs were confirmed by PCR-amplification of the sORF sequence with GoTaq Green Mastermix (Promega, Cat# M7123) and subsequent subcloning the PCR products into pCR2.1 (Thermo Fisher Scientific). For each hESC clone, at least six pCR2.1 clones were Sanger sequenced. Oligonucleotide sequences for sgRNA cloning are provided in Figure 4—source data 2A.

**PCR genotyping of CRISPR clones**

Four days after transfer of single cell-derived clones into 96-wells, cell culture supernatants containing dead cells were collected from each well prior to the daily media change. Cell debris was then pelleted and used for gDNA extraction with 10–20 μl QuickExtract DNA Extraction Solution (Lucigen, Cat# QE09050) according to the manufacturer’s instructions. 1 μl DNA was then PCR-amplified with GoTaq Green Mastermix (Promega, Cat# M7123) and locus-specific primers that anneal either within or outside of the excised genomic DNA. PCR products generated with ‘inside’ primers were visualized on a 2% agarose gel, PCR bands generated with primers flanking the deletion were gel-purified and submitted for Sanger sequencing (see Figure 4—source data 2B for genotyping and sequencing primers).

For genotyping of sORF frameshift clones, PCR amplicons designed to encompass the Cas9 cut site were amplified and Sanger sequenced (Figure 4—source data 2B). If out-of-frame indels were apparent in the sequencing chromatogram, the sequenced PCR product was ligated into pCR2.1-TOPO via TOPO-TA cloning. A minimum of six clones were Sanger sequenced in order to determine the genotype at both alleles with high confidence.

**Generation of sORF translation reporter plasmids**

The four IncRNAs tested were PCR-amplified with KOD Xtreme DNA Hotstart Polymerase (Millipore) from their 5’ end up until the last codon of the sORF to be tested, omitting its stop codon (primer sequences are listed in Figure 4—source data 2D). cDNA was used as PCR template for LINC00261 and LHFPL3-AS2; RP11-834C11.4, and MIR7-3HG were amplified from a gBlock synthetic gene fragment (Integrated DNA Technologies; see Figure 4—source data 2F). The GFP coding sequence (without start codon; amplified from pRRLSIN.cPPT.PGK-GFP.WPRE; RRID: Addgene_12252) was then fused in-frame to the sORF via overlap extension PCR. The resulting fusion product was cloned into pRRLSIN.cPPT.PGK-GFP.WPRE (Addgene plasmid #12252, gift from Didier Trono) via BshTI and SalI restriction sites. To create the RP11-834C11.4-sORF-1XFLAG reporter construct in an analogous way, a gBlock synthetic gene fragment encompassing the FLAG-tagged sORF served as PCR template (Figure 4—source data 2F). The resulting PCR product was cloned into pRRLSIN.cPPT.PGK-GFP.WPRE via BshTI and Sall restriction sites.
Generation of LINC00261 wild type and ΔATG\textsuperscript{sORF1–7} expression plasmids

The LINC00261 wild type cDNA was PCR-amplified from pENTR/D-TOPO-LINC00261 (gift from Leo Kurian) with KOD Xtreme DNA Hotstart Polymerase (Millipore, Cat# 71975). The resulting PCR product was inserted into pRRLSIN.cPPT.PGK-GFP.WPRE via its appended BshTISalI cloning sites. Full-length LINC00261 ΔATG\textsuperscript{sORF1–7} was assembled through overlap extension PCR from the following three fragments and subsequently cloned into pRRLSIN.cPPT.PGK-GFP.WPRE via appended BshTISalI cloning sites: (i) a 1,248 bp PCR product amplified from a synthetic gene construct (Genewiz; see Figure 4—source data 2F for sequence) in which the ATG start codons of sORFs 1–6 had been mutated (ATG → AAG/ATT/AGG/AAG/ATA/AGG), and (ii–iii) 3,111 bp/610 bp PCR fragments (amplified from the LINC00261 cDNA) in which the sORF7 start codon was mutated (ATG → AAG). The obtained plasmids were sequence-verified by Sanger sequencing.

Immunofluorescence staining

H1 hESC-derived cells grown as monolayer on Matrigel-coated coverslips were washed twice with PBS and then fixed with 4% paraformaldehyde in PBS for 30 min at room temperature. Following three washes in PBS, samples on coverslips were permeabilized and blocked with Permeabilization/Blocking Buffer (0.15% (v/v) Triton X-100% and 1% normal donkey serum in PBS) for 1 hr at room temperature. Primary and secondary antibodies were diluted in Permeabilization/Blocking Buffer. Sections were incubated overnight at 4˚C with primary antibodies, and then secondary antibodies for 30 min at room temperature. The following primary antibodies were used: rabbit anti-OCT4 (Cell Signaling Technology, Cat# 2890, RRID: AB_2167725, 1:500), goat anti-SOX17 (Santa Cruz Biotechnology, Cat# AF1924, RRID:AB_355060, 1:250), goat anti-FOXA2 (Santa Cruz Biotechnology, Cat# sc-6554, RRID:AB_2262810, 1:250), goat anti-GATA6 (Santa Cruz Biotechnology, Cat# sc-9055, RRID:AB_2108768, 1:50), guinea pig anti-insulin (Dako, Cat# A0564, RRID:AB_10013624). Secondary antibodies (1:1000) were Cy3- or Alexafluor488-conjugated antibodies raised in donkey against guinea pig, rabbit or goat (Jackson ImmunoResearch Laboratories, Cat# 706-545-148, RRID:AB_2340472, Cat# 711-545-152, RRID:AB_2313584, Cat# 705-165-147, RRID:AB_2307351). Images were acquired on a Zeiss Axio-Observer-Z1 microscope with a Zeiss AxioCam digital camera, and figures prepared with Adobe Photoshop/Illustrator CS5.

Flow cytometry analysis

For intracellular flow cytometry, single cells were washed three times in FACS buffer (0.1% (w/v) BSA (Thermo Fisher Scientific in PBS) and then fixed and permeabilized with Cytofix/Cytoperm Fixation/Permeabilization Solution (BD Biosciences) for 20 min at 4˚C, followed by two washes in BD Perm/Wash Buffer. Cells were next incubated with either PE-conjugated anti-SOX17 antibody (BD Biosciences; Cat# 561591, RRID:AB_10717121), or PE-conjugated anti-INS antibody (Cell Signaling Technology, Cat# 8508S, RRID:AB_11179076) in 50 ml BD Perm/Wash Buffer for 60 min at 4˚C. Following three washes in BD Perm/Wash Buffer, cells were analyzed on a FACSCanto II (BD Biosciences) cytometer.

Insulin content measurements

To measure total insulin content of endocrine cell stage control and lncRNA KO cells, adherent cultures were enzymatically detached from a 24-well at day 16 of differentiation. Upon quenching with FACS buffer (0.1% (w/v) BSA (Thermo Fisher Scientific in PBS), the cells were pelleted and extracted over night at 4˚C in 100 ml acid-ethanol (2% HCl in 80% ethanol). Insulin was measured by Insulin ELISA (Alpco, Cat# 80-INSHU-E10.1) and normalized to total protein, as quantified with a BCA protein assay (Thermo Fisher Scientific, Cat# 23227).

Quantitative reverse transcription PCR (qRT-PCR)

Total RNA was isolated from hESC-derived cells and HEK293T cells using either TRIzol (Thermo Fisher Scientific, Cat# 15596018) or the RNAeasy Mini Kit (Qiagen, Cat# 15596018), respectively. Upon removal of genomic DNA (TURBO DNA-free Kit, Thermo Fisher Scientific, Cat# AM1907 or RNase-free DNase Set, Qiagen, Cat# 79254) cDNA was synthesized using the iScript cDNA Synthesis Kit (Bio-Rad, Cat# 1708890). PCR reactions were run in triplicate with 6.25–12.5 ng cDNA per
reaction using the CFX96 Real-Time PCR Detection System (BioRad). TATA-binding protein (TBP) was used as endogenous control to calculate relative gene expression using the \( \Delta \Delta Ct \) method. Primer sequences are provided in Figure 4—source data 2C.

**Transient transfection of HEK293T cells with polyethylenimine (PEI)**

Two hours prior to transfection, fresh pre-warmed DMEM medium (Corning, Cat# 45000–312) was added to each well. Transfection mix was prepared by combining PEI (Polysciences Cat# 23966–1) and plasmid DNA (4:1 ratio; 4 \( \mu \)g PEI per 1 \( \mu \)g DNA) in Opti-MEM Reduced Serum Medium (Thermo Fisher Scientific, Cat# 31985062) followed by brief vortexing. After five minutes, the transfection complex was added dropwise to the cells.

**Lentivirus preparation and ectopic LINC00261 expression**

Lentiviral particles were prepared by co-transfecting HEK293T cells (using PEI) with the pCMVR8.74/pMD2.G helper plasmids (Addgene plasmids #22036/12259, RRID:Addgene_22036 and RRID:Addgene_12259, gift from Didier Trono) and with pRRLSIN.cPPT.PGK-GFP.WPRE transfer plasmid (RRID: Addgene_12252), in which the GFP ORF had been replaced with the 4.9 kb LINC00261 cDNA. Virus-containing supernatant was collected for two consecutive days and concentrated by ultracentrifugation for 2 hr at 19,400 rpm using an Optima L-80 XP Ultracentrifuge (Beckman Coulter).

To express LINC00261 (wild type) and LINC00261 (\( \Delta ATG^{SORF1-7} \)) in HEK293T cells, the cells were plated in 6-well plates and transduced with lentivirus the following day. Two days post infection, the cells were passaged for RNA half-life measurements.

**LINC00261 RNA half-life measurement**

HEK293T cells transduced with either LINC00261 (wild type) or LINC00261 (\( \Delta ATG^{SORF1-7} \)) lentivirus were seeded in six 24-wells. 48 hr after plating, cells from one well were collected for RNA isolation as the ‘0 hr’ time point. To the remaining five wells, 100 \( \mu \)l growth media supplemented with 10 \( \mu \)g/ml actinomycin D (Sigma-Aldrich Cat# A9415) were added to inhibit transcription. At 2, 4, 6, 8, and 9 hr following actinomycin D addition, samples were collected for RNA isolation. Total RNA was then reverse transcribed and analyzed by qPCR, where the abundance of each time point was calculated relative to the abundance at the 0 hr time point (\( \Delta Ct \)). The half-life was then determined by non-linear regression (One phase decay; GraphPad Prism).

**Single molecule RNA fluorescence in situ hybridization (smRNA FISH)**

H1-derived PP2 stage cells (control and LINC00261 KO) were cultured on Matrigel-coated 12 mm diameter coverslips in a 24-well plate. Following 10 min fixation in 1 mL Fixation Buffer (3.7% (v/v) formaldehyde in PBS) at room temperature, the cells were washed twice in PBS and subsequently permeabilized in 70% (v/v) ethanol for one hour at 4°C. Following a five minute wash in Stellaris RNA FISH Wash Buffer A (LGC Biosearch Technologies, Cat# SMF-WA1-60; 1:5 diluted concentrate, with 10% (v/v) formamide added), the coverslips were incubated in a humidified chamber at 37°C for 14 hr with probes diluted in Stellaris RNA FISH Hybridisation Buffer (LGC Biosearch Technologies, Cat# SMF-HB1-10; with 10% (v/v) formamide added) to 125 nM. After a 30 min wash at 37°C in Wash Buffer A, the cells were counter-stained with Hoechst 33342 (Thermo Fisher Scientific) for 15 min and washed in RNA FISH Wash Buffer B (LGC Biosearch Technologies, Cat# SMF-WB1-20) for 5 min at room temperature. The coverslips were mounted in Vectashield Mounting Medium (Vector Laboratories, Cat# H-1000) and imaged on an UltraView Vox Spinning Disk confocal microscope (PerkinElmer) using a 100X oil objective.

**In vitro transcription/translation of lncRNAs**

Synthetic gene constructs containing complete transcript isoforms (including the predicted 5’ and 3’ UTR) of four translated lncRNAs (RP11-834C11.4, LINC00261, MIR7-3HG, and LHFPL3-AS2) were produced by Genewiz (constructs available upon request). Microproteins were translated in vitro from 0.5 \( \mu \)g linearized plasmid DNA using the TnT Coupled Wheat Germ Extract system (Promega; Cat# L4140) in the presence of 10 mCi/mL [\( ^{35} \)S]-methionine (Hartmann Analytic) according to manufacturer’s instructions. 5 \( \mu \)l lysate was denatured for 2 min at 85°C in 9.6 \( \mu \)L Novex Tricine SDS Sample Buffer (2X) (Thermo Fisher Scientific; Cat# LC1676) and 1.4 \( \mu \)L DTT (500 mM). Proteins
were separated on 16% Tricine gels (Thermo Fisher Scientific; Cat# EC66955BOX) for 1 hr at 50 V followed by 3.5 hr at 100 V and blotted on PVDF-membranes (Immobilon-PSQ Membrane, Merck Millipore; Cat# ISEQ00010). Incorporation of $^{35}$S-methionine into newly synthesized proteins enabled the detection of translation products by phosphor imaging (exposure time of 1 day).

**In vivo translation assays**

Reporter plasmids were transfected into HEK293T cells using PEI, and 36 hr post transfection live cells were imaged on an EVOS Cell Imaging System (Thermo Fisher Scientific) equipped with a 20X objective. Additional constructs were generated that served as negative controls (no GFP fluorescence): 1) a LINC00261-sORF3-GFP construct with a single ’T’ insertion inside sORF3, causing a frame-shift, 2) a LINC00261-sORF2-GFP construct with a stop codon preceding the GFP coding sequence, and 3) a LINC00261-sORF1-GFP construct with a frame-shift mutation within the GFP coding sequence.

**Stranded mRNA-seq library preparation for lncRNA KOs**

Total RNA from PP2 cells differentiated with the Rezania et al., 2012 protocol was isolated and DNase-treated using either TRIzol (Thermo Fisher Scientific), or the RNAeasy Mini kit (Qiagen) according to the manufacturer’s instructions. RNA integrity (RIN > 8) was verified on the Agilent 2200 TapeStation (Agilent Technologies), and 400 ng RNA was used for multiplex library preparation with the KAPA mRNA HyperPrep Kit (Roche; Cat# KK8581). All libraries were evaluated on TapeStation High Sensitivity DNA ScreenTapes (Agilent Technologies; Cat# 5067–5584) and with the Qubit dsDNA High Sensitivity (Life Technologies; Cat# Q10212) assays for size distribution and concentration prior to pooling the multiplexed libraries for single-end 1 × 51 nt or 1 × 75 sequencing on the HiSeq 2500 or HiSeq 4000 System (Illumina). Libraries were sequenced to a depth of > 20M uniquely aligned reads.

**Cell fractionation and ribo-minus RNA-seq**

H1 hESCs were differentiated to the PP2 stage with the Rezania et al., 2012 protocol, then nuclear and cytosolic RNA was isolated with the Paris Kit (Thermo Fisher Scientific). Unfractionated total RNA was set aside as a control. All samples were DNasel-treated prior to further processing (TURBO DNA-free Kit; Thermo Fisher Scientific). rRNA-depleted total RNA-seq libraries were prepared with TruSeq Stranded Total RNA Library Prep Gold (Illumina; Cat# 2002059), and sequencing was performed on a HiSeq4000 instrument.

**Alignment of lncRNA KO mRNA-seq samples and processing for gene expression analysis**

Using the Spliced Transcripts Alignment to a Reference (STAR) aligner (STAR 2.5.3b; Dobin et al., 2013), sequence reads were mapped to the human genome (hg38/GRCh38) with the Ensembl 87 annotations in 2-pass mapping mode, allowing for up to six mismatches. Cufflinks (part of the Cufflinks version 2.2.1 suite; Roberts et al., 2011; Trapnell et al., 2010), was then used to quantify the abundance of each transcript cataloged in the Ensembl 87 annotations in reads per kilobase per million mapped reads (RPKM). For plotting expression values, a pseudocount of 1 was added to all RPKM values prior to log2-transformation. Genes with RPKM ≥ 1 across two replicates were deemed expressed. Differential gene expression was tested using the DESeq2 v1.10.1 Bioconductor package (Love et al., 2014) with default parameters. Input count files for DESeq2 were created with htsseq-count from the HTSeq Python library (Anders et al., 2015). Genes with a > 2 fold change and an adjusted p-value of <0.01 were considered differentially expressed.

The chromosomal localization of genes differentially expressed upon LINC00261 KO was visualized with the RCircos package in R (https://cran.r-project.org/web/packages/RCircos/index.html).

**LncRNA classifications**

The following transcript biotypes were grouped into the ‘lncRNA’ classification: 3’ overlapping ncRNA, antisense, bidirectional promoter IncRNA, lincRNA, macro IncRNA, non coding, processed transcript, sense intronic, sense overlapping, TEC.
LncRNAs with ≥1 RPKM during all differentiation stages of CyT49 hESCs (ES, DE, FG, GT, PP1, PP2) were categorized as constitutively expressed ('constitutive'), whereas lncRNAs with <1 RPKM throughout differentiation were considered 'never expressed'. LncRNAs expressed in at least one of the stages (but not in all five stages) were referred to as dynamically expressed ('dynamic'). Furthermore, for each lncRNA, its maximum RPKM value was determined across 38 tissues/cell types (see ‘Gene-gene correlations and GO enrichment’ section below). Log₂-transformed maximum expression values (RPKM + pseudocount of 1) were graphed as boxplots for different gene sets using the ggplot2 R package (https://cran.r-project.org/web/packages/ggplot2/index.html).

To determine the subcellular localization of lncRNAs, first all lncRNAs expressed in the nuclear and/or cytosolic RNA fraction (RPKM ≥ 1 in two biological replicates) of H1-derived PP2 stage cells were selected. Among these expressed lncRNAs, those with ≥1 RPKMcytosol and <1 RPKMnucleus were classified as 'cytosol enriched'. Conversely, lncRNAs with <1 RPKMcytosol and ≥1 RPKMnucleus were termed ‘nucleus enriched’. LncRNAs expressed in both fractions (≥1 RPKMcytosol and ≥1 RPKMnucleus) were tagged with 'both'.

**Assignment of lncRNAs to their nearest coding gene using GREAT**

GREAT (Genomic Regions Enrichment of Annotations Tool 3.0.0; McLean et al., 2010) was run with the ‘Single nearest gene’ within 1000 kb option to assign the nearest coding genes to the following sets of lncRNAs: i) DE-transcribed lncRNAs, ii) PP2-transcribed lncRNAs that are not transcribed at the DE stage (non-transcribed control set for i)), iii) PP2-transcribed lncRNAs, and iv) lncRNAs transcribed at the DE stage but not transcribed in PP2 cells (non-transcribed control set for iii)). The log₂-transformed RPKM values of the lncRNA-associated coding genes were then graphed as boxplots using ggplot2. The corresponding absolute coding-to-lncRNA inter-gene distances were visualized as cumulative frequency plots.

**Gene-gene correlations and GO enrichment**

Pearson correlations were calculated among all genes across a catalog of 38 tissues/cell types derived from all three germ layers (16 Illumina BodyMap 2.0 tissues, other publicly available data sets (see ‘Data sources’ below), and EndoC-βH1 RNA-seq data generated in our lab). Scatter plots of the log₂-transformed RPKM values for lncRNAs/neighbor TFs and histograms of the Pearson correlation coefficients were plotted in R using ggplot2.

Spearman correlations were calculated to test for expression coregulation among all genes expressed (RPKM ≥ 1) in a minimum of ten out of 38 tissues. The resulting correlation matrix was then used to calculate the Euclidean distance followed by hierarchical clustering. The resulting heatmap was subdivided into ten clusters. Cluster visualization was done using heatmap.3 (https://raw.githubusercontent.com/obigriffith/biostar-tutorials/master/Heatmaps/heatmap.3.R) from gplots v3.0.1 (http://cran.r-project.org/web/packages/gplots/index.html). GO enrichment (Ashburner et al., 2000; The Gene Ontology Consortium, 2019) and KEGG pathway (Kanehisa et al., 2017) analyses to assign functional annotation to all ten clusters were performed with gProfiler v0.6.4 (Reimand et al., 2016) using g:Profiler archive revision 1741 (Ensembl 90, Ensembl Genomes 38).

**Alignment and processing of ChIP-seq samples**

All sequence reads were filtered to include only those passing the standard Illumina quality filter, and then aligned to the Homo sapiens reference genome (hg38/GRCh38) using Bowtie version 1.1.1 (Langmead et al., 2009). The following parameters were used to select only uniquely aligning reads with a maximum of two mismatches:

\[-k1 -m1 -l50 -n2 –best –strata\]

SAMtools (Li et al., 2009) was then used to filter reads with a MAPQ score less than 30 and to remove duplicate reads. Finally, replicate ChIP-seq and input BAM files were merged and sorted. The HOMER makeUCSCfile function (Heinz et al., 2010) was used to create a bedGraph formatted file for viewing in the UCSC Genome Browser.
Ribosome profiling and matching RNA-seq

Ribosome profiling was performed on PP2 cells obtained from six independent differentiations of H1 hESCs with the *Rezania et al., 2014* protocol, yielding an average of 89% PDX1-positive cells. Ribosome footprinting and sequencing library preparation was done with the TruSeq Ribo Profile (Mammalian) Library Prep Kit (Illumina, Cat# RPYS12116, currently out of production) according to the TruSeq Ribo Profile (Mammalian) Reference Guide (version August 2016). In short, 50 mg of PP2 aggregates were washed twice with cold PBS and lysed for 10 min on ice in 1 mL lysis buffer (1 × TruSeq Ribo Profile mammalian polysome buffer, 1% Triton X-100, 0.1% NP-40, 1 mM dithiothreitol, 10 U ml⁻¹ DNase I, cycloheximide (0.1 mg/ml) and nuclease-free H₂O). Per sample, 400 μL of lysate was further processed according to manufacturer’s instructions. Final library size distributions were checked on the Bioanalyzer 2100 using a High Sensitivity DNA assay (Agilent Technologies), multiplexed and sequenced on an Illumina HiSeq 4000 producing single end 1 × 51 nt reads. Ribo-seq libraries were sequenced to an average depth of 85M reads.

Total RNA was isolated using TRIzol Reagent (Thermo Fisher Scientific) from the exact same cell cultures processed for ribosome profiling (10% of the total number of cells). Total RNA was DNase treated and purified using the RNA Clean and Concentrator−25 kit (Zymo Research). RIN scores (RIN = 10 for all six samples) were measured on a BioAnalyzer 2100 using the RNA 6000 Nano assay (Agilent Technologies). Poly(A)-purified mRNA-seq library preparation was performed according to the TruSeq Stranded mRNA Reference Guide (Illumina), using 500 ng of total RNA as input. Libraries were multiplexed and sequenced on an Illumina HiSeq 4000 producing paired-end 2 × 101 nt reads.

Alignment of Ribo-seq and matched mRNA-seq samples

Prior to mapping, ribosome-profiling reads were clipped for residual adapter sequences and filtered for mitochondrial, ribosomal RNA and tRNA sequences (*Figure 2—source data 1*). Next, all mRNA and ribosome profiling data were mapped to the Ensembl 87 transcriptome annotation of the human genome hg38 assembly using STAR 2.5.2b (*Dobin et al., 2013*) in 2-pass mapping mode. To avoid mRNA-seq mapping biases due to read length, the 2 × 101 nt mRNA-seq reads were next trimmed to 29-mers, and those mRNA reads were processed and mapped with the exact same settings as the ribosome profiling data. For the mapping of 2 × 101 nt RNA-seq reads six mismatches per read were allowed (default is 10), whereas two mismatches were permitted for the Ribo-seq and trimmed mRNA-seq reads. To account for variable ribosome footprint lengths, the search start point of the read was defined using the option `--seedSearchStartLmaxOverLread`, which was set to 0.5 (half the read, independent of ribosome footprint length). Furthermore, `--outFilterMulti-mapNmax` was set to 20 and `--outSAMmultNmax` to 1, which prevents the reporting of multimapping reads.

Detecting actively translated reading frames

Canonical ORF detection using ribosome profiling data was performed with RiboTaper v1.3 (*Calviello et al., 2016*) with standard settings. For each sample, we selected only the ribosome footprint lengths for which at least 70% of the reads matched the primary ORF in a meta-gene analysis. Following the standard configuration of RiboTaper, we required ORFs to have a minimum length of 8aa, evidence from uniquely mapping reads and at least 21 P-sites. The final list of translation events was stringently filtered requiring the translated gene to have an average RNA RPKM ≥ 1 and to be detected as translated in all six profiled samples. Furthermore, we required the exact ORF to be detected independently in at least 4 out of 6 samples.

Translational efficiency estimates

Translational efficiency (TE) estimations were calculated as the ratio of Ribo-seq over mRNA-seq DESeq2 normalized counts, yielding independent gene-specific TEs for each of the six individual replicate differentiations. For this, mRNA-seq and Ribo-seq based expression quantification was calculated for (annotated and newly detected) coding sequences (CDSs/ORFs) only, using RNA reads trimmed to footprint sizes as described above.
Data sources

The following datasets used in this study were downloaded from the GEO and ArrayExpress repositories:

- **RNA-seq**: Illumina BodyMap 2.0 expression data from 16 human tissues (GSE30611); polyA mRNA RNA-seq from BE2C (GSE93448), GM12878 (GSE33480), 293T (GSE34995), HeLa (GSE33480), HepG2 (GSE9322), HUVEC (GSE33480), Jurkat (GSE93435), K562 (GSE33480), MiaPaCa-2 (GSE43770), Panc1 (GSE93450), PFSK-1 (GSE93451), U-87 MG (GSE90176); CyT49 hESC/DE/GT/PP1/PP2/CD142+ progenitors/CD200+ polyhormonal cells/in vivo matured endocrine cells/pancreatic islets (E-MTAB-1086).

- **ChIP-seq**: H3K4me3/H3K27me3 in CyT49 hESC/DE/GT/PP1/PP2/CD142+ progenitors/CD200+ polyhormonal cells/in vivo matured endocrine cells/pancreatic islets (E-MTAB-1086).

Statistical analyses

Statistical analyses were performed using Microsoft Excel, GraphPad Prism (7.05), and R (v.3.5.0). Statistical parameters such as the value of n, mean, standard deviation (S.D.), standard error of the mean (S.E.M.), significance level (*p<0.05, **p<0.01, ***p<0.001 and ****p<0.0001), and the statistical tests used are reported in the figures and figure legends. The “n” refers to the number of independent pancreatic differentiation experiments analyzed (biological replicates), or the number of genes/transcripts and sORFs detected.

Statistically significant gene expression changes were determined with DESeq2.

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**Author contributions**

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Additional files
Supplementary files
• Transparent reporting form

Data availability
All mRNA-seq and Ribo-seq datasets generated for this study have been deposited at GEO under the accession number GSE144682.

The following dataset was generated:

| Author(s)                     | Year | Dataset title                                           | Dataset URL                                                                 | Database and Identifier               |
|-------------------------------|------|---------------------------------------------------------|----------------------------------------------------------------------------|---------------------------------------|
| Gaertner B, van Heesch S,     | 2020 | The role of long noncoding RNAs during pancreas development | http://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE144682                | NCBI Gene Expression Omnibus, GSE144682 |
| Schneider-Lunitz V, Schulz    |      |                                                         |                                                                            |                                       |
| F, Blachut S, Nguyen S, Wong  |      |                                                         |                                                                            |                                       |
| R, Matta I, Hubner N, Sander  |      |                                                         |                                                                            |                                       |
| M                             |      |                                                         |                                                                            |                                       |

The following previously published datasets were used:

| Author(s)                     | Year | Dataset title                                           | Dataset URL                                                                 | Database and Identifier               |
|-------------------------------|------|---------------------------------------------------------|----------------------------------------------------------------------------|---------------------------------------|
| Khrebtukova I                 | 2011 | Illumina BodyMap 2.0                                    | http://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE30611                  | NCBI Gene Expression Omnibus, GSE30611 |
| ENCODE project consortium     | 2012 | RNA-seq from ENCODE/Caltech                             | http://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE33480                  | NCBI Gene Expression Omnibus, GSE33480 |
| ENCODE Project Consortium     | 2012 | polyA mRNA RNA-seq from BE2C (ENCSR000BYK)              | http://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE34995                  | NCBI Gene Expression Omnibus, GSE34995 |
| Huelga SC, Vu AQ, Arnold JD,  |      |                                                         |                                                                            |                                       |
| Liang TY, Liu PP, Yan BY,     |      |                                                         |                                                                            |                                       |
| Donohue JP, Shue L, Hoon S,   |      |                                                         |                                                                            |                                       |
| Brenner S, Ares M, Yeo GW     |      |                                                         |                                                                            |                                       |
| ENCODE Project Consortium     | 2016 | polyA mRNA RNA-seq from HepG2 (ENCSR329MHM)             | http://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE90322                  | NCBI Gene Expression Omnibus, GSE90322 |
| ENCODE Project Consortium     | 2017 | polyA mRNA RNA-seq from Jurkat clone E61 (ENCSR000BXX)  | http://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE93435                  | NCBI Gene Expression Omnibus, GSE93435 |
| Sherman MH, Yu RT, Engle DD,  | 2014 | Vitamin d receptor-mediated stromal reprogramming suppresses | http://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE93488                  | NCBI Gene Expression Omnibus, GSE93488 |

[Decision letter and Author response](https://doi.org/10.7554/eLife.58659.sa1)
[Author response](https://doi.org/10.7554/eLife.58659.sa2)
Ding N, Atkins AR, Tiriac H, Collisson EA, Connor F, Van Dyke T, Kozlov S, Martin P, Tseng TW, Dawson DW, Donahue TR, Mamsune A, Shimosegawa T, Apte MV, Wilson JS, Ng B, Lau SL, Gunton JE, Wahl GM, Hunter T, Drebin JA, O’Dwyer PJ, Liddle C, Tuveson DA, Downes M, Evans RM

pancreatitis and enhances pancreatic cancer therapy

acc=GSE43770

GSE43770

ENCODER Project Consortium 2017 polyA mRNA RNA-seq from Panc1 (ENCSR000BYM)

http://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE93450

NCBI Gene Expression Omnibus, GSE93450

ENCODER Project Consortium 2017 polyA mRNA RNA-seq from PFSK-1 (ENCSR000BYN)

http://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE93451

NCBI Gene Expression Omnibus, GSE93451

ENCODER Project Consortium 2016 polyA mRNA RNA-seq from U-87 MG (ENCSR000BYO)

http://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE90176

NCBI Gene Expression Omnibus, GSE90176

Xie R, Everett LJ, Lim HW, Patel NA, Schug J, Kroon E, Kelly OG, Wang A, D’Amour KA, Robins AJ, Won KJ, Kaestner KH, Sanders M 2013 ChIP-seq and RNA-seq of coding RNA of the progression of human embryonic stem cells to beta cells to characterize the epigenetic programs that underlie pancreas differentiation

https://www.ebi.ac.uk/arrayexpress/experiments/E-MTAB-1086/

ArrayExpress, E-MTAB-1086

References

Aguet F, Barbeira AN, Bonazzola R, Brown A, Castel SE, Jo B, Kasela S, Kim-Hellmuth S, Liang Y, Oliva M, Parsana PE, Flynn E, Fresard L, Gaamzon ER, Hamel AR, He Y, Hormozdiari F, Mohammadi P, Munoz-Aguirre M, Park Y, et al. 2019. The GTEx consortium atlas of genetic regulatory effects across human tissues. bioRxiv. DOI: https://doi.org/10.1101/787903

Amaral PP, Leonardi T, Han N, Vire E, Gascoigne DK, Arias-Carrasco R, Bu¨ scher M, Pandolfini L, Zhang A, Pluchino S, Maracaja-Coutinho V, Nakaya HI, Hemberg M, Shiekhattar R, Enright AJ, Kouzarides T. 2018. Genomic positional conservation identifies topological anchor point RNAs linked to developmental loci. Genome Biology 19:32. DOI: https://doi.org/10.1186/s13059-018-1405-5, PMID: 29540241

Anders S, Pyl PT, Huber W. 2015. HTSeq—a Python framework to work with high-throughput sequencing data. Bioinformatics 31:166–169. DOI: https://doi.org/10.1093/bioinformatics/btu638, PMID: 25260700

Arnes L, Akerman I, Balderes DA, Ferrer J, Sussel L. 2016. ??linc1 encodes a long noncoding RNA that regulates islet β-cell formation and function. Genes & Development 30:502–507. DOI: https://doi.org/10.1101/gad.273821.115, PMID: 26944677

Artner I, Bianchi B, Raum JC, Guo M, Kaneko T, Cordes S, Sieweke M, Stein R. 2007. MaB is required for islet beta cell maturation. PNAS 104:3853–3858. DOI: https://doi.org/10.1073/pnas.0700013104, PMID: 17360442

Ashburner M, Ball CA, Blake JA, Botstein D, Butler H, Cherry JM, Davis AP, Dolinski K, Dwight SS, Eppig JT, Harris MA, Hill DP, Issel-Tarver L, Kasarskis A, Lewis S, Matese JC, Richardson JE, Ringwald M, Rubin GM, Sherlock G. 2000. Gene ontology: tool for the unification of biology the gene ontology consortium. Nature Genetics 25:25–29. DOI: https://doi.org/10.1038/75556, PMID: 10802651

Balhhti M, Böttcher A, Lackert H. 2019. Modelling the endocrine pancreas in health and disease. Nature Reviews Endocrinology 15:155–171. DOI: https://doi.org/10.1038/s41574-018-0132-z, PMID: 30504925

Bazini AA, Johnstone TG, Christiano R, Mackowiak SD, Obermayer B, Fleming ES, Vejar CE, Lee MT, Rajewsky N, Walther TC, Giraldez AJ. 2014. Identification of small ORFs in vertebrates using ribosome footprinting and evolutionary conservation. The EMBO Journal 33:981–993. DOI: https://doi.org/10.1002/embj.201488411, PMID: 24705786

Cabili MN, Dunagin MC, McLanahan PD, Biaesch A, Padovan-Merhar O, Regev A, Rinn JL, Raj A. 2015. Localization and abundance analysis of human IncRNAs at single-cell and single-molecule resolution. Genome Biology 16:20. DOI: https://doi.org/10.1186/s13059-015-0586-4, PMID: 25630241
Gaertner L, Mukherjee N, Wyler E, Zauber H, Hirsekorn A, Selbach M, Landthaler M, Obermayer B, Ohler U. 2016. Detecting actively translated open reading frames in ribosome profiling data. *Nature Methods* **13**:165–170. DOI: https://doi.org/10.1038/nmeth.3688, PMID: 26657557.

Carlevaro-Fita J, Rahim A, Guigó R, Vardy LA, Johnson R. 2016. Cytoplasmic long noncoding RNAs are frequently bound to and degraded at ribosomes in human cells. *RNA* **22**:867–882. DOI: https://doi.org/10.1016/j.rna.2015.12.011, PMID: 27909285.

Chen J, Brunner AD, Cogan JZ, Nuñez JK, Fields AP, Adamson B, Izhak DN, Li JY, Mann M, Leonetti MD, Weissman JS. 2020. Pervasive functional translation of noncanonical human open reading frames. *Science* **367**:1140–1146. DOI: https://doi.org/10.1126/science.aay0262, PMID: 32139545.

Clark MB, Johnston RL, Inostroza-Ponta M, Fox AH, Fortini E, Moscat P, Dinger ME, Mattick JS. 2012. Genome-wide analysis of long noncoding RNA stability. *Genome Research* **22**:885–898. DOI: https://doi.org/10.1101/gr.131037.111, PMID: 22407655.

Daneshvar K, Pondick JV, Kim BM, Zhou C, York SR, Macklin JA, Abualteen A, Tan B, Sigova AA, Marcho C, Tremblay KD, Mager J, Choi MY, Mullen AC. 2016. DIGIT is a conserved long noncoding RNA that regulates GSC expression to control definitive endoderm differentiation of embryonic stem cells. *Cell Reports* **17**:353–365. DOI: https://doi.org/10.1016/j.celrep.2016.09.017, PMID: 27705785.

Dobin A, Davis CA, Schlesinger F, Drenkow J, Zaleski C, Jha S, Batut P, Chaisson M, Gingeras TR. 2013. STAR: ultrafast universal RNA-seq aligner. *Bioinformatics* **29**:15–21. DOI: https://doi.org/10.1093/bioinformatics/bts635, PMID: 23104886.

El-Brolosy MA, Kontarakis Z, Rossi A, Kuene C, Günther S, Fukuda N, Kiški K, Boeziio GLM, Takacs CM, Lai SL, Fukuda R, Gerri C, Giraldes AJ, Stainier DYR. 2019. Genetic compensation triggered by mutant mRNA degradation. *Nature* **568**:193–197. DOI: https://doi.org/10.1038/s41586-019-1064-z, PMID: 30944477.

Heinz S, Benner C, Spann N, Bertolino E, Lin YC, Laslo P, Cheng JX, Murre C, Singh H, Glass CK. 2010. Simple combinations of lineage-determining transcription factors prime cis-regulatory elements required for macrophage and B cell identities. *Molecular Cell* **38**:576–589. DOI: https://doi.org/10.1016/j.molcel.2010.05.004, PMID: 20513432.

Ji Z, Song R, Regev A, Struhl K. 2015. Many lncRNAs, 5'UTRs, and pseudogenes are translated and some are likely to express functional proteins. *eLife* **4**:e08890. DOI: https://doi.org/10.7554/eLife.08890, PMID: 26687005.

Jiang W, Liu Y, Liu R, Zhang K, Zhang Y. 2015. The IncRNA DEANR1 facilitates human endoderm differentiation by activating FOXA2 expression. *Cell Reports* **11**:137–148. DOI: https://doi.org/10.1016/j.celrep.2015.03.008, PMID: 25843708.

Kanehisa M, Furumichi M, Tanabe M, Sato Y, Morishima K. 2017. KEGG: new perspectives on genomes, pathways, diseases and drugs. *Nucleic Acids Research* **45**:D353–D361. DOI: https://doi.org/10.1093/nar/gkw1092, PMID: 27899662.

Klattenhoff CA, Scheuermann JC, Surface LE, Bradley RK, Fields PA, Steinhauer ML, Ding H, Butty VL, Torrey L, Haas S, Abro R, Tabebordbar M, Lee RT, Burge CB, Boyer LA. 2013. Braveheart, a long noncoding RNA required for cardiovascular lineage commitment. *Cell* **152**:570–583. DOI: https://doi.org/10.1016/j.cell.2013.01.003, PMID: 23352431.

Kurian L, Aguierre A, Sancho-Martinez I, Benner C, Hishida T, Nguyen TB, Reddy P, Nivet E, Krause MN, Nelles DA, Esteban CR, Campistol JM, Yeo GW, Belmonte JCI. 2015. Identification of novel long noncoding RNAs underlying vertebrate cardiovascular development. *Circulation* **131**:1278–1290. DOI: https://doi.org/10.1161/CIRCULATIONAHA.114.013303, PMID: 25739401.

Langmead B, Trapnell C, Pop M, Salzberg SL. 2009. Ultrafast and memory-efficient alignment of short DNA sequences to the human genome. *Genome Biology* **10**:R25. DOI: https://doi.org/10.1186/gb-2009-10-3-r25, PMID: 19261174.

Levy A. 2019. How evolution builds genes from scratch. *Nature* **574**:314–316. DOI: https://doi.org/10.1038/s41586-019-03061-x, PMID: 31619796.

Li H, Handsaker B, Wysoker A, Fennell T, Ruan J, Homer N, Marth G, Abecasis G, Durbin R, 1000 Genome Project Data Processing Subgroup. 2009. The sequence alignment/map format and SAMtools. *Bioinformatics* **25**:2078–2079. DOI: https://doi.org/10.1093/bioinformatics/btp352, PMID: 19505943.

Lin MF, Jungreis I, Kellis M. 2011. PhyloCSF: a comparative genomics method to distinguish protein coding and non-coding regions. *Bioinformatics* **27**:i275–i282. DOI: https://doi.org/10.1093/bioinformatics/btr209, PMID: 21685081.

Lin N, Chang KY, Li Z, Gates K, Rana ZA, Dang J, Zhang D, Han T, Yang CS, Cunningham TJ, Head SR, Duester G, Dong PD, Rana TM. 2014. An evolutionarily conserved long noncoding RNA TUNA controls pluripotency and neural lineage commitment. *Molecular Cell* **53**:1005–1019. DOI: https://doi.org/10.1016/j.molcel.2014.01.021, PMID: 24530304.

Love MI, Huber W, Anders S. 2014. Moderated estimation of fold change and dispersion for RNA-seq data with DESeq2. *Genome Biology* **15**:R50. DOI: https://doi.org/10.1186/s13059-014-0550-8.

Luo S, Lu JY, Liu L, Yin Y, Chen C, Han X, Wu B, Xu R, Liu W, Yan P, Shao W, Lu Z, Li H, Na J, Tang F, Wang J, Zhang YE. 2016. Divergent IncRNAs regulate gene expression and lineage differentiation in pluripotent cells. *Cell Stem Cell* **18**:637–652. DOI: https://doi.org/10.1016/j.stem.2016.01.024, PMID: 26996597.

Makarewich CA, Olson EN. 2017. Mining for micropeptides. *Trends in Cell Biology* **27**:685–696. DOI: https://doi.org/10.1016/j.tcb.2017.04.006, PMID: 28528987.

Marchese FP, Raimondi I, Huarte M. 2017. The multidimensional mechanisms of long noncoding RNA function. *Genome Biology* **18**:206. DOI: https://doi.org/10.1186/s13059-017-1348-2, PMID: 29084573.
Martinez TF, Chu Q, Donaldson C, Tan D, Shokhirev MN, Saghatelian A. 2020. Accurate annotation of human protein-coding small open reading frames. *Nature Chemical Biology* **16**:458–468. DOI: https://doi.org/10.1038/s41589-019-0425-0, PMID: 31819274

McLean CY, Bristor D, Hiller M, Clarke SL, Schaar BT, Lowe CB, Wenger AM, Bejerano G. 2010. GREAT improves functional interpretation of cis-regulatory regions. *Nature Biotechnology* **28**:495–501. DOI: https://doi.org/10.1038/nbt.1630, PMID: 20436461

Morán I, Akerman I, van de Bunt M, Xie P, Benazra M, Nammo T, Arnes L, Nakić N, García-Hurtado J, Rodríguez-Seguí S, Pasquali L, Sauty-Colace C, Beucher A, Scharffmann R, van Arensbergen J, Johnson PR, Berry A, Lee C, Harkins T, Gmyr V, et al. 2012. Human β cell transcriptome analysis uncovers lncRNAs that are tissue-specific, dynamically regulated, and abnormally expressed in type 2 diabetes. *Cell Metabolism* **16**:435–448. DOI: https://doi.org/10.1016/j.cmet.2012.08.010, PMID: 23040067

Mudge JM, Jungreis I, Hunt T, Gonzalez JM, Wright JC, Kay M, Davidson C, Fitzgerald S, Seal R, Tweedie S, He L, Waterhouse RM, Li Y, Bruford E, Choudhary JS, Frankish A, Kellis M. 2019. Discovery of high-confidence human protein-coding genes and exons by whole-genome PhyloCSF helps elucidate 118 GWAS loci. *Genome Research* **29**:2073–2087. DOI: https://doi.org/10.1101/gr.246642.118, PMID: 31537640

Neumann P, Jaé N, Knau A, Glaser SF, Fouani Y, Rossbach O, Krüger M, John D, Bindreeff A, Grote P, Boon RA, Dimmeler S. 2018. The IncRNA GATA6-AS epigenetically regulates endothelial gene expression via interaction with LOXL2. *Nature Communications* **9**:237. DOI: https://doi.org/10.1038/s41467-017-02431-1, PMID: 29339785

Olexiouk V, Crappé J, Verbruggen S, Verhegen K, Martens L, Menschaert G. 2016. sORFs.org: a repository of small ORFs identified by ribosome profiling. *Nucleic Acids Research* **44**:D324–D329. DOI: https://doi.org/10.1093/nar/gkw1175

Prensner JR, Enache OM, Luria V, Krug K, Clauser KR, Dempster JM, Karger A, Wang L, Stumbrahte K, Wang VM, Botta G, Lyons NJ, Goodale A, Kalani Z, Fritchman B, Brown A, Alan D, Green T, Yang X, Jaffe JD, et al. 2020. Non-canonical open reading frames encode functional proteins essential for Cancer cell survival. *bioRxiv*. DOI: https://doi.org/10.1101/2020.03.981001

Quinlan AR, Hall IM. 2010. BEDTools: a flexible suite of utilities for comparing genomic features. *Bioinformatics* **26**:841–842. DOI: https://doi.org/10.1093/bioinformatics/btp133, PMID: 20110278

Raj A, Wang SH, Shim H, Harpak A, Li YI, Engelmann B, Stephens M, Gilad Y, Pritchard JK. 2016. Thousands of novel translated open reading frames in humans inferred by ribosome footprint profiling. *eLife* **5**:e13328. DOI: https://doi.org/10.7554/eLife.13328, PMID: 2732982

Ramos AD, Andersen RE, Liu SJ, Nowakowski TJ, Hong SJ, Gertz C, Salinas RD, Zarabi H, Kriegstein AR, Lim DA. 2015. The long noncoding RNA pmyk regulates neuronal differentiation of embryonic and postnatal neural stem cells. *Cell Stem Cell* **16**:439–447. DOI: https://doi.org/10.1016/j.stem.2015.02.007, PMID: 25800779

Reimand J, Arak T, Adler P, Kolberg L, Reisberg S, Peterson H, Vilo J. 2016. G.profiler-a web server for functional interpretation of gene lists (2016 update). *Nucleic Acids Research* **44**:W63–W69. DOI: https://doi.org/10.1093/nar/gkw199, PMID: 27098042

Rezania A, Bruin JE, Riedel MJ, Mojibian M, Asadi A, Xu J, Gauvin R, Karanu F, O’Neil JJ, Ao Z, Warnock GL, Kieffer TJ. 2012. Maturation of human embryonic stem cell-derived pancreatic progenitors into functional islets capable of treating pre-existing diabetes in mice. *Diabetes* **61**:2016–2029. DOI: https://doi.org/10.2337/db11-1711, PMID: 22740171

Rezania A, Bruin JE, Arora P, Rubin A, Batushansky I, Asadi A, O’Dwyer S, Quiskamp N, Mojiabian M, Albrecht T, Yang YH, Johnson JD, Kieffer TJ. 2014. Reversal of diabetes with insulin-producing cells derived in vitro from human pluripotent stem cells. *Nature Biotechnology* **32**:1121–1133. DOI: https://doi.org/10.1038/nbt.3033, PMID: 25211370

Roberts A, Pimentel H, Trapnell C, Pachter L. 2011. Identification of novel transcripts in annotated genomes using RNA-Seq. *Bioinformatics* **27**:2325–2329. DOI: https://doi.org/10.1093/bioinformatics/btr355, PMID: 21697122

Romer AJ, Susel L. 2015. Pancreatic islet cell development and regeneration. *Current Opinion in Endocrinology & Diabetes and Obesity* **22**:255–264. DOI: https://doi.org/10.1097/MED.0000000000000174, PMID: 26087337

Ruiz-Orera J, Meseguer X, Subirana JA, Alba MM. 2014. Long non-coding RNAs as a source of new peptides. *eLife* **3**:e03523. DOI: https://doi.org/10.7554/eLife.03523, PMID: 25233276

Ruiz-Orera J, Verdaguer-Grau P, Villanueva-Canas JL, Meseguer X, Alba MM. 2018. Translation of neutrally evolving peptides provides a basis for de novo gene evolution. *Nature Ecology & Evolution* **2**:890–896. DOI: https://doi.org/10.1038/s41559-018-0506-6, PMID: 29560078

Schiesser JV, Wells JM. 2014. Generation of β cells from human pluripotent stem cells: are we there yet? *Annals of the New York Academy of Sciences* **1311**:124–137. DOI: https://doi.org/10.1111/nyas.12369, PMID: 24611778

Shi J, Ma H, Wang H, Zhu W, Jiang S, Dou R, Yan B. 2019. Overexpression of LINC00261 inhibits non-small cell lung Cancer cells progression by interacting with miR-522-3p and suppressing wnt signaling. *Journal of Cellular Biochemistry* **120**:18378–18387. DOI: https://doi.org/10.1002/jcb.29149, PMID: 3190356

Singer RA, Arnes L, Cui Y, Wang J, Gao Y, Guney MA, Burnum-Johnson KE, Rabadran R, Ansong C, Orr G, Sussel L. 2019. The long noncoding RNA paupar modulates PAX6 regulatory activities to promote alpha cell development and function. *Cell Metabolism* **30**:1091–1106. DOI: https://doi.org/10.1016/j.cmet.2019.09.013, PMID: 31607563
Sosa-Pineda B, Chowdhury K, Torres M, Oliver G, Gruss P. 1997. The Pax4 gene is essential for differentiation of insulin-producing beta cells in the mammalian pancreas. Nature 386:399–402. DOI: https://doi.org/10.1038/386399a0, PMID: 9121556

Sun M, Gadad SS, Kim DS, Kraus WL. 2015. Discovery, annotation, and functional analysis of long noncoding RNAs controlling Cell-Cycle gene expression and proliferation in breast Cancer cells. Molecular Cell 59:698–711. DOI: https://doi.org/10.1016/j.molcel.2015.06.023, PMID: 26236012

Swarr DT, Herriges M, Li S, Morley M, Fernandes S, Sridharan A, Zhou S, Garcia BA, Stewart K, Morrisey EE. 2019. The long noncoding RNA falcor regulates Foxa2 expression to maintain lung epithelial homeostasis and promote regeneration. Genes & Development 33:656–668. DOI: https://doi.org/10.1101/gad.320523.118, PMID: 30923168

Tani H, Torimura M, Akimitsu N. 2013. The RNA degradation pathway regulates the function of GAS5 a non-coding RNA in mammalian cells. PLOS ONE 8:e55684. DOI: https://doi.org/10.1371/journal.pone.0055684, PMID: 23383264

The Gene Ontology Consortium. 2019. The gene ontology resource: 20 years and still GOing strong. Nucleic Acids Research 47:D330–D338. DOI: https://doi.org/10.1093/nar/gky1055, PMID: 30395331

Trappell C, Williams BA, Pertea G, Mortazavi A, Kwan G, van Baren MJ, Salzberg SL, Wold BJ, Pachter L. 2010. Transcript assembly and quantification by RNA-Seq reveals unannotated transcripts and isoform switching during cell differentiation. Nature Biotechnology 28:511–515. DOI: https://doi.org/10.1038/nbt.1621, PMID: 20436464

van Heesch S, van Iterson M, Jacobi J, Boymans S, Essers PB, de Bruijn E, Hao W, Machnes AW, Cuppen E, Simonis M. 2014. Extensive localization of long noncoding RNAs to the cytosol and mono- and polyribosomal complexes. Genome Biology 15:R6. DOI: https://doi.org/10.1186/gb-2014-15-1-r6, PMID: 24393600

van Heesch S, Witte F, Schneider-Lunitz V, Schulz JF, Adami E, Faber AB, Kirchner M, Maatz H, Blachut S, Sandmann CL, Kanda M, Worth CL, Schaefer S, Calviello L, Merriott R, Patone G, Hume M, Wyler E, Obermayer B, Mücke MB, et al. 2019. The translational landscape of the human heart. Cell 178:242–260. DOI: https://doi.org/10.1016/j.cell.2019.05.010, PMID: 31155234

Wang ZK, Yang L, Wu LL, Mao H, Zhou YH, Zhang PF, Dai GH. 2017. Long non-coding RNA LINC00261 sensitizes human Colon cancer cells to cisplatin therapy. Brazilian Journal of Medical and Biological Research 51:e6793. DOI: https://doi.org/10.1590/1414-431x20176793, PMID: 29267503

Wang H, Sha L, Huang L, Yang S, Zhou Q, Luo X, Shi B. 2019. LINC00261 functions as a competing endogenous RNA to regulate BCL2L11 expression by sponging miR-132-3p in endometriosis. American Journal of Translational Research 11:2269–2279. PMID: 31105834

Wong WK, Jiang G, Sørensen AE, Chew YV, Lee-Maynard C, Liuwanta D, Williams L, O’Connell PJ, Dalgaard LT, Ma RC, Hawthorne WJ, Joglekar MV, Hardikar AA. 2019. The long noncoding RNA MALAT1 predicts human pancreatic islet isolation quality. JCI Insight 5:e129299. DOI: https://doi.org/10.1172/jci.insight.129299

Yan D, Liu W, Liu Y, Luo M. 2019. LINC00261 suppresses human Colon cancer progression via sponging miR-324-3p and inactivating the wnt/b-catenin pathway. Journal of Cellular Physiology 234:22648–22656. DOI: https://doi.org/10.1002/jcp.28831, PMID: 31183860