Cystic fibrosis (CF) is the main genetic cause of death among the Caucasian population. The disease is characterized by abnormal fluid and electrolyte mobility across secretory epithelia. The first manifestations occur within hours of birth (meconium ileus), later extending to other organs, generally affecting the respiratory tract. It is caused by mutations in the cystic fibrosis transmembrane conductance regulator (CFTR) gene. CFTR encodes a cyclic adenosine monophosphate (cAMP)-dependent, phosphorylation-regulated chloride channel required for transport of chloride and other ions through cell membranes. There are more than 2,000 mutations described in the CFTR gene, but one of them, phenylalanine residue at amino acid (aa) position 508 (p.F508del), a recessive allele, is responsible for the vast majority of CF cases worldwide. Here, we present the results of the application of genome-editing techniques to the restoration of CFTR activity in p.F508del patient-derived induced pluripotent stem cells (iPSCs). Gene-edited iPSCs were subsequently used to produce intestinal organoids on which the physiological activity of the restored gene was tested in forskolin-induced swelling tests. The seamless restoration of the p.F508del mutation resulted in normal expression of the mature CFTR glycoprotein, full recovery of CFTR activity, and a normal response of the repaired organoids to treatment with two approved CF therapies: VX-770 and VX-809.

INTRODUCTION

Cystic fibrosis (CF) is the most frequent genetic disease among the Caucasian population, affecting ca. 90,000 individuals worldwide. The first manifestations occur in early childhood and affect several organs, resulting in defects in the pancreas, liver, intestine, vas deferens, sweat ducts, and airways. The most severe and life-threatening consequences of CF occur in the lung. Three decades ago, it was shown that mutations in the cystic fibrosis transmembrane conductance regulator (CFTR) gene, inherited in an autosomal-recessive fashion, are the underlying cause of CF disease. More than 2,000 different mutations have been described across the CFTR locus, but up to date, only 346 genetic variants have been identified as pathogenic, disease-causing mutations (https://www.cfr2.org/). CFTR acts as an anion channel at the apical surface of secretory epithelial tissues regulating electrolyte and fluid transport across the plasma membrane. Disruption of CFTR activity leads to acidification of luminal pH and generation of a thick, sticky mucus at the affected epithelia. Chronic pulmonary infections produced by bacterial colonization of the abnormal mucus layer still represent the primary cause of mortality. However, thanks to improved symptomatic treatments and higher success rates in lung transplantations, CF is becoming a multi-systemic disease affecting lung, intestine, pancreas, and liver.

The deletion of the CTT triplet in exon 11 of the CFTR gene is the most prevalent CF-causing mutation, representing 70% of all patients’ alleles. The resulting trafficking-deficient CFTR protein lacks the phenylalanine residue at amino acid (aa) position 508 (p.F508del). Despite tremendous efforts in the last years, there is still no cure for all CF patients, including p.F508del patients. Only two types of small molecule treatments for p.F508del patients are presently available on the market. Unfortunately, the strongest possible intervention, consisting of a combination of both a potentiator to stimulate CFTR channel activity (ivacaftor) and a corrector to promote CFTR trafficking to the apical plasma membrane (tezacaftor or lumacaftor), has only modest effects on patients’ lung function. Most recently, the US Food and Drug Administration (FDA) has approved a promising triple-combination therapy (ivacaftor-tezacaftor-lexacaftor)
Seamless Correction of the CFTR Gene by TALEN-Mediated Homologous Recombination in CF iPSCs

To obtain a seamless correction of the p.F508del mutation in patient-derived CF-iPSCs, we devised a strategy based on Transcription Activator-Like Effector Nuclease (TALEN)-mediated homologous recombination (HR), followed by the total removal of the selection cassette with a piggyBac (PB) transposase system. This approach guarantees the absence of any vector fragment in the patient’s genome after the whole procedure is completed. For this purpose, we first designed a pair of TALENs that could recognize a target site nearby the p.F508del mutation (Figure 1A). The specificity of these TALENs was initially determined in K562 and HeLa cell lines with the Surveyor nuclease assay. A high cleavage efficiency of about 50% confirmed their functionality in both cell types (Figure S1). Then, a donor vector was designed containing a functional allele of the CFTR gene. The genetic defect in p.F508del iPSCs was corrected by introducing a CTT triplet in exon 11 of the CFTR gene at the precise position where its absence causes the mutant phenotype (Figure 1A). The targeting vector contains a transposon-based, double-selection puromycin-(delta)thymidine kinase (purotk) cassette driven by a phosphoglycerate kinase (PGK) promoter and flanked by PB-specific inverted terminal repeat (ITR) sequences. Once the PB transposase recognizes those sites, it efficiently catalyzes the seamless excision of the cassette.

RESULTS

Seamless Correction of the CFTR Gene by TALEN-Mediated Homologous Recombination in CF iPSCs

Targeted repair of the p.F508del allele has been carried out on two iPSC lines, independently derived in our lab from the same CF patient (IMDEA001-1-F). The deletion of the CTT triplet in the targeted iPSC line was previously confirmed by PCR-directed mutagenesis and MboI digestion. CF iPSCs were nucleofected with both TALEN plasmids and the donor vector at a ratio of 1:1:1 and seeded onto antibiotic-resistant feeders. Following 2 weeks of puromycin selection, resistant clones were isolated, and expanded, and analyzed by PCR. To identify positive clones, both 5′ and 3′ ends of the recombination site were screened with combinations of primers that bind to the selection cassette and to genomic sites located outside the recombination arms (Figures 1B and 1C). To eliminate possible recombination events 5′ to the p.F508del site that would give rise to the integration of the targeting construct into the CFTR locus, but without repairing the CTT deletion, the 5′ PCR products were digested with BglII. The newly
The generated BgIII site is located at 28 bp 3' of the p.F508del site, so that successful BgIII digestion would very likely indicate the correct integration of the CTT triplet and therefore, the repair of the gene. 71 clones identified as having undergone homologous recombination were subjected to BgIII analysis and all except one (clone 111; Figure 1B) showed two bands of 1,019 and 306 bp, corresponding to the size of the fragments expected from digesting the repaired allele (Figure 1B). The absence of BgIII digestion in clone 111 indicated that this clone lacked the BgIII site and very likely, the corrected CFTR sequence. This was most probably due to a recombination event that occurred 3' of the BgIII restriction site.

The next step of the procedure consisted of the excision of the recombination cassette. For that purpose, four recombined clones from L1 (L1-27, L1-43, L1-65, L1-104) and five from L2 (L2-18, L2-47, L2-61, L2-64, L2-87) were transfected with the hyperactive PB transposase. Fialuridine (2'-deoxy-2'-fluoro-beta-D-arabinofuranosyl-5-iodouracil or FIAU)-resistant clones were isolated, expanded, and analyzed by PCR with primers located outside the selection cassette. To ensure specific amplification of the edited allele, the 5' primer was designed to bind to the BgIII restriction site. PCR screening demonstrated the removal of the PB-based selection cassette in approximately 53% of the analyzed clones in both previously edited CF iPSC lines (Figure 1C).

After CFTR correction and cassette excision, one subclone from L2-64 (L2-64.12) was randomly selected for further characterization. A 2.2-kb DNA fragment spanning the complete genomic region involved in HR was amplified by PCR using primers P1 and P4. PCR amplicons were cloned into the pGEM-T Easy Vector and out of 98 analyzed clones in CF iPSC line 2 (L2), an overall correction efficiency of approximately 10% (Figure 1D).

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sequenced. The detection of both wild-type and CFTR-repaired alleles confirmed CFTR repair in heterozygosity at the endogenous CFTR locus. Figure 1E shows the edited sequence around the p.F508del mutation site. The addition of the CTT triplet and the presence of the eight silent mutations originally incorporated into the targeting vector are highlighted. Alignment of the complete sequenced region with a CFTR p.F508del template is shown in Figure S2.

Characterization of CFTR-Corrected iPSCs

Once the presence of the corrected allele was verified, the repaired CF iPSC line L2-64.12 was further characterized regarding genome integrity, stemness, and pluripotency. G-band analysis indicated a normal diploid karyotype without chromosomal aberrations (Figure 2A). To detect potential genomic imbalances at a higher resolution, array comparative genomic hybridization (aCGH) was performed by comparing on the one side the genomes of repaired and nonrepaired CF iPSCs and on the other side the genomes of repaired CF iPSCs and parental primary fibroblasts (Figure S3). All in all, only five copy number variations (CNVs) could be identified across the genome. Although two CNVs emerged during reprogramming and iPSC generation, three CNVs were generated in the course of HR-directed repair. The observed modifications could have their origin in the direct activity of the genetic material added to the cells or just be a consequence of having spent a lengthy time in culture. To test possible off-target effects of the TALENs in the L2-64.12 line, we decided to study the stability of the loci predicted by the PROGNOS software as the most likely affected by nonspecific TALEN activity. CFTR, plus the top 15 sites ranked as possible off-target regions were selected and analyzed by the Surveyor assay (Figure 2B).

Stemness of the repaired iPSC line was then tested by immunofluorescent analysis with a panel of diagnostic antibodies: aSOX2, aOCT4, aNANOG, and aTRA-1-60 (Figure 2C). In all cases, the corresponding antigens were strongly expressed, indicating that the L2-64.12 line had maintained its stem cell properties during the repair process. Human pluripotent stem cells have the capacity to differentiate into three embryonic germ layers: ectoderm, mesoderm, and endoderm. To test whether our newly corrected iPSC clone L2-64.12 retained this ability, embryoid bodies (EBs) were produced, plated on gelatin-coated dishes for 20 days, and analyzed by immunostaining. The presence of ectodermal lineages was

| Source | Chromosome | Genomic Location | Chromosomal Region | Targeted Allele |
|--------|------------|------------------|-------------------|-----------------|
| 1      | 7          | intronic          | YTRF              |                 |
| 2      | 4          | intronic          | YTRF              |                 |
| 3      | 8          | intronic          | PANGA_60A4        |                 |
| 4      | 13         | intronic          | ZMT14             |                 |
| 5      | 17         | intronic          | KBR11             |                 |
| 6      | 13         | intronic          | LEMO120           |                 |
| 7      | 9          | intronic          | NPEC2             |                 |
| 8      | 2          | intronic          | KCM64             |                 |
| 9      | 9          | intronic          | MARM62            |                 |
| 10     | 9          | intronic          | CF70              |                 |
| 11     | 6          | intronic          | MAX460            |                 |
| 12     | 6          | intronic          | CM64              |                 |
| 13     | 3          | intronic          | TAMAS07           |                 |
| 14     | 3          | intronic          | APT12A            |                 |
| 15     | 8          | intronic          | WRY               |                 |
| 16     | 4          | intronic          | TRN2             |                 |
conferred by the expression of NESTIN, Paired box protein 6 (PAX6), and Neuron-specific class III beta-tubulin (TUJ1), whereas GATA Binding Protein 4 (GATA4), α-ACTININ, and smooth muscle actin (SMA) illustrated the formation of mesoderm and transcription factor SOX-17 (SOX17), Hepatocyte nuclear factor 3-beta (FOXA2), and α-fetoprotein (AFP) of endoderm (Figure 2D).

Directed Differentiation of the CFTR Mutant and Corrected iPSCs toward CFTR-Expressing Intestinal Organoids

Native CFTR is expressed in multiple organs, such as lungs, pancreas, liver, and gastrointestinal tract. iPSC differentiation toward lung epithelial cells is very complex and shows high variability in differentiation efficiencies. Recently, it has been demonstrated that CFTR-expressing HIOs, derived from rectal biopsies, could be used to monitor CFTR function in a robust assay with a simple readout. Based on this observation, we have developed a novel, stepwise, three-dimensional (3D) differentiation protocol to generate HIOs from iPSC (manuscript in preparation). Intestinal organoids, produced according to the new protocol, were used to study the restoration of CFTR function after HR-directed CFTR repair. During development, the first precursor of intestinal epithelium to appear is the multipotent definitive endoderm (DE). Thus, CF-corrected iPSCs were initially differentiated in vitro into DE by treating them with the signaling molecules ACTIVIN-A and Wingless-Type MMTV integration site family, member 3A (WNT3A). In order to ensure high differentiation efficiencies, we incorporated a DMSO (dimethylsulfoxide) pretreatment step in our differentiation protocol prior to DE generation. DE cells at day 3 of differentiation showed a typical expression pattern of endoderm transcription factors, such as FOXA2, SOX17, and GATA4. The activation of endodermal markers was accompanied by the downregulation of the pluripotent stem cell markers OCT4 and NANOG (data not shown). The efficiency of DE derivation from CF-corrected iPSCs was about 80%. The next differentiation step consisted of the production of hindgut endoderm (HGE), the direct precursor population of the more posterior digestive tract. Exposure to high concentration of WNT3A agonists for 5 days resulted in the formation of HGE lineages with the propensity to differentiate into intestinal epithelial cells. A large percentage of HGE cells expressed the hindgut marker Caudal Type Homeobox 2 (CDX2), whereas the generation of the anterior foregut lineage was completely blocked, as demonstrated by the lack of FOXA2/SOX2 coexpression. At that point, floating 3D HGE spheroids were embedded into Matrigel drops and cultured in the presence of intestinal growth factors. The sequential maturation of the hindgut spheroids into mature intestinal organoids could be observed during the following weeks. The resulting HIOs contained crypt-like structures and an internal lumen lined by columnar epithelium, closely resembling the morphology of native intestinal tissue. Although HIOs expressed intestinal markers, such as Kruppel Like Factor 5 (KLF5), CDX2, and VILLIN, all attempts made at this point to detect restoration of CFTR function by the FIS assay in CFTR-corrected, iPSC-derived HIOs failed (data not shown). This suggested very low to undetectable levels of CFTR expression that could be explained either by the failure in restoring the CFTR gene or by an incomplete maturation of the organoids.

The functionality of the FIS assay has been demonstrated in primary HIO obtained by crypt isolation during rectal biopsies, but not in iPSC-derived organoids. We, therefore, introduced several
Figure 3C). In addition, the growth rate was very similar to that of the mesenchymal layer that is normally found in iPSC-derived HIO cultures. This new protocol was able to grow without support of the mesenchymal layer. HIOs derived from CFTR-corrected and mutant iPSCs following DE differentiation showed an intestinal phenotype of biopsy-derived HIOs (Figure 3C; manuscript in preparation). The intestinal phenotype of HIOs was confirmed by positive E-cadherin (red) with Villin (green) but not with the basolateral marker E-cadherin (green/purple) at the apical surface in CFTR-homozygous HIOs (top row). Colocalization of CFTR (red) with Villin (green) but not with the basolateral marker E-cadherin (green/purple) at the apical surface in CFTR-corrected HIOs (middle and bottom rows). Nuclei were counterstained with DAPI (blue). Scale bars, 50 μm (top and middle rows); scale bar, 20 μm (bottom row).

To determine whether functional rescue of p.F508del mutant CFTR is entirely dependent on CFTR function, we first tested if CFTR activation by forskolin-induced elevation of cyclic AMP (cAMP) levels was able to induce fluid influx into the organoid’s lumen, thus augmenting their surface area.

To allow quantification of organoid expansion in the course of FIS assays, corrected and mutant HIOs were previously labeled with calcein green, a cell-permeant dye that exclusively marks living cells. Upon 180 min of forskolin treatment, corrected and mutant HIOs increased their surface area to 177% (±9 SD) and 103% (±1 SD), respectively (Figures 5A–5C), where 100% represents the original surface prior to treatment. These expansion rates are in good concordance with previous findings, where primary wild-type and p.F508del mutant HIOs were used in FIS assays over a period of 60 min.13. FIS response in human organoids was reported to be entirely dependent on CFTR function13, so to demonstrate CFTR dependency in our iPSC-derived HIO culture system, iPSC-derived organoids were pretreated for 16 h, with the specific CFTR inhibitor CFTRinh-17231, before starting the FIS assay. In fact, forskolin-induced expansion in CFTR-corrected HIOs was completely blocked in the presence of CFTRinh-172 (102% ± 2% SD; Figures 5A and 5C), whereas as expected, the negligible FIS response in CFTR mutant HIOs did not significantly change (96% ± 1% SD; Figures 5A and 5B). This inhibition was reversible after washing out CFTRinh-172 (data not shown).

To examine whether functional rescue of p.F508del mutant CFTR is feasible with a combined treatment of chemical CFTR modulators, we treated iPSC-derived HIOs simultaneously with the CFTR potentiator (VX-770) and the corrector (VX-809; preincubated for 16 h). The double treatment markedly increased the FIS response in p.F508del mutant HIOs (swelling rate: 133% ± 7% SD; Figures 5A and 5B), albeit to a lesser extent than what was observed in CFTR-corrected HIOs without VX-770/VX-809 treatment, suggesting that HR-directed CFTR repair is considerably more efficient in restoring CFTR function than the clinically approved double treatment VX-770/VX-809 (Orkambi). CFTR-corrected HIOs showed an even higher increase in surface area (206% ± 14% SD; Figures 5A and 5C) when treated with VX-770/VX-809, thus validating the impact of CFTR modulators on CFTR function in CF-corrected HIOs. Finally, organoid swelling was completely abolished in VX-770/VX-809-treated, CFTR-corrected HIOs, whereas only immature, core-glycosylated CFTR (B band) was expressed in mutant HIOs (Figure 4A). The CFTR expression pattern was similar to healthy and p.F508del primary HIOs, respectively. Moreover, corrected CFTR protein was found to be expressed on the apical membrane that lines the inner lumen of HIOs, as evidenced by colocalization of CFTR and the apical marker VILIN (Figure 4B).

Figure 4. Restoration of CFTR Protein Expression in Human Intestinal Organoids Derived from CFTR-Corrected iPSCs
(A) Western blot analysis. C band, mature, complex glycosylated CFTR; B band, immature CFTR. Calu-3 cells and primary rectal wild-type organoids (HUB WT) serve as positive control for the detection of the mature CFTR C band, whereas primary rectal CFTR p.F508del homozygous organoids (HUB 508/508) only express the immature B band of the CFTR protein. HIO L2 and HIO L2-64.12 intestinal organoids derived from mutant and corrected CF iPSCs, respectively. Note the inverse ratio between the immature and glycosylated forms of CFTR in the two types of organoids. α-tubulin is the protein-loading internal control. (B) Immunofluorescence analysis of CFTR expression. No CFTR protein is detected in iPSC-derived CFTR p.F508del homozygous HIOs (top row). Colocalization of CFTR (red) with Villin (green) but not with the basolateral marker E-cadherin (green/purple) at the apical surface in CFTR-corrected HIOs (middle and bottom rows). Nuclei were counterstained with DAPI (blue). Scale bars, 50 μm (top and middle rows); scale bar, 20 μm (bottom row).

Modifications to the maturation step of the original differentiation protocol in order to generate iPSC-derived HIOs that more closely resembled biopsy HIOs (Figure 3C; manuscript in preparation). HIOs derived from CFTR-corrected and mutant iPSCs following this new protocol were able to grow without support of the mesenchymal layer that is normally found in iPSC-derived HIO cultures (Figure 3C). In addition, the growth rate was very similar to that observed in biopsy-derived HIOs.29. The intestinal phenotype of iPSC-derived HIOs was confirmed by immunostaining, showing the expression of the intestinal transcription factors SOX9 and CDX2, as well as the detection of the enterocyte marker VILIN. The epithelial origin of HIOs was confirmed by positive E-cadherin (E-CAD) staining (Figure 3D). Successful differentiation of the original CF mutant iPSC line, L2, into intestinal organoids was equally verified by immunofluorescence studies at DE, HGE, and HIO stages (Figure S5). Most importantly, CFTR protein expression, a prerequisite for the recovery of CFTR activity, was detected mainly as the complex-glycosylated form (C band) in CFTR-corrected HIOs, whereas only immature, core-glycosylated CFTR (B band) was expressed in mutant HIOs (Figure 4A). The CFTR expression pattern was similar to healthy and p.F508del primary HIOs, respectively. Moreover, corrected CFTR protein was found to be expressed on the apical membrane that lines the inner lumen of HIOs, as evidenced by colocalization of CFTR and the apical marker VILIN (Figure 4B).

Functional Restoration of CFTR Channel Activity in iPSC-Derived, CFTR-Corrected Intestinal Organoids
To determine whether heterozygous CFTR repair of the p.F508del allele could support functional recovery of CFTR channel activity, we first tested if CFTR activation by forskolin-induced elevation of cyclic AMP (cAMP) levels was able to induce fluid influx into the organoid’s lumen, thus augmenting their surface area.
(104% ± 6% SD) and mutant (99% ± 2% SD) HIOs when CFTR activity was blocked by CFTRinh-172 exposure (Figures 5A–5C).

Taken together, these results strongly indicate that heterozygous CFTR correction by TALEN-mediated footprint-free gene targeting resulted in full restoration of CFTR function in iPSC-derived HIOs, a result compatible with similar observations made on iPSC-derived, 2D-cultured epithelial cells.32

DISCUSSION

Here, we describe the correction, by gene targeting in patient-derived iPSCs, of the CFTR p.F508del mutation. Gene-editing protocols are often associated with undesired and potentially harmful modifications of the genome; to minimize the possibility of this type of events, a strategy based on an antibiotic selection step, followed by a triple serial DNA screening, was devised. In addition, a new procedure to obtain intestinal organoids from iPSC-edited cells was validated, and the complete functional restoration of CFTR activity in CF-corrected, iPSC-derived intestinal organoids demonstrated using the CFTR-dependent FIS assay.

After transfection, antibiotic selection eliminated all cellular clones in which the antibiotic-resistant cassette had not been incorporated into the genome. The first PCR screening step was designed to identify cellular clones with HR-driven integration of the selection cassette into the targeted locus. Validation at both 5′ and 3′ recombination sites was necessary to avoid false positives due to the fragmentary integration of the recombination cassette. The second screening exploited a newly created, silent BglII restriction site. Due to its close proximity to the p.F508del mutation, successful digestion of the amplicon comprising the site was indicative of a very likely, correct cointegration of the restoring CTT triplet. Finally, the third screening step was carried out to confirm piggyBac-catalyzed excision of the selection cassette and therefore, a seamless repair of the CFTR gene. Due to the silent mutations introduced into the recombination cassette, allele-specific PCRs could be performed to identify recombinant CF-repaired clones in which the selection cassette had been properly cut out by PB transposase.

Our TALEN-mediated HR approach resulted in the correction of the p.F508del mutation in two iPSC lines derived from one CF patient with efficiencies between 9.3% and 10.2%. These efficiencies are superior or similar to correction rates reported in other nuclease-assisted, genome-editing studies, where the p.F508del mutation has been repaired.33–36 It should be noted that the efficiency of HR-driven correction depends on many factors, such as cell type, cell line, cutting efficiency of designer nuclease, nuclease retargeting, proximity of the double-strand breaks to the repair site, sequence identity between...
donor template and target region, as well as the type of homology-directed repair (HDR) template\textsuperscript{37}. For instance, Lee et al.\textsuperscript{34} reported an overall repair efficiency of <1% in a CF lung epithelial cell line. This low level of repair was most probably due to the long distance (203 bp) between the p.F508del mutation and the Zinc Finger Nuclease (ZFN) target site, as well as to sequence variation between donor plasmid and target region. Another study used short DNA fragments and TALENs to correct the CTT deletion in CF iPSCs, with correction rates as low as 0.1%, although the efficiency increased up to 10% after six cycles of enrichment of the recombinant cells\textsuperscript{35}. More recently, two other studies claimed to have obtained higher HR-driven repair efficiencies at the p.F508del locus by employing CRISPR/Cas9-mediated, gene-editing approaches in combination with a donor vector\textsuperscript{33} or single-stranded oligodeoxynucleotides (ssODNs)\textsuperscript{35} as repair templates. In the first study, editing efficiency of 16.7% was observed in a CF iPSC line, whereas the correction rate in the second study ranged from 2.5% to 22%, depending on the cell type used. Although higher HR rates can be achieved by electroporating CRISPR/Cas9 as a recombinant ribonucleoprotein\textsuperscript{36}, we have opted to utilize the TALEN technology to enhance CF gene correction in iPSCs, since the CRISPR/Cas9 system, in spite of recent efforts to minimize off-targeting, seems still to be associated with increased off-target effects\textsuperscript{38,39}. On the contrary, TALENs generally show a higher level of specificity, which relies on their ability to recognize longer target sequences (30–36 bp)\textsuperscript{40}, the limitations imposed by the required distance between target sequences\textsuperscript{41} and on the fact that they only act if both members of the pair bind to the target sequence on opposite strands.

Although small DNA fragments, including ssODNs, are nowadays widely used for HR-based gene correction of small fragments, such as single-base pair mutations, like the CTT deletion in the CFTR gene,\textsuperscript{35,36,42} this elegant and drug selection-free approach might require laborious screening analysis in order to identify targeted clones\textsuperscript{43}. Additional, more serious disadvantages of ssODN-based gene editing are the generation of indel mutations at HR sites by the microhomology-mediated end-joining mechanism\textsuperscript{44} and the random insertion of ssODN throughout the genome with a frequency of up to 18%\textsuperscript{45}. A recent study by Vaidyanathan et al.\textsuperscript{46} has reported recombination frequencies of around 40% at the CFTR locus in upper-airway basal stem cells from CF patients. They achieved this mark by adenovirus-associated (AAV) transduction of CRISPR/Cas9 and ssODNs. However, the results should be considered in light of new evidence suggesting the possibility of undesired rearrangements produced by ssODNs, both at the target site and at distant locations\textsuperscript{37,46}. In addition, therapies based on viral vectors, including AAVs, might generate immune responses against the virus and are not applicable to individuals with pre-existing neutralizing antibodies\textsuperscript{47,48}. For all of these reasons and despite the attractiveness of the ssODN-mediated HR system, we opted instead for an ex vivo PB transposon-based gene-targeting strategy. Similar to previous reports\textsuperscript{33,30}, we obtained integration-free-corrected iPSC clones with an excision efficiency of the drug-resistant cassette of 53% on already-selected edited iPSC lines. In several CFTR gene-targeting studies, seamless correction of the p.F508del allele was not employed, thereby leaving behind donor DNA remnants, such as the complete selection cassette\textsuperscript{27,24} or a residual loxP site after Causes Recombination enzyme (CRE)-mediated excision of the donor vector\textsuperscript{37}, in the genome of repaired CF iPSCs. iPSCs harboring those unwanted genetic modifications have a limited application, not only in clinical settings but also for disease modeling in basic research and drug-screening programs.

Functional restoration of CFTR channel activity after gene editing has been previously demonstrated in iPSC-derived epithelial cells by using 2D differentiation monolayer systems\textsuperscript{32,33,36,42}. However, due to variable differentiation efficiencies, iPSC differentiation often resulted in the derivation of heterogeneous populations in terms of CFTR expression, thereby impeding unambiguous readouts during functional assays. We, therefore, developed a novel iPSC-derived 3D organoid model that allows the precise assessment of CFTR activity in CFTR mutant and repaired HIOs by the FIS assay. Importantly, the readout of this technique has been shown to be completely dependent on CFTR in primary HIO\textsuperscript{12}, thus serving as an ideal method to measure the recovery of CFTR function following CFTR gene correction. Before starting the FIS assay, CFTR expression and its localization at the plasma membrane were confirmed in CFTR-corrected HIOs by immunostaining. In addition, detection of mature, complex, glycosylated CFTR protein by western blot strongly indicated the restoration of correct CFTR processing upon CFTR gene repair and a very likely functional recovery of the gene, since only this processed form of CFTR is able to reach the plasma membrane and exert its function as an anion channel. On the contrary, p.F508del mutant HIOs only expressed the immature core-glycosylated form of CFTR that is rapidly degraded in the endoplasmic reticulum (ER) before reaching the plasma membrane. These results are in concordance with recent findings showing the localization of mature CFTR at the plasma membrane in primary wild-type but not in p.F508del homozygous HIOs\textsuperscript{15}.

Finally, functional testing of CFTR protein activity in FIS assays demonstrated that p.F508del correction in one of the two mutant alleles is enough for the complete rescue of CFTR function, as indicated by similar levels of organoid swelling to those observed in wild-type\textsuperscript{13} or CFTR-corrected, primary HIOs\textsuperscript{17}. Interestingly, heterozygous gene correction seems to be far more efficient in terms of restoring CFTR activity than the combinatorial treatment of homozygous p.F508del samples with the CFTR potentiator (VX-770) and corrector (VX-809) together, which is consistent with a previous report using primary HIOs\textsuperscript{17}. It has been described that CF patients harboring p.F508del in homozygosity differ significantly in their response to Orkambi (VX-770/VX-809) treatment, suggesting that the CF patient in our study could belong to the low responder group\textsuperscript{6,71}. Patient-specific differences were also observed in primary HIOs with identical CF-causing mutations, further indicating that individual genetic background, in addition to CFTR status, might affect the degree of FIS response in vitro and successful therapy in vivo\textsuperscript{14,15}. Given the influence of genetic modifiers on CFTR functional tests and more importantly, on the course of the disease and the response of patients...
to treatment\(^4\), CFTR-repaired HIOs and their original mutant counterparts represent an optional isogenic platform to use in disease modeling and drug screening. Although previous gene-targeting studies have validated the restoration of CFTR activity in iPSC-derived lung epithelial cells using a variety of techniques, such as using chamber\(^32,36\), whole-cell patch-clamp\(^33\) and iodide efflux assays\(^32\), the combination of the HIO and FIS assay is presumably the easiest and most robust way to validate CFTR activity in a complex, polytropic tissue model. In addition, the possibility of expanding iPSC-derived HIOs to an unlimited number, thanks to the maintenance of an intestinal stem-cell subpopulation, makes it very attractive for high-throughput screening in 384-well plates when combined with automated live microscopy. Indeed, pharmaceutical companies have already started to integrate intestinal organoids in their drug-discovery pipeline\(^34\). It is important to keep in mind, however, that until exhaustive comparisons are performed, it cannot formally be ruled out that drug responses of certain CFTR variants can vary in a tissue-specific manner due to the recruitment of different cis-regulatory elements nearby or within the CFTR locus. In this regard, it has been demonstrated that CFTR expression is differentially regulated in lung and intestinal cells through cell type-specific enhancer elements that are located in intergenic and intronic regions\(^18\).

In summary, this report demonstrates, to our knowledge, for the first time, the integration of an efficient method for seamless CFTR correction in CF patient-derived iPSCs with a functional intestinal organoid swelling test to determine the biological activity of the resulting protein. The linkage of the two methods was made possible by a novel and efficient protocol for the production of intestinal organoids from edited cells and culture procedures.

**Cells and Culture Procedures**

For gene-targeting experiments, we used an iPSC line (IMEDEA001-F) recently generated in our lab from a CF patient\(^7\). Human iPSCs were cultured in KnockOut DMEM (Invitrogen, USA), supplemented with 20% KnockOut Serum Replacement (KOSR; Invitrogen, USA), 2 mM GlutaMAX, 0.1 mM β-mercaptoethanol, 1% nonessential amino acid (NEAA; Invitrogen, USA), and 10 ng/mL basic fibroblast growth factor (bFGF; PeproTech, UK). iPSC passage was carried out using recombinant trypsin (TrypLE Select; Invitrogen, USA) and the final addition of 10 μM rho-associated protein kinase (ROCK) inhibitor Y-27632 (Sigma, Spain). Mature, intestinal iPSC-derived organoids and rectal organoids were maintained in intestinal growth media and passed 1:3 to 1:6 every 3–8 days, as described previously\(^18,25\). HUB wild-type and HUB p.F508del homozygous rectal organoid lines were purchased from Hubrecht Organoid Technology (https://huborganoids.eu/).

**Generation of CFTR-Specific TALENs and Detection of Nuclease Activity**

TALEN expression vectors designed to target and cleave 55–57 bp 3’ of the p.F508del mutation, were synthesized by Transposagen (USA). The length of the recognition sequence of each TALEN is 17 bp (left binding site: 5’-ACAGAAGGTCATCAA-3’; right binding site: 5’-AGGTAAGAAACTATGTG-3’) and is separated by a 16-bp spacer. The expression of the TALEN genes is under control of the cytomegalovirus (CMV) promoter.

The cleavage activity of the newly generated CFTR TALENs was analyzed in two highly transfectable cell lines, K562 and HeLa. 1 × 10⁶ K562 and HeLa cells were transiently transfected with 2 μg of each TALEN vector for 48 h. Genomic DNA from the TALEN-transfected cell populations was isolated and the target site amplified by PCR using the ZFN-forward (Fw)/P6 primers (see Table S1). To assess the frequency of indels generated after cleavage by TALENs, the Surveyor nuclease assay (Integrated DNA Technologies [IDT], Belgium) was used. For this purpose, PCR amplicons were denatured and slowly renatured. Resulting heteroduplexes were cleaved by the Surveyor nuclease, whereas homoduplexes were left intact. The heterogeneous DNA population was run on a microchip device (Bioanalyzer 2100; Agilent, USA), and the efficiency of TALEN cleavage (percentage of indels) was quantified by densitometry as a ratio between cleaved and noncleaved DNA bands.

**Targeting Vector Design**

Two vectors were produced. The first one (Seq_1) was designed to be used, prior to gene-targeting experiments, as a template to optimize the PCR reactions used in the screening for recombinant clones. For this purpose, specific genomic regions, immediately adjacent but external to the 5’ and 3’ recombination arms, were added to the vector to allow primer binding. The length of the 5’ and 3’ homology arms was 916 and 920 bp, respectively. In order to prevent retargeting of the recombinated allele by TALENs, five silent point mutations were introduced into the 5’ recombination arm. In addition, a new silent BglII restriction site was also created to facilitate the detection of recombinant clones by PCR, followed by digestion of the amplification product with the enzyme (Figure 1A). Primers specific for the recombinant allele were also designed (Table S1). The final targeting vector, pMC3.1, was obtained by modifying Seq_1 in three cloning steps. The elimination of the external primer-binding regions attached to the homology arms was performed by EcoRI and MluI digestions. Finally, the double-selection cassette PB:PGKpuroΔtk from pMCS-AAT-PB:PGKpuroΔtk\(^35\) was cloned via NheI/NotI digestion/ligation in between the homology arms. The pMCS-AAT-PB:PGKpuroΔtk plasmid was provided by the Sanger Institute under a Material Transfer Agreement (MTA).

**Electroporation of iPSCs**

Nucleofection was performed with the P3 Primary Cell 4D-Nucleofector Kit (Lonza, Walkersville, MD, USA), according to the manufacturer’s instructions. To avoid feeder cell contamination, iPSCs were expanded in Matrigel-coated p100 culture plates prior to
nucleofection. Adherent iPSCs were detached by using TrypLE (Gibco, USA), centrifuged at 200 × g, and resuspended in nucleofection buffer 3. For homologous recombination experiments, 2 μg of TALEN and donor plasmids were transfected per 1 × 10^6 cells in a 100-μL cuvette using the CB-150 program. In piggyBac excision experiments, 5 μg of piggyBac transposase (pCMV-HAhyPBase) per million cells was employed in nucleofection assays. Nucleofected iPSCs were seeded in the presence of 10 μM ROCK inhibitor onto irradiated HAF-1/W3R puromycin-resistant feeder cells, kindly provided by Dr. Chen from Johns Hopkins University School of Medicine (Baltimore, MD, USA).56

Isolation of Recombined Clones and PCR Screening

After selection with 1 μg/mL puromycin (Sigma, Spain) or 250 nM FIAU (Sigma, Spain) for 12–18 days, individual resistant colonies were manually picked and transferred to 48-well plates previously seeded with irradiated feeder cells. After 7–10 days, iPSC clones were trypsinized (TrypLE) and duplicated in either Matrigel-coated or feeder cell-containing 48-well plates. Genomic DNA was extracted from clones grown in feeder-free, Matrigel-coated plates using a standard protocol, whereas clones grown on irradiated feeders were maintained and later expanded in the case of successful recombination. Shortly, genomic DNA from 48-well plates was isolated by proteinase K digestion overnight (O.N.) at 55°C, precipitation of DNA with isopropanol, washing with 70% ethanol, and resuspension in Tris-EDTA buffer. The first round of the screening consisted of the identification of recombinant clones. For this purpose, both 5’ and 3’ ends of the recombination site were analyzed by PCR. One primer of each primer pair was located in the genomic region outside the corresponding recombination arm; the other one was situated within the selection cassette (P1/P2 and P3/P4 pairs; Table S1). To confirm positive recombinant clones, the amplicons generated by 5’ PCR were subsequently digested with BgIII. The absence of random integration of the recombination cassette into the genome was verified by PCR using primers specific for the selection cassette (Reint-Fw/Reint-Rv; Table S1).

Excision of Recombination Cassette by piggyBac Transposase

Following the identification of corrected clones, the selection cassette was removed by nucleofection of 5 μg of pCMV-HAhyPBase per 1 × 10^6 iPSCs. Selection with FIAU (250 nM) was started 24 h after nucleofection and resistant clones picked, expanded, and duplicated, as described above. To identify clones with successful excision of the selection cassette, the correct size of the CFTR target site was confirmed by PCR using primers specific for the edited allele (P5/P6; Table S1).

Sequencing

Genomic DNA from donor cells was isolated, as described above. For sequencing, a 2,250-bp long fragment comprising the whole CFTR-targeted area, including the p.F508del mutation site, was amplified by PCR with the P1 and P4 primers (Table S1). PCR products were cleaned using the Zymoclean Gel DNA Recovery Kit (Zymo Research, CA, USA) and subcloned into the pGEM-T Easy Vector, according to the manufacturer’s recommendations (Promega, USA). Sequencing of the CFTR alleles was performed by Secugen (Madrid, Spain) using the same amplification primers.

Prediction and Analysis of Off-Target Sites

Prediction of potential off-target sites after p.F508del correction was done with PROGNOS software from Dr. Gang Bao’s laboratory.7 Parameters, such as targeted region and repeat variable domains (RVDs) for both TALENs as well as variable length of spacer (10–30 bp), were introduced to predict off-target sequences throughout the genome. The 16 most likely candidates were PCR amplified in the CFTR-correlated clone using primers designed by PROGNOS (Table S1). Subsequently, the amplicons were analyzed by the Surveyor nuclease assay (IDT, Belgium), as previously described.

Generation of Embryoid Bodies

To produce EBs, iPSC colonies were mechanically collected and transferred into a 15-mL conical tube. Once the colonies settled at the bottom of the tube, media were discarded, and iPSCs were cultured in bacteriological Petri dishes with DMEM plus 10% fetal bovine serum (FBS), 1% nonessential amino acids, and 2 mM GlutaMAX for 10 days. Cavitation was then assessed and EBs mechanically disrupted and plated onto gelatin-coated, 24-well plates. Media were changed twice weekly, and after 10–20 days of spontaneous differentiation, cells were used for immunofluorescence studies.

Differentiation toward HIOs

iPSCs were harvested by trypsinization and replated for 1 h on gelatin-coated dishes to induce the depletion of human fibroblast feeders. Then, the nonadhered iPSC suspension was centrifuged; resuspended in iPSC growth media, supplemented with 10% FBS (Sigma, Spain), and plated on tissue-culture plates (100,000 cells/cm²), previously coated with growth factor-reduced Matrigel (BD Biosciences, Spain). To improve differentiation efficiency, cells were treated on the following day with 1% DMSO (Sigma, Spain) for 24 h.25 Subsequently, differentiation into definitive endoderm was performed, as previously described, with slight modifications25. Briefly, cells were washed three times with PBS (Biowest, France) and cultured in RPMI 1640, supplemented with GlutaMAX (2 mM; Life Technologies, USA), 1% NEAA (Life Technologies, USA), 2 mM Glutamine (Life Technologies, USA), 1% penicillin/streptomycin (Life Technologies, USA), 0.2% FBS (Biowest, France), 100 ng/ml human activin A (R&D Systems, UK), and 25 ng/mL WNT3A (R&D Systems, UK). Following 24 h of endoderm differentiation, WNT3A was withdrawn, and cells were differentiated only in the presence of activin A. At day 2 of endoderm induction, FBS concentration was shifted to 2%.

For hindgut specification, early definitive endodermal cells from day 3 were exposed to high doses of WNT3A. Briefly, cells were washed twice with basal differentiation media (Biowest, France) and cultured in RPMI 1640, supplemented with 1xB27, GlutaMAX (2 mM; Life Technologies, USA), 1% NEAA (Life Technologies, USA), 2 mM Glutamine/streptomycin (Life Technologies, USA), 2% FBS (Biowest, France), and 6 μM CHIR99021 (STEMCELL Technologies, France) for 5 days. Floating 3D hindgut spheroids were embedded in the
Matrigel matrix (Corning, NY, USA) and cultured in the presence of R-spondin1, Noggin, epidermal growth factor (EGF), and WNT3A (PeproTech, UK) to allow further intestinal maturation. Finally, a novel differentiation protocol was employed to generate intestinal organoids that were able to grow without a mesenchymal niche (manuscript in preparation), thus facilitating their use in FIS assays.

**FIS Assay**

FIS assays were performed as recently described but with minor modifications. Briefly, human iPSC-derived intestinal organoids from a 5- to 7-day-old culture were harvested, mechanically broken by pipetting, resuspended in Matrigel, seeded as 4 µL drops in a 96-well culture plate (Nunc), and cultured for 24 h in intestinal growth media. Each drop contained approximately 40–80 organoids. 50 µM CFTRinh-172 (Sigma, Spain) and 3 µM VX-809 (Selleck Chemicals, USA) were added to media after seeding for CFTR inhibition and correction, respectively. After 16 h pretreatment and following preincubation for 30 min with media after seeding for CFTR inhibition and correction, respectively.

**Immunofluorescence**

Undifferentiated iPSCs, differentiated cells, and iPSC-derived organoids were washed with PBS, fixed for 20 min with 4% paraformaldehyde, and washed again with PBS. Immunocytochemistry was performed for Nanog (1:150, rabbit immunoglobulin [Ig]G polyclonal; Abcam, Spain), Oct4 (1:100, rabbit IgG polyclonal; Santa Cruz Biotechnology, USA), Tra-1-60 (1:100, mouse IgM; Millipore, USA), AFP (1:100, rabbit IgG; Dako, Denmark), Nestin (1:500, rabbit IgG; Sigma, Spain), TuJ1 (1:500, rabbit IgG; Covance, UK), Sox17 (1:100, goat IgG; R&D Systems, UK), α-actinin (1:200, mouse IgM; Sigma, Spain), SMA (1:200, mouse; Sigma), CFTR (clone MM13-4 or M3A7, 1:100; Merck Millipore, Spain), E-CAD (1:200, rabbit; Cell Signaling Technology), SOX9 (1:100, goat; R&D Systems, UK), Villin (1:100, goat; Santa Cruz Biotechnology, USA), FOXA2 (1:100, goat IgG; Santa Cruz Biotechnology, USA), CDX2 (1:100, mouse IgG; DCS ImmunoLine, Germany), and GATA4 (1:100, mouse IgG; Santa Cruz Biotechnology, USA). After permeabilization in 0.2% Triton X-100/100 mM glycine/PBS buffer for 30 min at room temperature (RT), cells were blocked with 5% BSA in PBS for 1 h at RT. Cells were incubated with primary antibodies overnight at 4°C in 2% BSA in PBS, followed by three washing steps with PBS. Alexa Fluor 555 (1:500, donkey anti-mouse IgG or donkey anti-rabbit IgG; Invitrogen, USA), Alexa Fluor 488 (1:500, donkey anti-rabbit IgG, or donkey anti-rat IgG; Invitrogen, USA) were used as secondary antibodies, incubated 1 h at RT with 2% BSA in PBS. After washing with PBS, cells were stained with 4’,6-diamidino-2-phenylindole (DAPI; 5 min, 1 µg/mL), washed three times, and mounted with Dako fluorescent-mounting medium.

**Western Blot Analysis**

Cells were lysed in radioimmunoprecipitation assay (RIPA) buffer (150 mM NaCl, 1% [v/v] Nonidet P-40 [NP-40], 1% [w/v] SDS, and 50 mM Tris HCl, pH 8.0) in the presence of protease inhibitor cocktail (P8340; Sigma, Spain), incubated for 30 min at 37°C, and passed through a 22G needle to reduce viscosity. Protein samples were quantified by a modified Lowry method, and 70 µg of total protein diluted 1:1 in sample buffer (100 mM Tris HCl, 10% [v/v] β-mercaptoethanol, 4% [w/v] SDS, 0.02% [w/v] bromophenol blue, and 20% [v/v] glycerol) was separated on an 8% SDS-PAGE. Proteins were transferred to a polyvinylidene fluoride (PVDF) membrane, following a wet transfer standard protocol. Membrane was blocked with 5% (w/v) skim milk in Tris-buffered saline (TBS)-TWEEN (TBST) for 1 h at RT, washed three times with TBST, and incubated overnight at 4°C with a cocktail of three primary CFTR antibodies: anti-CFTR monoclonal antibody (mAb) 450 (R-domain 696–712), anti-CFTR mAb 570 (R-domain 731–742 aa), and anti-CFTR mAb 596 (nucleotide-binding domain 2 [NBD2] 1204–1211 aa) (Cystic Fibrosis Foundation, https://www.cff.org), diluted 1:500 in TBST 2% (w/v) skim milk. Membrane was washed and incubated 1 h at RT in the dark with the secondary antibody (Thermo Scientific, Spain) in TBST 2% (w/v) skim milk. After repeated washes, membrane was developed using SuperSignal West Pico PLUS Chemiluminescent Substrate (Thermo Scientific, Spain). As an internal loading control, α-tubulin (Sigma, Spain) was used.

**Karyotyping**

A standard optimized G-band technique with minor modifications was employed to determine the karyotype of iPSC lines. Briefly, actively proliferating cells were treated with 10 µg/mL Colcemid (Sigma, Spain) for 3 h, trypsinized, washed in PBS, and incubated in hypotonic PBS (74.85 mOsm/kg H2O) for 20 min, before immersed them in Carnoy’s fixative (cold methanol/acetic acid 3:1; both from BDH) for 30 min. Nuclei were then centrifuged at 500 × g for 2 min and resuspended in fresh Carnoy’s to wash residual PBS. Fixed nuclei were spread and Gbanded by Biobanco del Sistema Sanitario Público de Andalucía, Granada (Spain).

**Array CGH Analysis**

Array CGH analysis was performed by Genetabi Diagnostics (Spain). Genomic DNA was isolated from repaired and nonrepaired CF-human iPSC (hiPSC) and from parental fibroblasts by standard techniques. Whole-genome array CGH analysis was carried out using 0.5 µg of genomic DNA and a 60-K oligonucleotide array (Agilent Technologies, Santa Clara, USA; G4450A), according to the manufacturer’s instructions. Image quantification, hybridization quality control, and CNV detection were accomplished using Agilent Feature Extraction version (v.)11.5 and Agilent Workbench v.7.0. CNVs identified in the samples were visualized using the University of California, Santa Cruz (UCSC), Genome Browser (http://genome.ucsc.edu).
SUPPLEMENTAL INFORMATION
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AUTHOR CONTRIBUTIONS
A.F. designed and performed experiments, analyzed the data, and wrote the manuscript. S.V.-D., J.M.M.-F., A.S.-G., S.A., and J.L.C. performed experiments and interpreted results. S.V.-D. contributed to writing the manuscript. M.C., A.d.P., A.E., J.L.P., and M.V. provided intellectual input. D.B. designed and coordinated the research, analyzed the data, and wrote the manuscript. All authors reviewed and approved the final manuscript.

CONFLICTS OF INTEREST
S.V.-D., A.d.P., A.E., J.L.P., M.V., and D.B. are founders and shareholders of Karuna Good Cells Technologies S.L.

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