Multiomic Profiling Identifies cis-Regulatory Networks Underlying Human Pancreatic β Cell Identity and Function

Graphical Abstract

Highlights

- Comprehensive multiomic maps of EndoC-βH1 human β cell line and primary islets
- Sequence motifs enriched in EndoC-specific enhancers reflect its precursor state
- Identification of regulatory hubs preserved between EndoC-βH1 and human islets
- Identified SNP alleles (including T2D GWAS) altering cis-regulatory signatures

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In Brief

EndoC-βH1 is becoming an important cellular model to study genes and pathways governing human β cell identity and function, but its (epi)genomic similarity to primary human islets is unknown. Lawlor et al. complete and compare extensive EndoC and primary human islet multiomic maps to identify shared and distinct genomic circuitry.
SUMMARY

EndoC-βH1 is emerging as a critical human β cell model to study the genetic and environmental etiologies of β cell (dys)function and diabetes. Comprehensive knowledge of its molecular landscape is lacking, yet required, for effective use of this model. Here, we report chromosomal (spectral karyotyping), genetic (genotyping), epigenomic (ChIP-seq and ATAC-seq), chromatin interaction (Hi-C and Pol2 ChIA-PET), and transcriptomic (RNA-seq and miRNA-seq) maps of EndoC-βH1. Analyses of these maps define known (e.g., PDX1 and ISL1) and putative (e.g., PCSK1 and mir-375) β cell-specific transcriptional cis-regulatory networks and identify allelic effects on cis-regulatory element use. Importantly, comparison with maps generated in primary human islets and/or β cells indicates preservation of chromatin looping but also highlights chromosomal aberrations and fetal genomic signatures in EndoC-βH1. Together, these maps, and a web application we created for their exploration, provide important tools for the design of experiments to probe and manipulate the genetic programs governing β cell identity and (dys)function in diabetes.

INTRODUCTION

Type 2 diabetes (T2D) is a complex disease characterized by elevated blood glucose levels. Ultimately, T2D results when pancreatic islets are unable to produce and secrete enough insulin to compensate for insulin resistance in peripheral tissues of the body. Individual genetic variation combined with dietary and environmental stressors contribute to disease risk and pathogenesis (Lawlor et al., 2017b; Mohlke and Boehnke, 2015). Genome-wide association studies have identified hundreds of genetic loci associated with T2D and related traits, but extensive work remains to identify the causal or functional variants, define their target genes, and determine the roles of these genes in β cell identity and function. Several studies have employed (epi) genomic and transcriptomic profiling of human islets (van de Bunt et al., 2015; Fadista et al., 2014; Varshney et al., 2017), purified β cells (Ackermann et al., 2016; Blodgett et al., 2015), and single-cell populations (Lawlor et al., 2017a; Segerstolpe et al., 2016; Xin et al., 2016) to identify changes in transcriptional regulation and gene expression associated with β cell (dys)function and T2D. However, the molecular and physiologic consequences
of these alterations and their causal link to β cell failure and T2D pathogenesis remain largely undefined.

With the recent creation of an immortalized human β cell line, EndoC-iH1 (Ravassard et al., 2011), islet researchers now possess a tool to experimentally interrogate the molecular mechanisms governing human β cell identity and (dys)function. Since the initial report of their creation, studies utilizing EndoC-iH1 to build insights into human β cell regulation and function have grown steadily. These studies have demonstrated that the physiology (e.g., response to glucose and insulin secretion) of EndoC-iH1 cells resembles that of their primary islet counterparts (Andersson et al., 2015; Krizhanovskii et al., 2017; Oleson et al., 2015; Teraoku and Lenzen, 2017) and that EndoC-iH1 can be used to identify novel genes involved in human insulin secretion (Ndiaye et al., 2017). To motivate further functional studies of human β cell molecular biology and guide the development of cellular models (e.g., for small molecule screening; Tsonkova et al., 2018), extensive characterization of the EndoC-iH1 molecular landscape is needed. Here, we completed multilomic profiling of EndoC-iH1 cells to extensively map (1) chromosomal (spectral karyotyping), (2) 3D epigenomic and/or chromatin looping (Hi-C [Belton et al., 2012] and ChIA-PET [Li et al., 2014]), (3) histone modification (ChiP-seq), (4) chromatin accessibility (ATAC-seq) (Buensrostro et al., 2013), (5) genetic (dense genotyping and imputation), and (6) transcriptomic (RNA-seq and miRNA-seq) signatures of EndoC-iH1. With these high-resolution maps, we sought to (1) identify gene regulatory programs central to human β cell identity and function; (2) nominate putative functional variants, putative molecular mechanisms, and target genes underlying T2D, glucose, and insulin genetic associations; and (3) build a publicly available web application for interactive, intuitive exploration of these data. By comparing these multilomic profiles to those generated from human islets in this study (Hi-C) and parallel studies (Khetan et al., 2018), we identified shared and unique cis-regulatory elements (cis-REs) and gene expression features. Taken together, these data, the insights gleaned from their analysis, and the research support provided by the web application serve as a high-content resource to enable and guide future functional assessment and molecular studies of β cell (dys)function.

RESULTS

Chromosomal and Genetic Heterogeneity in EndoC-iH1
To pursue a precise, comprehensive characterization of the regulatory networks that govern EndoC-iH1 and/or islet β cell identity and function, we first investigated the somatic genomic complement and stability of this cell line using spectral karyotyping (SKY) (Figure 1A). SKY analysis of 14 EndoC-iH1 metaphase spreads revealed that the number of chromosomes was pseudodiploid (n = 46–48) (Figure S1). Nearly all metaphases (n = 13/14) had a normal XY sex complement, with only one having a missing Y chromosome, (metaphase 2; Table S1).

The most common autosomal aberrations in EndoC-iH1 included chromosome 20 gains (n = 11/14 metaphases) and chromosome 10 losses (n = 10/14). Both of these were independently detected as copy-number changes by comparative genomic hybridization (CGH) analysis of the cell line (Univercell Biosolutions, 2011). As summarized in Figure 1B and Table S1, we also noted recurrent 10;17 (11/14 metaphases), 7;18 (10/14 metaphases), 3;17 (7/14 metaphases), and 3;21 (7/14 metaphases) chromosomal translocations as well as rarer events including 12;22 (metaphase 1) and 3;5 (metaphase S2.5) translocations and chromosome 12 losses (2/14 metaphases; Table S1). Together, these results emphasize that although EndoC-iH1 is largely diploid, vigilance and caution are warranted when interpreting studies of genes or cis-REs on chromosomes 3, 7, 10, 17, 18, 20, and 21. We advise investigators to specifically assess copy-number variation at loci of interest, particularly in the regions identified herein as unstable or variable among the population.

Delineation of T2D- and Related-Metabolic-Trait-Associated GWAS SNP Genotypes in EndoC-iH1
Genome-wide association studies (GWASs) have identified hundreds of index and linked SNPs representing putative causal variants at hundreds of loci (Mahajan et al., 2018) associated with T2D genetic risk and changes in associated quantitative traits (e.g., fasting glucose, insulin, and proinsulin levels). We completed dense genotyping and imputation of EndoC-iH1 (STAR Methods) to determine the genotypes at ~2.5 million sites genome-wide (minor allele frequency, MAF >1%), including disease-associated SNPs. First, we overlapped EndoC-iH1 genotypes with National Human Genome Research Institute/European Bioinformatics Institute (NHGRI/EBI) GWAS catalog (STAR Methods) single lead SNPs associated with glucose levels (fasting glucose), insulin levels (fasting insulin and proinsulin levels), type 1 diabetes (T1D), or T2D (MacArthur et al., 2017). EndoC-iH1 exhibited homozygous non-risk genotypes at >50% of these SNPs (Figure 1C; Table S2). For ~20% of analyzed GWAS loci, EndoC-iH1 possessed a heterozygous genotype, including rs10830963 at the MTNR1B locus (chr11) and rs11920090 at the SLC2A2 locus (chr9) (Figure 1D; Table S2). Overlap with T2D-associated SNPs (n = 6,725 index and linked [R2 >0.8] SNPs, representing 403 unique signals) reported in the most recent meta-analysis (Mahajan et al., 2018) revealed a similar genotype distribution, in which EndoC-iH1 was heterozygous for ~30% (n = 119/403; Table S2) of T2D signals. These unique signals represent attractive candidates for (epi)genome editing to experimentally determine T2D-associated allelic effects on cis-RE use in human β cells.

The EndoC-iH1 Epigenome and Transcriptome Largely Resemble Those of Primary Islets but Retain Fetal or Progenitor Islet Cell Signatures
To identify the genome-wide location of EndoC-iH1 cis-REs (Figure 2A), we generated chromatin accessibility maps using ATAC-seq and defined chromatin states (ChromHMM) by completing and integrating ChiP-seq profiles for multiple histone modifications. ATAC-seq identified 127,894 open chromatin sites in EndoC-iH1. Qualitative comparison of EndoC-iH1 open chromatin and chromatin state maps to those in primary human islets (Khetan et al., 2018; Varshney et al., 2017) revealed that the genomic architecture for well-known islet-specific loci such as PCSK1 (Figure 2A), PDX1, and NKX6-1 was remarkably
similar in both, suggesting EndoC-βH1 cells effectively recapitulate β cell cis-regulatory landscapes.

We further compared each ATAC-seq dataset to those from primary islets, sorted β or α cells, and other primary cell types, including adipocyte, skeletal muscle, peripheral blood mononuclear cells (PBMCs), and CD4+ T cells (STAR Methods). We identified a total of 269,701 open chromatin regions (OCRs) across all cell types analyzed (STAR Methods). Among all studied cell types, EndoC-βH1 ATAC-seq profiles most resembled those of β cells (Figure 2B; Spearman R = 0.67), islets (Figure 2B; Spearman R = 0.64), and α cells (Figure 2B; Spearman R = 0.62) (Spearman’s test p values < 2.225 e−308).

Next, we compared OCRs and chromatin states to determine where and to what extent EndoC-βH1 chromatin states recapitulated those of human islets, their constituent cell types, or other metabolic tissues. EndoC-βH1, islet, and β cell OCRs were commonly enriched for binding sites of transcription factors (TFs) implicated in islet cellular identity and function (Figure 2C; group III: FOXA2, FOXO1, RFX, NKX6-1, and PDX1). EndoC-βH1 OCRs also showed exclusive enrichment of sequence motifs that correspond to TFs reported to regulate pluripotency and pancreatic progenitor states (Figure 2C; group IV: HNF6, SOX2, and OCT4), perhaps reflecting the fetal origin and/or derivation of EndoC-βH1. At EndoC-βH1 ATAC-seq OCRs, promoter annotations (from ChromHMM; STAR Methods) were widely conserved between EndoC-βH1 and other cell types, including islets as expected (Figure S2A; centered Pearson correlation >0.95; STAR Methods). In contrast, enhancers, which often encode cell-specific transcriptional regulatory elements (Heinz et al., 2015), at EndoC-βH1 ATAC-seq OCRs were most comparable between islet and EndoC-βH1 (Figure S2A, black point; centered Pearson correlation ~0.71).

To further assess similarities and differences between islet and EndoC-βH1 epigenomes, we investigated the proportions and features of chromatin states that were preserved or disparate between them. Unsurprisingly, a large proportion of promoters (11,907/19,482; 61%) were preserved (Figure 2D) and contained motifs for a variety of TFs from the ETS family (e.g., ELK4, ETS, and ELF1; Table S3) with established roles in cellular differentiation, proliferation, and apoptosis (Findlay et al., 2013). Regions annotated as repressed in both islets and EndoC-βH1 were enriched for CTCF and BORIS binding.
motifs (Table S3), DNA-binding proteins known to bind and establish transcriptional insulators at chromatin territory boundaries. 16,351 out of 51,325 putative enhancers (defined via ChromHMM) were shared between islets and EndoC-bH1 (Figure 2D, blue box) and showed strong enrichment for general (ATF3, AP-1, and JUN) TFs (Figure 2E, blue dots; Table S3) relative to all enhancer regions. Interestingly, we observed a substantial number of EndoC-bH1 enhancers that were annotated as quiescent or repressed in islets (n = 19,380) (Figure 2D, orange box). Relative to all enhancers, these sites were enriched for sequence motifs of TFs controlling pluripotency (OCT2 and NANOG) (Sokolik et al., 2015; Tantin, 2013), pancreatic development or lineage specification (HNF6 and ISL1) (Zhang et al., 2009), and β cell fate determination (PDX1 and NKX6-1) (Thompson and Bhushan, 2017) (Figure 2E, orange dots; Table S3). Based on these findings, it is possible that these regions may represent fetal or developmental cis-REs that are active in the fetal-derived EndoC-bH1 and inactive in adult islets composed of mature β cells. Nonetheless, a significant number of cis-REs (n = 16,351) are conserved between EndoC-bH1 and human islets.

Next, we measured EndoC-bH1 gene expression using RNA-seq and compared it to RNA-seq profiles of islets and other cell types and/or tissues (Figure S2B). As anticipated, the EndoC-bH1 transcriptome most strongly correlated with transcriptomes of islets (R = 0.87) and primary β cells (R = 0.86) among all tissues or cells tested. Of the 27,564 protein coding and/or large intergenic noncoding RNA (lincRNA) genes considered, 11,554...
were expressed in EndoC-βH1 and 12,231 genes were expressed in islet (with 10,473 genes expressed in both). Similarly, EndoC-βH1 small non-coding RNA (miRNA) profiles resembled human islets more than other profiled tissues (adipose, skeletal muscle; Figure S2C) in principal-component analysis (PCA). In particular, PC1 loadings were highly correlated with key islet miRNAs, including miR-375 (Table 1), a critical regulator of β cell mass and identity (Eliasson, 2017), while PC2 stratified primary tissue (adipose, skeletal muscle, and islet) from immortalized cells (EndoC-βH1). Consistent with the PCA, miRNA expression levels in EndoC-βH1 and islets were highly correlated (R = 0.779; Figure S2D), and the vast majority of the most highly expressed miRNAs in EndoC-βH1 have been reported previously to be enriched in primary human β cells relative to whole islets (Table 1) (van de Bunt et al., 2013). Together, the chromatin accessibility, chromatin state, gene expression, and small RNA expression analyses reveal substantial conservation between the transcriptional regulatory and gene expression landscapes of EndoC-βH1 and primary islets.

### Table 1. Top 25 Expressed miRNAs in EndoC-βH1 Cells

| miRNA     | EndoC-βH1 Rank | EndoC-βH1 Enrichment / Depletion |
|-----------|----------------|----------------------------------|
| hsa-miR-375 | 1              | 162076.925 2.84                  |
| hsa-miR-127-3p | 2           | 90512.35 –                     |
| hsa-miR-27b-3p | 3             | 85043.195 3.09                    |
| hsa-miR-192-5p |4            | 70126.5525 2.01                    |
| hsa-miR-192-5p_1 | 5          | 37621.0275 –                      |
| hsa-miR-182-5p | 6           | 36662.2475 2.15                    |
| hsa-miR-22-3p | 7             | 29623.8 2.87                     |
| hsa-miR-191-5p | 8           | 25424.1575 3.02                    |
| hsa-miR-26a-2-5p | 9          | 15578.9175 2.94                    |
| hsa-miR-26a-1-5p | 10         | 15496.7075 2.94                    |
| hsa-miR-141-3p | 11          | 13656.6825 2.84                    |
| hsa-miR-92b-3p | 12          | 13615.6425 2.30                    |
| hsa-miR-30d-5p | 13          | 11108.9275 0.75                    |
| hsa-miR-654-3p | 14          | 11107.385 –                      |
| hsa-miR-148a-3p | 15         | 10651.885 2.22                    |
| hsa-miR-200b-3p | 16         | 10266.2575 1.48                    |
| hsa-miR-92a-2-3p | 17         | 10161.165 2.30                    |
| hsa-miR-381 | 18            | 9342.8825 1.08                     |
| hsa-miR-25-3p | 19            | 9230.2225 1.98                     |
| hsa-miR-181a-1-5p | 20         | 8702.47 1.81                       |
| hsa-miR-181a-2-5p | 21        | 8702.3125 1.81                    |
| hsa-miR-21-5p | 22            | 8439.755 0.21                      |
| hsa-miR-183-5p | 23            | 8338.08 1.58                       |
| hsa-miR-92b-3p | 24            | 7531.295 4.63                      |
| hsa-miR-125a-5p | 25          | 6743.8375 1.24                    |

Table consisting of the top 25 expressed miRNAs in EndoC-βH1 cells. Each miRNA rank was determined by its corresponding expression level (reads per million mapped miRNA [RPMMM]). β cell enrichment / depletion was determined by dividing the counts of each miRNA in β cells (from van de Bunt et al., 2013) by the counts in islets.

Hi-C Profiling of EndoC-βH1 and Human Islets Reveals β Cell-Specific Chromatin Looping Domains

Next, we sought to determine spatial chromatin organization and identify chromatin domains both in EndoC-βH1 and islets using Hi-C. We generated Hi-C maps with ~6 billion reads each for EndoC-βH1 and human islet cells. The maps have 1.9 billion contacts and 1.5 billion contacts, respectively. Using Juicer (Durand et al., 2016a) (STAR Methods), we identified 9,100, 2,580, and 9,448 Hi-C loops in EndoC-βH1, human islet, and GM12878 (a human lymphoblastoid cell line) (Rao et al., 2014), respectively. The reduced number of Hi-C loops identified in primary islet may be attributed to lower unique read depth (~3.7 billion unique reads in islet cells versus ~4.7 billion unique reads in EndoC-βH1 (Table S1)). Together, this represents 19,428 independent DNA loops. Aggregate peak analyses (APAs) (Rao et al., 2014) (Figure 3A, top plots) revealed that chromatin looping sites (anchors) were comparable in EndoC-βH1, islets, and GM12878 for the majority of (>90%) the total chromatin loops (n = 19,428/ 21,128). Consistent with previous studies (Rao et al., 2014; Vietri Rudan et al., 2015), CTCF and CTCFL DNA-binding motifs were overwhelmingly enriched (p < 1e-229 and p < 1e-114, respectively) among all Hi-C anchor sequences (Table S4), verifying that general 3D chromatin structures and loops are preserved between different mammalian tissues and cell types. Importantly, however, we detected 1,078 and 117 chromatin loops that were exclusively present in EndoC-βH1 and islet, respectively, compared to GM12878 (Figure 3A, bottom plots).

To further study cell-specific loops, we subdivided EndoC-βH1 and GM12878 differential Hi-C loops into three classes based on the cell type specificity of the ATAC-seq OCRs they bring into physical proximity (Figure 3B): (A) loops between two non-specific OCRs, (B) loops between two cell-specific OCRs, or (C) loops between one cell-specific OCR and one non-specific OCR. Class B/C loops were classified as cell-specific and further studied. Comparison of EndoC-βH1-specific (n = 315) and GM12878-specific (n = 308) loops revealed a strong bias for cell-specific TF binding at anchor sites (Figure 3C). In EndoC-βH1-specific anchors, we observed enrichment for TFs involved in β cell differentiation and function (NKX6-1, FOXA2, and FOXA1) (Thompson and Bhushan, 2017) as well as OCT4, a key regulator for early embryo development (Le Bin et al., 2014; Wu and Schöler, 2014), while GM12878-specific anchors were enriched for TFs necessary for B cell proliferation and activation (MEF2C and NFAT) (Herglotz et al., 2016; Peng et al., 2001). Furthermore, genes adjacent to EndoC-βH1-specific anchors (STAR Methods) were most enriched (hypergeometric false discovery rate [FDR]-adjusted p value < 0.05) for islet-associated gene ontology (GO) terms, including insulin secretion, glucose homeostasis, and neuronal or endocrine development (Figure S3A; complete results are shown in Table S4). For several genes affiliated with these GO terms, such as SLC30A8, which encodes a zinc efflux transporter involved in zinc ion sequestering and insulin secretion (Mitchell et al., 2016), we observed striking similarities in Hi-C contact frequencies between islet and EndoC-βH1 cells (Figure 3D). In contrast, we observed far fewer chromatin loops and large spans of polycomb-repressed and/or quiescent chromatin for this locus in GM12878 cells (Figure 3E).
Approximately 50% (4,543/9,100) of EndoC-βH1 loop anchors overlapped EndoC-βH1 ATAC-seq OCRs, 44% (n = 1,987/4,543) of which occurred between promoter and enhancer elements (Figure S3B). Of these 4,543 loops, 587 were specifically present in EndoC-βH1 yet absent in GM12878 cells (EndoC-βH1 specific); the remaining 3,956 were captured in both EndoC-βH1 and GM12878 (non-specific). We observed a substantially higher proportion (64%; 376/587) of EndoC-βH1-specific loops that overlapped EndoC-βH1 stretch enhancers (Parker et al., 2013) (Fisher’s exact test p value < 4.23 e-42) compared to that of nonspecific loops (34%; 1,358/3,956). To examine the functional specificity of these loops, we overlapped chromatin state (ChromHMM) information from EndoC-βH1 and 27 other tissue or cell types for all EndoC-βH1 Hi-C loops. For each cell type, we determined the percentage of Hi-C anchors that contained the same chromatin state as EndoC-βH1 (STAR Methods). Islets had the highest percentage of chromatin states identical to EndoC-βH1 at Hi-C anchor sites among all tested tissues or
cell types, especially promoter-enhancer loops (Figure S3C; orange line plot). These findings enumerate regions of cell-specific chromatin looping associated with islet development and function and indicate that EndoC-βH1 forms cell-type-specific chromatin domains or territories highly similar to primary human islets.

**EndoC-βH1 Pol2 ChIA-PET Identifies β Cell cis-Regulatory Hubs**

To map functional cis-regulatory networks, we completed RNA polymerase Pol2 ChIA-PET (Li et al., 2017b) in EndoC-βH1, identifying 25,336 putative Pol2-mediated chromatin interactions. We further filtered these interactions, retaining only those for which both interacting sites (ChIA-PET anchors) overlapped EndoC-βH1 ATAC-seq OCRs, resulting in 16,756 putative cis-regulatory interactions (STAR Methods). As shown in Figure 4A, the overwhelming majority of Pol2-mediated chromatin interactions linked active enhancer and active promoter chromatin states to themselves and each other. Importantly, ChIA-PET detected EndoC-βH1-specific interactions (Figure 4B; compare EndoC-βH1, GM12878, and K562 ChIA-PET tracks) coinciding with those previously reported in targeted 4C-seq analyses of human islets (Pasquali et al., 2014), including the ISL1 (Figure 4B; n = 8 sites denoted by asterisks) and PDX1 (Figure S4A; n = 9 sites) loci.

In addition to replicating interactions previously studied by 4C-seq in human islets, Pol2 ChIA-PET identified hundreds of additional promoter-promoter and promoter-enhancer interactions genome-wide (Figure 4C; STAR Methods). These include extensive Pol2 interactions in loci containing genes crucial for β cell identity and development such as PDX1, ISL1, NKX6-1, MAFB, and mir375 (Figure 4C, red text). As shown in Figures 4C and S4A, multiple interactions were detected between the PDX1 promoter and classically described essential PDX1 transcriptional enhancer sequences (n = 7/14 enhancer interactions), which contain binding sites for islet TFs such as FOXA2 (Gao et al., 2008; Gerrish et al., 2004). During embryonic development, C57BL/6 mouse pancreata displayed a transition in expression of MafB to MafA (Nishimura et al., 2006), suggesting that these two factors are tightly involved in β cell differentiation and function. The high degree of connectivity in MAFB (versus that of MAF) may therefore reflect the fetal or naive state of EndoC-βH1 cells. mir375, a small non-coding RNA, possessed multiple connections to active promoter and enhancer elements (Figure 4C; red text), consistent with its role as a post-transcriptional regulator of genes involved in β cell development or differentiation and insulin secretion or exocytosis (Eliaison, 2017).

Interestingly, INS1M1, a gene necessary for pancreatic endocrine cell differentiation (Osipovich et al., 2014), harbored the most connections in EndoC-βH1 (n = 97 total interactions, n = 7 between active promoters and enhancers; Figure S4B). Other genes linked by ChIA-PET interactions are involved in insulin processing and secretion, including PCSK1, one of the prohormone convertases that catalyzes (pro)insulin processing; RIMBP2, whose protein mediates formation of a complex for polarized accumulation and exocytosis of insulin granules (Fan et al., 2017); RGS7, a critical regulator of muscarinic-stimulated insulin secretion (Wang et al., 2017); and CDC42, which is essential for second-phase insulin secretion (Wang et al., 2007). Additionally, genes implicated in the protection and management of stress were highly connected in EndoC-βH1 ChIA-PET interactions. Notable candidates were ZFAND2B, whose induction helps protect against human β amyloid peptide toxicity or accumulation in a C. elegans transgenic Alzheimer’s disease model (Hassan et al., 2009); SUSD4, a complement inhibitor and tumor suppressor that modulates endoplasmic reticulum stress; and CD59, which is required for mediating exocytosis events facilitating insulin secretion (Blom, 2017; Krus et al., 2014). Finally, TSHZ1, a PDX1 target gene whose expression levels were notably lower in human islet donors with T2D (Raum et al., 2015), harbored five links to active enhancer elements in EndoC-βH1, suggesting that perturbation of the cis-regulatory networks identified herein may contribute to T2D pathogenesis.

Finally, we sought to study to what extent the putative cis-regulatory networks detected in EndoC-βH1 may be preserved in islets and other cell types. Due to limited availability of ChIA-PET data in human islets and other relevant tissues, we decided to use the chromatin interaction sites determined by EndoC-βH1 ChIA-PET and compare the functional annotations (ChromHMM state annotations) at these loci across 27 different tissue or cell types. Overall, aggregate counts of these chromatin state interactions for each cell type were most similar between EndoC-βH1 and islet (Figure 4D, green bar plots; STAR Methods). ChIA-PET interactions between regions annotated as active promoters in EndoC-βH1 were similarly annotated as active promoters across the 27 other cell or tissue types (Figure 4D, Act. promoter × Act. promoter, red line plot; STAR Methods). In contrast, the majority of active enhancers interacting in EndoC-βH1 were marked as active enhancers only in human islets (Figure 4D, Act. Enhancer × Act. Enhancer, yellow line plot). Multidimensional scaling of all cell or tissue chromatin state annotations at EndoC-βH1 ChIA-PET interacting sites reaffirmed high similarity between EndoC-βH1 and islets (Figure S4C), consistent with strong conservation of active enhancer state annotations, as previously observed (Figure 4D, line plots). These results suggest that these interactions may represent β cell cis-regulatory hubs. Indeed, anchors for 41% of ChIA-PET interactions (6,904/16,756) overlapped islet stretch enhancers, suggesting that these interactions may encompass key islet functional chromatin domains.

**Integration of EndoC-βH1 Genotype and 3D Genomic Interaction Maps Identifies Allelic Imbalance at β Cell-Specific cis-REs**

We and others have demonstrated that genetic variants, including those associated with T2D and other quantitative measures of islet (dys)function, can alter cis-RE use (chromatin accessibility quantitative trait loci [caQTL]) (Khetan et al., 2018) and target gene expression (expression quantitative trait loci [eQTL]; van de Bunt et al., 2015; Fadista et al., 2014). Recently, approaches have been used to assess allelic effects on these molecular features at heterozygous sites within a single sample. We applied these allelic imbalance (AI) analyses in EndoC-βH1 to identify genetic variants that alter β cell cis-REs and target gene expression. To identify instances of AI in EndoC-βH1, we examined the allelic bias of ~2 million heterozygous SNPs (STAR Methods) within OCRs (ATAC-seq peaks), active enhancer
elements (H3K27ac peaks), or expressed genes (RNA-seq; Figure 5A). Less than 10% of all SNPs occurring in OCRs and enhancer elements showed significant AI (Figure 5A; part I; FDR <10%). Approximately 25% of SNPs exhibited gene expression AI (Figure 5A). When considering variants with adequate coverage in both EndoC-βH1 ATAC-seq OCRs and H3K27ac-marked enhancer regions (n = 1,734 SNPs), we noted a positive correlation (R = 0.2) in the corresponding AI ratios (Figure S5A), suggesting the potential for coordinate regulation of chromatin accessibility and histone modification at these

Figure 4. RNA Polymerase 2 ChIA-PET Identifies Chromatin Interactions in EndoC-βH1
(A) Heatmap showing the chromatin states of EndoC-βH1 ChIA-PET interaction nodes.
(B) Example of a Pol2 ChIA-PET interaction between active enhancer (blue box) and active promoter (green box) cis-REs in the ISL1 locus on chromosome 5. Asterisks under EndoC-βH1 ChIA-PET interactions (red) indicate interacting sites in the ISL1 locus detected in human islet 4C-seq analyses (Pasquali et al., 2014).
(C) ChIA-PET network connectivity of gene promoters in EndoC-βH1 containing at least three interactions with other regulatory elements. For each gene, the number of connections between other regulatory elements (e.g., active enhancer and weak enhancer) and the proportion of total links in which the chromatin states are EndoC-βH1-specific (blue) or identical in both human islet and EndoC-βH1 (green) are shown in bar plots on the right. The remaining proportions that are neither EndoC-βH1 specific nor common to islet and EndoC-βH1 are not shown. Red font denotes loci containing genes crucial for β cell identity and development.
(D) Top: Bar plot illustrating the proportions of chromatin states at the Pol2 ChIA-PET interacting sites (nodes) shared between EndoC-βH1, islets, and additional Epigenomics Roadmap tissues and cell lines. Bottom: Heatmap demonstrating the chromatin states of EndoC-βH1 Pol2 ChIA-PET interacting sites (nodes) in islets (left) or stomach smooth muscle (right). See also Figure S4.
cis-regulatory sites. In total, 119 out of 403 T2D-associated signals (~30%) overlapped EndoC-βH1 cis-REs (Table S2); 34 out of 119 of these unique signals (~29%) were heterozygous in EndoC-βH1 and potentially amenable to allelic analyses. GREGOR (Schmidt et al., 2015) enrichment analysis of these same 403 signals identified significant overlap (p value < 1 e-7) in EndoC-βH1 ATAC-seq (n = 67 SNPs) OCRs. Moreover, 24,102 out of 126,013 of T2D-associated 99% genetic credible set SNPs (19%) overlapped significant allelic bias in gene expression (RNA-seq) in a direction consistent with the H3K27ac allelic bias. Although the exact role of SAMD5 in human islets has not been described, expression of this gene is high in adult β cells but absent in adult α cells (Lawlor et al., 2017a; Segerstolpe et al., 2016; Wang et al., 2016). SAMD5 has been recently identified as

Figure 5. Allelic Effects on EndoC-βH1 Transcriptional Regulatory Features

(A) EndoC-βH1 genotype information was integrated with ATAC-seq, H3K27ac, and RNA-seq data to identify sequence variants altering cis-RE accessibility and/or activity (ATAC-seq and H3K27ac) or mRNA levels (RNA-seq) in EndoC-βH1. Pie charts summarize the proportions of variants exhibit significant AIs (blue; FDR < 10%) in each of the corresponding sequencing profiles.

(B) Cartoon representation of approach to identify systematic allelic effects on EndoC-βH1 cis-regulatory networks.

(C) Multibeam view highlighting allelic effects on the SAMD5 locus cis-regulatory network in EndoC-βH1. A variant site exhibiting significant Al in H3K27 acetylation (denoted by blue arrow) is linked (red ChIA-PET interaction) to the transcription start site (TSS) of SAMD5. Within the SAMD5 locus, five transcribed SNPs exhibited significant allelic bias in gene expression (RNA-seq) in a direction consistent with the H3K27ac allelic bias.

(D) Left: Bar plots summarizing the proportions of variants with ATAC-seq/H3K27ac imbalance (blue bars; FDR < 10%) that overlap ChIA-PET interacting loci. Right: Pie charts specifying the chromatin state (ChromHMM) annotations of the overlapping variants. See also Figure S5 and Table S5.
These data identify SAMDS5 as one of the most highly connected loci in EndoC-H1 (Figures 4C and S4B; n = 35 ChIA-PET connections between the gene and active enhancers; n = 5 interactions to other active promoters) and highlight a potential cis-regulatory hub for fetal β and islet cell development. Further manipulation of the cis-regulatory network in this locus may provide greater insight into its putative roles in islet cell differentiation and function.

Overall, 2,500 out of 5,515 (~45%) and 8,794 out of 43,492 (~20%) of heterozygous SNPs passing coverage thresholds (STAR Methods) in ATAC-seq and H3K27ac cis-REs, respectively, overlapped a ChIA-PET anchor (Figure 5D, bar plots). For both datasets, <5% of SNPs demonstrated significant AI. AI SNPs were present in active promoter and enhancer regions of the genome (Figure 5D, pie charts marked with blue arrows). Enhancers exhibited enriched AI compared to promoters. For example, in the ATAC-seq data, 49.5% of AI SNPs occurred in enhancers compared to 30.9% of non-AI SNPs (Fisher’s exact test p value = 0.000282 for comparison of enhancer versus promoter counts). Similarly, 54.6% of H3K27ac AI SNPs occurred in enhancers compared to 43% of non-AI SNPs (Fisher’s exact test p value = 3.343e-05 for comparison of enhancer versus promoter counts) (Figure 5D).

Enhancers govern cell-type-specific gene expression patterns (Heinz et al., 2015). We identified 50 and 185 enhancer SNPs with significant ATAC-seq and H3K27ac AI, respectively (Figure 5D, yellow portion of pie charts marked with blue arrows). We used ChIA-PET interactions to link enhancers showing AI to promoters of the genes they might regulate (Table S5). This identified enhancer-promoter links for 21 out of 50 (42%) ATAC-seq and 91 out of 185 (~49%) H3K27ac AI enhancer SNPs, respectively. To assess if observed allelic effects on EndoC-H1 cis-regulatory networks extended to primary islets, we examined the eQTL direction of effect of these SNPs on steady-state expression of their predicted target genes in human islets (Varshney et al., 2017). Islet eQTL Z scores (STAR Methods) were most correlated to the H3K27ac allelic ratio at SNPs where the human islet eQTL target gene (i.e., the gene whose expression in human islets is influenced by a genetic variant) matched that of the EndoC-H1 ChIA-PET target gene (i.e., the gene linked to the enhancer by EndoC-H1 ChIA-PET as depicted in Figure 5B) (n = 42/91) (Figure S5B, red points; R = 0.32), as opposed to those genes in the locus that were not linked to the enhancer by ChIA-PET (Figure S5B, gray points; R = 0.17).

Importantly, these analyses suggest that integrated EndoC-H1 omics analyses provide molecular insights into diabetes genetics (e.g., GWAS). For example, T2D-associated index and 99% credible set SNP rs57235767, for which the “C” risk allele exhibited reduced EndoC-H1 H3K27ac counts, exhibited consistent downregulation of the ChIA-PET-predicted target gene C11orf54 (Figure S5B, asterisk) expression in islet cohorts (Fadista et al., 2014; Varshney et al., 2017). Similarly, the T2D risk allele of rs3807136, a 99% credible set SNP in linkage disequilibrium (r² > 0.8) with the index and 99% credible set SNP rs2268382, displayed a higher proportion of H3K27ac counts in EndoC-H1 and increased expression of the predicted target gene, CEP41. These human islet SNP-gene interactions (that are also recapitulated in EndoC-H1 cells) represent high-priority targets for (epi)genome modification and should assist efforts to decrypt the genes contributing to T2D pathogenesis.

**DISCUSSION**

Here, we report extensive multiomic mapping and integrated analysis of cytogenetic (karyotyping), large-scale chromatin structural conformation (H3C), cis-regulatory networks (ChIA-PET), histone mark (ChIP-seq), chromatin accessibility (ATAC-seq), genetic (genotyping), and gene expression (RNA-seq) information in EndoC-H1 human β cells. For convenient and interactive browsing of this data, we have created an R shiny web application (Chang et al., 2018) available at https://shinyapps.jax.org/endoc-islet-multi-omics/. These data and the browser application should serve as a resource for future studies to explore the complex β cell regulatory programs uncovered in this study and guide targeted studies of regulatory networks, genes, and pathways of interest.

SKY revealed chromosomal heterogeneity among individual cells in the EndoC-H1 population. These included copy number variation, such as chromosome 20 gain and chromosome 10 loss, that has been identified independently by array CGH analyses, as well as previously unappreciated structural alterations, including chromosome 10:17 and 3:21 translocations, that are frequent within the population. Consistent with SKY, we observed enhanced contact frequency between chromosomes 3 and 21 in EndoC-H1 Hi-C maps (Juicebox [Durand et al., 2016b]; http://aidenlab.org/ juicebox/?juiceboxURL=http://bit.ly/2NxWcDp), suggesting that these two technologies may complement one another to identify cell line abnormalities. As evidenced by other less prevalent chromosomal aberrations among the population, it is possible that this cell line may continue to evolve with continued passaging. Thus, caution should be taken, and these aberrations should be considered, in future (epi)genome editing or EndoC-H1 molecular and functional experiments involving genes or regulatory elements on these chromosomes.

Overall, comparative analyses of omics profiles indicate substantial similarity among EndoC-H1, islet, and primary β cell transcriptomes (Figure 2B; Pearson R > 0.86). EndoC-H1 open chromatin profiles were modestly correlated with islet (R = 0.64) and primary β (R = 0.67) cells, highlighting potential drift between the cell line and primary islet cells at the level of chromatin accessibility. This includes ~19,000 putative EndoC-H1 enhancers annotated as quiescent or polycomb repressed in human islets. Interestingly, these sites contain potential binding sites for TFs with important roles in β cell development and pancreatic precursor fates and functions (e.g., NKX6-1, PDX1, ISL1, and HNF6) (Thompson and Bhushan, 2017) and pluripotency (e.g., NANOG and OCT2) (Sokolik et al., 2015; Tantin, 2013). Thus, these discordant features may reflect the fetal nature of EndoC-H1 cells, their transformed state, or both.

Using Hi-C to map higher-order chromatin structure in EndoC-H1 and a corresponding map from a human islet donor, we defined islet and β cell chromatin domains and territories. Consistent with previous findings (Rao et al., 2014), the overall spatial chromatin organization was similar across EndoC-H1,
islet, and GM12878 cells (Figure 3A; top panel), and all Hi-C anchors were enriched for TFs with general roles in chromatin organization (e.g., CTCF and BORIS, Table S4). Importantly, however, Hi-C analyses also identified ~1,078 islet and β cell-specific chromatin domains, several of which were evident in both EndoC-βH1 and primary human islets (Figure 3). These cell-specific chromatin territories were enriched for β cell-specific TFs (Figure 3C) and brought into close physical proximity genes linked to islet-associated biological process GO terms (Figure S3A) compared to those of GM12878. We and others have linked a drive islet and vide valuable insights into important transcriptional hubs that compared to those of 27 other cell or tissue types. Thus, the advanced expression or target genes. Using this approach, we linked 91 out of 185 works in EndoC-βH1, which further refined chromatin territories to reveal functional EndoC-βH1 cis-regulatory networks. In addition to validating chromatin interactions previously reported in 4C-seq analyses at select loci in human islets, Pol2 ChIA-PET identified hundreds of interactions genome-wide between active promoter and enhancer regions potentially involved in the regulation and transcription of dozens of β cell-specific loci (Figures 4A–4C). Comparison of Pol2 ChIA-PET interaction locations in EndoC-βH1, GM12878, and K562 revealed that the overwhelming majority of interactions at these loci were unique to EndoC-βH1. Due to high cell input requirements (~100 million cells) for current ChIA-PET library construction protocols (Li et al., 2017b), we were unable to validate these findings in human islets. However, consistent with our previous observations between islet and EndoC-βH1 Hi-C maps (Figure S3C), we noted that chromatin states of ChIA-PET interaction nodes in EndoC-βH1 were most conserved in islet (Figure 4D, Figure S4C) compared to those of 27 other cell or tissue types. Thus, the cis-regulatory programs we define for EndoC-βH1 should provide valuable insights into important transcriptional hubs that drive islet and β cell identity and function. GWASs have identified hundreds of loci that contribute to genetic risk of T2D and other quantitative measures of islet dysfunction, such as glucose, insulin, and proinsulin levels (Fuchsberger et al., 2016; Mahajan et al., 2018). We and others have linked a subset of SNPs and loci to altered cis-RE activity and steady-state islet gene expression (van de Bunt et al., 2015; Fadista et al., 2014; Khetan et al., 2018; Varshney et al., 2017). For the majority of loci, challenges remain to define the (1) causal or functional SNP, (2) determine its molecular effect on cis-RE activity, and (3) identify the putative target gene(s). By combining our dense genotype and 3D chromatin interaction (ChIA-PET) networks in EndoC-βH1, we sought to identify SNPs with imbalanced expression or cis-RE use and link them physically with their target genes. Using this approach, we linked 91 out of 185 (H3K27ac imbalanced) and 21 out of 50 (ATAC-seq imbalanced) SNPs to potential target genes (e.g., rs2294805 to SAMD5 in Figure 5B). Our ability to assess chromatin interactions between diabetes-associated SNPs and their target genes was modest. Nonetheless, we identified two candidate SNPs (rs57235767, rs3807136) that demonstrated consistent directions-of-effect on measures of cis-RE activity (e.g., H3K27ac) and target gene expression in EndoC-βH1 and human islets (Figure S5B) (Varshney et al., 2017). Several factors could underlie the modest frequency of diabetes-associated GWAS SNPs linked by Pol2 ChIA-PET interactions: (1) limited sensitivity of ChIA-PET technology, (2) condition or disease specificity of GWAS SNP effects on cis-RE use or activity, or (3) condition-specific Pol2 interactions between cis-REs and their target genes. In summary, this study provides an integrated multiomic analysis of EndoC-βH1, a human pancreatic β cell line with increasing utility and importance to the β cell and diabetes communities. Integrated analysis of chromatin interaction and gene expression information identified chromosomal territories and cis-regulatory networks governing β cell identity and function. Overall, comparison of EndoC-βH1 (epigenomic and 3D chromatin profiles with those of human islets verified common signatures of gene expression, TF binding, and cis-RE use. These analyses also highlighted genomic discrepancies between EndoC-βH1 and their primary cell counterparts, possibly reflecting the fetal or embryonic origin of the cell line and/or its transformed state. Integration of EndoC-βH1 cis-regulatory maps with genome-wide genotype information nominated target genes and identified SNP allelic effects on transcriptional regulatory networks, including a subset of T2D-associated SNPs. Together, the data and tools provided here should serve as helpful guides for rational design of targeted and hypothesis-driven studies of candidate genes, pathways, or cis-REs to determine their roles in β cell (dys)function and diabetes.

STAR★METHODS

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SUPPLEMENTAL INFORMATION

Supplemental Information includes five figures and five tables and can be found with this article online at https://doi.org/10.1016/j.celrep.2018.12.083.
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AUTHOR CONTRIBUTIONS

M.L.S., S.C.J.P., M.R.E., F.S.C., D.U., Y.R., A.P.A., E.L.A., and P.S. designed the experimental and data analysis strategies. A.D. and E. Pak performed SKY. M.L.S., S.C.J.P., M.R.E., F.S.C., D.U., Y.R., A.P.A., E.L.A., and P.S. designed the generation and sequencing of EndoC-M.S.S., R.K., M.R.E., E. Piecuch, X.L., O.L., E.L.A., H.G., and Y.R. contributed to the generation and sequencing of EndoC-M-iH1 and human islet libraries. N.L., AT., E.J.M., PO., S.C.J.P., A.V., M.K., P.S., M.R.E., P.S.C., S.R., M.S.S., and N.N. performed bioinformatics analyses of the data. Genotyping and imputation of EndoC-M-iH1 was performed by M.E. and N.N. N.L., N.E., N.C.J.P., and M.L.S. wrote the manuscript, and N.L., E.J.M., N.N., P.S., and M.K. designed the corresponding figures. All authors reviewed the manuscript prior to submission.

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REFERENCES

Ackermann, A.M., Wang, Z., Schug, J., Naji, A., and Kaestner, K.H. (2016). Integration of ATAC-seq and RNA-seq identifies human α cell and β cell signature genes. Mol. Metab. 5, 233–244.
Andersson, L.E., Valtat, B., Bagge, A., Sharoyko, V.V., Nicholls, D.G., Ravassard, P., Scharffmann, R., Spégel, P., and Mulder, H. (2015). Characterization of stimulus-secretion coupling in the human pancreatic EndoC-M-iH1 β cell line. PLoS ONE 10, e0120879.
Belton, J.-M., McCord, R.P., Gibcus, J.H., Naumova, N., Zhan, Y., and Dekker, J. (2012). Hi-C: a comprehensive technique to capture the formation of genomes. Methods 58, 268–276.
Blodgett, D.M., Nowsosiekta, A., Afik, S., Pechhold, S., Cura, A.J., Kennedy, N.J., Kim, S., Kucukural, A., Davis, R.J., Kent, S.C., et al. (2015). Novel observations from next-generation RNA sequencing of highly purified human adult and fetal islet cell subsets. Diabetes 64, 3172–3181.
Blom, A.M. (2017). The role of complement inhibitors beyond controlling inflammation. J. Intern. Med. 282, 116–128.
Bolger, A.M., Lohse, M., and Usadel, B. (2014). Trimmomatic: a flexible trimmer for Illumina sequence data. Bioinformatics 30, 2114–2120.
Buenrostro, J.D., Giresi, P.G., Zaba, L.C., Chang, H.Y., and Greenleaf, W.J. (2013). Transposition of native chromatin for fast and sensitive epigenomic profiling of open chromatin, DNA-binding proteins and nucleosome position. Nat. Methods 10, 1213–1218.
Cardinale, V., Wang, Y., Carpino, G., Cui, C.-B., Gatto, M., Rossi, M., Berloco, P.B., Cantafiora, A., Wauthier, E., Furth, M.E., et al. (2011). Multipotent stem/progenitor cells in human biliary tree give rise to hepatocytes, cholangiocytes, and pancreatic islets. Hepatology 54, 2159–2172.
Chang, W., Cheng, J., Alaire, J.J., Xie, Y., McPherson, J. (2018). shiny: Web Application Framework for R. R package version 1.2.0. https://CRAN.R-project.org/package=shiny.
Das, S., Forer, L., Schönherr, S., Sidore, C., Locke, A.E., Kwong, A., Vrieze, S.I., Chew, E.Y., Levy, S., McGue, M., et al. (2016). Next-generation genotyping imputation service and methods. Nat. Genet. 48, 1284–1287.
Delaneau, O., Marchini, J., and Zagury, J.-F. (2011). A linear complexity phasing method for thousands of genomes. Nat. Methods 9, 179–181.
Dobin, A., Davis, C.A., Schlesinger, F., Drenkow, J., Zaleski, C., Jha, S., Batut, P., Chaisson, M., and Gingeras, T.R. (2013). STAR: ultrafast universal RNA-seq aligner. Bioinformatics 29, 15–21.
Durand, N.C., Shamim, M.S., Machol, I., Rao, S.S.P., Huntley, M.H., Lander, E.S., and Aiden, E.L. (2016a). Juicer provides a one-click system for analyzing loop-resolution Hi-C experiments. Cell Syst. 3, 95–98.
Durand, N.C., Robinson, J.T., Shamim, M.S., Machol, I., Mesirov, J.P., Lander, E.S., and Aiden, E.L. (2016b). Juicebox provides a visualization system for Hi-C contact maps with unlimited zoom. Cell Syst. 3, 99–101.
Elalsson, L. (2017). The small RNA miR-375 - a pancreatic islet abundant miRNA with multiple roles in endocrine β cell function. Mol. Cell. Endocrinol. 456, 95–101.
ENCOD Project Consortium (2012). An integrated encyclopedia of DNA elements in the human genome. Nature 489, 57–74.
Ernst, J., and Kellis, M. (2017). Chromatin-state discovery and genome annotation with CHROMHMM. Nat. Protoc. 12, 2478–2492.
Fadista, J., Vikman, P., Laakso, E.O., Mollet, I.G., Esquerra, J.L., Taneera, J., Storm, P., Osmark, P., Ladenvall, C., Prasad, R.B., et al. (2014). Global genomic and transcriptomic analysis of human pancreatic islets reveals novel genes influencing glucose metabolism. Proc. Natl. Acad. Sci. USA 111, 13924–13929.
Fan, F., Matsunaga, K., Wang, H., Ishizaki, R., Kobayashi, E., Kiyonari, H., Mckunoto, Y., Okunishi, K., and Izumi, T. (2017). Exophilin-8 assembles secretory granules for exocytosis in the actin cortex via interaction with RIM-BP2 and myosin-VIlla. eLife 6, e26174.
Findlay, V.J., LaRue, A.C., Turner, D.P., Watson, P.M., and Watson, D.K. (2013). Understanding the role of ETS-mediated gene regulation in complex biological processes. Adv. Cancer Res. 119, 1–61.
Fuchsberger, C., Flannick, J., Teslovich, T.M., Mahajan, A., Agarwala, V., Gaulton, K.J., Ma, C., Fontanillas, P., Moutsianas, L., McCarthy, D.J., et al. (2016). The genetic architecture of type 2 diabetes. Nature 536, 41–47, advance online publication.
Gao, N., LeLay, J., Vatamaniuk, M.Z., Rieck, S., Friedman, J.R., and Kaestner, K.H. (2008). Dynamic regulation of Pdx1 enhancers by Foxa1 and Foxa2 is essential for pancreas development. Genes Dev. 22, 3435–3448.
Gerrish, K., Van Velkinburgh, J.C., and Stein, R. (2004). Conserved transcriptional regulatory domains of the pdx-1 gene. Mol. Endocrinol. 18, 533–548.
Hartley, S.W., and Mullikin, J.C. (2015). QoRTs: a comprehensive toolset for quality control and data processing of RNA-Seq experiments. BMC Bioinformatics 16, 224.
Hassan, W.M., Merin, D.A., Fonte, V., and Link, C.D. (2009). AIP-1 ameliorates β-amyloid peptide toxicity in a Caenorhabditis elegans Alzheimer’s disease model. Hum. Mol. Genet. 18, 2739–2747.
Heinz, S., Benner, C., Spann, N., Bertolino, E., Lin, Y.C., Laslo, P., Cheng, J.X., Murre, C., Singh, H., and Glass, C.K. (2010). Simple combinations of lineage-determining transcription factors prime cis-regulatory elements required for macrophage and B cell identities. Mol. Cell 38, 576–589.
Heinz, S., Romanoski, C.E., Benner, C., and Glass, C.K. (2015). The selection and function of cell type-specific enhancers. Nat. Rev. Mol. Cell Biol. 16, 144–154.
Khetan, S., Kursawe, R., Youn, A., Lawlor, N., Jillette, A., Marquez, E.J., Ucar, D., and Stitzel, M.L. (2018). Type 2 diabetes-associated genetic variants regulate chromatin accessibility in human islets. Diabetes D., and Stitzel, M.L. (2018). Type 2 diabetes-associated genetic variants regulate chromatin accessibility in human islets. Diabetes.

Khetan, S., Kursawe, R., Youn, A., Lawlor, N., Jillette, A., Marquez, E.J., Ucar, D., and Stitzel, M.L. (2018). Type 2 diabetes-associated genetic variants regulate chromatin accessibility in human islets. Diabetes.

Krizhanovskii, C., Kristinsson, H., Elkins, A., Wang, X., Gavali, H., Bergsten, P., Scharfmann, R., and Welsh, N. (2017). EndoC-J1H cells display increased sensitivity to sodium palmitate when cultured in DMEM/F12 medium. Islets 9, e1269995.

Krus, U., King, B.C., Nagaraj, N., Fajardo, S., Jin, J., Buda, P., Garcia-Vaz, E., Gomez, M.F., Ottosson-Laasko, E., Storm, P., et al. (2014). The complement inhibitor CD59 regulates insulin secretion by modulating exocytotic events. Cell Metab.

Lawlor, N., George, J., Boisset, M., Kursawe, R., Sun, L., Kycia, I., Robson, P., and Stitzel, M.L. (2017a). Single cell transcriptomes identify human islet cell signatures and reveal cell-type-specific expression changes in type 2 diabetes. Genome Res 27, 208–222.

Lawlor, N., Khetan, S., Ucar, D., and Stitzel, M.L. (2017b). Genomics of islet (dys)function and type 2 diabetes. Trends Genet. 33, 244–255.

Le Bin, G.C., Munoz-Descalzo, S., Kurowski, A., Leitch, H., Lou, X., Mansfield, W., Etsene-Dumeau, C., Grabole, N., Mulas, C., Niwa, H., et al. (2014). Oct4 is required for lineage priming in the developing inner cell mass of the mouse blastocyst. Development 141, 1001–1010.

Li, H. (2013). Aligning sequence reads, clone sequences and assembly contigs with BWA-MEM. arXiv. doi: 1303.3979v2.

Li, H., Handsaker, B., Wysoker, A., Fennell, T., Ruan, J., Homer, N., Marth, G., Abecasis, G., and Durbin, R. (2011). 1000 Genome Project Data Processing Subgroup (2009). The Sequence Alignment/Map format and SAMtools. Bioinformatics 25, 2078–2079.

Li, G., Ruan, X., Auerbach, R.K., Sandhu, K.S., Zheng, M., Wang, P., Poh, H.M., Goh, Y., Lim, J., Zhang, J., et al. (2012). Extensive promoter-centered chromatin interactions provide a topological basis for transcription regulation. Cell 148, 84–98.

Li, G., Cai, L., Chang, H., Hong, P., Zhou, Q., Kulikova, E.A., Kolchanov, N.A., and Ruan, Y. (2014). Chromatin interaction analysis with paired-end tag (ChIA-PET) sequencing technology and application. BMC Genomics 15, S11.

Li, G., Chen, Y., Snyder, M.P., and Zhang, M.Q. (2017a). ChIA-PET2: a versatile and flexible pipeline for ChIA-PET data analysis. Nucleic Acids Res. 45, e4.

Li, X., Luo, O.J., Wang, P., Zheng, M., Wang, D., Piechuch, E., Zhu, J.J., Tian, S.Z., Tang, Z., Li, G., and Ruan, Y. (2017b). Long-read ChIA-PET for base-resolution mapping of haplotype-specific chromatin interactions. Nat. Protoc. 12, 899–915.

Loh, P.-R., Palamara, P.F., and Price, A.L. (2016). Fast and accurate long-range phasing in a UK Biobank cohort. Nat. Genet. 48, 811–816.

Lun, A.T.L., Perry, M., and Ing-Simmons, E. (2016). Infrastructure for genomic interactions: Bioconductor classes for HI-C, ChIA-PET and related experiments. F1000Res. 5, 950.

MacArthur, J., Bowling, E., Cerezo, M., Gil, L., Hall, P., Hestings, E., Junkins, H., McMahon, A., Milano, A., Morales, J., et al. (2017). The new NHGRI-EBI Catalog of published genome-wide association studies (GWAS Catalog). Nucleic Acids Res. 45 (D1), D896–D901.

Mahajan, A., Tailun, D., Thumer, M., Robertson, N.R., Torres, J.M., Rayner, N.W., Payne, A.J., Steinhorsdottir, V., Scott, R.A., Granup, N., et al. (2018). Fine-mapping type 2 diabetes loci to single-variant resolution using high-den-
Ramirez, F., Dündar, F., Diehl, S., Grüning, B.A., and Manke, T. (2014). deepTools: a flexible platform for exploring deep-sequencing data. Nucleic Acids Res. 42, W187-91.

Rao, S.S.P., Huntley, M.H., Durand, N.C., Stamenova, E.K., Bochkov, I.D., Robinson, J.T., Sanborn, A.L., Machol, I., Omer, A.D., Lander, E.S., and Eiden, E.L. (2014). A 3D map of the human genome at kilobase resolution reveals principles of chromatin looping. Cell 159, 1665–1680.

Raum, J.C., Soleimanpour, S.A., Groff, D.N., Coré, N., Fassano, L., Garratt, A.N., Dai, C., Powers, A.C., and Stoffers, D.A. (2015). Tshz1 regulates pancreatic β-cell maturation. Diabetes 64, 2905–2914.

Ravassard, P., Hazhouz, Y., Pechherty, S., Bricout-Neveu, E., Armanet, M., Czerinchow, P., and Scharfmann, R. (2011). A genetically engineered human pancreatic β cell line exhibiting glucose-inducible insulin secretion. J. Clin. Invest. 121, 3589–3597.

Roadmap Epigenomics Consortium; Kundaje, A., Meuleman, W., Ernst, J., Blenkly, M., Yen, A., Heravi-Moussavi, A., Kheradpour, P., Zhang, Z., Wang, J., Ziller, M.J., et al. (2015). Integrative analysis of 111 reference human epigenomes. Nature 518, 317–330.

Ross-Innes, C.S., Stark, R., Teschedorff, A.E., Holmes, K.A., Ali, H.R., Dunning, M.J., Brown, G.D., Gojis, O., Ellis, I.O., Green, A.R., et al. (2012). Differential oestrogen receptor binding is associated with clinical outcome in breast cancer. Nature 481, 389–393.

Rozowsky, J., Abyzov, A., Wang, J., Alves, P., Raha, D., Harmanaci, A., Leng, J., Bjornson, R., Kong, Y., Kitabayashi, N., et al. (2011). AlleleSeq: analysis of allele-specific expression and binding in a network framework. Mol. Syst. Biol. 7, 522.

Schmidt, E.M., Zhang, J., Zhou, W., Chen, J., Mohlke, K.L., Chen, Y.E., and Willer, C.J. (2015). GREGOR: evaluating global enrichment of trait-associated variants in epigenetic features using a systematic, data-driven approach. Bioinformatics 31(7), 2601–2606.

Schultz, M.D., He, Y., Whitaker, J.W., Harirhan, M., Muckel, E.A., Leung, D., Rajagopal, N., Nery, J.R., Unich, M.A., Chen, H., et al. (2015). Human body epigenome maps reveal noncanonical DNA methylation variation. Nature 523, 212–216.

Scott, L.J., Erdos, M.R., Huyghe, J.R., Welch, R.P., Beck, A.T., Wolford, B.N., Chines, P.S., Didion, J.P., Narisu, N., Strüwing, H.M., et al. (2016). The genetic regulatory signature of type 2 diabetes in human skeletal muscle. Nat. Commun. 7, 11764.

Segerstolpe, Å, Palasanta, A., Eliasson, P., Andersson, E.-M., Andréasson, A.-C., Sun, X., Picelli, S., Sabirish, A., Claussen, M., Bjursell, M.K., et al. (2016). Single-Cell Transcriptome Profiling of Human Pancreatic Islets in Health and Type 2 Diabetes. Cell Metab. 24, 593–607.

Sokolik, C., Liu, Y., Bauer, D., McPherson, J., Broeker, M., Heimberg, G., Qi, L.S., Sivak, D.A., and Thomson, M. (2015). Transcription factor competition allows embryonic stem cells to distinguish authentic signals from noise. Cell Syst. 1, 117–129.

Stitzel, M.L., Sethupathy, P., Pearson, D.S., Chines, P.S., Song, L., Erdos, M.R., Welch, R., Parker, S.C.J., Boyle, A.P., Scott, L.J., et al.; NISC Comparative Sequencing Program (2010). Global epigenomic analysis of primary human pancreatic islets provides insights into type 2 diabetes susceptibility loci. Cell Metab. 22, 443–455.

Tang, Z., Luo, O.J., Li, X., Zheng, M., Zhu, J.J., Szalaj, P., Trzaskoma, P., Magalska, A., Wlodarczyk, J., Ruszczycki, B., et al. (2015). CTCF-mediated human 3D genome architecture reveals chromatin topology for transcription. Cell 163, 1611–1627.

Tantin, D. (2013). Oct transcription factors in development and stem cells: insights and mechanisms. Development 140, 2857–2866.

Teraoku, H., and Lenzen, S. (2017). Dynamics of insulin secretion from EndoC-β1 cell pseudoislets in response to glucose and other nutrient and nonnutrient secretagogues. J. Diabetes Res 2017, 2309630.

The Broad Institute (2013). Picard-Tools: A set of command line tools (in Java) for manipulating high-throughput sequencing (HTS) data and formats such as SAM/BAM/CRAM and VCF. http://broadinstitute.github.io/picard/.
**STAR METHODS**

**KEY RESOURCES TABLE**

| REAGENT or RESOURCE | SOURCE | IDENTIFIER |
|---------------------|--------|------------|
| Antibodies          |        |            |
| H3K27ac-human antibody | Abcam | Abcam Cat# ab4729; RRID:AB_2118291 |
| H3K4me3-mouse antibody | Abcam | Abcam Cat# ab12209; RRID:AB_442957 |
| ENCODE Project Antibody validation H3K4me1 | Abcam | Abcam Cat# ab8895; RRID:AB_306847 |
| ENCODE Project Antibody validation H3K36me3 | Abcam | Abcam Cat# ab9050; RRID:AB_306966 |
| H3K27me3-celegans antibody | Abcam | Abcam Cat# ab6002; RRID:AB_305237 |
| H3K79me2-human antibody | Abcam | Abcam Cat# ab3594; RRID:AB_303937 |
| H3K9me3-human,H3K9me3-mouse antibody | Abcam | Abcam Cat# ab8898; RRID:AB_306848 |
| Rabbit Anti-CTCF Polyclonal Antibody, Unconjugated | Abcam | Abcam Cat# ab70303; RRID:AB_1209546 |
| RNA Polymerase II 8WG16 Monoclonal Antibody, anti-RNA Polymerase II | Covance Research Products | Covance Research Products Cat# MMS-126R; RRID:AB_10013665 |
| Biological Samples |        |            |
| Human pancreatic islet | NDRI | UNOS ID: ADAC418 |
| Critical Commercial Assays |        |            |
| SKY Probe | Applied Spectral Imaging INC | Cat# FPRPR0028 |
| HumanOmní2.5–4v1_H BeadChip Array | Illumina | Cat# WG-311-2514 |
| TruSeq ChIP Sample Prep Kit | Illumina | Cat# IP-202-1024 |
| TruSeq Stranded mRNA Kit | Illumina | Cat# 20020595 |
| TruSeq Small RNA Library Kit | Illumina | Cat# RS-200-0036 |
| Deposited Data |        |            |
| Raw and analyzed data | This paper | NCBI BioProject: PRJNA480287; European Variation Archive: ERZ674947; NCBI GEO: GSE118588 |
| Islet ATAC-seq and RNA-seq | Khetan et al., 2018 | NCBI SRA: SRP117935 |
| Purified beta and alpha cell ATAC-seq and RNA-seq | Ackermann et al., 2016 | NCBI GEO: GSE76268 |
| Peripheral mononuclear blood cells (PBMC) ATAC-seq | Ucar et al., 2017 | European Genome-phenome Archive: EGAS00001002605 |
| PBMC RNA-seq | ENCODE Project Consortium, 2012 | NCBI GEO: GSE90275 |
| Skeletal muscle ATAC-seq | Scott et al., 2016 | dbGap: phs001068.v1.p1 |
| Skeletal muscle RNA-seq | ENCODE Project Consortium, 2012 | NCBI GEO: GSE78611 |
| GM12878 and CD4+ T ATAC-seq | Buenrostro et al., 2013 | NCBI GEO: GSE47753 |
| GM12878 RNA-seq | Rozowsky et al., 2011 | NCBI GEO: GSE30400 |
| CD4+ T RNA-seq | Schultz et al., 2015 | NCBI GEO: GSE18927 |
| Adipocyte RNA-seq | ENCODE Project Consortium, 2012 | NCBI GEO: GSE93486 |
| GM12878 Hi-C | Rao et al., 2014 | NCBI GEO: GSE63525 |
| K562 RNA Pol2 CHIA-PET | Li et al., 2012 | NCBI GEO: GSE39495 |
| GM12878 RNA Pol2 CHIA-PET | Tang et al., 2015 | NCBI GEO: GSE72816 |
| Experimental Models: Cell Lines |        |            |
| EndoC-betaH1: Cell line | EndoCells/INSERM | RRID: CVCL_L909 |
| Software and Algorithms |        |            |
| Michigan Imputation Server | Das et al., 2016 | https://imputationserver.sph.umich.edu/index.html |
| Eagle v2.3 | Loh et al., 2016 | https://data.broadinstitute.org/alkesgroup/Eagle/ |
| SHAPEIT v2.r790 | Delaneau et al., 2011 | http://mathgen.stats.ox.ac.uk/genetics_software/shapeit/shapeit.html |

(Continued on next page)
CONTACT FOR REAGENT AND RESOURCE SHARING

Further information and requests for resources and reagents should be directed to and will be fulfilled by the Lead Contact, Michael Stitzel (michael.stitzel@jax.org).

EXPERIMENTAL MODEL AND SUBJECT DETAILS

EndoC-βH1 cell culture and processing

EndoC-βH1 cells provided by EndoCells/INSERM were cultured and passaged as previously described (Ravassard et al., 2011). Briefly, cells were seeded at a density of approximately 600,000 cells/cm² on tissue culture-treated plates pre-coated overnight with extracellular matrix (Sigma) and fibronectin (Sigma) in EndoC-βH1 complete medium. Cells were passaged approximately every 7 days. Cells were harvested at various passages and distinct sites (e.g., NHGRI, JAX-GM) for karyotyping, genotyping, ATAC-seq, ChIP-seq, RNA-seq, Hi-C, and Pol2 ChIA-PET analyses.

Human islet acquisition and procurement

The single human pancreatic islet sample (obtained from a cadaveric donor; UNOS ID: ADAC418; sex: Female; age: 43 years old) used in this study was obtained from the National Disease Research Interchange (NDRI) and processed according to NHGRI institutional review board-approved protocols. The islet was shipped overnight from the distribution center. On receipt, we pre-warmed the islet to 37°C in shipping media for 1–2 h before harvest; ~50,000 islet equivalents (IEQs) were harvested for Hi-C.
METHOD DETAILS

Spectral karyotyping (SKY)
Spectral karyotyping of EndoC-βH1 was completed to identify structural and numerical chromosome aberrations using standard procedures as previously described. In brief, EndoC-βH1 cells were cultured to 80% confluence. Metaphase spreads were prepared from these cells after mitotic arrest with Colcemid (0.015 μg/mL, 16 to 18 hours) (GIBCO, Gaithersburg, MD), hypotonic treatment (0.075 mol/L KCl, 20 minutes, 37° C), and fixation with methanol–acetic acid (3:1). Commercial SKY probe and software (Applied Spectral Imaging INC, Carlsbad, CA) was used to identify and visualize the individually colored chromosomes obtained from two slides’ worth of metaphase spreads from the same passage.

Genotyping, imputation, and phasing of EndoC-βH1
EndoC-βH1 was genotyped with the HumanOmni2.5–4v1_H BeadChip Array (Illumina, San Diego, CA, USA). We mapped the Illumina array probe sequences to the hg19 genome assembly and excluded likely problematic ones as described in (Varshney et al., 2017).

We applied the following filtering criteria to remove additional SNP probes prior to pre-phasing of the array genotypes: 1) we assessed allele frequency of the SNPs using combined genotypes of EndoC-βH1 and 163 other samples that were genotyped on similar chips; and 2) we removed SNPs with an alternate allele frequency difference with 1000G EUR samples > 20%, or palindromic SNPs with a minor allele frequency > 20%, genotype missingness > 2.5%, Hardy-Weinberg p value < 10⁻⁴. At the end, a total of 1,851,388 SNPs were used in pre-phasing and imputation.

We performed pre-phasing and imputation separately on autosomal and chrX markers using the Michigan Imputation Server (Das et al., 2016). We used Eagle v2.3 (Loh et al., 2016) for autosomal chip marker pre-phasing and SHAPEIT v2.r790 (Delaneau et al., 2011) for chrX markers. We subsequently used minima3 (Das et al., 2016) for imputation of missing genotypes using the Haplotype Reference Consortium (HRC version, hrc.r1.1.2016) panel (McCarthy et al., 2016).

GWAS SNP pruning
Lists of reference SNP identifiers were obtained from the NHGRI-EBI Catalog of SNPs (https://www.ebi.ac.uk/gwas/; accessed January 19th, 2017) for Type 2 diabetes, Type 1 diabetes, fasting glucose traits, fasting insulin traits, and proinsulin level categories. For each disease category, GWAS SNPs were pruned using PLINK version 1.9 (Purcell et al., 2007) to identify SNPs in high linkage disequilibrium (LD) (R² > 0.8) the SNP with the least significant p value was discarded. T2D-associated SNPs from (Mahajan et al., 2018) were obtained and pruned using the same methodology described above.

ATAC-seq
EndoC-βH1 ATAC-seq libraries were prepared as previously described (Varshney et al., 2017) and sequenced on an Illumina NextSeq 500 with 2 x 125 bp cycles. Raw sequence fastq files for adipocyte tissue, bulk islet (Khetan et al., 2018), islet beta and alpha (GSE76268) (Ackermann et al., 2016), peripheral mononuclear blood cells (PBMC) (Ucar et al., 2017), skeletal muscle (Scott et al., 2016), GM12878 and CD4+ T cells (GSE47753) (Buenrostro et al., 2013) were obtained from their corresponding studies.

Paired-end ATAC-seq reads were quality trimmed using Trimmomatic version 0.33 (Bolger et al., 2014) and parameters “TRIMMING:3 SLIDINGWINDOW:4:15 MINLEN:36.” Trimmed reads were aligned to human genome (hg19) using BWA version 0.7.12 (Li, 2013), specifically using the bwa mem –M option. Duplicate reads were removed using “MarkDuplicates” from Picard-tools version 1.95 (The Broad Institute, 2013). Quality of aligned reads were examined using Qualimap version 2.2.1 and default parameters “bamqc –bam –gd hg19.” (Okonechnikov et al., 2016). After preprocessing and quality filtering, peaks were called on alignments with MACS version 2.1.0 (Zhang et al., 2008) using the parameters “–g ‘hs’-nomodel-keep-dup all–broad–broad-cutoff 0.05 -f BAMPE.” Peaks located in blacklisted regions of the genome were removed. Remaining overlapping peaks from all cell types were merged with BEDTools version 2.26.0 (Quinlan and Hall, 2010) to generate a single peak set (n = 269,701). Raw read counts in these peaks for each cell type were determined using the R package DiffBind version 2.4.8 (Ross-Innes et al., 2012). Spearman rank-order correlation was calculated for cell types using the merged peaks with deepTools version 2.4.2 (Ramirez et al., 2014) and parameters “multiBamSummary BED-file –BED.”

ChIP-seq
CTCF, H3K27ac, H3K27me3, H3K36me3, H3K4me1, H3K79me2, H3K4me3, H3K9me3 ChIP-seq was performed as previously described (Stitzel et al. 2016) and sequenced on an Illumina HiSeq 2000 using 2 x 100 bp cycles. Harmonized ChromHMM (Ernst and Kellis, 2017) states for EndoC-βH1 and NIH Roadmap cells/tissues were determined as previously described: “Chromatin states were determined by applying the ChromHMM (version 1.10) hidden Markov model algorithm at 200-bp resolution to five chromatin marks and input. We ran ChromHMM with a range of possible states and selected a 13-state model, because it most accurately captured information from higher-state models and provided sufficient resolution to identify biologically meaningful patterns in a reproducible way” (Varshney et al., 2017).
Transcription factor motif enrichment analysis

“findMotifsGenome.pl” (HOMER version 4.6 (Heinz et al., 2010)) script with parameters “hg19 -size 200” was used to determine TF motifs enriched in ATAC-seq OCRs for each cell type (Figure 2C). In each analysis, all merged OCRs (n = 269,701) were provided as background (e.g., EndoC-βH1 called OCRs (foreground) versus all merged OCRs (background)). The same parameters were used to identify enriched motifs in either “Enhancers in Both EndoC-βH1 and Islet” (n = 16,351) or “Enhancers in EndoC-βH1 Only” (n = 19,380) compared to all enhancers (n = 51,325) (Figure 2D). The same HOMER script and parameters were also used to identify enriched motifs in EndoC-βH1 (n = 315) versus GM12878 (n = 308) cell-specific Hi-C loops.

Similarity of cell/tissue type chromatin state (ChromHMM) annotations at EndoC-βH1 ATAC-seq OCRs

EndoC-βH1 ATAC-seq OCRs (n = 127,894) were overlapped with chromatin state (ChromHMM) annotations from EndoC-βH1, human islet, adipocyte, skeletal muscle, GM12878, and PBMC cells provided in (Varshney et al., 2017). Next, only OCRs that intersected a ChromHMM annotation from all tissue/cell types (n = 127,887/127,894) were retained (union set). Within a tissue/cell type, or instances where multiple chromatin state elements intersected an EndoC-βH1 OCR, annotations were prioritized as follows: promoter, enhancer, transcription, repressed, or low signal. At each OCR, cell/tissue ChromHMM annotations were compared to those of EndoC-βH1 and assigned a binary classification (1 = the annotations were the same, 0 = the annotations were different). Aggregated counts of pairwise chromatin state annotations based on EndoC-βH1 OCRs were then computed for each tissue/cell type, and a resulting similarity matrix was calculated using the “simil” function within the proxy version 0.4 R package (Meyer and Buchta, 2018).

RNA-seq

Total RNA was extracted and purified from EndoC-βH1 using Trizol as previously described (Varshney et al., 2017). All sequencing was performed on an Illumina NextSeq 500 with 2 x 101 bp cycles. Raw fastq files for human islets (Khetan et al., 2018), islet beta and alpha (Ackermann et al., 2016), PBMC (GSE90275), skeletal muscle (GSE78611), adipocyte (GSE93486) (ENCODE Project Consortium, 2012), GM12878 (GSE30400) (Rozowsky et al., 2011), and CD4+ T cell (GSE18927) (Schultz et al., 2015) were obtained from the associated databases. Paired-end RNA-seq reads were trimmed using Trimmomatic with the same parameters as used for ATAC-seq reads. Trimmed reads were aligned to human genome (hg19) using STAR version 2.5.3 (Dobin et al., 2013) with default parameters and expression levels of all genes were determined using QoRTs version 1.2.42 (Hartley and Mullikin, 2015) with default parameters and Gencode v19 transcript annotations. A total of 27,564 protein-coding genes and long intergenic non-coding RNAs (lincRNAs) were considered in the study.

miRNA-seq

Total RNA was extracted and purified from 2000-3000 islet equivalents (IEQ) or 2 x 106 EndoC-βH1 cells using Trizol (Life Technologies). RNA quality was confirmed with Bioanalyzer 2100 (Agilent); islet samples with RNA integrity number (RIN) greater than 6.5 were prepared for miRNA sequencing; EndoC-βH1 cells RNA RIN scores were > 9.0. miRNA libraries were prepared at the NIH Intramural Sequencing Core (NISC) from 1 μg total RNA using Illumina’s TruSeq Small RNA Library Kit according to the manufacturer’s guidelines, except a 10% acrylamide gel was used for better separation of library from adapters. Libraries were pooled in groups of about 8 for gel purification. Single-end 51 base sequencing was performed on Illumina HiSeq 2500 sequencers in Rapid Mode using version 2 chemistry. Data was processed using RTA version 1.18.64 and CASAVA 1.8.2. All resulting data was processed with miRquant 2.0 (Kanke et al., 2016).

Hi-C

Hi-C libraries were generated as described in (Rao et al., 2014) and analyzed using the Juicer Tools version 1.75 pipeline (Durand et al., 2016a). We sequenced 6,065,763,792 Hi-C read pairs in EndoC-βH1 cells, yielding 1,909,699,446 Hi-C contacts; we also sequenced 6,009,242,588 Hi-C read pairs in islet cells, yielding 1,516,995,339 Hi-C contacts. Loci were assigned to A and B compartments at 500 kB resolution. Loops were annotated using HiCUPs at 5kb and 10kb resolutions with default Juicer parameters. This yielded a list of 9,100 loops in EndoC-βH1, annotations were prioritized as follows: promoter, enhancer, transcription, repressed, or low signal. At each OCR, cell/tissue ChromHMM annotations were compared to those of EndoC-βH1 and assigned a binary classification (1 = the annotations were the same, 0 = the annotations were different). Aggregated counts of pairwise chromatin state annotations based on EndoC-βH1 OCRs were then computed for each tissue/cell type, and a resulting similarity matrix was calculated using the “simil” function within the proxy version 0.4 R package (Meyer and Buchta, 2018).

ChIA-PET

EndoC-βH1 RNA Polymerase 2 (Pol2) ChIA-PET libraries were generated and sequenced reads were processed and analyzed according to the protocol in (Li et al., 2017b). ChIA-PET interactions were identified using ChIA-PET2 (Li et al., 2017a) using the “bridge
Corresponding ChIA-PET interactions for K562 (GSE39495) and GM12878 (GSE72816) cells were obtained from Gene Expression Omnibus. ChIA-PET and Hi-C loops were further filtered using the Bioconductor package InteractionSet_1.8.0 (Lun et al., 2016) to retain only those in which both interacting sites (anchors) overlapped OCRs. ChIA-PET anchors were annotated to the nearest gene. To assess the physical connectivity between genes and their putative regulatory regions as captured by ChIA-PET interactions, the number of distinct links between anchors annotated to each gene were counted and categorized by their chromatin state and regulatory function. Counting was carried out both between each gene promoter and all linked regulatory regions (Figure 4C), and also between all annotated anchors regardless of their chromatin state and all linked regulatory regions (Figure S4B). For example, consider anchors A1 and A2, both overlapping enhancer regions annotated to gene $i$. These anchors respectively link to anchors B1, located in a TSS, and B2, in an enhancer. In this scenario, a connectivity degree of two would be computed for gene $i$, corresponding to an enhancer-TSS and a TSS-TSS interaction, respectively.

The functional specificity of EndoC-$\beta$H1 ChIA-PET interactions was investigated by overlapping interaction anchors on ChromHMM chromatin states computed from EndoC-$\beta$H1 data as well as 27 other tissue/cell types (Varshney et al., 2017), and calculating the rate of conservation of chromatin states of both anchors. For example, a rate of 80% enhancer-enhancer conservation would mean that 8 out of 10 interactions of this type in EndoC-$\beta$H1 are also found in another cell type. The resulting proportions were computed for all interactions combined, and for specific relevant regulatory interactions (Figure 4D, line plots). In addition, aggregated counts of pairwise chromatin state interactions based on EndoC-$\beta$H1 ChIA-PET interactions were computed for the same cell types as above, and pairwise distances ($D$) between the resulting 29 count matrices were computed and plotted as scaled similarity values relative to EndoC-$\beta$H1 (i.e., $1-D/D_{\text{max}}$), so that $D = 0$ for EndoC-$\beta$H1 interactions and $D = D_{\text{max}}$ for the most divergent cell type (Figure 4D, bar plot). These same methods were used to determine the functional specificity of EndoC-$\beta$H1 Hi-C interactions.

**ATAC-seq allelic bias analysis**

All allelic bias analyses were performed using WASP (van de Geijn et al., 2015) (version 0.2.2 after GitHub commit 5a52185 and bug fix in pull request #67). For the EndoC-$\beta$H1 ATAC-seq allelic bias analyses, after original BWA mapping, reads were filtered to properly-paired, high-quality autosomal reads using SAMTools (v. 1.3.1; flags -f 3 -F 4 -F 8 -F 256 -F 2048 -q 30) (Li et al., 2009). Remapping and filtering as part of the WASP pipeline utilized the same parameters. As the last step of the WASP pipeline, duplicate removal was performed using WASP’s rmdup_pe.py script. In order to avoid double-counting SNPs covered by both reads in a pair, overlapping read pairs were clipped using bamUtil’s clipOverlap (https://genome.sph.umich.edu/wiki/BamUtil_clipOverlap; v. 1.0.14). The four replicate libraries were then merged using SAMTools merge.

For each SNP, we determined the number of reads containing each allele (requiring base quality of at least 20). We excluded SNPs with total coverage less than 10, as well as SNPs in regions blacklisted by the ENCODE Consortium because of poor mappability (wgEncodeDacMapabilityConsensusExcludable.bed and wgEncodeDukeMapabilityRegionsExcludable.bed). Allelic bias testing was performed using a two-tailed binomial test, using an adjusted expectation for the null to account for residual reference bias as described in (Scott et al., 2016). Briefly, for each of the 16 reference-alternate allele pairs (e.g., AG and GA are separate allele pairs), we calculated the expected fraction of reference alleles (fracRef) under the null as the sum of the reference allele counts divided by the sum of the total allele counts for SNPs of that allele pair. To prevent SNPs of high coverage from biasing the expected fracRef, we down-sampled SNPs with coverage in the top 25th percentile to the median coverage, and used these downsampled reference and total allele counts when calculating the expected fracRef. We used the observed allele-pair specific fracRef as the true fracRef under the null hypothesis of no allelic bias in the binomial test. Multiple testing correction was performed using the Benjamini-Hochberg correction (FDR < 10%).

**ChIP-seq allelic bias analysis**

For the EndoC-$\beta$H1 ChIP-seq allelic bias analyses, paired-end libraries were processed as follows. Adapters were trimmed using cta (v. 0.1.2) and reads mapped using BWA mem (-M flag; v. 0.7.12). Reads were filtered to properly-paired, high-quality autosomal reads using SAMTools (flags -f 3 -F 4 -F 8 -F 256 -F 2048 -q 30). Single-end libraries were mapped using BWA aln (v. 0.7.12; default parameters) and filtered using SAMTools (flags -F 4 -F 256 -F 2048 -q 30). Remapping and filtering as part of the WASP pipeline utilized the same parameters as for the original mapping. For paired-end libraries, overlapping read pairs were clipped using bamUtil’s clipOverlap. Replicates were merged using SAMTools merge. Allele counting and allelic bias testing was performed as described above for ATAC-seq.

**RNA-seq allelic bias analysis**

For the EndoC-$\beta$H1 RNA-seq allelic bias analyses, after original STAR mapping, reads were filtered to properly-paired, high-quality autosomal reads using SAMTools (v. 1.3.1; flags -f 3 -F 4 -F 8 -F 256 -F 2048 -q 255). Remapping and filtering as part of the WASP pipeline utilized the same parameters. In order to avoid double-counting SNPs covered by both reads in a pair, overlapping read pairs were clipped using bamUtil’s clipOverlap. Allele counting and allelic bias testing was performed as described above for ATAC-seq.

**Comparison of islet eQTL and EndoC-$\beta$H1 biased SNPs allelic effect**

Human islet eQTL data were obtained from (Varshney et al., 2017). For EndoC-$\beta$H1 biased (H3K27ac) enhancer SNPs that were also linked (via ChIA-PET chromatin interaction) to a target gene ($n = 91/185$ SNPs in Figure 5C), all corresponding islet eQTL SNP-gene...
pairs were retrieved. For 42/91 SNPs linked to target genes, the H3K27ac allelic effect bias was calculated assuming that the EndoC-
βH1 effect allele was the same as the islet eQTL effect allele. Allelic effect bias was calculated by dividing the effect allele coverage (either reference or alternate allele) by the total coverage of the SNP. A scalar value of 0.5 was subtracted from this value to determine whether the effect allele had an increased (positive value), decreased (negative value), or no (zero) bias in H3K27ac coverage. Randomly selected eQTL SNP-gene pairs that did not have corresponding connections (via ChIA-PET chromatin interaction) to a target gene were considered as a null/background set.

**QUANTIFICATION AND STATISTICAL ANALYSIS**

Visualization of all Hi-C, ChIA-PET, chromatin state (ChromHMM), ATAC-seq, and RNA-seq data examined in this study (Figures 1, 2, 3, 4, and 5) were produced with R using ggplot2_3.0.0 (Wickham, 2016), pheatmap_1.0.10 (Kolde, 2018), Sushi_1.18.0 (Phanstiel et al., 2014), as well as Juicer Tools (Figure 3) (Durand et al., 2016a) and Juicebox (Figure 3) (Durand et al., 2016b).

**DATA AND SOFTWARE AVAILABILITY**

The accession numbers for EndoC-βH1 Hi-C, ChIA-PET, ChIP-seq, ATAC-seq, RNA-seq, miRNA-seq as well as human islet Hi-C data reported in this paper are: NCBI BioProject accession PRJNA480287, NCBI Gene Expression Omnibus accession GSE118588. The accession number for EndoC-βH1 genotype data is: European Variation Archive accession ERZ674947 and project PRJEB27824. The R shiny web application to visualize and interact with the multi-omic data generated in this study is freely available at https://shinyapps.jax.org/endoc-islet-multi-omics/. Interactive Hi-C maps are available at aidenlab.org/juicebox.
Supplemental Information

Multiomic Profiling Identifies

cis-Regulatory Networks Underlying Human Pancreatic β Cell Identity and Function

Nathan Lawlor, Eladio J. Márquez, Peter Orchard, Narisu Narisu, Muhammad Saad Shamim, Asa Thibodeau, Arushi Varshney, Romy Kursawe, Michael R. Erdos, Matt Kanke, Huiya Gu, Evgenia Pak, Amalia Dutra, Sheikh Russell, Xingwang Li, Emaly Piecuch, Oscar Luo, Peter S. Chines, Christian Fuchberger, NIH Intramural Sequencing Center, Praveen Sethupathy, Aviva Presser Aiden, Yijun Ruan, Erez Lieberman Aiden, Francis S. Collins, Duygu Ucar, Stephen C.J. Parker, and Michael L. Stitzel
SKY EndoC-βH1 Metaphase S2.1
46,XY,der(17)t(10;17),der(17)t(3;17),der(17)t(3;17)

SKY EndoC-βH1 Metaphase S2.7
48,XY,t(3;21),-10,+der(17)t(10;17),+20

SKY EndoC-βH1 Metaphase 7
47,XY,t(7;18),-10,t(10;17),t(3;17),der(17)t(3;17),+20

Common event to multiple metaphases
Figure S1, related to Figure 1: Representative spectral karyotypes (SKY) of EndoC-βH1 cells at metaphase. Specific metaphases shown are S2.1 (top), 7 (middle), and S2.7 (bottom). Common structural or numerical chromosomal aberrations that are evident in multiple metaphases are indicated in green.
A Similarity of cell type chromatin states to EndoC-βH1

B Correlation of EndoC-βH1 and other tissue RNA-seq profiles

C PCA of EndoC-βH1 and other tissue miRNA profiles

D Correlation of EndoC-βH1 and Islet miRNA profiles

Chromatin State (chromHMM) Annotation

Tissue
- Adipose
- EndoC-βH1
- Islet
- Skeletal

PC1: 34.3% variance
PC2: 23.4% variance
Figure S2, related to Figure 2: Comparison of EndoC-ßH1 and islet epigenomic and transcriptomic profiles.
(A) Similarity (centered Pearson correlation) of human islet, adipocyte, skeletal muscle, GM12878, and PBMC chromatin state annotations (ChromHMM) to those of EndoC-ßH1 at EndoC-ßH1 OCRs. The similarity matrix was calculated using the “simil” function within the proxy version 0.4 R package (Meyer and Buchta, 2018).
(B) Spearman correlation between EndoC-ßH1 RNA-seq and the corresponding RNA-seq from islets, sorted alpha or beta cells (Ackermann et al., 2016), and other cell types and tissues. PBMC=peripheral blood mononuclear cells, GM12878 = lymphoblastoid cell line, CD4T = CD4+ T immune cell, skeletal = skeletal muscle, Alpha = primary islet alpha cells, Beta = primary islet beta cells.
(C) Principal component analysis (PCA) of miRNA-seq profiles from EndoC-ßH1 and 5 representative human islet, skeletal muscle, and adipose tissue samples. (D) Scatter plot illustrating the resemblance of miRNA expression levels between EndoC-ßH1 and human islets. RPMMM = reads per million mapped miRNA, R = Pearson R.
A Pathway analysis of genes adjacent to EndoC-βH1 anchors

B EndoC-βH1 Hi-C loop locations

C Specificity of EndoC-βH1 Hi-C looping locations
Figure S3, related to Figure 3: Enrichment analysis and annotation of EndoC-ßH1 Hi-C loops.

(A) Biological process gene ontology terms enriched in genes adjacent to EndoC-ßH1-specific Hi-C anchors. Enrichment analysis was performed with GREAT (McLean et al., 2010) using the single gene whose transcription start site (TSS) was nearest to each anchor. Results with an adjusted (Benjamini & Hochberg) hypergeometric p-value < 0.05 were regarded as statistically significant. (B) Frequency of EndoC-ßH1 Hi-C loop anchors and their corresponding chromatin state (ChromHMM) annotations. (C) (Top) Bar plot depicting the average similarity of chromatin state (ChromHMM) annotations for each cell type at EndoC-ßH1 Hi-C loop anchor positions. Overlaid line plots highlight the relative similarity for various chromatin state interactions (e.g. active promoter vs. active promoter). (Bottom) Select heat maps showing the frequency of human islet (left) and GM12878 (right) chromatin states at EndoC-ßH1 loop anchors.
Network connectivity of EndoC-βH1 anchor genes
MDS based on dissimilarity with EndoC-βH1 chromHMM states
Figure S4, related to Figure 4: Annotation and evaluation of EndoC-ßH1 RNA Pol2 ChIA-PET interactions.

(A) Genome-wide view of RNA polymerase 2-mediated (Pol2 ChIA-PET) chromatin interactions around the PDX1 locus on chromosome 13 in EndoC-ßH1. Asterisks under EndoC-ßH1 (red) ChIA-PET interactions indicate interacting sites/anchors identified by targeted 4C-seq analyses in this locus in human islets (Pasquali et al., 2014). (B) Connectivity of EndoC-ßH1 ChIA-PET interactions when considering nodes with 6 or more links to other regulatory elements. (Left) Circular plots depict the number of links that occur with corresponding regulatory elements (e.g., active promoter, active enhancer). (Right) bar plots illustrate the proportion of interacting nodes that exhibit the active regulatory element chromatin states exclusively in EndoC-ßH1 (blue) or identical chromatin states in both EndoC-ßH1 and islet (green). (C) Multidimensional scaling (MDS) plot based on pairwise Chi-square distances of vectors of proportions of chromatin states in EndoC-ßH1, islets, and additional Epigenomics Roadmap cell and tissue types at EndoC-ßH1 defined ChIA-PET interacting nodes.
A  Correlation of cis-regulatory element allelic imbalance

N = 1734 SNPs, R = 0.2

FDR < 0.1
ATAC, n=71
H3K27ac, n=50
Both, n=13

B  Cis-regulatory element imbalance at human islet eQTL SNP-gene pairs
Figure S5, related to Figure 5: Assessment of cis-regulatory element allelic imbalance in EndoC-ßH1 and islets.

(A) Correlation of cis-regulatory element allelic imbalance for SNPs with heterozygous genotypes in EndoC-ßH1. For a given heterozygous SNP (n = 1,734 total with 20X coverage in EndoC-ßH1 ATAC-seq and H327ac data), allelic imbalance ratios were calculated and plotted. Colored points signify SNPs with significant allelic imbalance (FDR < 10%) in ATAC-seq (red), H3K27ac (blue), or both (purple) datasets. (B) Correlation of human islet eQTL SNP-gene pair direction of effect (z-score) from (Varshney et al., 2017) and their corresponding H3K27ac effect allele bias deviation from 0.5 in EndoC-ßH1. Red points indicate eQTL SNP-gene pairs that are also linked by an EndoC-ßH1 ChIA-PET interaction. Asterisks indicate eQTL SNPs that are also diabetes-associated GWAS SNPs.