A TALEN Genome-Editing System for Generating Human Stem Cell-Based Disease Models

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SUMMARY

Transcription activator-like effector nucleases (TALENs) are a new class of engineered nucleases that are easier to design to cleave at desired sites in a genome than previous types of nucleases. We report here the use of TALENs to rapidly and efficiently generate mutant alleles of 15 genes in cultured somatic cells or human pluripotent stem cells, the latter for which we differentiated both the targeted lines and isogenic control lines into various metabolic cell types. We demonstrate cell-autonomous phenotypes directly linked to disease—dyslipidemia, insulin resistance, hypoglycemia, lipodystrophy, motor-neuron death, and hepatitis C infection. We found little evidence of TALEN off-target effects, but each clonal line nevertheless harbors a significant number of unique mutations. Given the speed and ease with which we were able to derive and characterize these cell lines, we anticipate TALEN-mediated genome editing of human cells becoming a mainstay for the investigation of human biology and disease.

INTRODUCTION

The study of human disease has been facilitated by the ability to identify the gene mutations responsible; at the same time, it has been hampered by the lack of an inexhaustible supply of easily accessible tissues from patients bearing those mutations. Another limitation is that many gene mutations that would be informative for disease biology if they could be studied in isolated cells are incompatible with human life (i.e., embryonic lethal). Classical gene-targeting technology via homologous recombination has proven to be an invaluable tool of experimental biology through its use in mouse embryonic stem cells for generating germline knockout and knockin mice; however, its use in mammalian systems has been limited primarily to studies in mice. In many cases, mice do not faithfully phenocopy human physiology and disease, e.g., cholesterol metabolism, coronary artery disease, and human hepatitis C virus (HCV) infection. The emergence of genome editing with engineered nucleases, as well as human pluripotent stem cell (hPSC) technology and differentiation protocols to obtain a variety of cell and tissue types in vitro, now make it possible to rapidly interrogate the effects of genetic modification in otherwise isogenic human model systems.

Transcription activator-like effector nucleases (TALENs) are a new class of engineered nucleases that, due to their modular domain structure, have proven more straightforward to design and construct for performing genome editing than other types of nucleases (Bogdanove and Voytas, 2011). TALENs are typically designed as a pair that binds to genomic sequences flanking a target site and generates a double-strand break, which is repaired by the cell using either homology-directed repair (HDR) or the error-prone process of nonhomologous end-joining (NHEJ) (Christian et al., 2010; Li et al., 2011; Miller et al., 2011; Hockemeyer et al., 2011). NHEJ can be exploited to introduce small insertions or deletions (indels) resulting in frameshift mutations that effectively knock out a protein-coding gene. An exogenously introduced double-stranded DNA or single-stranded DNA oligonucleotide (ssODN) can serve as a repair template for HDR to incorporate an alteration into the genome (Soldner et al., 2011). In principle, TALEN pairs can be generated de novo with standard molecular biology techniques in a matter of

days (Cermak et al., 2011; Sanjana et al., 2012). To demonstrate the utility, efficiency, and rapidity of TALEN technology in generating human cellular models with which to derive new biological insights, we created mutations in 15 genes and performed detailed phenotypic analysis of four genes for which novel roles in disease biology have emerged in recent years—*APOB*, *SORT1*, *AKT2*, and *PLIN1*.

RESULTS

Modular Assembly and Use of TALENs for Efficient and Rapid Genome Editing

The DNA-binding domain of a TALEN comprises an array of 33- to 35-amino-acid monomers that are "coded" to recognize and bind specific DNA base pairs (bp) in a 1:1 fashion (Moscou and Bogdanove, 2009; Boch et al., 2009). We built upon previously described modular Golden Gate methodologies to allow assembly of multiple DNA fragments in an ordered fashion (Li et al., 2011; Cermak et al., 2011), such that a single ligation of preassembled tetramers and trimers generates TALENs that recognize any 15 bp recognition site in the genome (Figure 1; Figure S1 available online). This assembly method requires only 1–2 days for completion and is not prone to errors that complicate methods that rely on PCR amplification of monomers. Furthermore, we have developed a set of optimized vectors and methods for the delivery of TALENs into mammalian cells and, in particular, hPSCs. In brief, we transfect or electroporate TALEN pairs into cells and then subject them to fluorescence-activated cell sorting (FACS) 48 hr posttransfection based on fluorescent-marker expression. We replate the sorted cells at low density and allow them to recover and grow for 1 week, resulting in the formation of distinct single colonies. Colonies are expanded, genomic DNA purified, and mutations analyzed by PCR, agarose-gel screening, and Sanger sequencing (Figure 1 and Figure S1). The entire process from start to finish can be completed in less than one month.

Utilizing these methods, we generated TALEN pairs to target 16 distinct sites in 15 genes in human somatic cell lines, human embryonic stem cell (hESC) lines, or human induced pluripotent stem cell (iPSC) lines; the alterations included a variety of knockout mutations as well as a knockin missense mutation and a functional frameshift mutation (Table 1). We observed that the efficiency of mutation varied by genomic location as well as among different cell lines, with indels from NHEJ occurring in roughly 2% to 34% of clones screened and the efficiency of knockin by HDR occurring at a frequency of 1.6%. We then proceeded to perform detailed phenotypic analyses of cells harboring mutations in four human disease-related genes—*APOB*, *SORT1*, *AKT2*, and *PLIN1*.

APOB Is Required for HCV Replication

APOB, which encodes apolipoprotein B, the core protein of very-low-density lipoprotein and low-density lipoprotein (LDL) particles that transport cholesterol and triglycerides from the liver to other tissues via the bloodstream, has been suggested as playing a critical role in HCV infection. In HCV models using cultured human HuH-7 hepatoma cells, RNA interference resulting in partial knockdown of *APOB* expression has been reported to reduce HCV secretion, albeit not HCV replication (Huang et al., 2007; Nahmias et al., 2008); however, another report has sug-

gested that apolipoprotein E, but not apolipoprotein B, is necessary for HCV production (Jiang and Luo, 2009). Thus, the importance of *APOB* and precise points of interaction with the HCV lifecycle remain to be determined. We sought to address this question by generating *APOB* knockout HuH-7 cells.

The human *APOB* gene encodes a 512 kDa protein termed apoB-100. We designed a TALEN pair targeting a site in exon 13 (Figure 2A); frameshift mutations at the site would generate truncated proteins about 12.5% of the size of apoB-100 (apoB-12.5). We transfected a clonal line of HuH-7 with high expression of CD81 (a coreceptor for HCV entry, HuH-7 [CD81^{hi}]) with the *APOB* TALEN pair. Following FACS with a cotranslated fluorescent marker, replating of sorted cells at a limiting dilution, and expansion of single clones, we found that of 126 screened clones, indels were present in nine clones (Figure S2A), of which four had exon 13 frameshift mutations in both alleles (Figure 2A). Compared to wild-type controls from the same set of screened clones, *APOB* knockout cells had no detectable intracellular apoB protein, no secreted apoB mass in the media, and <3% *APOB* messenger RNA (mRNA) expression, consistent with nonsense-mediated mRNA decay (Figures 2B, 2C, and 2D).

We infected *APOB*$^{-/-}$ and wild-type cells with the tissue-culture-infectious HCV strain JFH-1. The *APOB*$^{-/-}$ cells had significantly lower intracellular HCV RNA levels (74% reduction, p = 0.006), with minimal detectable HCV core protein (Figures 2B and 2E). Reintroduction of apoB-100 protein into the *APOB*$^{-/-}$ cells by adding LDL particles to the media, allowing for cellular LDL uptake, resulted in partial restoration of HCV core protein levels, arguing that the HCV replication defect was the result of loss of *APOB* function rather than an off-target effect of the TALENs (i.e., mutagenesis at other sites in the genome) (Figure 2B). Together, these data suggest that apoB-100 is integral to the HCV viral life cycle and that *APOB*-targeting therapeutics (e.g., mipomersen) may have efficacy in treating HCV-infected patients.

Isogenic Disease Models in hPSCs

We found that the karyotype of the HuH-7 cells was severely abnormal (Figure S2B)—fortuitously, it harbored two *APOB* alleles, in contrast to *SORT1*, with at least five alleles—highlighting the disadvantages of cultured tumor cell lines for rigorous genetic studies. hPSCs offer several advantages: they can maintain stable genomes with normal karyotypes while propagated in culture (Figure S2B), preserving correct gene dosage; they can be differentiated into a variety of cell types, extending studies beyond a single cell type; and they can yield human cell types that are not available as cultured cell lines, e.g., adipocytes and motor neurons.

These advantages are mitigated by the significant variability in differentiation capacity and phenotypic characteristics among different hPSC lines, particularly among iPSC lines. This variability is attributed to differences in genetic background, in epigenetic state, and in derivation of the cell lines and adaptation to culture, among other factors. In this variability lies the potential for confounding of any phenotypic differences observed among differentiated cell lines generated to serve as disease models or controls—a significant weakness of studies in which a few iPSC lines from patients are compared with a few iPSC lines from healthy individuals, as has been the case with most published
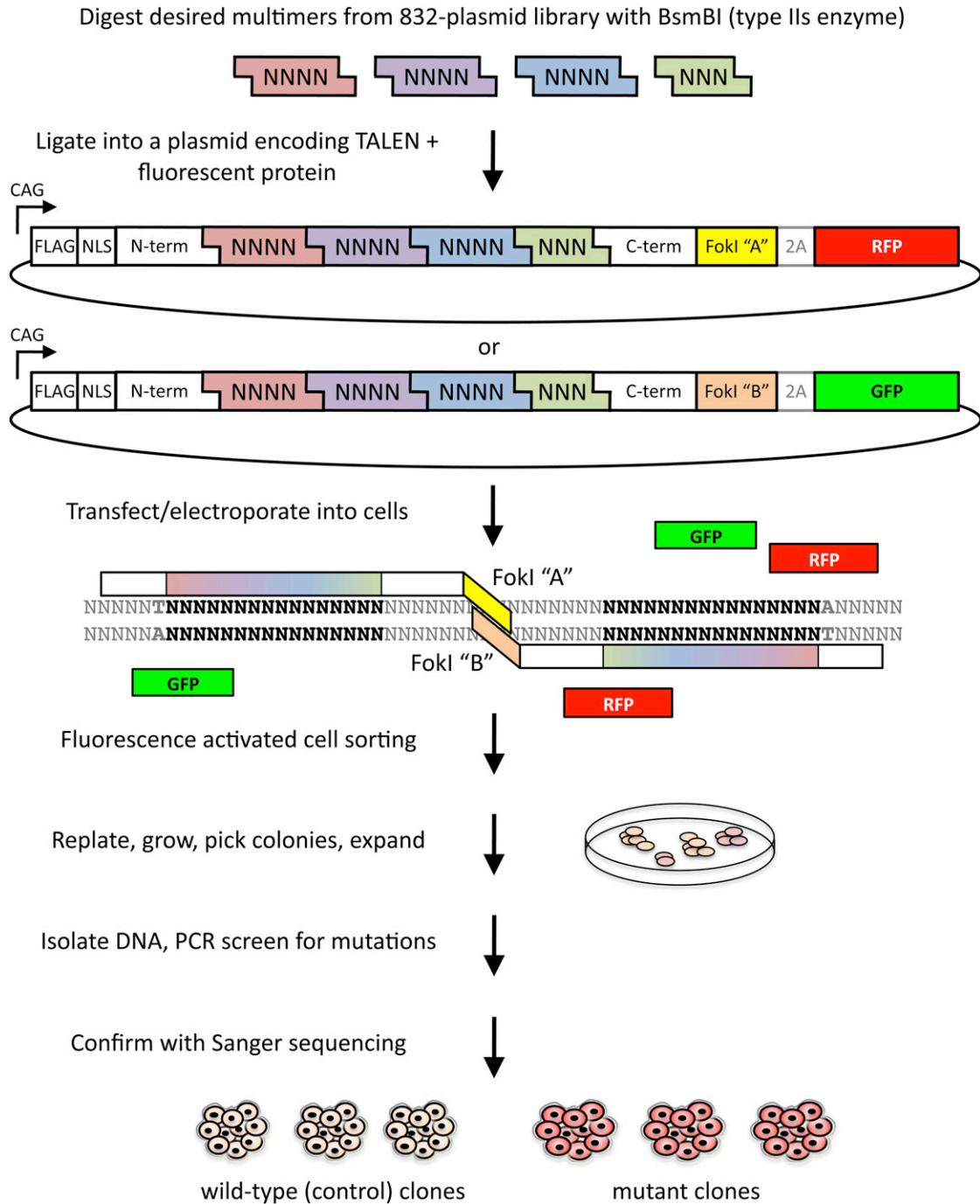
Figure 1. Schematic of System for Efficient and Rapid Genome Editing with TALENs
GFP, green fluorescent protein; RFP, red fluorescent protein; NLS, nuclear localization signal; N-term; N-terminal doman; C-term, C-terminal domain. See also Figure S1.

studies to date, because any observed differences cannot be reliably attributed to the effects of disease mutations. We demonstrated this cell line-to-cell line variability by differentiating two hESC lines, HUES 1 and HUES 9, into hepatocyte-like cells (HLCs) using an adapted protocol (Si-Tayeb et al., 2010) (Figure S3). We found that there were significant differences in the amounts of apoB and albumin secreted by the

two cell lines and retained in the media (Figure S4A); when apoB mass was normalized to albumin mass, there was a 2-fold difference between the two lines (p = 0.0001).

Using genome editing to generate isogenic cell lines that differ only with respect to a single mutation of interest provides a superior study design, because the cell lines have the same origin and would thus be matched in genetic background, epigenetic state,

Table 1. Targeting Efficiency of TALENs at 16 Loci in 15 Genes in Various Cell Types

Gene	Target Sequence (Flanked by Underlined TALEN Binding Sites)	Cell Line[a]	Number of Clones Screened	Number of Mutants	Efficiency of Mutation
AKT2	TCCCTTCCTGCCTCATTTCAGGTGAATACATCAAGACCTGGAGGCCA	HUES 9	192	17	8.9%
AKT2 E17K	TCCCTTCCTGCCTCATTTCAGGTGAATACATCAAGACCTGGAGGCCA	HUES 9	192	3[b]	1.6%
ANGPTL3	TCAAAACTTGAAAGCCTCCTAGAAGAAAAAATTCTACTTCAACAAAA	HUES 9	424	84	19.8%
APOB	TAAGCGACTGGCTGCCTATCTTATGTTGATGAGGAGTCCTTCACA	HuH-7	126	9	7.1%
ATGL	TTGCTGGCGTTGCCCCCCGCCCGCCCCCACGGCCCAGAGGACAAGGA	HUES 9	192	28	14.6%
C6orf106	TGGAGGGCATGGACGTAGACCTGGACCCGGAGCTGATGCAGAAGTTCA	HUES 9	216	13	6.0%
CIITA	TAACAGCGATGCTGACCCCCTGTGCCTCTACCACTTCTATGACCAGA	BJRiPS	292	37	12.7%
CELSR2	TGCTGGCTCGGCTGCCCTGAGGTTGCTCAATCAAGCACAGGTTTCAA	HUES 1	506	18	3.5%
CFTR	TGGAATCACACTGAGTGGAGGTCAACGAGCAAGAATTTCTTTAGCA	CF-RiPS	140	3	2.1%
GLUT4	TGGTCCTTGCTGTGTTCTCTGCGGTGCTTGGCTCCCTGCAGTTTGGGTA	HUES9	155	52	33.5%
LINC00116	TCAGAGAGGACACTGCAGTTGTCCGTGCTAGTAGCCTTCGCTTCTGGA	HUES 9	88	26	29.5%
NLRC5	TGGAGCTGTCTTGTGAGGCTGCTCACCAAAGACCCAGAATGGCTGAA	BJ-RiPS	83	6	7.2%
PLIN1	TGACAACGTGGTGGACACAGTGGTGCATTACGTGCCGGTGAGTACCA	HUES 9	293	70	23.8%
PLIN1	TGACAACGTGGTGGACACAGTGGTGCATTACGTGCCGGTGAGTACCA	BJ-RiPS	439	29	6.6%
SORT1 (exon 2)	TGATGATCTCAGAGGCTCAGTATCCTTGTCCTGGGTTGGAGATAGCA	HUES 1	576	128	22.2%
SORT1 (exon 3)	TGGTAATTATGACTTTTGGACAGTCCAAGCTATATCGAAGGTGAGATCA	HUES 9	192	21	10.9%
TRIB1	TCTTCAAGCAGATTGTCTCCGCCGTCGCCCACTGCCACCAGTCAGCCA	HUES 9	169	41	24.2%
TTN	TTGAGGCTCCTTAATGATGACAGAAGAGAAGGCTTCTCTGGGTTCA	BJ-RiPS	250	17	6.8%

TALENs, transcription activator-like effector nucleases.
[a]HUES 1 and HUES 9 are human embryonic stem cell lines (Cowan et al., 2004); BJ-RiPS and CF-RiPS are induced pluripotent stem cell lines (Warren et al., 2010); and HuH-7 cells are cultured human hepatocellular carcinoma cells.
[b]Successfully inserted E17K knockin mutation using single-stranded DNA oligonucleotide.

differentiation capacity, derivation and adaptation to culture, etc. This would minimize confounding of the experiment and allow for more confidence in concluding that any phenotypic differences are secondary to the mutation. For these reasons, our subsequent studies were all performed in genome-edited hESCs.

SORT1 Mediates Diverse Cellular Functions in Hepatocytes, Adipocytes, and Neurons

SORT1 (encoding sortilin) was recently discovered through genome-wide association studies to regulate human blood LDL cholesterol levels and risk for coronary artery disease, via the modulation of the hepatic secretion of apoB-100-containing particles into the bloodstream; however, conflicting studies in humans and mice disagree about the direction of the effect of sortilin on apoB secretion (Musunuru et al., 2010; Kjolby et al., 2010). Human genetic studies have found that SNPs associated with increased hepatic SORT1 expression are also associated with decreased blood LDL cholesterol levels (Musunuru et al., 2010). Knockdown and overexpression of Sort1 in mouse liver suggested that sortilin functions to decrease hepatocyte apoB secretion (Musunuru et al., 2010). In contrast, a study of Sort1 knockout mice suggested that sortilin increases hepatocyte apoB secretion (Kjolby et al., 2010).

We targeted exon 2 in the hESC line HUES 1 and, in a single round of TALEN targeting, generated three clones that were compound heterozygous for frameshift mutations (out of 576 clones screened) and confirmed that they lacked sortilin protein

(Figures 3A and 3B). In parallel, we targeted exon 3 in the hESC line HUES 9 and obtained two knockout clones (out of 192 clones screened). We differentiated two SORT1−/− and two wild-type HUES 1 clones or two SORT1−/− and two wild-type HUES 9 clones into HLCs using an adapted protocol (Si-Tayeb et al., 2010) (Figure S3). Measuring the levels of apoB as well as albumin and apoA-I (reference controls) secreted from the HLCs and retained in the media, we found that knockout cells had significantly increased apoB mass (HUES 1: 117% increase in apoB/albumin ratio, $p = 0.04$; HUES 9: 65% increase in apoB/albumin ratio, $p = 0.05$) (Figure 3C and Figure S4B). We infected knockout HUES 1 HLCs or HUES 9 HLCs with a lentivirus expressing the SORT1 complementary DNA (cDNA) or a control lentivirus and found that reconstitution of SORT1 to the levels observed in wild-type HUES 1 or HUES 9 HLCs resulted in normalization of the apoB mass (Figure 3D and Figure S4C), confirming that the observed differences in apoB mass are specific to SORT1 function and not the result of off-target effects. We found that secreted levels of additional hepatic proteins—ANGPTL4, ANGPTL6, HGF, and FGF-19—did not differ among the various experimental conditions (Figure S4D), nor did mRNA levels of APOB and other lipid-related genes such as HMGCR, LDLR, and SREBP1 (Figure S4E). Our data suggest that, in humans, sortilin acts in hepatocytes to reduce apoB-containing particle levels in the blood, resulting in lower cholesterol levels and reduced risk of coronary artery disease—consistent with human genetic studies (Musunuru et al., 2010) and, notably,
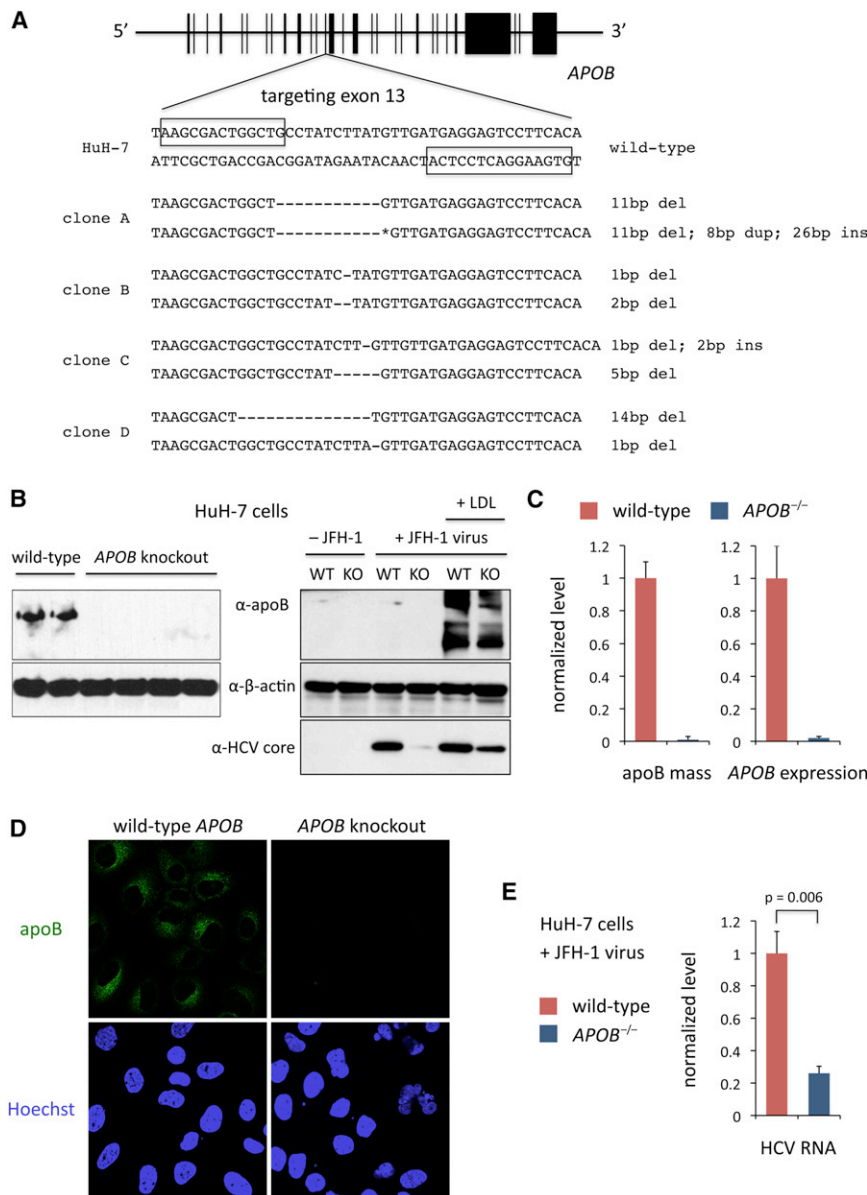
Figure 2. APOB Is Important for HCV Replication

(A) Generation of APOB knockout HuH-7 clones with TALENs targeting exon 13. Boxes indicate the TALEN binding sites. Deletions, insertions, and duplications in the two alleles of each clone are indicated. The 26 bp insertion and 8 bp duplication (asterisk) in clone A are 5′-GAGTCGCTTCT CCGGGAGATAAGTCA-3′ and 5′-GACTGGCT-3′, respectively.

(B) Left panel, western blot using whole-cell lysates from two wild-type (WT) and four knockout (KO) HuH-7 cell lines (clones A–D). Right panel, western blot from a wild-type clone and a knockout clone (clone A) infected with or without JFH-1 virus and incubated with or without LDL particles. The same wild-type clone and knockout clone (clone A) were used for all subsequent experiments.

(C) Left panel, apoB ELISAs performed on conditioned media from cells; values are normalized to the level in the wild-type clone. Right panel, APOB mRNA expression by qRT-PCR from whole-cell lysates; expression is indicated as the fold change of $2^{-\Delta\Delta Ct}$ with reference to 18S ribosomal RNA (rRNA), normalized to the level in the wild-type clone.

(D) Immunocytochemistry for apoB.

(E) HCV RNA levels by qRT-PCR from clones infected with JFH-1 virus; expression is indicated as the fold change of $2^{-\Delta\Delta Ct}$ with reference to GAPDH, normalized to the level in the wild-type clone.

The error bars show SEM from experiments with biological replicates, N = 3. p values were calculated with an unpaired t test. See also Figure S2.

p = 0.002), confirming that the loss of insulin response in the knockout adipocytes was specific to SORT1 function and not the result of off-target effects (Figure 3E and Figure S5B). Thus, SORT1 appears to be critical for insulin-responsive glucose uptake in human adipocytes and may play a role in insulin sensitivity in humans.

contradicting the results reported from Sort1 knockout mice (Kjolby et al., 2010).

SORT1 has also been suggested as playing an important role in regulating blood glucose levels by modulating insulin-dependent translocation of the fat- and muscle-specific glucose transporter, Glut4, to the plasma membrane via the formation and transport of Glut4 storage vesicles, based on studies in cultured mouse 3T3-L1 cells (Shi and Kandror, 2005). We differentiated two SORT1−/− and two wild-type HUES 1 clones into white adipocytes using a recently published protocol (Ahfeldt et al., 2012), and we observed a substantial increase in glucose uptake in wild-type adipocytes upon treatment with insulin (63% increase, p = 0.009), but not in SORT1−/− adipocytes (Figure S5A). We infected the knockout adipocytes with a SORT1 or control lentivirus and found that reconstitution of SORT1 restored insulin-responsive glucose uptake (60% increase,

Finally, SORT1 has also been implicated in the viability and function of neurons (Nykjaer and Willnow, 2012). In motor neurons, sortilin has been found to regulate neuronal survival during a temporally and spatially specific period of programmed cell death. Specifically, induction of motor-neuron cell death by the proform of brain-derived neurotrophic factor (proBDNF) has been reported to be dependent on the presence of sortilin (Teng et al., 2005; Taylor et al., 2012). We differentiated two SORT1−/− and two wild-type HUES 9 clones into TUJ1+ ISL-1+ motor neurons using an adapted protocol (Di Giorgio et al., 2008; Chambers et al., 2009) and observed that although both SORT1−/− and wild-type hPSCs generated similar numbers of motor neurons (Figure S5C), wild-type motor neurons exhibited a substantial reduction after 3 days of proBDNF treatment (23% reduction, p = 0.004), whereas SORT1−/− motor neurons were unaffected (Figures 3F and 3G).
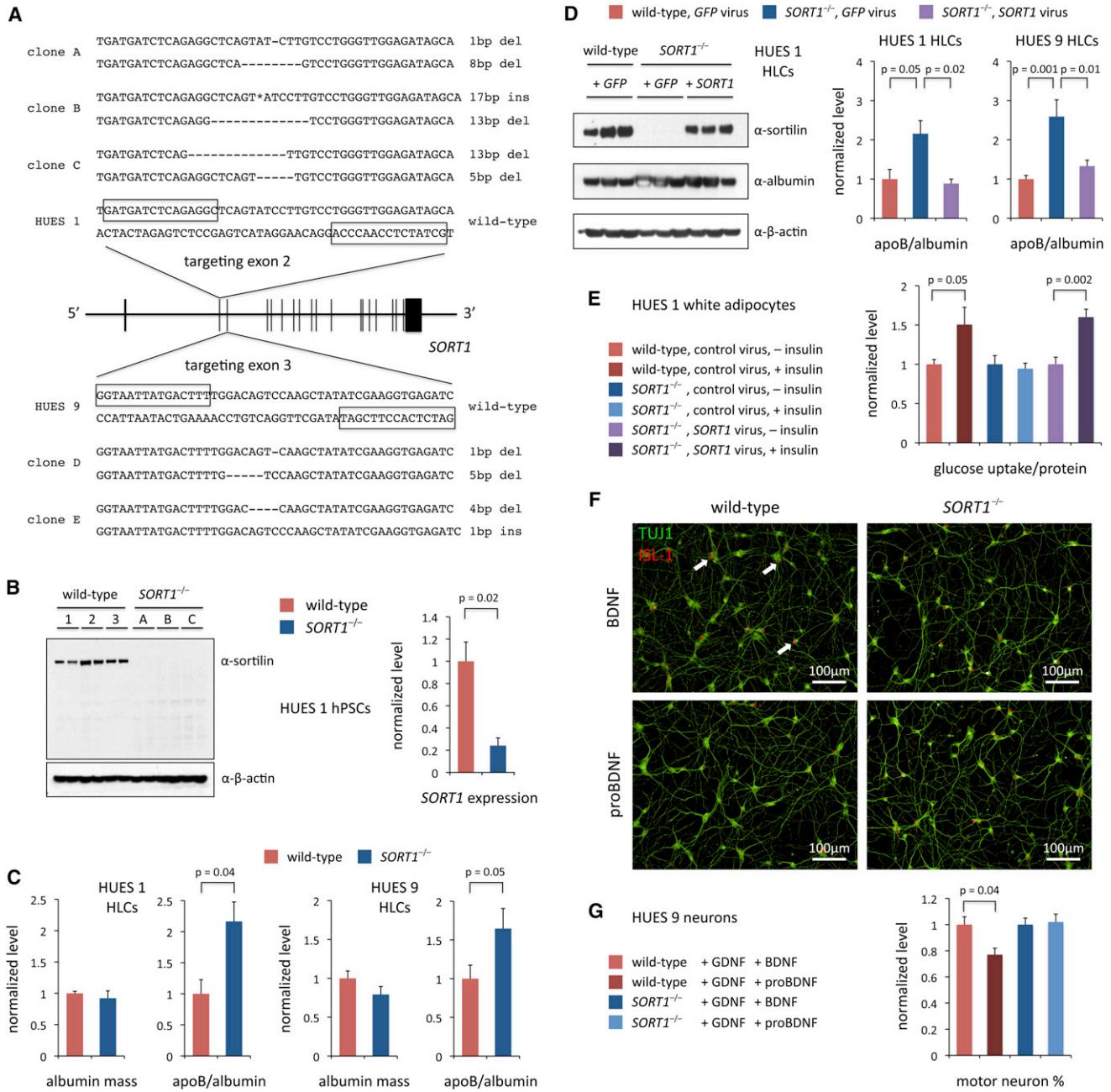
Figure 3. *SORT1* Reduces Hepatocyte-Secreted ApoB Mass, Is Important for Insulin-Responsive Glucose Transport in Adipocytes, and Mediates proBDNF-Induced Motor-Neuron Death

(A) Generation of *SORT1* knockout hPSC clones with TALENs targeting exon 2 or exon 3. Boxes indicate the TALEN binding sites. Deletions and insertions in the two alleles of each clone are indicated. The 17 bp insertion (asterisk) in clone B is 5′-TGCTATCTCCAACCAGG-3′.

(B) Western blot for sortilin and qRT-PCR for *SORT1* mRNA in wild-type and knockout HUES 1 clones (clones A–C); mRNA expression is indicated as the fold change of $2^{-\Delta\Delta Ct}$ with reference to 18S rRNA, normalized to the mean level in the wild-type clones.

(C) Albumin and apoB mass measured by ELISA in media collected from wild-type and knockout HLCs (two clones each; A and B for HUES 1, D and E for HUES 9), normalized to mean levels in wild-type HLCs. N = 3 for HUES 1, N = 6 for HUES 9.

(D) Western blots of lysates and ELISAs in media from wild-type and knockout HLCs (one clone each, A for HUES 1; two clones each, D and E for HUES 9) infected with *SORT1*- or *GFP*-expressing lentivirus, normalized to mean levels in wild-type HLCs. N = 2 for HUES 1, N = 6 for HUES 9.

(E) Ratios of glucose uptake to total protein content in wild-type and knockout HUES 1 adipocytes (one clone each; clone A) infected with *SORT1*-expressing or control lentivirus and treated with or without insulin, all normalized to the mean ratio in wild-type adipocytes without insulin. N = 6.

(F) Immunocytochemistry for TUJ1 and ISL-1 in wild-type and knockout HUES 9 motor neurons. Arrows indicate representative double-positive cells.

(G) Counts of wild-type and knockout HUES 9 (two clones each, D and E) motor neurons (TUJ1 and ISL-1 double-positive cells) treated with BDNF versus proBDNF. N = 12. GDNF, glial cell-derived neurotrophic factor.

The error bars show SEM from experiments with biological replicates. p values were calculated with an unpaired t test. See also Figures S3, S4, and S5.
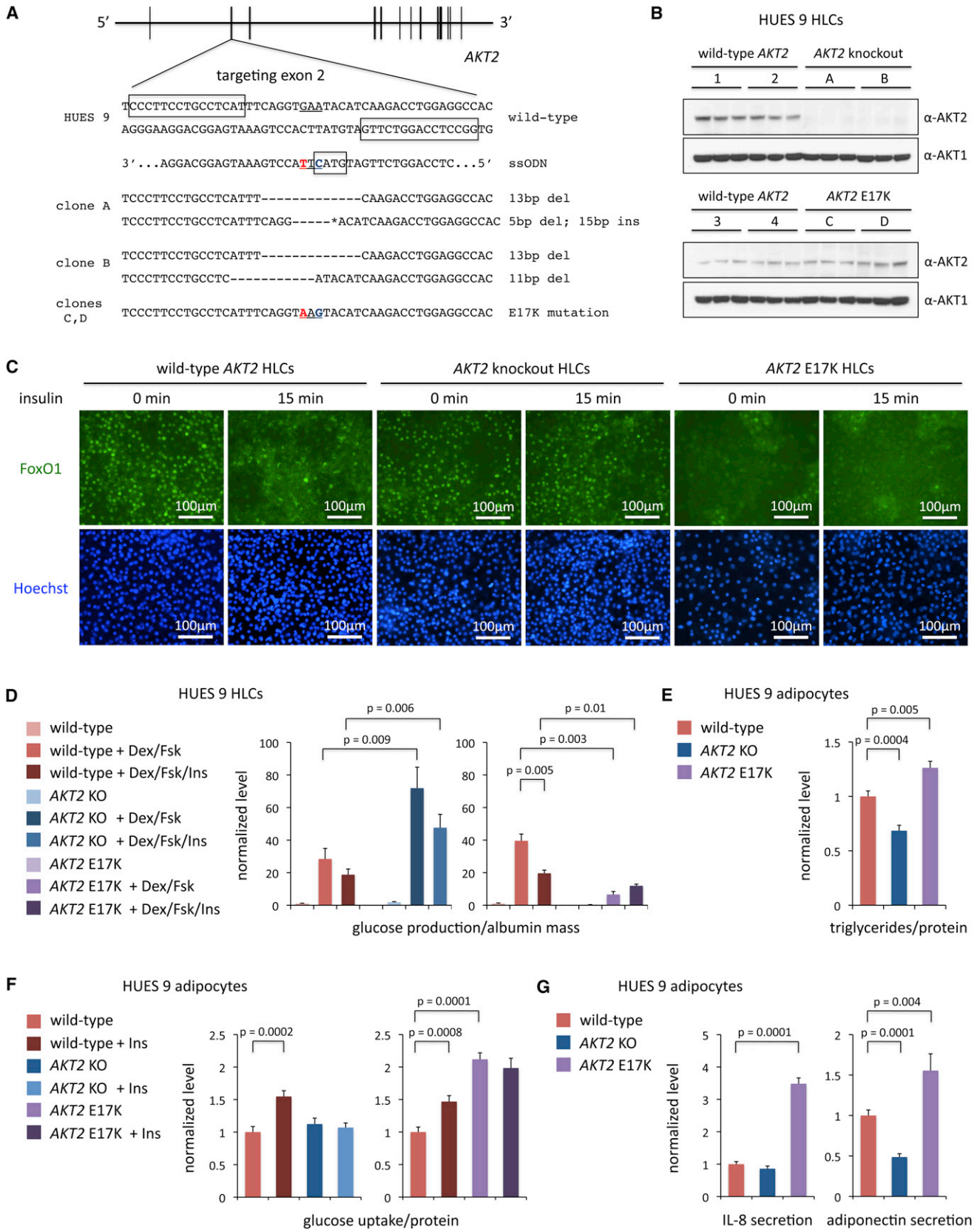
These data agree with the reported requirement of *SORT1* for proBDNF-induced programmed cell death in human motor neurons.

AKT2 Regulates Insulin Signaling and Glucose Metabolism in Hepatocytes and Adipocytes

The human *AKT2* gene (encoding RAC-β serine/threonine-protein kinase, also known as AKT2 or PKBβ) has also been implicated in the regulation of insulin sensitivity. Loss of function of *AKT2* in humans has been reported to result in severe insulin resistance, as well as decreased body fat and partial lipodystrophy, attributed to reduced adipocyte differentiation (George et al., 2004; Agarwal and Garg, 2006); also, *Akt2* knockout mice are resistant to the effects of insulin on glucose metabolism in liver and muscle, and they manifest lipoatrophy (Cho et al., 2001; Garofalo et al., 2003). Recently, three patients with severe hypoglycemia, hypoinsulinemia, and increased body fat were reported to bear a missense mutation in *AKT2*, p.Glu17Lys (E17K) (Hussain et al., 2011). Although the function of the mutant AKT2 E17K protein was assessed by heterologous overexpression studies in cultured cell lines (HeLa and 3T3-L1) and interpreted as being activated, the inability to study a physiologically relevant phenotype (e.g., glucose metabolism) in physiologically relevant tissues (e.g., human liver) precluded the conclusion that the *AKT2* mutation was causal for the metabolic disorder in the patients.

We sought to unequivocally establish a dominant, activated function of the AKT2 E17K mutant on glucose metabolism by generating an allelic series of isogenic hPSC lines with wild-type *AKT2*, knockout of *AKT2*, or a single *AKT2^{E17K}* allele. We designed TALENs to target the site of the E17K mutation in the second coding exon (Figure 4A). In one round of targeting of HUES 9 cells with the TALEN pair alone, we obtained 17 clones with indels (out of 192 clones screened), none of which was compound heterozygous for frameshift mutations. A second round of TALEN targeting with a clone with one frameshift allele yielded two clones compound heterozygous for frameshift mutations (out of 96 clones screened). In parallel, we coelectroporated wild-type HUES 9 cells with the TALEN pair and a 67 nt antisense ssODN harboring the E17K missense variant, yielding three *AKT2^{E17K}* heterozygous clones (out of 192 clones screened) (Figure S6A).

We differentiated the allelic series of hPSC clones (two clones each) into HLCs. No AKT2 protein was apparent in the knockout cells, with comparable levels of AKT2 observed in the wild-type and E17K cells (Figure 4B). We assessed the regulation of the FoxO1 transcription factor, an AKT2 substrate that upon phosphorylation is translocated from the nucleus to the cytoplasm. In wild-type HLCs, FoxO1 was predominantly nuclear at baseline and cytoplasmic after insulin stimulation; in *AKT2^{−/−}* HLCs, it was predominantly nuclear both at baseline and after stimulation; and in *AKT2^{E17K}* HLCs, it was predominantly cytoplasmic both at baseline and after stimulation (Figure 4C). We assessed glucose production in the allelic series of HLCs and found that with all three genotypes, addition of dexamethasone and forskolin to the media dramatically increased glucose production; the further addition of insulin decreased glucose production in the wild-type HLCs, but not in the mutant HLCs (Figure 4D). In all media conditions, glucose production was significantly higher in *AKT2^{−/−}* HLCs and lower in *AKT2^{E17K}* HLCs compared to wild-type HLCs. Similar trends were observed in the mRNA expression levels of two genes involved in gluconeogenesis, *G6PC* and *PCK1* (Figure S6B).

We also differentiated the *AKT2* allelic series of hPSC clones into white adipocytes and found that *AKT2^{−/−}* adipocytes had significantly decreased triglyceride content (32% reduction, p = 0.0004) and *AKT2^{E17K}* adipocytes had significantly increased triglyceride content (26% increase, p = 0.005) (Figure 4E), consistent with the fat-related phenotypes observed in patients with *AKT2* mutations. We observed a substantial increase in glucose uptake in wild-type adipocytes upon treatment with insulin (∼50% increase in two different experiments) but, as with *SORT1^{−/−}* adipocytes, we observed no significant change in glucose uptake in *AKT2^{−/−}* adipocytes with insulin (Figure 4F). In contrast, *AKT2^{E17K}* adipocytes displayed higher levels of glucose uptake at baseline compared to wild-type adipocytes (111% increase, p = 0.0001); upon treatment with insulin, there was no further increase in glucose uptake, presumably because the cells were in a constitutively active state with respect to insulin signaling (Figure 4F). In the same vein, *AKT2^{E17K}* adipocytes displayed substantially increased secretion of inflammatory adipokines such as interleukin-18 (IL-8) (Figure 4G), MCP1, and PAI-1 (Figure S6C). Finally, *AKT2^{−/−}* and *AKT2^{E17K}*

Figure 4. *AKT2* E17K Is a Dominant, Activating Mutation

(A) Generation of *AKT2* knockout and E17K knockin hPSC clones with TALENs targeting exon 2. The underline indicates the codon encoding the E17K change. Boxes in the wild-type sequence indicate the TALEN binding sites. The box in the ssODN sequence indicates the RsaI restriction site created by the synonymous mutation in blue; the nucleotide in red indicates the missense mutation for E17K. Deletions and insertions in the two alleles of each clone are indicated. The 15 bp insertion (asterisk) in clone A is 5′-GACCTCCAGGTCCTG-3′.

(B) Western blot for AKT2 and AKT1 in wild-type, knockout, and E17K HUES 9 HLCs (A and B for knockout; C and D for E17K).

(C) Immunocytochemistry for FoxO1 in HLCs (clones A and C) at baseline and after 15 min of insulin stimulation.

(D) Ratios of glucose production to secreted albumin mass (ELISA) in wild-type, knockout, and E17K HUES 9 HLCs (two clones each; A and B for knockout; C and D for E17K) treated with or without dexamethasone (Dex) and forskolin (Fsk) with or without insulin (Ins), all normalized to the mean ratios in wild-type HLCs without additives. Left panel, N = 4; right panel, N = 2.

(E) Ratios of triglyceride content to total protein content in wild-type, knockout, and E17K HUES 9 adipocytes (two clones each; A and B for knockout; C and D for E17K), normalized to the mean ratio in wild-type adipocytes. N = 5.

(F) Ratios of glucose uptake to total protein content in wild-type, knockout, and E17K HUES 9 adipocytes (two clones each; A and B for knockout; C and D for E17K) treated with or without insulin, all normalized to the mean ratio in wild-type adipocytes without insulin. N = 6.

(G) IL-8 and adiponectin mass measured by ELISA in media collected from wild-type, knockout, and E17K HUES 9 adipocytes (two clones each; A and B for knockout; C and D for E17K), normalized to mean levels from wild-type adipocytes. N = 4.

The error bars show SEM from experiments with biological replicates. p values were calculated with an unpaired t test. See also Figures S3 and S6.
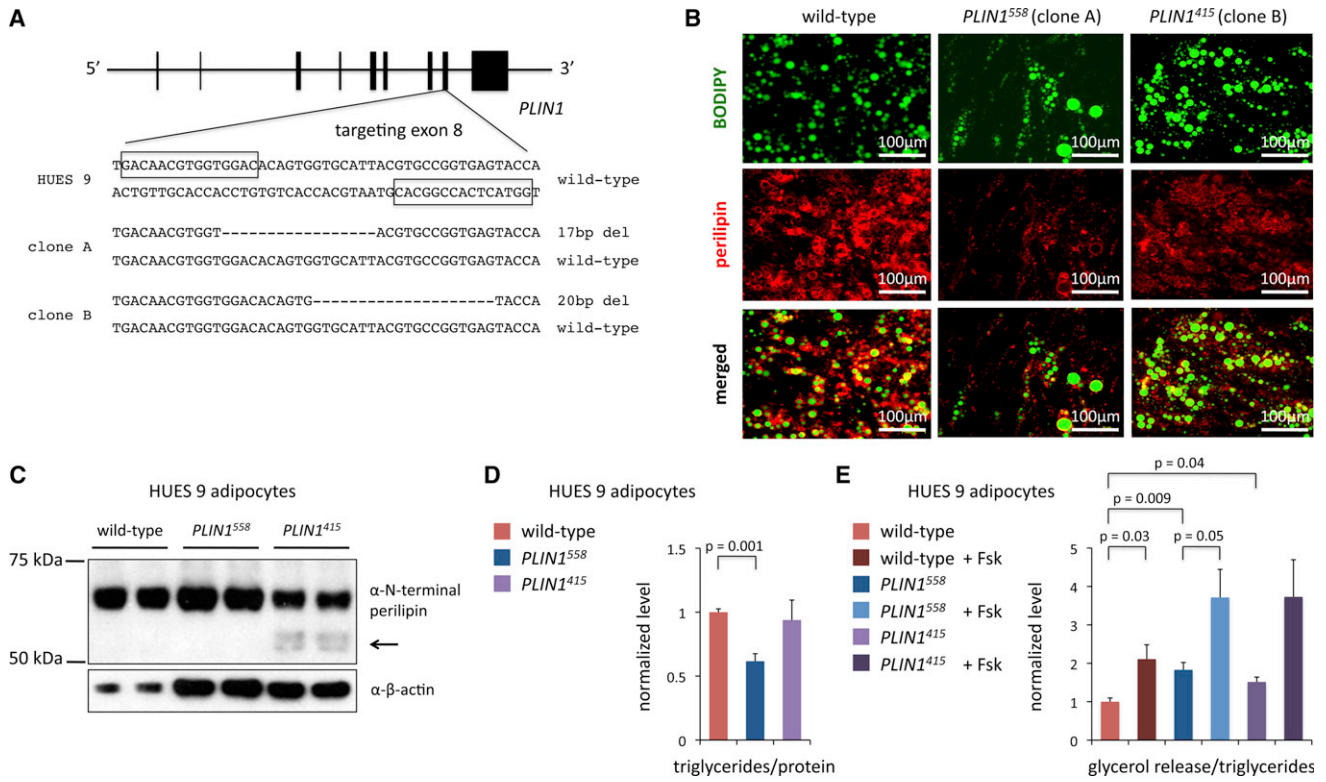
Figure 5. *PLIN1* Frameshift Mutations Produce Dominant-Acting and Truncated Proteins
(A) Generation of *PLIN1* frameshift mutant hPSC clones with TALENs targeting exon 8. Boxes indicate the TALEN binding sites. Deletions in the one allele of each clone are indicated.
(B) Immunocytochemistry for perilipin and BODIPY in wild-type, *PLIN1*[558] (clone A), and *PLIN1*[415] (clone B) adipocytes.
(C) Western blot for the N terminus of perilipin in wild-type, *PLIN1*[558], and *PLIN1*[415] adipocytes; the arrow indicates a truncated protein.
(D) Ratios of triglyceride content to total protein content in wild-type, *PLIN1*[558], and *PLIN1*[415] adipocytes. N = 3.
(E) Ratios of lipolysis activity (as measured by glycerol release) to total triglyceride content in wild-type, *PLIN1*[558], and *PLIN1*[415] adipocytes treated with or without 10 μM forskolin. N = 4.
The error bars show SEM from experiments with biological replicates. p values were calculated with an unpaired t test. See also Figure S7.

adipocytes showed decreased and increased secretion of adiponectin, respectively (Figure 4G).

The opposing effects of the knockout and *AKT2*[E17K] alleles, in addition to indicating that the effects were specific to *AKT2* function and not the result of off-target effects, establish that E17K is indeed a dominant, activating mutation in *AKT2* and causal for the hypoglycemia and increased body fat observed in the three patients.

PLIN1 Frameshift Mutations Dominantly Alter Lipolysis in Adipocytes
PLIN1 encodes the protein perilipin, the most abundant protein coating lipid droplets in adipocytes, where it is required for droplet formation and maturation, optimal triglyceride storage, and the release of free fatty acids from the droplet (Brasaemle et al., 2009). Frameshift mutations in *PLIN1* have recently been identified in patients with a novel autosomal-dominant subtype of partial lipodystrophy (Gandotra et al., 2011). The frameshift mutations found in patients result in a C-terminal elongation of perilipin, with a significantly altered amino acid sequence. Mice lacking *Plin1* exhibit elevated levels of basal lipolysis in adipocytes (Tansey et al., 2001; Zhai et al., 2010), which has been

suggested as the mechanism by which patients harboring the frameshift mutations develop lipodystrophy (Gandotra et al., 2011). Mechanistic studies of these disease-causing mutations have been limited to the overexpression of mutant and wild-type human cDNAs in mouse 3T3-L1-derived adipocytes, with the conclusion being that wild-type *PLIN1*, but not mutant *PLIN1*, is able to inhibit basal lipolysis (Gandotra et al., 2011).

We designed TALENs to target the site of one of the naturally occurring patient-specific mutations (Val398fs) in the eighth coding exon of *PLIN1* (Figure 5A). In a single round of targeting of HUES 9 cells we identified 70 mutant clones (out of 293 clones screened). We characterized two mutant clones, one of which harbors a frameshift mutation that elongates perilipin to a length of 558 amino acids (designated *PLIN1*[558]; wild-type perilipin has 522 amino acids)—very similar to the effect of the naturally occurring Val398fs mutation—and the other of which has a frameshift resulting in a C-terminal truncation of the protein (415 amino acids; designated *PLIN1*[415]) (Figure S7A).

We differentiated the allelic series of hPSCs—wild-type, *PLIN1*[558], and *PLIN1*[415]—into white adipocytes and observed a substantial reduction in the number of lipid-droplet-containing cells, as well as smaller lipid droplets in *PLIN1*[558] adipocytes

compared to either wild-type or $PLIN1^{415}$ cell lines (Figure 5B and Figure S7B). We confirmed the presence of perilipin protein in the adipocytes via western blot analysis (Figure 5C and Figure S7C). We found that the $PLIN1^{558}$ adipocytes had significantly reduced triglyceride content (38% reduction compared to wild-type adipocytes, p = 0.0009), whereas the $PLIN1^{415}$ adipocytes had similar triglyceride content to wild-type adipocytes (Figure 5D). We also measured basal and forskolin-stimulated lipolysis and found that both $PLIN1^{558}$ and $PLIN1^{415}$ adipocytes had increased basal lipolysis compared to wild-type adipocytes (83% increase, p = 0.008, and 52% increase, p = 0.04, respectively) (Figure 5E). Interestingly, both the $PLIN1^{558}$ and $PLIN1^{415}$ mutations resulted in increased lipolysis, though the effect was more marked with the $PLIN1^{558}$ mutation; furthermore, there were significant differences in the expression levels of adipocyte-specific genes between the $PLIN1^{558}$ and $PLIN1^{415}$ cells, underscoring that the two different frameshifts (one leading to elongation of perilipin, the other to truncation) have distinct functional consequences (Figure S7D). Together, these data point to the C-terminal elongated form of perilipin, via a frameshift similar to naturally occurring mutations in lipodystrophy patients, acting in a dominant fashion to alter lipolysis and reduce triglyceride storage and lipid-droplet formation in human adipocytes.

TALENs Exhibit Minimal Off-Target Effects, but Sequence Variants Abound

Because the extent of off-target effects of TALENs (i.e., mutagenesis at other sites in the genome) in hPSCs remains to be defined, we performed exome sequencing of six cell lines: the parental HUES 1 cell line (clone X); the three *SORT1* knockout HUES 1 clones (clones A–C in Figure 3A); a control HUES 1 clone that had been grown in parallel with the *SORT1* knockout clones (i.e., had been exposed to the *SORT1* exon 2 TALEN pair but retained two wild-type alleles; clone W); and a clone that had been targeted in the *CELSR2* gene with a different TALEN pair (clone Y). It should be noted that TALENs virtually always induce indels by NHEJ rather than single-nucleotide variants (SNVs). Restricting our analysis to novel DNA sequence variants not found in the parental HUES 1 cell line, we identified the known on-target indels in *SORT1* in clones A–C; otherwise, we identified just two indels in the exome, a 4 bp deletion in clone C in the coding sequence of *LARP6*, resulting in a predicted frameshift mutation, and a 1 bp deletion in clone Y in an intron near an exon-intron boundary of *LUC7L3* (Table 2). Neither of these sites is flanked by sequences resembling predicted TALEN binding sites, arguing against (but not ruling out) the indels being TALEN-mediated off-target effects. More noteworthy were the 35 Sanger-sequencing-confirmed SNVs we discovered across the five experimental and control clones (Table 2). None of these SNVs lay near predicted off-target TALEN binding sites (see next paragraph). It is more likely that these SNVs represent intrinsic and perhaps unavoidable heterogeneity among single-cell clones of the original pool of HUES 1 cells. Arguing in favor of this interpretation, several of the SNVs were shared by both experimental and control clones, implying a common clonal origin within the original pool of HUES 1 cells. The functional significance of these SNVs is unclear, but the majority resulted in missense mutations, and at least one

lay in a well-established disease gene (*DMD*, responsible for Duchenne muscular dystrophy).

We also performed whole-genome sequencing of the same six cell lines. Because the sequencing was performed at low coverage (6–12× coverage on average), it was not possible to perform de novo genome assembly and ascertain all sequence variation among the genomes. Instead, we used the sequencing data to interrogate the sites in the genome at which one or the other TALEN of the *SORT1* exon 2 TALEN pair would be most likely to bind based on a weighted TAL monomer-nucleotide association probability matrix developed by Doyle et al. (2012), and would thus be most likely to induce an off-target sequence change. About 100,000 potential off-target genomic sites were identified; we screened all of these sites for evidence of nearby indels. Besides the known *SORT1* indels in clones A–C, we identified no indels meeting our criteria.

Thus, although we are not able to completely rule out TALEN off-target effects, we conclude that off-target indels rarely occur based on the results of the exome and whole-genome sequence analyses. However, extrapolating to the entire genome, we expect that each clonal cell line harbors hundreds of SNVs that distinguish it from other cell lines derived from the same pool of parental cells. Thus, it may be virtually impossible to derive truly isogenic cell lines, even with the minimized manipulation of cells entailed by our genome-editing system.

DISCUSSION

In our studies we have used human model systems to generate strong evidence that apoB-100 is critical for HCV replication in human hepatocytes; that sortilin reduces apoB secretion by human hepatocytes, facilitates insulin-mediated glucose uptake by human adipocytes, and is necessary for proBDNF-mediated motor-neuron apoptosis; that *AKT2* E17K is a gain-of-function mutation that leads to reduced glucose production in human hepatocytes and increased triglyceride content in human adipocytes; and that *PLIN1* frameshift mutations increase basal lipolysis in human adipocytes. More generally, these findings highlight the various types of studies to which genome editing in human cells may be applied for obtaining novel biological insights.

We note that genome-editing technology is rapidly advancing, and we anticipate that improvements in the engineering of TALENs will continue to make genome editing more rapid and efficient. Indeed, since we established our system, high-throughput automated assembly methods have been reported (Reyon et al., 2012; Briggs et al., 2012), as well as the characterization of TAL monomers with improved nucleotide-binding specificity (Streubel et al., 2012; Cong et al., 2012). Although our specific TALEN assembly platform does not incorporate these latest advancements, in principle any up-to-date assembly platform that is paired to a delivery methodology similar to ours should be able to achieve efficient genome editing on a timescale of less than a month.

Whatever the assembly platform, our studies suggest that TALENs incur a low burden of off-target effects but that there is nevertheless significant clone-to-clone genetic variation in the form of SNVs; even if not secondary to TALEN use, they cannot be ignored. The ease and rapidity of TALEN-mediated

Table 2. Clonal Sequence Variants Detected by Exome Sequencing and Their Predicted Effects on Protein Products

Chr	Pos	Ref	Alt[a]	Gene	Clone A[b]	Clone B	Clone C	Clone W	Clone Y
1	21014224	G	T	KIF17	–	–	S432Y	–	–
1	90178297	G	A	LRRC8C	–	–	–	P56P	–
1	149921554	C	T	OTUD7B	–	–	–	–	Q367Q
2	109347284	T	C	RANBP2	–	–	–	F39F	–
2	113820048	C	A	IL36RN	–	–	–	–	L88I
2	173330424	C	T	ITGA6	–	intronic	–	–	–
3	132319348	T	C	CCRL1	–	–	–	–	V36A
3	193002674	G	T	ATP13A5	intronic	–	intronic	intronic	–
4	113568377	GAAGA	G	LARP7	–	–	frameshift[c]	–	–
6	152652970	C	A	SYNE1	–	–	–	–	A4284S
7	2260574	C	T	MAD1L1	–	–	–	–	G47D
7	22179660	G	C	RAPGEF5	L601V	–	–	–	–
8	15508246	C	T	TUSC3	–	–	R117C	–	–
9	33354142	G	A	NFX1	E130K	–	–	–	–
10	75541848	C	T	CHCHD1	–	S5S	–	–	–
11	124095534	G	A	OR8G2	–	–	G46E	–	–
12	80752469	G	T	OTOGL	–	C2026F	–	–	–
12	101750754	G	T	UTP20	R1862I	–	R1862I	R1862I	–
12	105445861	G	T	ALDH1L2	–	intronic	–	–	–
14	21967435	C	A	METTL3	–	intronic	–	–	–
15	43827249	G	T	PPIP5K1	–	–	P1071T	–	–
15	48805891	C	A	FBN1	–	–	intronic	–	–
15	90631649	C	A	IDH2	–	–	–	–	W155L
16	20043135	C	A	GPR139	–	S328S	–	–	–
17	11998899	G	A	MAP24K4	R145Q	–	R145Q	R145Q	–
17	48817653	AT	A	LUC7L3	–	–	–	–	intronic[d]
17	73567174	G	T	LLGL2	–	L712L	–	–	–
17	79226178	C	A	SLC38A10	–	–	–	A588S	–
18	30992086	G	T	C18orf34	–	–	–	–	intronic
19	9028296	G	T	MUC16	–	–	–	–	L12166M
19	56552245	G	T	NLRP5	intronic	–	intronic	intronic	–
20	60498685	G	T	CDH44	–	E425D	–	–	–
22	37325592	C	A	CSF2RB	D100E	–	D100E	D100E	–
22	39078278	A	T	TOMM22	–	intronic	–	–	–
X	32328257	G	T	DMD	–	A1897D	–	–	–
X	48558634	C	T	SUV39H1	–	–	T106T	–	–
X	135956509	C	A	RBMX	–	–	–	–	S323I

Chr, chromosome; Pos, position; Ref, reference allele; Alt, alternate allele.

[a]Alternate allele compared to reference allele in parental HUES 1 cell line (clone X), confirmed by Sanger sequencing.

[b]Clones A, B, and C are SORT1^{−/−} clones in HUES 1; clone W is a wild-type (SORT1^{+/+}) clone in HUES 1 that was exposed to SORT1 transcription activator-like effector nucleases (TALENs), from the same pool as clones A–C; clone Y is a wild-type (SORT1^{+/+}) clone that was exposed to a different set of TALENS (targeting CELSR2).

[c]The surrounding sequence is CAGAA GAGAA GAAAA AGAAA AAGAA **GAAGA** AAGGC CGAAT GAAAA AGGAA GACAA.

[d]The surrounding sequence is ATAAT CACAG ATAAT TTATA CAATT AT**AT**T TTTTC CCCCA GGTCC GTGTG AAAAA.

genome editing allows for rigorous study designs that can alleviate any concerns about off-target effects or other potential confounding by clonal sequence variation. As we have demonstrated with SORT1, it is straightforward to (1) generate multiple distinct mutant cell lines with each TALEN pair, (2) use distinct TALEN pairs to target different sites in a gene, (3) generate mutant clones in different cell lines with different genetic backgrounds, and (4) perform reconstitution experiments in knockout clones. Having used all of these approaches, we are able to conclude with great confidence that the observed cellular phenotypes are indeed related to SORT1 function. We suggest that using at least one of these approaches should become de rigeur for future genetic studies in order to minimize confounding by clonal sequence variation.

Given the ability to use TALENs to readily insert specific gene variants into cells, the current enthusiasm for the generation and comparison of "disease" iPSC lines from patients with genetic disorders and "control" iPSC lines from unmatched healthy individuals should shift to the use of genome editing to engineer isogenic cell lines with and without disease mutations. The time required to recruit a patient for the donation of tissue from which to make iPSCs (assuming such a patient is readily accessible, which may not be the case for rare disorders), to perform reprogramming for deriving iPSC clones, to perform quality control for identifying clones that are pluripotent and that will readily differentiate into the desired cell type, and then to undertake differentiation and phenotypic studies—in the absence of isogenic control cell lines—is a minimum of six months and usually longer. Within a shorter time frame, we have found it to be quite feasible to use TALENs to edit a well-characterized and prevalidated (with respect to differentiation capacity) hPSC line and yield both mutant cell lines and isogenic control cell lines, allowing for a more rigorous study design, and to undertake differentiation and phenotypic studies, without any need for patient contact.

The potential advantages offered by prevalidated wild-type cell lines notwithstanding, there are many disorders for which the genetic background (i.e., modifier genes) plays a significant role in determining whether disease mutations result in clinical phenotypes. In these cases, it will be important to use iPSC lines from patients with clinically apparent diseases in order to have cell lines with the correct genetic backgrounds for complete disease penetrance (whereas wild-type cell lines may have nonpermissive genetic backgrounds). Genome editing with TALENs could be readily applied to patient-specific iPSC lines to "cure" disease mutations and generate appropriate isogenic control lines. Indeed, the most robust study design possible may be to assess both the effect of inserting a disease mutation into a wild-type cell line, thereby testing the sufficiency of the mutation for disease, and the effect of removing a disease mutation from a patient-specific iPSC line, thereby testing the necessity of the mutation for disease. Certainly the rapidity and efficiency of genome editing with TALENs should make it feasible to test the effects of a disease mutation in a variety of genetic backgrounds.

Finally, genome editing potentially allows for the interrogation of a large number of DNA sequence variants, such as those now emerging from next-generation sequencing studies of human populations, on a single genetic background. Creating a robust allelic series of isogenic cell lines represents an approach that hitherto has only been possible in nonmammalian organisms. Such studies will represent a significant advance in our ability to dissect genotype-phenotype relationships and thereby better elucidate human biology and disease.

EXPERIMENTAL PROCEDURES

TALEN Construction

TALEN genomic binding sites were chosen to be 15 bp in length or, in a few cases, 13 bp in length such that the target sequence between the two binding sites was between 14 and 18 bp in length; each binding site was anchored by a preceding T base in position "0" as has been shown to be optimal for naturally occurring TAL proteins (Moscou and Bogdanove, 2009; Boch et al., 2009). A library of 832 tetramer or trimer TAL repeats was constructed using methods based on the PCR-based protocol of Zhang et al. (2011); these multimers were

designed to have complementary sticky ends when digested out of library plasmids with the type IIs restriction enzyme BsmBI. As outlined in Figure 1, multimers were assembled into an array and subcloned into a full-length TALEN harboring, in order, an N-terminal FLAG tag, a nuclear localization signal, the N-terminal portion of the TALE PthXo1 from the rice pathogen *X. oryzae* pv. oryzae (a kind gift of Dr. Daniel Voytas, University of Minnesota) lacking the first 176 amino acids (after Miller et al., 2011), the engineered TAL repeat array, the following 63 amino acids from the corresponding C-terminal portion of PthXo1 (after Miller et al., 2011), and one of two enhanced FokI domains. The FokI domains used were obligate heterodimers with both the Sharkey (Guo et al., 2010) and ELD:KKR (Doyon et al., 2011) mutations for enhancing cleavage activity, engineered by PCR. Each TALEN was in a plasmid with the CAG promoter for optimal expression in hPSCs, with the TALEN being coexpressed with a fluorescent marker (enhanced green fluorescent protein, mCherry [Clontech], or turbo red fluorescent protein [Evrogen]) via an intervening viral 2A sequence. The generic TALEN protein sequences are shown in Figure S1A. All reagents, protocols, and plasmid sequences needed for generating TALENs and performing genome editing by the methods described in this manuscript will be available to academic researchers through Addgene (http://www.addgene.org/TALEN_genome_editing_collection).

Cell Culture, Transfection or Electroporation, and Sorting

HuH-7 (CD81hi) cells were grown in adherent culture in Dulbecco's modified Eagle's medium High Glucose containing glutamine and pyruvate (Invitrogen) and supplemented with 10% fetal bovine serum, penicillin, and streptomycin. Transfection of the plasmids expressing the *APOB* TALEN pair into HuH-7 cells was performed using FuGENE 6 (Roche) in 10 cm tissue-culture plates according to manufacturer instructions. HUES 1 and HUES 9 cells (Cowan et al., 2004) were grown in feeder-free adherent culture in chemically defined mTeSR1 (STEMCELL Technologies) supplemented with penicillin and streptomycin on plates precoated with Geltrex matrix (Invitrogen). The cells were disassociated into single cells with Accutase (Invitrogen), and 10 million cells were electroporated with 50 μg of the TALEN pair (25 μg of each plasmid), or with a mix of 30 μg of the TALEN pair (15 μg of each plasmid) and 30 μg of the ssODN (5′-CAGGA AGTAC CGTGG CCTCC AGGTC TTGAT GTACT **T**ACCT GAAAT GAGGC AGGAA GGGAG GGAGA GA-3′) in a single cuvette and replated as previously described (Schinzel et al., 2011). The cells were collected from the culture plates 48 hr posttransfection or postelectroporation by trypsin or Accutase treatment, respectively, and resuspended in PBS. Cells expressing green and/or red fluorescent markers were collected by FACS (FACSAria II; BD Biosciences) and replated on 10 cm tissue-culture plates at 15,000 cells/plate to allow for recovery in growth media.

Isolation of Targeted Clonal Cell Populations

Post-FACS, the cells were allowed to recover for 7–10 days, after which single colonies were manually picked, dispersed, and replated individually to wells of 96-well plates. Colonies were allowed to grow to near confluence over the next 7 days, at which point they were split using trypsin (for HuH-7 cells) or Accutase (for hESCs) and replica-plated for the creation of a working stock and a frozen stock. The working stock was grown to confluence. Genomic DNA was extracted in 96-well format from working stocks in lysis buffer (10 mM Tris [pH 7.5], 10 mM EDTA, 10 mM NaCl, 0.5% Sarcosyl) containing proteinase K at 56°C overnight in a humidified chamber. Genomic DNA was precipitated by the addition of 95% ethanol containing 75 mM NaCl for 1 hr at room temperature. The DNA was then washed two times in 70% ethanol, allowed to dry at room temperature, and then resuspended in nuclease-free water.

Genotyping at the TALEN target site was then performed for each sample by PCR amplication (94°C, 30 s; 56°C, 30 s; 68°C, 30 s) using FastStart Taq (Roche) and a primer pair designed to yield small amplicons (~150–200 bp) around the target site. Amplicons were subjected to electrophoresis on 2.5% agarose gels to discriminate clones with indels, with positive clones having a band or bands visibly shifted in size from the baseline (see Figures S1B and S6A for examples); for *AKT2* E17K candidate clones, the amplicons were digested with RsaI for 1 hr and subjected to electrophoresis, with positive clones displaying cleavage products (Figure S6A). For a subset of the potentially positive clones, PCR amplicons were subcloned using the TOPO TA Cloning Kit (Invitrogen) and subjected to numerous sequence reads for confirmation of the presence of mutant alleles; in a similar fashion, a subset of the

potentially negative clones were confirmed to be wild-type. Clones with confirmed compound heterozygous mutant alleles (or the *AKT2* E17K mutation) or confirmed to be wild-type were retrieved from the frozen stocks and expanded for further experiments. When no compound heterozygous clones were identified, a heterozygous clone with one mutant allele was expanded and subjected to a second round of TALEN targeting.

Differentiation of hPSCs into HLCs, White Adipoctyes, and Motor Neurons

Differentiation was performed following the protocols of Si-Tayeb et al. (2010), Ahfeldt et al. (2012), and Di Giorgio et al. (2008) and Chambers et al. (2009). Details are given in Supplemental Experimental Procedures.

ELISAs, Immunocytochemistry, and Western Blot Analysis

These procedures were performed using standard methods. Details are given in Supplemental Experimental Procedures.

Glucose Production, Glucose Uptake, Triglyceride Content, and Lipolysis Assays

Glucose production and glucose uptake were measured using protocols adapted from Hagiwara et al. (2012) and Ahfeldt et al. (2012), respectively. Details of the various procedures are given in Supplemental Experimental Procedures.

Quantitative RT-PCR

Quantitative RT-PCR (qRT-PCR) was performed using standard methods. Details and oligonucleotide sequences are given in Supplemental Experimental Procedures.

HCV Infection of HuH-7 Cells

Details are given in Supplemental Experimental Procedures.

Exome and Whole-Genome Sequence Analyses

Exome sequencing and whole-genome sequencing were performed as previously described (Gnirke et al., 2009; Stransky et al., 2011). Procedural and analytical details are given in Supplemental Experimental Procedures.

SUPPLEMENTAL INFORMATION

Supplemental Information includes seven figures and Supplemental Experimental Procedures and can be found with this article online at http://dx.doi.org/10.1016/j.stem.2012.11.011.

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