



Adeno-associated virus–targeted disruption of the *CFTR* gene in cloned ferrets

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Somatic cell gene targeting combined with nuclear transfer cloning presents tremendous potential for the creation of new, large-animal models of human diseases. Mouse disease models often fail to reproduce human phenotypes, underscoring the need for the generation and study of alternative disease models. Mice deficient for *CFTR* have been poor models for cystic fibrosis (CF), lacking many aspects of human CF lung disease. In this study, we describe the production of a *CFTR* gene–deficient model in the domestic ferret using recombinant adeno-associated virus–mediated gene targeting in fibroblasts, followed by nuclear transfer cloning. As part of this approach, we developed a somatic cell rejuvenation protocol using serial nuclear transfer to produce live *CFTR*-deficient clones from senescent gene-targeted fibroblasts. We transferred 472 reconstructed embryos into 11 recipient jills and obtained 8 healthy male ferret clones heterozygous for a disruption in exon 10 of the *CFTR* gene. To our knowledge, this study represents the first description of genetically engineered ferrets and describes an approach that may be of substantial utility in modeling not only CF, but also other genetic diseases.

Introduction

Cystic fibrosis (CF) is the most common autosomal recessive condition affecting white individuals. Although the *CFTR* gene – which encodes an epithelial chloride channel that is defective in CF – was cloned nearly 2 decades ago, progress toward treatments has been hindered by the lack of an animal model that reproduces the life-threatening lung infections observed in CF patients. While certain CF mouse models demonstrate phenotypic alterations in response to bacterial agarose bead challenge in the lung (1), they fail to develop the natural progression of spontaneous disease seen in humans. This is most likely due to species-specific expression of alternative chloride channels in their airways (2, 3). The domestic ferret (*Mustela putorius furo*) has been considered as an alternative species for modeling CF, given the high degree of similarity between ferret and human lung biology and known utility as a model for other types of lung infection, such as SARS (4) and influenza (5) virus.

The domestic ferret is an attractive alternative species to model CF for several reasons. First, ferrets and humans share a remarkably similar airway cytoarchitecture (6–8), a feature not shared by humans and mice. Second, the expression pattern of the *CFTR* gene is extremely similar in ferret and human airways (9, 10), with the highest levels in submucosal glands. Bioelectric and pharmacologic properties of the *CFTR* chloride channel in ferret airway epithelia are also similar to those seen in human airway epithelia (11). In contrast to those of mice, ferret and human tracheobronchial airways contain abundant submucosal glands that express high levels of *CFTR* (9, 10, 12); these glands have been shown to be critical for

airway innate immunity in the ferret (13), as predicted for humans (14, 15). Gene transfer to ferret airway epithelia with several adeno-associated virus (AAV) serotypes has also been shown to be very closely conserved to that seen in human airway epithelia (16); again, a conservation not shared with mice (3). In addition, the ferret has a 42-day gestation time and reaches sexual maturity in 5–6 months (17), making it one of the more rapidly reproducing species for animal modeling by somatic cell nuclear transfer (SCNT). Together, these studies point to the potential of the ferret to be a good species on which to model CF lung disease in humans.

Since the birth of Dolly (18), the concept of combining SCNT with gene targeting in somatic cells has held tremendous potential for the development of new animal models of human disease such as CF. However, to date, this approach has failed to deliver on such potential due to technical challenges. A few laboratories have successfully applied SCNT with gene-targeting technology in livestock for agricultural and biomedical (organ transplantation and protein production) applications (19–22). However, to our knowledge, no report has yet to apply this application to generate better non-rodent disease models. The development of robust gene-targeting technologies that are compatible with SCNT have also been rate limiting, slowing progress in this area. Additionally, efficient nuclear transfer (NT) cloning procedures for potentially useful species such as ferret have lagged behind those for larger species such as sheep (8%–10%) (18, 23), cattle (10%–20%) (24, 25), and pigs (5.5%) (26). These cloning efficiencies of 5%–20% have been sufficient to generate gene-targeted sheep, cattle, and pigs (19–22). While ferret cloning from highly reprogrammable somatic cumulus cells has been reported (27), SCNT methods with fibroblasts (the best cell type for gene targeting) have yet to be developed. Thus, further optimization of fibroblast-based SCNT cloning procedures are needed to facilitate the development of genetic models in the ferret.

Nonstandard abbreviations used: AAV, adeno-associated virus; CF, cystic fibrosis; NT, nuclear transfer; rAAV, recombinant AAV; SCNT, somatic cell nuclear transfer.

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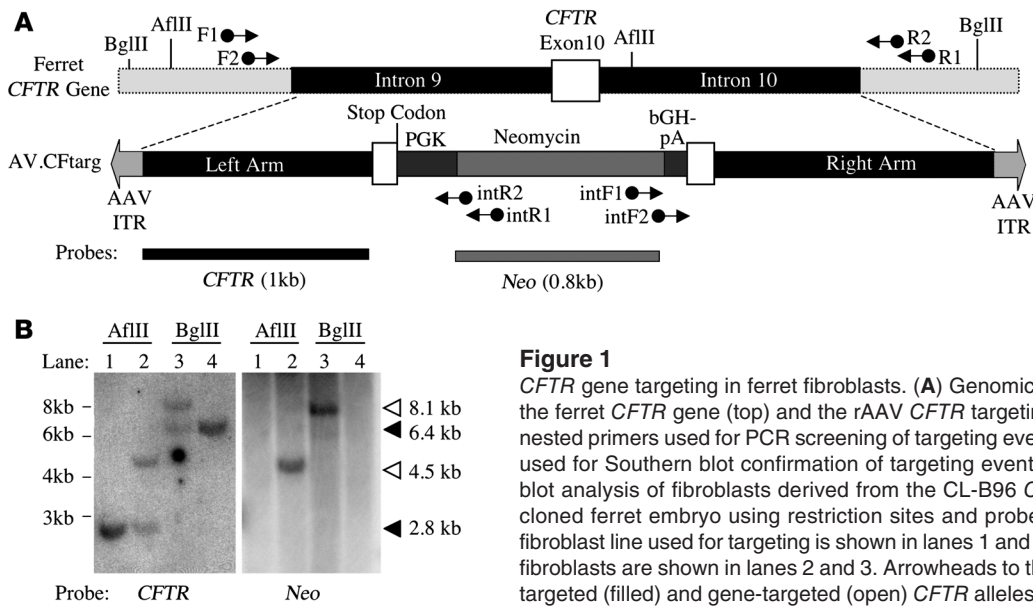


Figure 1

CFTR gene targeting in ferret fibroblasts. (A) Genomic fragment containing exon 10 of the ferret *CFTR* gene (top) and the rAAV *CFTR* targeting vector (bottom). Arrows mark nested primers used for PCR screening of targeting events. Restriction sites and probes used for Southern blot confirmation of targeting events are also shown. (B) Southern blot analysis of fibroblasts derived from the CL-B96 *CFTR* gene-targeted 21-day NT cloned ferret embryo using restriction sites and probes marked in A. The originating fibroblast line used for targeting is shown in lanes 1 and 4, while the *CFTR* gene targeted fibroblasts are shown in lanes 2 and 3. Arrowheads to the right of blots indicate the non-targeted (filled) and gene-targeted (open) *CFTR* alleles.

Here we report the production of cloned domestic ferrets heterozygous for a disrupted *CFTR* gene. Critical to this success was the development of recombinant AAV (rAAV) gene-targeting methods in fetal ferret fibroblasts and a serial NT cloning technique to rejuvenate senescent gene-targeted cells prior to cloning of full-term ferrets. Using this serial cloning technique, 8 healthy *CFTR* gene-disrupted male ferrets were obtained from 11 recipient jills. This study has demonstrated the feasibility of AAV-mediated genetic manipulation in the ferret using SCNT. Such an approach may also be of utility in genetic manipulation and disease modeling in other species.

Results

rAAV-mediated targeting of the CFTR gene in ferret fetal fibroblasts. rAAV has been previously shown to facilitate homologous recombination between its viral DNA and cellular genomic DNA of infected fibroblasts (28). Therefore, we chose to use this virus to target the *CFTR* gene in ferret fibroblasts. A ferret BAC library (CHORI-237) was constructed by the BACPAC Resources Center, Children’s Hospital Oakland Research Institute, and used to isolate an approximately 150-kb genomic fragment containing exon 10 of the *CFTR* gene. Genomic sequences from this BAC clone were used to generate a rAAV targeting vector harboring a PGK promoter-driven neomycin resistance gene cassette flanked by sequences encompassing *CFTR* exon 10 and the adjacent introns (Figure 1). Male fetal fibroblasts derived from 28-day fetuses were used for *CFTR* gene targeting with rAAV serotype-2 (rAAV2). Fibroblasts were infected with rAAV2 virus at a multiplicity of infection of 100,000 particles per cell and subsequently serially diluted into 96-well plates and placed under G418 selection. During optimization of this procedure, we found that the number of cells seeded into each well and the timing of G418 selection after plating were the most critical variables to the subcloning process, which targeted 15%–30% of wells with surviving clones by 15 days. Typically, seeding 200–500 cells per well and initiation of selection (300 µg/ml G418) at day 2 following passaging into 96-well plates was optimal.

Following replica plating of each primary 96-well plate of selected fibroblast clones, a single plate was used for PCR screening of

flanking sequences outside each targeting arm of the vector (Figure 1A). In total, approximately 500 clones were typically screened in a single experiment. Although the efficiency of gene targeting ranged from 0.5%–2% depending on the experiment, we found that the largest hurdle was the rapid senescence of PCR-positive targeted clones once they were expanded to 24-well plates. Initially, this was an obstacle that prevented direct confirmation of gene targeting by Southern blotting prior to SCNT.

Rejuvenation of PCR-positive CFTR-targeted senescent fibroblasts by SCNT. *CFTR*-targeted ferret fibroblast clones senesce rapidly during expansion following the initial round of PCR screening; therefore, it was necessary to develop methods for rejuvenating and expanding these candidate *CFTR*-targeted clones for Southern blot confirmation of the *CFTR*-targeting events. To this end, we performed several rounds of SCNT on 3 independent PCR-positive *CFTR*-targeted clones and successfully cloned six 21-day ferret fetuses by NT (Table 1). Each of these NT fetuses was dissociated with trypsin, and primary fetal fibroblasts were generated under selection with G418. DNA was then derived from each of these primary fibroblast lines and used for Southern blotting. From these cultures, one of the 3 original senescent fibroblast lines (CL-B96) gave rise to secondary NT-derived fetal fibroblasts with a “clean” *CFTR* gene-targeting event and no other integrations, as shown by Southern blotting (Figure 1B). The remaining 2 senescent fibroblast lines gave rise to NT-derived fetal fibroblasts that were neomycin resistant but had an insertion of the rAAV vector at a non-homologous site in the genome. NT-derived fetal fibroblasts from the CL-B96 rejuvenated fibroblast clone were expanded to passage 5 for SCNT cloning of live-birth ferrets.

SCNT cloning of CFTR-targeted ferrets from fetal fibroblasts. We have reported methods of SCNT cloning of ferrets using cumulus cells (27). However, in initial pilot studies, we observed that these previous methods were not compatible with efficient cloning from fetal fibroblasts. Thus, ferret SCNT protocols were optimized to resolve the major deficiencies preventing successful ferret cloning with fetal fibroblasts. Major variables included the age of recipient oocytes, alterations to media designed to reduce the stress



Table 1
Rejuvenation of senescent *CFTR* gene–targeted ferret fibroblasts by SCNT

Senescent PCR-positive <i>CFTR</i> -targeted fibroblast clones	No. of NT embryos transferred	No. of recipients pregnant (%)	No. of 21-day fetuses (%)	No. of Neo-resistant secondary cell lines	Southern blot positive for <i>CFTR</i> targeting
CL-58	230	1/6 (16.7)	2 (0.9)	2	0
CL-B96	130	1/3 (33.3)	1 (0.8)	1	1
CL-124/590	143	1/3 (33.3)	3 (2.1)	3	0

during embryo manipulation, and optimization of the timing for oocyte activation to more efficiently promote nuclear remodeling. Using this optimized method for ferret fibroblasts, we observed a doubling in oocyte implantation as compared with the previous methods developed for cumulus cells (Table 2). This enhanced level of implantation also led to normal fetal development in 2.2% of implanted reconstructed embryos, whereas fetal development failed to occur when we used the previous methods employing cumulus cells (Table 2).

Using this improved method of SCNT with ferret fibroblasts, we proceeded to clone live-birth ferrets from the NT-rejuvenated, *CFTR*-targeted fibroblasts (CL-B96). In total, 11 recipient jills were each adoptively transferred with 35–60 reconstructed NT embryos derived from the CL-B96 line and allowed to develop to term (Table 3). Fourteen live pups were born by natural birth or C-section from these 11 jills. Twelve of these 14 pups appeared to be healthy at birth. In 3 cases however, the jills only nursed one pup in the litter or failed to nurse at all. Two abandoned pups were rescued by transfer to foster jills. However, it was not possible to do this in all cases of parental neglect, since surrogate jills were not always available. Two of the 14 pups born appeared weak and died; however, it was unclear whether the jills were lactating and surrogates were not available at the time. Southern blotting confirmed that all 8 of the surviving NT cloned ferrets (Figure 2A) were heterozygous for a single targeted allele of the *CFTR* gene (Figure 2B). Furthermore, these ferrets all remained healthy and had preweaning growth rates similar to those of noncloned pups (Figure 2C).

Discussion

Combining rAAV-mediated gene targeting with SCNT cloning lays the foundation for numerous other applications in disease modeling with ferrets and other species (see the companion article in this issue; ref. 29). In previous reports of the generation of gene-targeted animals in pig (20, 21), cow (22), and sheep (19), linear fragments have been used to facilitate gene targeting with non-viral gene transfer methods. Important differences between these

previous approaches and our current strategy are worth noting. First, gene-targeting experiments in pig and sheep fibroblasts have utilized expression from the endogenous promoter of the target gene to facilitate positive selection by insertion of an internal ribosome entry site upstream to the resistance marker gene (19–21). This was not feasible for targeting the *CFTR* locus, since this gene is not expressed in fibroblasts. A second strategy used to target the bovine gene encoding immunoglobulin- μ required an alternative approach, since this gene is not expressed in fibroblasts (22). In this context, a diphtheria toxin A gene was used as a negative marker to select against random integration events and led to a targeting efficiency of approximately 0.5%. This efficiency of targeting appears to be similar to that obtained in our experiments to target the ferret *CFTR* allele using rAAV.

Selection of gene-targeted fibroblasts can lead to rapid senescence. This was indeed the case in the ferret, where *CFTR*-targeted fibroblast clones rarely expanded beyond 1×10^6 cells. Factors affecting senescence appear to be linked to neomycin selection and stress induced by this process, as serial dilution cloning of nonselected fibroblast could easily be expanded to greater than 5×10^7 cells. This phenomenon may be one reason why there are few reports of gene-targeted animals produced by SCNT. Reversal of this phenotype in fibroblasts by NT, as described in this report, indicates that the causes of selection-based senescence are not permanent. Cell rejuvenation provides an alternative strategy to more efficiently produce gene-targeted animal models. Interestingly, senescence of *CFTR*-targeted fibroblasts was not a major obstacle in the cloning of CF pig models (see the companion article in this issue; ref. 29), suggesting that species-specific factors likely influence the biology of selection-induced senescence.

The creation of both ferret and pig CF models provides new opportunities for dissecting the pathophysiology of CF and testing of new therapies not previously approachable in mouse models of this disease. Furthermore, the fact that AAV-mediated gene targeting was successfully used to genetically engineer both ferret and pig models suggests that this technology may be generally applicable to modeling in any species. It remains unknown

Table 2
Optimization of SCNT procedures with ferret fibroblasts

Procedure	No. of oocytes fused (%)	No. of oocytes cleaved (%)	No. of embryos transferred	No. of recipients pregnant (%)	No. of implantations (%)	No. of fetuses (%)
Original ^A	100 (90.9)	81 (73.6)	110	0/3 (0)	4 (3.6)	0 (0)
Optimized ^B	262 (94.9)	224 (81.2)	276	4/6 (66.7%)	18 (6.5)	6 (2.2)

^AMethods used for cumulus cell–based SCNT as previously reported (27). ^BMethods developed in the current report for fibroblast-based SCNT.



Table 3
Cloning of *CFTR*-targeted ferrets by SCNT

Recipient jill	Reconstructed oocytes transferred	Live births	Pups surviving	Percent efficiency ^A
1	60	1 ^B	1	1.7
2	45	0	0	0
3	38	2 ^C	1	5.3
4	41	2	2	4.9
5	45	1	1	2.2
6	38	1	0	2.6
7	46	3 ^C	1	6.5
8	35	1	0	2.9
9	49	2 ^B	2	4.1
10	35	0	0	0
11	40	1 ^C	0	2.5
Total (n = 11)	472	14	8	3.0 ± 0.6

^APercent efficiency of transferred oocytes that gave rise to live births.

^BOne abandoned pup was transferred to a foster jill. ^CDespite the fact all pups were born healthy, the jill only nursed one pup and neglected others or failed to nurse all pups in the litter.

whether homozygous *CFTR* gene-disrupted ferrets will contract lung disease identical to that of humans. However, regardless of their phenotype, it is likely the field will learn much about *CFTR* biology from this model, as has been the case for CF mice. Under the proper light cycle, ferrets can reach sexual maturity in approximately 5–6 months. Hence, with a gestation time of 42 days, *CFTR*-deficient ferrets could be available in approximately 1–1.5 years. Since ferrets are a preferred model for other devastating infectious human lung diseases, such as H5N1 influenza (5) and SARS virus (4), the ability to generate genetically engineered ferrets may also be of significant utility to pandemic viral disease research.

Methods

Animals. Ferrets were purchased from Marshall Farms. Female sable jills (virgin, 6–7 months of age) and albino jills (primipara, 9–12 months of age) were in estrus when delivered. Vasectomized male ferrets (albino, 12 months of age) were used for mating to induce follicular oocyte maturation in oocyte donor jills and to induce pseudopregnancy in NT embryo surrogate jill recipients. All ferrets were housed in separate cages under controlled temperature (20–22°C) and long daylight cycle (16 hours light/8 hours dark). Ferret chow was obtained from Marshall Farms. The use of animals in this study was carried out according to a protocol approved by the University of Iowa Institutional Animal Care and Use Committee and conformed to or exceeded NIH standards.

Collection of fetal ferret fibroblast. Fetal ferret fibroblasts were obtained from 28-dpc fetuses derived from a sable (female) × sable (male) mating (Marshall Farms) as previously described (30). Each fetus was treated individually. Karyotype analysis was performed on each embryo line, and only male fibroblasts with a normal chromosomal profile were used for SCNT and *CFTR* gene targeting with rAAV.

Cloning ferret *CFTR* genomic DNA. Ferret genomic DNA was extracted from internal organs collected from an E28 female fetus and used to construct a ferret genomic BAC library through the BACPAC Resources Center (<http://bacpac.chori.org>) at Children's Hospital Oakland Research Institute. The average length of the ferret genomic DNA inserts in this library is approximately 150 kbp. BAC clones encompassing ferret *CFTR* exon 10 were isolated from this library after screening with the *CFTR* exon 10 probe, synthesized according to the partial ferret *CFTR* cDNA sequence (gb:S82688) (9).

BAC DNA from the *CFTR*-positive clone was prepared with the ΨCLONE BAC DNA isolation kit (Princeton Separations). Sequencing of the *CFTR* exon 10 and adjacent introns was initiated with 2 primers located inside the *CFTR* exon 10 region (primers: e10F: 5'-TGATGATTATGGGAGAGTTGGAGCC-3' and e10R: 5'-GCATGCTTTGATGACACTCCTG-3'). Primer walking was used to sequence approximately 2 kb on each side of exon 10 to generate a contig for cloning of the targeting vector. Based on the obtained *CFTR* genomic sequence, primers for subcloning were designed, and two 2.0-kb *CFTR* fragments containing exon 10 and flanking intronic sequence were retrieved from BAC DNA by PCR with AccuPrime Pfx SuperMix (Invitrogen). The PCR products were cloned into the pBlunt4PCR vector with Topo cloning kit (Invitrogen) and confirmed by sequencing. The resultant plasmids were designated as pTopo-Left and pTopo-Right, encompassing exon 10 and left-arm or right-arm introns, respectively.

Generation of *CFTR*-targeting proviral vector. To construct the AAV targeting vector centered on *CFTR* exon 10, a 0.97-kb left homologous arm and a 1.23-kb right homologous arm were retrieved by PCR from pTopo-Left and pTopo-Right, respectively. The primer set for the left arm was: 5'-ccatcgatGGCACCCCTGTGTTATCTTTCT-3' (forward) and 5'-ccggctacctatcagATCCAGGAAAAGTGGAGCAG-3' (reverse). The right-arm set was: 5'-ccatcgatgcccgcgagctcGCCTGGCACCATCAAAGAAAAC-3' (forward) and 5'-ggactagtggatccGATGGCCTTTCCTTTGGATGGA-3' (reverse) (lowercase letters indicate restriction enzyme sequences introduced for cloning). The reverse primer of the left arm and forward primer of the right arm are located at the center of exon 10. The 2 PCR products together with a 1.7-kb PGK promoter-driven neomycin resistance expression cassette were assembled and finally cloned into an AAV2 proviral plasmid, giving rise to vector harboring 2.3-kb ferret *CFTR* genomic DNA with a neomycin cassette inserted at the center of exon 10. The rAAV2 targeting virus (AV.CFtag) was produced as previously described using a triple plasmid transfection procedure in 293 cells and purified over an iodixanol cushion followed by ion exchange HPLC (31).

Screening for *CFTR*-targeted fibroblast clones. Targeting was initiated by infecting ferret primary fibroblasts derived from a male E28 fetus with AV.CFtag at a multiplicity of infection of 100,000 particles per cell. On day 1 following infection, fibroblasts were subsequently serially diluted into twenty 96-well plates at 200–500 cells per well. These seeding densities allowed approximately 15%–30% of wells to give rise to G418-resistant clones. Selection was initiated on day 2 following replating by the addition of 300 µg/ml G418 to the media, and cells were cultured for an additional 15 days. Typically, this screening gave rise to approximately 500 G418-resistant clones, which were subsequently expanded into 3 replica 96-well plates. Once these replica plates reached confluence, a single plate was used for PCR screening of flanking genomic sequences outside each targeting arm of the vector and anchored sequences within the vector. Nested PCR screening was then performed for the predicted left-side homologous recombination event. Cells in the 96-well plates were directly lysed with 10 µl per well of lysis buffer (50 mM KCl, 1.5 mM MgCl₂, 10 mM Tris, pH 8.5, 0.5% NP40, 0.5% Tween-20), and 10% of the cell lysate (1 µl) was used for PCR with the first-round PCR primer set: F1 (5'-TGGTTTCAAGGGAATGGGGTC-3') and intR1 (5'-AAGCGAAGGAGCAAAGCTGCTA-3'). Two percent of the first-round PCR products was then used as template for the second-round PCR with primers: F2 (5'-GGTGCAGGAGGTGTTTTGTCATAGA-3') and intR2 (5'-GCTAAAGCGCATGCTCCAGACT-3'). The positive clones were further confirmed by another nested PCR reaction against the right arm of the integration site using the first-round primer set: intF1 (5'-CGGACCCTATCAGGACATAG-3') and R1 (5'-TACGAAATGCAGCAAGCGCC-3'); and the second-round nested primer set: intF2 (5'-AGGTGTCATTCTATTCTGGGG-3') and R2 (5'-CCCAGGATCCCTGAAACT-3'). The clones that were PCR positive for both the left and right

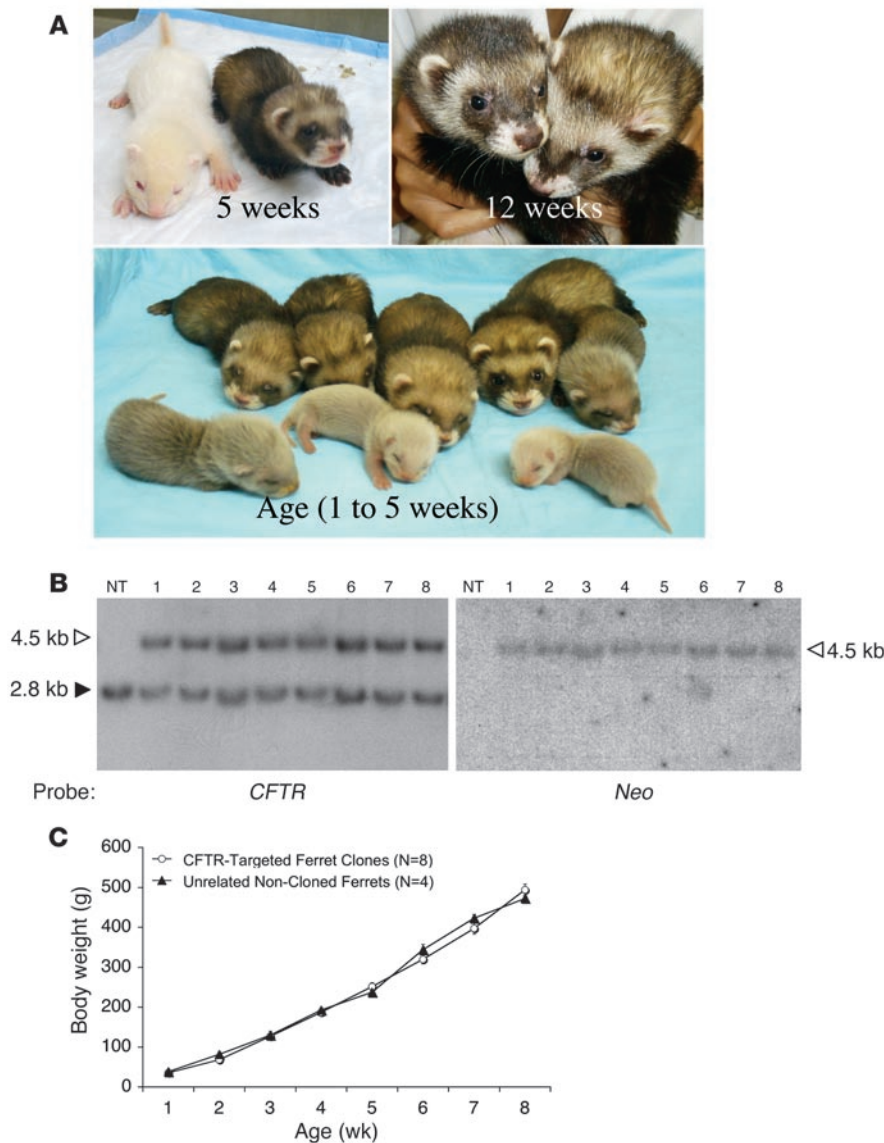


Figure 2

Cloning of *CFTR*-targeted ferrets. (A) Cloned *CFTR*-targeted ferrets. The top left panel shows the first clone (sable coat color) at 5 weeks of age with its albino noncloned foster sibling. The other panels show sable clones with ages indicated. (B) Southern blot analysis of ear fibroblast DNA from the 8 *CFTR*-targeted cloned ferrets (nos. 1–8) using *Afl*III-digested genomic DNA and the indicated *CFTR* and neomycin probes. NT, non-targeted unrelated ferret DNA. Arrowheads indicate the nontargeted (filled) and gene-targeted (open) *CFTR* alleles. (C) Prewearing growth rate of *CFTR*-targeted ferret clones as compared with unrelated noncloned ferrets. Results are shown as the mean \pm SEM for the indicated *n* in each group.

G4527; Sigma-Aldrich) plus 5 IU/ml of human chorionic gonadotrophin (hCG; C8554; Sigma-Aldrich). For enucleation, oocytes were transferred to mPBS medium containing 7.5 μ g/ml cytochalasin B (CB; C6762; Sigma-Aldrich) in the micromanipulation chamber. Using Nomarski optics, the first polar body and chromosome spindle were aspirated with a minimal volume of oocyte cytoplasm with a 20- μ m (inside diameter [ID]) PeizoDrill glass pipette (Humagen). A senescent fibroblast (diameter \geq 40 μ m) was inserted into the perivitelline space (PVS) of enucleated oocytes using another pipette (35 μ m ID). This larger-diameter pipette was specifically needed to accommodate the larger size of senescent fibroblasts. The NT-reconstructed embryos were transferred into fusion medium (0.3–0.26 M mannitol, 0.1 mM MgCl₂, 0.1 mM CaCl₂, 0.5 mM HEPES, 0.01% [wt/vol] BSA), placed between 2 parallel electrodes and subjected to an electrical pulse of 1 DC of 180 V/mm for 30 μ s from an ECM 2001 (BTX). The fused embryos were then activated with 5 mM iono-

mycin for 4 minutes, then 2 mM 6-dimethylaminopurine for 3 hours. Activated embryos were cultured for 24 hours and then were transferred into pseudopregnant albino recipient jills. A pseudopregnant state was achieved in surrogate albino virgin jills through mating with a vasectomized albino male 24 hours prior to embryo transfer. At E21 the fetuses were collected and fetal fibroblasts were established as described above, except 300 μ g/ml G418 was added to the media. Once expanded, these cells were then used for Southern blot screening of the *CFTR* gene targeting event.

Cloning of *CFTR*-targeted ferrets by SCNT. The procedure described for rejuvenating fibroblast lines by SCNT was also used for cloning live-birth ferrets from Southern blot–confirmed E21 *CFTR* gene–targeted fibroblasts (CL-B96 clone) with minimal modification. Briefly, the same pipette was used for both enucleating the oocyte and insertion of fibroblasts into the PVS of enucleated oocytes. All other NT procedures were identical to those described above. The reconstructed embryos were transferred into albino primipara pseudopregnant jills. If natural birth did not occur at 42 days gestation, the recipients were treated with prostaglandin (Lutalyse; Pfizer; 0.5 mg to 1 mg i.m.) to induce labor. If no kits were delivered within 3 hours, 0.3 ml of oxytocin was subsequently administered to the jill.

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If these treatments were unsuccessful to induce labor within 8 hours, a C-section was performed. If the recipients failed to produce enough milk to feed kits, the young were fostered onto another jill when available.

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