



ORIGINAL ARTICLE

Next generation diagnostics of cystic fibrosis and *CFTR*-related disorders by targeted multiplex high-coverage resequencing of *CFTR*

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ABSTRACT

Background Here we have developed a novel and much more efficient strategy for the complete molecular characterisation of the cystic fibrosis (CF) transmembrane regulator (*CFTR*) gene, based on multiplexed targeted resequencing. We have tested this approach in a cohort of 92 samples with previously characterised *CFTR* mutations and polymorphisms.

Methods After enrichment of the pooled barcoded DNA libraries with a custom NimbleGen SeqCap EZ Choice array (Roche) and sequencing with a HiSeq2000 (Illumina) sequencer, we applied several bioinformatics tools to call mutations and polymorphisms in *CFTR*.

Results The combination of several bioinformatics tools allowed us to detect all known pathogenic variants (point mutations, short insertions/deletions, and large genomic rearrangements) and polymorphisms (including the poly-T and poly-thymidine-guanine polymorphic tracts) in the 92 samples. In addition, we report the precise characterisation of the breakpoints of seven genomic rearrangements in *CFTR*, including those of a novel deletion of exon 22 and a complex 85 kb inversion which includes two large deletions affecting exons 4–8 and 12–21, respectively.

Conclusions This work is a proof-of-principle that targeted resequencing is an accurate and cost-effective approach for the genetic testing of CF and *CFTR*-related disorders (ie, male infertility) amenable to the routine clinical practice, and ready to substitute classical molecular methods in medical genetics.

INTRODUCTION

Cystic fibrosis (CF; MIM #219700) is one of the most common, life-threatening, autosomal recessive genetic disorders, with a carrier frequency in the Caucasian population of around 1 in 20–80 people.¹ Mutations in the CF transmembrane conductance regulator (*CFTR/ABCC7*; MIM #602421) gene determine the impairment of chloride transport in epithelial cells, mainly affecting lungs, digestive tract, sweat glands and vas deferens in men.² Although a major mutation (Δ F508) accounts for over two-thirds of CF alleles worldwide,³ a high level of allelic heterogeneity has been described within different CF populations,⁴ including single nucleotide variants (SNVs), short insertions and deletions (InDels) and large structural variants (SVs). Since the characterisation of *CFTR* more than 20 years ago,^{5–7} 1937 *CFTR* variants have been reported (Cystic Fibrosis Mutation

Database, <http://www.genet.sickkids.on.ca>). In addition to the classical CF phenotype, mild mutations in *CFTR* can cause other *CFTR*-related disorders (*CFTR*-RD), such as male infertility due to congenital bilateral absence of the vas deferens (CBAVD; MIM #277180), idiopathic chronic pancreatitis (MIM #167800), and bronchiectasis (MIM #211400) among others.⁸ Some of these mild alleles are common polymorphisms, such as poly-thymidine (poly-T) and poly-thymidine-guanine (poly-TG) tracts, associated with aberrant splicing of exon 10 of *CFTR*, being the most common mutation in CBAVD.⁹ Although *CFTR* is one of the most extensively studied human disease genes, its high allelic heterogeneity makes CF and *CFTR*-RD molecular diagnostics challenging.

The precise diagnosis of CF combines clinical evaluation (clinical features of CF phenotype and sweat test measurements) with *CFTR* molecular genetic studies. To date, the molecular characterisation of *CFTR* mutations in a given sample relies on commercial tests that screen for specific common mutations (reverse dot blot INNO-LIPA *CFTR* [Innogenetics], Cystic Fibrosis Genotyping Assay/OLA [Abbott], Elucigene CF-EU2 [Zeneca], xTAG Cystic Fibrosis 71 kit v2 [Luminex], among others). The detection rate of these panels varies depending on the mutations included (ranging from 4 to 70 *CFTR* mutations) and the molecular heterogeneity of each population. For many patients with common *CFTR* mutations that are present in these commercial panels, there is no need for additional studies, but the high heterogeneity of *CFTR* mutations in some CF populations and in *CFTR*-RD, often makes necessary the complete molecular screening of the 27 exons and the regulatory regions of *CFTR*, which is a costly and labour-intensive task.

As a first step towards the implementation of next-generation sequencing (NGS) approaches to molecular testing that can replace current low-throughput and time-consuming molecular methods, we assessed the efficacy of targeted resequencing for the molecular diagnosis of CF and *CFTR*-RD in a heterogeneous panel of 92 patients with CF and *CFTR*-RD, and CF carriers with known *CFTR* mutations.

MATERIALS AND METHODS

Detailed protocols are available in online supplementary materials.

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Subjects

High-quality genomic DNA from 92 unrelated samples, including patients with CF (n=45), CF carriers (n=27) and patients with *CFTR*-RD (n=20), were extracted from peripheral blood lymphocytes using standard protocols. The group of subjects with *CFTR*-RD included 12 patients with CBAVD, 5 patients with idiopathic bronchiectasis, and 3 patients with *CFTR*-related metabolic syndrome. All samples included in this study had previously undergone conventional *CFTR* screening,^{10 11} and all *CFTR* mutations were confirmed by Sanger sequencing, multiplex ligation-dependent probe amplification (MLPA) or quantitative PCR (qPCR). The samples selected for this study were recruited for diagnostic purposes between 1998 and 2011. For obvious reasons, it has been impossible to obtain the corresponding informed consents, although all samples were obtained with the purpose of *CFTR* mutation screening. For that, all samples were anonymised in order to ensure the protection of their identity and the list of confirmed mutations was not provided to the investigators performing the bioinformatics mutation analysis until the end of the variant prioritisation process.

In-solution capture and multiplexed resequencing of *CFTR*

Figure 1 summarises the mutation screening workflow that we have implemented in this study. Briefly, DNA from blood was sonicated to obtain fragments of approximately 200 bp. Then, fragments underwent end repair, A-tailing, and ligation to Illumina paired-end indexed adapters, as outlined in the DNA Truseq protocol (Illumina). Once the DNA libraries were indexed, they were PCR amplified and pooled before in-solution hybridisation to a custom NimbleGen SeqCap EZ Choice Library (Roche) of *CFTR* complementary oligonucleotide DNA baits. After stringent washing, the captured libraries were PCR amplified and sent for sequencing (24 libraries per lane) to generate 2×101 bp paired-end reads with a HiSeq 2000 instrument (Illumina). Finally, the resulting DNA sequences were aligned to

the human reference genome and sequence variants were detected and annotated as outlined in online supplementary materials.

Identification of CF and *CFTR*-RD mutations

In order to identify *CFTR* pathogenic mutations that could cause CF and *CFTR*-RD, we applied the following filtering steps¹²:

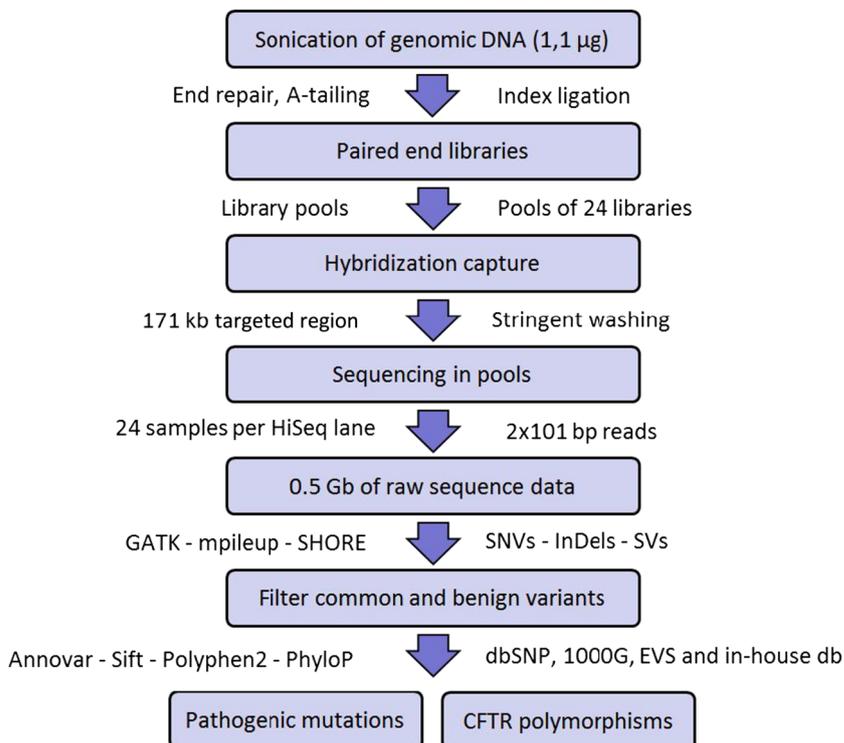
1. We required all candidate variants on both sequenced DNA strands and to account for ≥15% of total reads at that site.
2. Common polymorphisms (≥5% in the general population) were discarded by comparison with National Center for Biotechnology Information (NCBI) single nucleotide polymorphism (SNP) Database (dbSNP) build 132, the March 2010 release of the 1000 Genomes project (<http://www.1000genomes.org>), the Exome Variant Server (<http://evs.gs.washington.edu>) and an inhouse exome variant database to filter out common benign variants and recurrent artefact variant calls. However, since these databases contain known disease-associated mutations, all detected variants were compared with gene-specific mutation databases (<http://www.hgmd.cf.ac.uk> and <http://www.genet.sickkids.on.ca>).
3. Then, we screened for mutations that could give rise to premature protein truncating mutations, that is, stop mutations, damaging missense variants, splice sites, exonic deletions/insertions and large SVs.
4. Variants were ranked based upon evolutionary conservation and potential deleteriousness of the affected nucleotide using Sift,¹³ Polyphen2,¹⁴ PhyloP,¹⁵ and MutationTaster.¹⁶

RESULTS

***CFTR* enrichment**

We designed oligonucleotides to target the complete genomic sequence (the 27 exons plus all introns), and 10 kb of 5' and 3' flanking genomic regions of *CFTR* covering a total of 208 701 bp. After removal of repetitive sequences, 87% of the

Figure 1 Assay workflow to identify *CFTR* polymorphisms and pathogenic mutations.



targeted bases could be covered with capture baits for a total targeted region of 181 539 bp in 171 individual regions, with lengths ranging from 68 bp to 6689 bp (average 1062 bp). We included the untranslated region of *CFTR* to have a complete definition of the non-coding variability and to favour the detection and sizing of large SVs within the gene.

***CFTR* sequencing statistics**

On average, for each of the four HiSeq2000 (Illumina) lanes, 95.8% of the paired-end 2×101 bp reads could be assigned unambiguously to individual samples, according to their tags, receiving similar proportion of reads for each sample. The remaining 4.2% of unassigned reads were removed because of sequencing errors in their index tags. Therefore, the losses of sequence data associated with high sample multiplexing were minimal. On average, for every sample, 95% of high quality sequencing reads mapped to the reference genome. This resulted in an evenly distributed mean depth of coverage for *CFTR* of 231X (199X if the targeted regions are expanded by 150 bp at each end) with a coefficient of variation of 35%, across samples. In fact, 99.7% of all targeted bases were covered by at least 5 reads (the minimum that we require for variant calling) and 78.53% by at least 100 reads (table 1). For a comprehensive summary of the obtained sequencing results, see also online supplementary table S1.

To determine if coverage was substantially lower for any region, we calculated the proportion of base pairs that were captured by <50 reads. The proportion of these poorly covered regions accounted for 0.069 of *CFTR* targeted bases, and only 0.07% of the targeted bases were not covered by any read (table 1). As expected, these low-covered genomic regions are characterised by low complexity and a high GC content. Sequence targets with these two characteristics are usually refractory to enrichment, resulting in reduced coverage for these sites. However, as shown above, this was the case for only a very small proportion of all bases intended to be captured in this study. From these data we can conclude that all samples, regardless of the pool sizes in the precapture step, were uniformly covered at depths that in all cases exceed by far the minimum coverage required for a reliable variant calling (see online supplementary table S1). The minor differences between samples and pools were neutralised by the excessive overall *CFTR* coverage achieved by our assay. The sequence quality metrics of this data warrant a confident detection of variants in all samples.

Identification of CF and *CFTR*-RD mutations

The selection of the samples for this study was done with the idea to include as many different types of *CFTR* mutations as possible, to simulate a real-world diagnostics scenario, including SNVs, InDels, and large SVs, so that we could test the performance of our approach for all these types of genetic variations. To assess the sensitivity of our assay to detect pathogenic mutations, we blindly inspected all mapped sequence reads from the 92 samples with previously defined mutations in *CFTR*.

By using our multiplexed capture approach and automated variant calling pipeline, we were able to detect, before variant filtering and ranking, 115 SNVs (4 novel) and 28 InDels (19 novel) in *CFTR* per sample on average (table 1). Among these variants we identified several common *CFTR* polymorphisms (see online supplementary table S2). Then, we applied our variant prioritisation strategy to identify *CFTR* pathogenic mutations present in each sample. Using this strategy we detected 122 different pathogenic mutations on *CFTR* in their correct heterozygous/homozygous state across the 92 samples included

in the study (some variants were present in more than one sample). We correctly identified 58 missense, 14 nonsense, 23 splice site SNVs, 12 frameshift deletions, 2 frameshift insertions, and 3 inframe deletions (one of 84 nucleotides long), known to cause CF and *CFTR*-RD (see online supplementary table S3). In addition, we were also able to detect three different 5T pathogenic haplotypes, five large deletions, one duplication and one large genomic rearrangement (that includes one inversion and two deletions) involving various *CFTR* exons.

Intron 9 poly-TG and poly-T haplotypes and alternative splicing of *CFTR*

The 5T variant in intron 9 (c.1210–12T[5]) is the most common mutation associated with CBAVD.⁹ The penetrance of the 5T variant depends on the neighbouring TG sequence repeat.¹⁷ Thus, the definition of the TG-T (c.1210–34TG[11–13]T[5–9]) haplotype contributes to predict the most likely *CFTR*-RD phenotype of the carrier subject. However, the repetitiveness of its sequence at the nucleotide level makes difficult to determine the TG-T haplotype using standard variant calling algorithms (figure 2). In order to address this issue, we developed an in-house script that scans the very raw sequencing data of each sample for all possible combinations of c.1210–34TG[11–13]T[5–9]. By doing this, we were able to determine the exact TG-T haplotype of each sample, including three T5-TG11, eight T5-TG12 and two T5-TG13 haplotypes (see online supplementary table S4).

Characterisation of large structural changes in *CFTR*

Several of the unknown CF and *CFTR*-RD mutations in affected individuals may not have been identified yet because of the intrinsic low sensitivity of traditional PCR-based *CFTR* screening approaches for large SVs. It has been estimated that large genomic rearrangements of *CFTR*, which exhibit extensive allelic heterogeneity and are mainly caused by non-homologous recombination events, may account for up to 20% of the unidentified *CFTR* alleles in patients with CF and *CFTR*-RD.¹⁸ A major step-forward of NGS technologies with respect to classical molecular approaches is the possibility to detect large genomic rearrangements at the same time than SNVs and InDels, without the need for additional assays specific for large SVs, such as array-comparative genomic hybridisation, semi qPCR based methods, MLPA or quantitative multiplex PCR of short fluorescent fragments. In our study, the combination of paired-end mapping, split-read analysis, and normalised depth of coverage strategies allowed the blind identification of 7/7 (100% sensitivity) large SVs (5 deletions, one duplication and one complex rearrangement) in *CFTR* (figure 3). We were able to accurately identify the breakpoints of all of them, with a perfect concordance between the prediction of the algorithms and the validations for each of them (table 2).

Among the seven SVs analysed in this study we have also characterised in silico and validated by Sanger sequencing the breakpoints of a novel (ie, not previously reported to the public databases) *CFTR* 1899 bp deletion (chr7:117267155-117269054, hg19) that includes the loss of exon 22 (c.3469-420_3717+1230del1899), and all the breakpoints of a large genomic rearrangement previously reported as *CFTR*50kdel (legacy name).²¹ These two SVs were previously identified in their respective samples by means of MLPA and qPCR, but their breakpoints were not known. Thanks to the results of this study now we know that *CFTR*50kdel consists of a 85 kb inversion, with breakpoints chr7:117169862/117169876-117255003; containing two large deletions:

Table 1 Sequencing quality control parameters, coverage and detected variants by targeted resequencing of the *CFTR* gene using pools of 8, 12, 16 and 24 samples

Samples	Pool name Precapture pooling (number of samples)	All samples All	8A 8	8B 8	12A 12	12B 12	16A 16	16B 16	24A 24
Sequencing	QC-passed reads±%CV	11701689±35	19187383±14	17639.073±20	8430967±15	11654345±15	8614519±10	10449749±30	11779102±26
	% Mapped	94.9	97.13	96.36	94.53	96.36	96.4	94.23	92.58
	% Properly paired	92.98	96.03	95.05	92.49	95.06	95.04	92.07	89.73
Coverage	Mean coverage (X)±%CV	231±43	425±16	358±22	101±12	223±16	173±12	212±29	244±22
	Mean coverage extended 150 bp (X)±%CV	199±43	367±17	313±22	88±11	192±16	150±13	181±29	209±22
	% Enrichment	55.31	58.54	57.12	43.29	55.8	56.42	56.67	57.75
	% target bases covered=0×	0.07	0.03	0.01	0.12	0.1	0.09	0.07	0.06
	% target bases covered≥1×	99.93	99.97	99.99	99.88	99.9	99.91	99.93	99.94
	% target bases covered≥5×	99.7	99.84	99.86	99.41	99.72	99.76	99.67	99.72
	% target bases covered≥10×	99.39	99.76	99.78	98.55	99.49	99.56	99.29	99.45
	% target bases covered≥20×	98.48	99.63	99.58	95.92	98.82	98.92	98.23	98.69
	% target bases covered≥50×	93.13	98.86	98.38	79.25	95.1	94.74	92.49	94.78
	% target bases covered≥100×	78.53	96.34	93.79	43.14	83.23	77.56	78.29	83.64
CFTR variants	SNVs	115	124	89	121	119	124	121	104
	Novel SNVs	4	5	5	3	3	4	3	4
	Exonic SNVs	3	2	2	3	3	3	3	2
	Missense, nonsense and splice site SNPs	2	2	1	2	2	2	2	2
	InDels	28	29	29	28	30	31	28	25
	Novel InDels	19	19	19	19	20	21	17	16
	Frameshift and non-Frameshift InDels	1	1	1	1	1	1	1	1

CFTR, cystic fibrosis transmembrane regulator; CV, Coefficient of variation; InDels, insertions and deletions; SNV, single nucleotide variants; QC, Coefficient of variation.

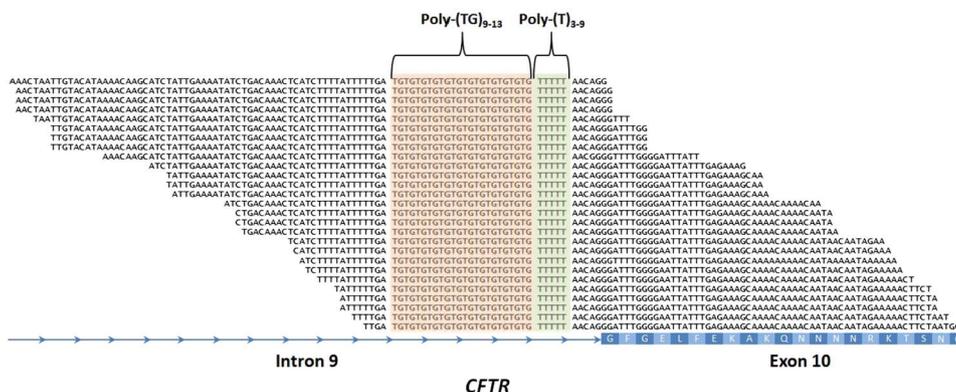


Figure 2 Detection of the intron 9 poly-TG-T haplotype involved in male infertility and other *CFTR*-RD. Example of a patient with *CFTR*-RD with the c.1210-34T[G12]T[5] haplotype. The centre of the alignment of the 100 nt NGS reads shows the poly-TG (in orange) and poly-T (in green) tracts. The *CFTR* intron 9 and exon 10 (with the amino acid sequence in white) are represented in the bottom in blue. poly-TG, poly-thymidine-guanine.

chr7:117169908-117180511(10.6 kb deletion of exons 4–8), and chr7:117216401-117254987(38.6 kb deletion of exons 12–21), 49.2 kb in total, which is remarkably close to the 50 kb deletion originally estimated by classical molecular methods²¹ (figure 4A,B). In addition, cDNA analysis evidenced an aberrant transcript showing a unique junction exon 3/22 indicating the loss of the entire inverted region (figure 4C). This is the first time that a *CFTR* large inversion is reported, and, to our knowledge, it is the most complex rearrangement ever characterised in *CFTR* (c.[274-1091_3468+236inv85141ins38; 274-1044_1116+111del10602insTATAT; 1585-11 392_3468+219del38585]).

Sensitivity and specificity of targeted resequencing of *CFTR*

The molecular diagnostic strategy for CF and *CFTR*-RD that we present here has blindly identified all previously known pathogenic *CFTR* variants in the 92 CF samples studied. This represents a mutation detection rate of 100% (122/122), with zero false-positive calls, and would have resulted in a positive molecular diagnosis in 91 of the 92 patients with CF and *CFTR*-RD and CF carriers (diagnostic rate of 98.9%), since for one of the patients with CF (sample 80) we were unable to identify his previously unknown second CF allele. As expected, for patients with *CFTR*-RD with only one previously known *CFTR* mutation our NGS strategy hasn't identified a second *CFTR* allele, as it is the case of three idiopathic bronchiectasis and four

patients with CBAVD. It is known that other genetic and environmental factors may contribute to these phenotypes,^{23–25} so the apparently missing *CFTR* alleles in these samples cannot be solely attributed to issues with the specificity or sensibility of our approach. Overall, the high success rate achieved in this study highlights the accuracy of this strategy as a molecular diagnostics tool for CF and *CFTR*-RD.

Precapture pooling and multiplexed sequencing reproducibility

Precapture pooling reduces substantially the library preparation time and, in combination with multiplexed sequencing, allows to exploit the full potential of NGS for clinical diagnostics. In order to assess how precapture multiplexing affects coverage and accuracy, we tested different pool sizes: two captures of 8, 12 and 16 samples each and one capture of 24 samples. All samples were marked with a specific index/tag, so that their individual identification was warranted at the end of the sequencing run. The sequence quality data and the variant calling results indicate that there were no sensitivity or specificity problems associated with the use of precapture pools of high number of samples (table 1). Thus, the major technical consequences of precapture pooling, which are the reduction in the input amount of the individual libraries and the addition of multiple barcodes, which may lead to less efficient blocking and

Figure 3 Detection of large structural variants in the *CFTR* gene by normalised depth of coverage analysis. Representation of the SVD-ZRPKM Values calculated by Conifer²⁹ for the 92 samples. Coloured peaks indicate the five largest structural variants identified in this study.

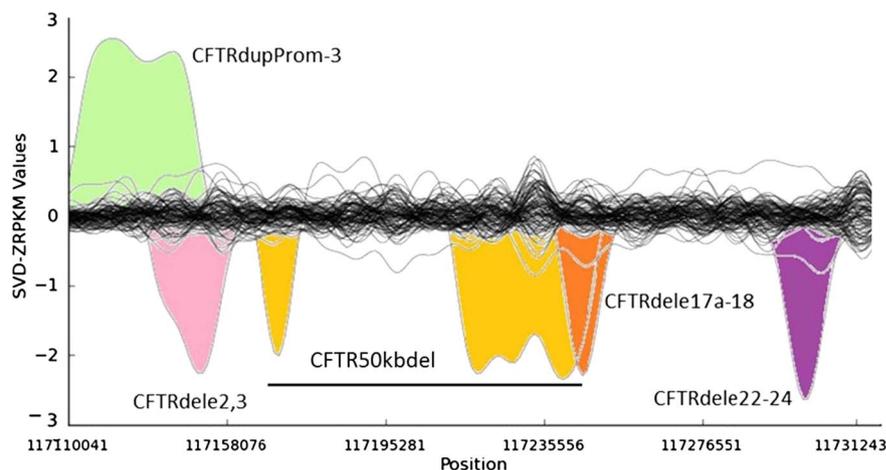


Table 2 Large structural variants identified in the *CFTR* by targeted resequencing

Sample	SV	Predicted breakpoints	Validated breakpoints	Reference
47FQ	CFTRdele20	chr7:117282464-117283245	chr7:117282468-117283248	18
69FQ	CFTRdele2,3	chr7:117138362-117159442	chr7:117138367-117159446	19
70FQ	CFTRdupProm-3	chr7:117113959-117149700	chr7:117113985-117149644	10
78FQ	CFTRdele22-24	chr7:117300851-117310305	chr7:117300852-117310305	20
83FQ	CFTR50kdbdel	INV chr7:117169861-117254986+DELS chr7:117170000-117182000+chr7:117217000-117255000	INV chr7:117169862/117169876-117255003+DELS chr7:117169908-117180511+chr7:117216401-117254987	21 and this study
88FQ	CFTRdele17a-18	chr7:117247975-117256874	chr7:117247980-117256878	22
93FQ	CFTRdele22	chr7:117267155-117269054	chr7:117267155-117269054	This study

CFTR, cystic fibrosis transmembrane regulator; SV, structural variant.

favour unspecific hybridisation,²⁶ have minor effects in the final variant calling process.

Reproducibility was determined by running four samples (two patients with CF and two patients with *CFTR*-RD) in duplicate on the same run, but captured in pools of different sample sizes (in two different precapture pools of 8 and 24 samples, respectively), and sequenced in independent HiSeq2000 lanes. We detected eight of eight pathogenic mutations in the replicated samples, yielding 100% reproducibility of mutation detection. We next assessed the reproducibility for all variant calls in the entire *CFTR* captured region (mean=138±47 SNVs and 32±11 InDels per sample). Across the four samples, reproducibility was 96.09% for SNVs and 71.62% for InDels, with an overall reproducibility of 91% for all variants in all four samples (table 3). Variant calls that did not replicate were all intronic or in intergenic regions, and almost all of them were located very close to the ends of the targeted regions of *CFTR* or in regions not covered by capture baits. This explains the observed low coverage of these unreplicated variants (mean=33.47X vs

144.58X of the replicated variants), and highlights the impact of the depth of coverage on the assay reproducibility.

Twenty-one out of the 122 pathogenic variants detected by our analysis were present in two or more individuals (see online supplementary table S3). This means a reproducibility of 100% for pathogenic variant calls between two or more samples (based on the results of 17.21% of the mutations included in this study). Since most of the samples bearing these mutations were multiplexed in independent precapture pools of different sample sizes, and also were run in different sequencer lanes, we can conclude that our approach offers great robustness and reproducibility in the detection of *CFTR* pathogenic variants. Although the coverage for a given mutation can vary significantly between samples, the proportion of reads supporting the non-reference allele was always maintained (see online supplementary table S3). Altogether, these results highlight the sensitivity and reproducibility of our assay, and support the use of a larger number of samples in precapture pools in future studies, when more index tags are available (24 when we planned this study).

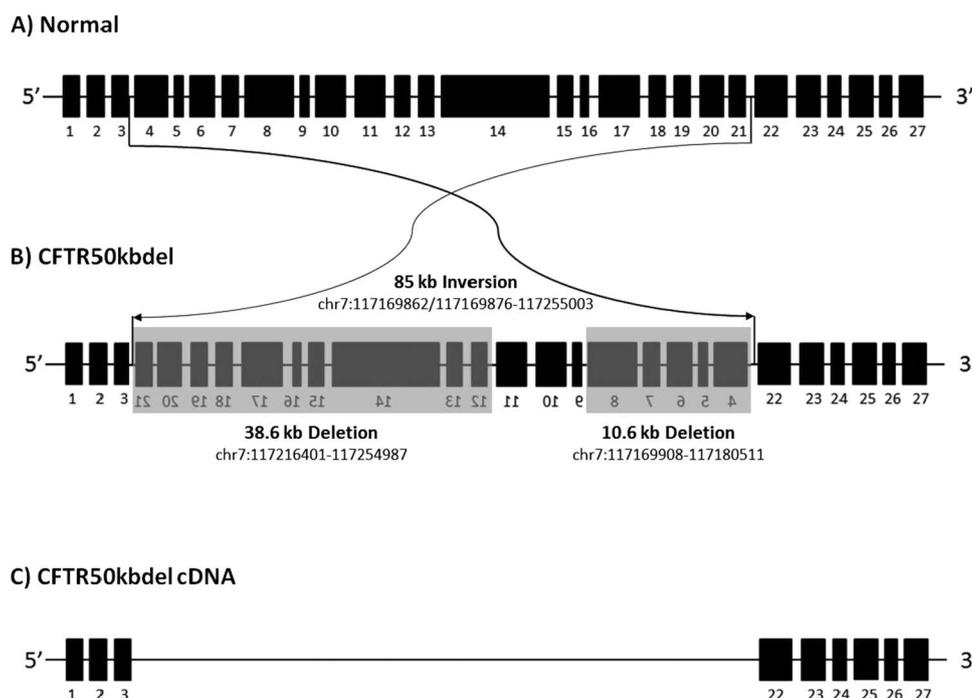


Figure 4 Schematic representation of the complex *CFTR*50kdbdel. (A) Normal structure of *CFTR*. Black boxes represent each of the 27 *CFTR* exons. (B) Diagram shows the complex architecture of the *CFTR*50kdbdel mutation. Arrows indicate the breakpoints of the 85 kb inversion. Grey areas indicate the two deleted regions. (C) cDNA sequence of *CFTR*50kdbdel, showing the loss of exons 4–21.

Table 3 Assay reproducibility of the identification of *CFTR* mutations by targeted resequencing

Sample	1		2		3		4		4 Samples	
	Reproducibility	Per cent								
SNPs	200/208	96.15	129/131	98.47	134/142	94.37	78/82	95.12	541/563	96.09
InDels	38/55	69.09	21/33	63.64	31/37	83.78	16/23	69.57	106/148	71.62
All Variants	238/263	90.49	150/164	91.46	165/179	92.18	94/105	89.52	647/711	91.00
Coverage	Mean	SD								
Matched	128.14	107.16	242.77	174.01	106.61	89.59	99.19	76.98	144.58	130.48
Unmatched	41.94	74.56	80.86	132.04	10.15	7.58	15.22	13.65	33.47	70.24

CFTR, cystic fibrosis transmembrane regulator; InDels, insertions and deletions.

DISCUSSION

Here we have implemented and tested a novel strategy for the molecular analysis of CF and *CFTR*-RD, based on pooled target enrichment and multiplexed NGS of *CFTR*. We have validated this new approach in a cohort of 92 samples with previously known pathogenic *CFTR* mutations. The different pools of simultaneously enriched *CFTR* samples were multiplexed in groups of 24 samples in four sequencer lanes. After mapping the sequencing reads to the reference genome and performing blind variant calling and filtering, our bioinformatics pipeline successfully retrieved all known pathogenic mutations in their correct heterozygous/homozygous state. With this approach we were able to identify a heterogeneous panel of *CFTR* mutations, including SNVs, InDels and large SVs. Our results (mutation detection rate of 100% and diagnostic rate of 98.91%) demonstrate the suitability of targeted resequencing for the routine clinical diagnosis of CF and *CFTR*-RD.

Clinical diagnostic tools must meet very stringent sensitivity and specificity parameters, while keeping their cost-effectiveness and time-effectiveness. The approach that we describe here represents a change in the paradigm for the molecular diagnostics of CF and *CFTR*-RD. Until now, the ideal strategy for *CFTR* screening consisted of three sequential steps:¹⁰ (1) genotyping by commercially available kits a small subset (30–50) of common *CFTR* mutations; (2) in case of not having identified the two *CFTR* alleles, complete screening of the coding portion and flanking regions of *CFTR* by scanning techniques, like denaturing gradient gel electrophoresis or single strand conformation polymorphism/heteroduplex among others, and subsequent Sanger sequencing; and if still insufficient, (3) screening by MLPA and/or array-comparative genomic hybridisation for large genomic rearrangements. The average cost per sample of this strategy is around €400 with an estimated turnaround time of 2–3 months for samples that have to undergo all three steps described above. We estimate that the approach that we present here has an overall cost of less than €200 per sample, which represents a 50% of cost savings per sample and makes the whole process eight times faster when compared with the techniques currently used for the molecular diagnosis of CF and *CFTR*-RD. In addition, our strategy offers a complete definition of the captured *CFTR*, without the need for stepwise testing anymore. We foresee that these differences will become even more significant because of the constantly dropping sequencing costs²⁷ and optimised library preparation and sequencing protocols. The complete process of library preparation, sequence enrichment, NGS and bioinformatics analysis could be completed within 14 days after reception of the DNA sample. The most time-consuming step was sequencing the *CFTR*-enriched DNA libraries on the HiSeq2000 (Illumina), which took

approximately 10 days. In addition to saving time in the process of library preparation with new capture strategies, using the most recent enrichment technologies such as Haloplex (Agilent), and optimising the bioinformatics, major time savings could be made by using the new generation of HiSeqs (Illumina) sequencers (series 2500), which have been recently reported to be able to generate up to 140 GB of sequence (2×100 bp) in approximately 24 h.²⁸ As an alternative that would reduce NGS costs, we propose the use of smaller, benchtop, personal sequencers such as the MiSeq (Illumina) or Ion Torrent (Ion Torrent Systems). The amount of sequence output of these instruments is approximately 10 times smaller than its bigger siblings, so they would be ideal for the analysis of batches of reduced numbers of samples (up to 10 samples per run).

The major drawback of capturing the complete genomic sequence of *CFTR* instead of focusing only on the coding regions is that more sequencing is needed to achieve similar coverage. However, the benefits of this approach are that no deep intronic mutations are missed, nor variants in the promoters or in the Untranslated regions (UTRs). In addition, this strategy has also proven its utility to detect large deletions, duplications and inversions, involving various *CFTR* exons, as well as to detect their breakpoints. The detection of variation in the untranslated regions of *CFTR* can also be used for the identification of alleles of clinical relevance, such as the 5T variant, which has variable penetrance and accounts for part of the phenotypic variability of *CFTR*-RD.¹⁷

In addition to the technical limitations inherent to hybrid capture, such as selection bias and uneven capture efficiency, the main limitation of the targeted resequencing approach is the impossibility to efficiently capture and sequence the repetitive and low-complexity, and GC-rich genomic segments of *CFTR* that are refractory to enrichment. However, the constant optimisation of the capture probes and NGS chemistries will gradually close the capture gaps (mainly due to uniqueness constraints, homopolymer runs, ambiguous bases or other factors that are known to cause issues in either oligonucleotide synthesis or hybridisation), and reduce enrichment variability between samples. But until then, this will require backup methods to assess the variability in these ‘dark’ regions, in the case of samples with clear CF or *CFTR*-RD phenotypes, but with no identified mutations in the captured fraction of *CFTR*, as in the case of sample 80 for which we were not able to find its previously unknown second CF allele. However, the major sources variability that could potentially affect the sensitivity and specificity in our study (such as variations in Guanine-cytosine (GC) content or differential hybridisation efficiency of the two alleles in a diploid genome) are neutralised by the high level of sequencing depth achieved.

The transition of NGS technologies from basic research to routine molecular diagnostics over the next years, will take advantage of the constant improvements in the reliability and robustness of these technologies, and of simplified bioinformatics analyses able to generate medical report-like outputs adapted to clinical laboratories. We are still in the process of defining the methods and guidelines for the application of NGS to clinical genetic diagnostics. In this initial phase, we still recommend that novel mutations are validated by Sanger sequencing before informing the patient.

In conclusion, this represents, to the best of our knowledge, the first study successfully using targeted NGS to detect pathogenic lesions in the CFTR gene. With the approach reported here we have been able to describe for the first time the break-points of a novel deletion and the most complex genomic rearrangement in CFTR. We have only had one false positive and zero spurious calls. Altogether, our assay shows a clear superiority with respect to traditional methods for CFTR screening and overcomes their technical limitations, making it their natural replacement in the diagnostic laboratories.

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Contributors This study was conceived and designed by DT, TC, LA, and XE. Selection of samples was performed by TC. NGS libraries were prepared by JG. The bioinformatics pipeline and the NGS analysis was performed by DT, CT, GE, and SO. Validation of SVs was performed by FS, MDR, and TC. The manuscript was written by DT, TC, and XE. All aspects of the study were supervised by TC and XE.

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