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Genetic spectrum of Chinese children with cystic fibrosis: comprehensive data analysis from the main referral centre in China

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ABSTRACT

Background and objectives Cystic fibrosis (CF) is a heterogeneous disease with a diverse genetic spectrum among populations. Few patients with CF of Chinese origin have been reported worldwide. The objective of this study is to characterise the genotypic features of CF in Chinese children.

Methods We recruited and characterised the genetic manifestations of 103 Chinese children with CF in Beijing Children's Hospital from 2010 to 2022. Whole-exome sequencing were performed to define the genotypes. Meanwhile, other 99 genetically confirmed patients with Chinese origin described in 45 references were also summarised.

Results 158 different variants including 23 novel observations were identified after sequencing. The majority of *CFTR* variants (82.3%) in Chinese have been observed only once or twice. 43.7% of the variants were only identified in patients of Chinese origin. The c.2909G>A(p.Gly970Asp), c.1766+5G>T and c.1657C>T(p.Arg553X) were the most frequent variants among Chinese patients, with allele frequency of 12.1%, 5.4% and 3.6%, respectively. The first two variants both showed significant Chinese ethnic tendency, while the latter one most likely came from Europeans for historical reasons. They also demonstrated significant differences in geographical distribution. c.1521_1523delCTT(p.F508del) was rarely observed in patients of pure Chinese origin, with an allele frequency of 1.8%. Two de novo variants (c.960dupA[p.Ser321IlefsX43] and c.2491-2A>G) and two deep-intronic variants (c.3718-2477C>T and c.3874-4522A>G) were identified, which were also quite rare among Chinese.

Conclusions The genetic spectrum of CF in Chinese is unique and quite different from that observed in Caucasians. The geographical distributions of the most frequent variants were reported for the first time.

Cystic fibrosis (CF) is the most frequent monogenic disease in Caucasian populations, with incidences ranging from 1/1800 to 1/25000.^{1,2} According to the American Cystic Fibrosis Foundation registry, there are currently more than 100 000 CF patients throughout the world. China is the world's most populated country with a population of approximately 1.4 billion. However, up to now, only approximately 110 CF patients of Chinese origin were reported in literature,³ and most of publications were case reports, without any epidemiological

WHAT IS ALREADY KNOWN ON THIS TOPIC

⇒ Cystic fibrosis (CF) is a rare disease in Chinese populations, with only approximately 110 CF patients of Chinese origin were reported in literature up to now. Furthermore, most of publications were case reports.

WHAT THIS STUDY ADDS

⇒ This is the largest study and most comprehensive analysis of genotypic features of CF in Chinese population to date. The genetic spectrum of CF in Chinese is unique and quite different from that observed in Caucasians. Meanwhile, the geographical distributions of the most frequent variants were reported for the first time in this study.

HOW THIS STUDY MIGHT AFFECT RESEARCH, PRACTICE OR POLICY

⇒ This study could expand our knowledge on the genotypic features of CF in Chinese population, which will greatly contribute to prevention, diagnosis and even future molecular genetic management of the disease in China.

data on the prevalence available. Interestingly, most of the cases have been diagnosed in the last 5 years, which suggests that the actual incidence of CF in China may be seriously underestimated. The contributing factors may include underdiagnosis, under-reporting, lack of national registries and variability in the frequency of mutation carriers. It has been identified that the profiles of CF transmembrane conductance regulator (*CFTR*) gene mutation spectrum vary widely among different populations based on their geographic and ethnic origins. Although CF in China is being increasingly recognised, there is still an urgent need to understand the genetic spectrum of the *CFTR* gene in Chinese patients, which will greatly contribute to prevention, diagnosis and even future molecular genetic management of the disease.

In this study, we describe 103 children with CF receiving care at the main referral centre in China. Meanwhile, other 99 genetically confirmed cases with Chinese origin reported worldwide from 1993 to 2022 were also summarised. To our knowledge, this is the largest study and most comprehensive



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analysis of genotypic features of CF in Chinese population to date.

STUDY DESIGN AND METHODS

Children referred to the Respiratory Department of Beijing Children's Hospital from January 2010 to May 2022 were enrolled in the study after meeting the diagnostic criteria of CF. CF was diagnosed based on the Consensus Guidelines from the Cystic Fibrosis Foundation²: at least one of the key clinical features highly suggestive of CF, including sinopulmonary, gastrointestinal, reproductive systems manifestations, as well as evidence of *CFTR* dysfunction, including elevated sweat test and/or presence of biallelic pathogenic variants.² Duplicate cases that had been reported in previous literatures were excluded.

In addition, we studied all the published CF cases of Chinese origin by searching China National Knowledge Infrastructure database, Wanfang database, PubMed, Embase, Cochrane Library, OVID medicine and SinoMed databases from January 1975 to January 2022. The search strategy included the following term keys: ('cystic fibrosis') AND ('Chinese' OR 'China'). Study types included clinical trials, meta-analyses, randomised controlled trials, case reports, case series or reviews. Original articles were included if they met the criteria including the patients were of Chinese origin, the presence of CF disease and complete data of both clinical manifestations and genetic sequencing. Incomplete cases or duplicate reports were excluded from our final analysis.

SWEAT CONDUCTIVITY MEASUREMENT

The Macroduct collection system and Sweat-Chek conductivity analyzer (Wescor Inc, Logan, Utah, USA) were used for CF sweat conductivity analysis as previously described.⁴⁻⁶ Based on a user's manual, sweat secretion was stimulated by pilocarpine iontophoresis. Following the stimulation, sweat was collected in a coiled plastic tubing collector cup for 30 min. Then, the sweat sample was transferred from the Macroduct tube to the take-up tube on the conductivity cell during the analysis. The test was repeated on two separate days. The average results were considered normal if values were below 60 mmol/L and intermediate if values were between 60 and 80 mmol/L.⁷ CF was very likely if values were equal to or above 80 mmol/L.

CFTR gene sequencing

Genomic DNA samples were extracted from peripheral blood leukocytes by using standard genomic DNA purification methods. The whole-exome sequencing, bioinformatics analysis and Sanger sequencing validation were performed according to their standard approach as previously described.⁸ Multiplex ligation-dependent probe amplification (MLPA) (MRC-Holland) was applied to detect the large deletions or duplications of *CFTR* gene. All exons, proximal introns and selected regions of deep introns were fully analysed.

RESULTS

Demographic data

A total of 103 patients (40 males and 63 females) from 100 Chinese families presented to Beijing Children's Hospital were recruited into this study. All 103 patients were of Chinese origin, and none of them had a family history of intermarriage with Caucasians. Patients no. 30-1/no. 30-2, no. 31-1/no.31-2 and No. 76-1/no.7622 were siblings respectively both suffering from CF. Only two children (case no. 30-1/no. 30-2) from the same family were the products of consanguineous marriage. The mean (\pm SD) ages at CF diagnosis were 7.6 (\pm 4.4) years.

Eighty-four children accepted sweat conductivity analysis; among them, 78 children had positive results and six of them showed intermediate values. The mean value was 114.1 mmol/L ranging from 60.0 to 168.0 mmol/L (online supplemental e-table 1).

In addition, other 99 Chinese patients from 94 different families described in 45 references were included in this study (Mainland China (82 cases),^{4 6 9-41} Taiwan area (nine cases),⁴²⁻⁴⁷ Hong Kong (five cases),⁴⁸ Australia (one case),⁴⁹ Canada (one case)⁵⁰ and USA (one case)).⁵¹ Overall, a total of 202 CF patients of Chinese origin constituted the cohort for the current analysis.

Genetic spectrum summarising

The *CFTR* gene variants identified in all the 103 patients (100 families) presented to Beijing Children's Hospital are listed in online supplemental e-table 1. Of note, cases 1~19⁴ and cases 20~31⁶ (except for patient no. 30-1 and no. 31-1) have been reported in 2016 and 2020, respectively, by the authors. The remaining cases have never been reported so far. Twenty-three novel observations were identified after sequencing (c.222delG[p.Arg75AspfsX16], c.298C>T[p.Leu100Phe], c.464C>G[p.Ala155Gly], c.940G>T[p.Gly314Trp], c.1064C>G[p.Pro355Arg], c.1219G>T[p.Glu407X], c.1265C>T[p.Ser422Phe], c.1347_1350delAGAA[p.Arg450AspfsX18], c.1368delT[p.Ala457LeufsX12], c.1393-?_1584+?del, c.1514delA[p.Asn505IlefsX22], c.1523_1534delTTGGT-GTTTCCT[p.Phe508_Ser511del], c.1772T>C[p.Val591Ala], c.1810A>C[p.Thr604Pro], c.2042A>T[p.Glu681Val], c.2058_2061delTT [p.Phe687X], c.2328dupA[p.Val777S-erfsX2], c.2489dupA[p.Glu831GlyfsX5], c.2909-?_3468+?del, c.3140-?_3367+?del, c.3469-12T>G, c.3469-2A>T and c.3659C>T[p.Thr1220Ile]).

The spectrum of *CFTR* variants detected in all Chinese patients with CF (194 families, 388 alleles) was summarised in online supplemental e-table 2. As a result, 373 mutated alleles were detected (detection rate: 96.1%). The identified variants included 56 missense, 31 nonsense, 27 frameshift, 21 splicing, 11 large insertion/deletion, 8 sequence variation and 4 in-frame insertion/deletion. Overall, 158 different variants were identified after sequencing. The majority of *CFTR* variants (82.3%, 130/158) in Chinese have been observed only once or twice. Approximately half of the variants (43.7%, 69/158) were only identified in patients of Chinese origin thus far, and to our knowledge, they have never been reported in Caucasians (online supplemental e-table 2). The c.2909G>A(p.Gly970Asp) was found to be the most frequent variant among Chinese CF patients with the highest allele frequency of 12.1% (47/388), followed by c.1766+5G>T (5.4% (21/388)) and c.1657C>T[p.Arg553X] (3.6% (14/388)), which were the second and third most common variants, respectively (online supplemental e-table 2, figure 1). They also showed significant differences in geographical distribution, that is, the c.2909G>A(p.Gly970Asp) variant was found in the Northern and Eastern China, while the c.1766+5G>T and the c.1657C>T(p.Arg553X) variants were most common observed in the Southern and Eastern coasts (figure 2). Meanwhile, c.1521_1523delCTT(p.F508del) was observed in six patients of pure Chinese origin, with an allele frequency of 1.8% (7/388). Interestingly, two de novo variants (c.960dupA[p.Ser321IlefsX43] and c.2491-2A>G) and two deep-intronic variants (c.3718-2477C>T and c.3874-4522A>G) were identified, which were also quite rare among Chinese.

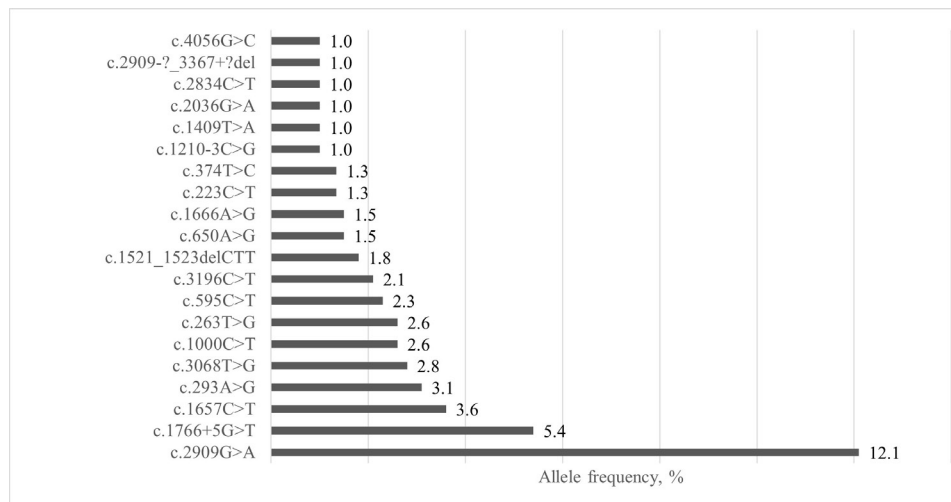


Figure 1 Frequency (>1%) of CFTR variants observed among 202 Chinese CF patients. CF, cystic fibrosis; CFTR, CF transmembrane conductance regulator.

DISCUSSION

Herein, to the best of our knowledge, we report results of the most comprehensive analysis to date of genotypic features associated with CF patients originating from across China. The first CF patient of Chinese origin was reported in 1975, which was diagnosed by sweat test.⁵² Then, it was not until 1993 that there was the first genetically confirmed case.⁴⁶ From 1993 to 2022, a total of 202 CF patients of Chinese origin have been diagnosed with definite CFTR variants. In addition, 158 different variants of CFTR gene were identified, including 23 novel observations (current report). Of these, only 45 variants are known to be CF causing in CFTR2.⁵³ In addition, 50 loss-of-function variants (nonsense, frameshift and large insertion/deletion) are likely to be CF causing. However, the pathogenic significance for the remaining 63 variants is unknown, and further functional elucidation is necessary.

The variant spectrum of CFTR among Caucasians in Western countries has been well established. c.1521_1523delCTT(p.

F508del) is the most frequent variant in Caucasians, accounting for approximately 70% of mutated alleles in general.⁵⁴ However, it is quite rarely seen in Asia, especially East Asia.⁵⁵ The c.1521_1523delCTT (p.F508del) was only observed in six patients of Chinese origin (one in homozygosity and five in compound heterozygosity), with an allele frequency of only 1.8%. No cases have ever been reported in other East Asian countries so far (except for mixed Asian-Caucasian parentage). By contrast, among Chinese population, the majority of CFTR variants (82.3%, 130/158) have been observed only once or twice. Approximately half of the variants (43.7%, 69/158) were only identified in patients of Chinese origin thus far. The c.2909G>A(p.Gly970Asp) and c.1766+5G>T variants were the most predominant observations, occurring in 12.1% and 5.4% of the alleles among all the reported Chinese patients, respectively. Notably, both variants show significant Chinese ethnic tendency, because to our knowledge, most cases with them were reported among Chinese. The c.1657C>T(p.Arg553X) was the

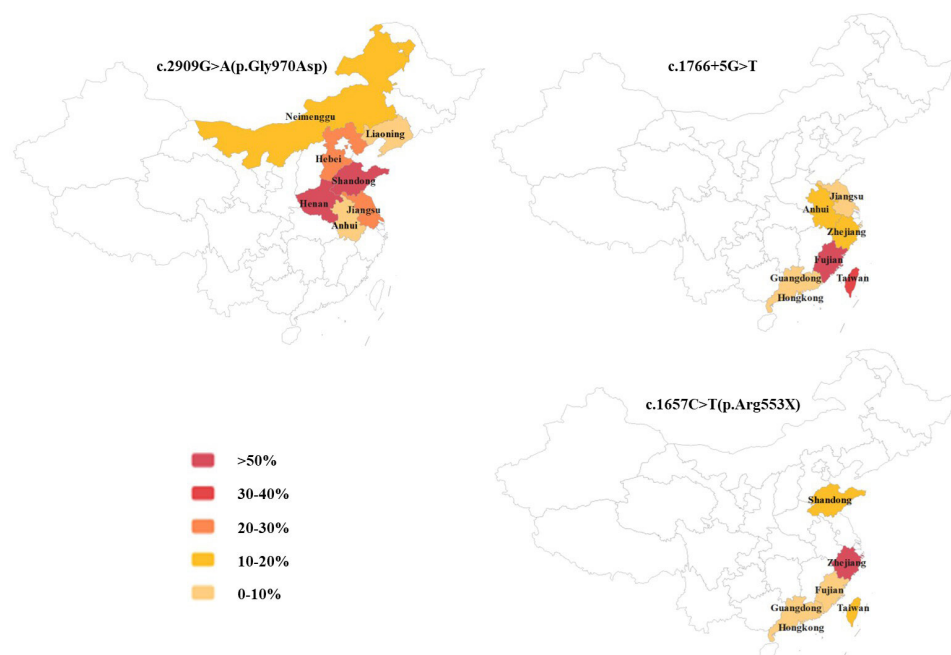


Figure 2 Frequency distribution of the top three most frequent variants among 202 Chinese patients with CF. CF, cystic fibrosis.

third most commonly observed variants, with an allele frequency of 3.6%. Interestingly, the c.1657C>T(p.Arg553X) was also present in the panel of 23 mutations proposed by the American College of Medical Genetics and Genomics.⁵⁶ It has been associated with Central European-derived populations, and the clinical consequence of this variant is known to be CF causing.⁵⁷ In terms of geographic distribution, the c.2909G>A(p.Gly970Asp) variant was found in the Northern and Eastern China, while the c.1766+5G>T and the c.1657C>T(p.Arg553X) variants were most common observed in the Southern and Eastern coasts (figure 2). We presume that the different distribution of these variants may be due to the different ethnic groups in different parts of China. Eastern coast is considered the most prosperous region of China's economy and trade, with more frequent population migrations from both Northern and Southern areas. That may be a possible explanation that the top three most frequent variants were all found in the Eastern area. The first description of c.1657C>T(p.Arg553X) among Chinese was made in homozygosity genotype in a native Taiwanese boy diagnosed with CF. Chen *et al*⁴⁵ proposed that the occurrence of c.1657C>T(p.Arg553X) variant in Taiwan area may correspond to the colonisation by the Dutch and Spanish 300 years ago. Interestingly, in this study, half of the c.1657C>T(p.Arg553X) variants were found in Zhejiang Province, which located on the southeastern coast of China. Historically, since China established diplomatic relations with European countries in the 1950s, Zhejiang people immigrated to Europe through Macao and Hong Kong. At present, Zhejiang migrants live throughout Europe. Based on this, we speculate that the origin of the c.1657C>T(p.Arg553X) variant most likely came from Europeans, which may be due to the intermarriage between Taiwanese and Europeans during the colonial period, or the wave of European economic immigration after the founding of new China.

Variants occurring de novo in the *CFTR* gene are extremely rare, with approximately 10 cases of de novo *CFTR* variants published to date.⁵⁸ There was an interesting finding that two de novo variants (c.960dupA[p.Ser321IlefsX43] and c.2491-2A>G) were identified among Chinese, which were confirmed after paternity test as well as *CFTR* gene screening for the biological parents. Furthermore, both de novo variants were found in Beijing Children's Hospital. The insertion c.960dupA(p.Ser321IlefsX43) has been reported in 2016,⁴ while the splicing variant c.2491-2A>G was a recent observation. Among previous reports, the only description of the c.2491-2A>G was made in an Irish CF patient, without any additional clinical data available.⁵⁹ In the present study, c.2491-2A>G was the first time observed as a de novo variant in a 11-year-old Chinese boy (case no. 68) with severe sinopulmonary manifestations, liver disease and pancreatic insufficiency (PI), who bore the c.2491-2A>G/c.3196C>T genotype disease. Compared with inherited variants, de novo variants are probably more deleterious because they have been subjected to less stringent evolutionary selection.⁶⁰ Most often, the de novo variants appeared on the paternal chromosome, and the same observation was also shown for c.2491-2A>G in our patient. Casals *et al*⁶¹ proposed that this tendency may reflect a higher mutation rate in paternal gametes. Nevertheless, the c.960dupA (p.Ser321IlefsX43) earlier found in our study was located on the maternal chromosome. The reason for this is unknown.

The *CFTR* genotype remains incomplete in 1% of CF cases, deep-intronic variants are putative candidates to fill this gap.⁶² A collection of variants in non-coding regions of the *CFTR* gene could help to assess their potential role as genetic factors that modify the phenotype. To our knowledge, only seven

deep-intronic disease-causing variants have been identified in the *CFTR* gene until now.⁶² Interestingly, two deep-intronic variants (c.3718-2477C>T[c.3717+12 191C>T] and c.3874-4522A>G) were the first time identified among Chinese patients. The c.3718-2477C>T has been associated with multiethnicity-derived populations, including Ashkenazi-Jewish, Southern European, Middle Eastern, Iranian and Indian.⁵⁷ In addition, it belongs to class V mutations in the *CFTR2* database, which result in reduced synthesis of *CFTR* protein with normal function at the epithelial cell membrane.^{57 63} In our report, the c.3718-2477C>T variant was found in two Chinese CF children (case no. 44 and case no. 83) with typical pulmonary features but without PI. The results of sweat conductivity test were intermediate (64 mmol/L) and weak positive (87 mmol/L), respectively, which were consistent with previous reports and may be considered milder CF phenotype. As far as c.3874-4522A>G was concerned, the ethnic origins were reported to be France, Iran and Laos.⁶² Moreover, this variant was considered CF causing associated with a large phenotypic spectrum, including *CFTR*-related disorders and typical CF.⁶² In the present study, c.3874-4522A>G was found in two Chinese children (case no. 39 and case no. 57) with severe sinopulmonary diseases from early childhood, including progressive bronchiectasis, recurrent airway *Pseudomonas aeruginosa*, lung function defect and sinusitis (both cases), and allergic bronchopulmonary aspergillosis (only for case no. 39). Due to the limited sequencing methods at the first time they presented to us, we only detected the c.2936A>C(p.Asp979Ala) and the c.1368delT(p.Ala457LeufsX12) variant on one allele, respectively. Three years later, we found the c.3874-4522A>G variant deeply located in intron 23 of the other allele and eventually supplemented the genetic data for both children.

In terms of the novel observations, 12 variants (c.222delG[p.Arg75AspfsX16], c.1219G>T[p.Glu407X], c.1347_1350delAGAA[p.Arg450AspfsX18], c.1368delT[p.Ala457LeufsX12], c.1393-?_1584+?del, c.1514delA[p.Asn505IlefsX22], c.1523_1534delTTGGTGTTCCT[p.Phe508_1521Ser511del], c.2058_2061delITT[p.Phe687X], c.2328dupA[p.Val777SerfsX2], c.2489dupA[p.Glu831GlyfsX5], c.2909-?_3468+?del and c.3140-?_3367+?del) are likely to be CF causing, and the clinical consequences of remaining 11 variants are uncertain. Clinical characterisation includes the presence of high frequency of sinopulmonary diseases and relatively low frequency of PI compared with Caucasians, which were consistent with previous reports on phenotype in individuals with CF of Chinese origin (online supplemental e-table 3). The geographical distribution of the novel variants showed no significant discrepancies between North and South in China.

The genetic spectrum of CF in Chinese is unique and quite different from that observed in Caucasians. Therefore, the Caucasian *CFTR* common mutation-screening panel is not applicable for Chinese patients. We recommend using the extensive *CFTR* gene sequencing (including all *CFTR* exons, their intronic boundaries and selected regions of deep introns) followed by MLPA analysis for effective diagnosis, although it is relatively expensive. If patients with CF are still incompletely genotyped, whole genome sequencing for the identification of unknown deep-intronic variants is advocated.

To date, 2110 variants of *CFTR* have been identified worldwide,⁶⁴ but the disease liability of only 401 variants has been ascertained in *CFTR2*.⁵³ Based on the nature of the molecular defects in *CFTR*, small molecules (*CFTR* modulators) can successfully restore activity to the mutant protein, thereby ameliorating disease manifestations.⁵⁵ Unfortunately, few studies on the molecular consequences of Chinese-specific *CFTR* variants have

been reported. The most frequent *CFTR* mutation in Chinese, c.2909G>A(p.Gly970Asp), was predicted to be a gating mutation with partial trafficking defect.⁶⁵ Furthermore, lumacaftor/ivacaftor therapy was proven efficacy in both *ex vivo*⁶⁵ and *in vivo*⁶⁶ studies of a patient with c.1521_1523delCTT(p.F508del)/c.2909G>A(p.Gly970Asp) genotype. These findings have significant implications for *CFTR* modulator use in or availability to the Chinese population. Future studies on functional consequences analysis of Chinese ethnicity-specific variants would be beneficial to take the first step to research on *CFTR* modulator therapies in China.

The limitation of the present study would be that the sweat chloride concentration test, which has been the gold standard for diagnosis of CF, was not available in China. However, the sweat conductivity measurement has been shown to have excellent correlation with the sweat chloride concentration.^{67,68} It is accurate, simple to perform and economical. Thus, the sweat conductivity measurement is used as an assistant diagnostic test for CF in China. As the patients in this study were enrolled over a period of decades, this could have caused unintended bias such as improvement in laboratory technology and lifestyle changes. First, sweat conductivity analysis is not readily available in all the paediatric centres in China, except Beijing Children's Hospital, which is considered the main referral centre for patients with CF from all over the country. Besides this, even in our centre, sweat conductivity analysis was not performed on all the patients enrolled because the testing facility was only available from 2014. Finally, MLPA analysis was only adopted in patients recruited from 2016 onwards also due to inaccessibility of this technique.

In conclusion, results presented herein describing the genetic spectrum of CF in Chinese is unique and quite different from that observed in Caucasians, consistent with our previous statement.⁴ The c.2909G>A(p.Gly970Asp), c.1766+5G>T and c.1657C>T(p.Arg553X) are the most frequent variants among Chinese CF patients studied. The geographical distributions of the most frequent variants were reported for the first time. These data demonstrate that it is very urgent and necessary to establish a national CF registry in China, which would be beneficial to compare genetic data intranationally and internationally from an epidemiological perspective, to evaluate the phenotype-genotype association in China and to enhance our understanding of CF pathogenic processes for improvement of disease management and prognosis.

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Contributors YS: conceptualised and designed the study, validated genetic studies, analysed the data and drafted the initial manuscript. XT and QC: validated genetic studies, carried out the initial analyses, and reviewed and revised the manuscript. HX and HLi: recruited and evaluated patients, and reviewed and revised the manuscript. JL and HY: designed the data collection instruments, coordinated and supervised data collection, interpreted the results, and reviewed and revised the manuscript. HLi and SZ: conceptualised and designed the study, interpreted the results and critically reviewed the manuscript for important intellectual content. SZ: responsible for the overall content as guarantor. All authors approved the final manuscript as submitted and agree to be accountable for all aspects of the work.

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Competing interests None declared.

Patient consent for publication Not applicable.

Ethics approval This study protocol was approved by the Ethics Committees of Beijing Children's Hospital, China (Approval no. (2022)-E-029-R). Informed written consent was obtained from all participants or parent/legal guardians. Participants gave informed consent to participate in the study before taking part.

Provenance and peer review Not commissioned; externally peer reviewed.

Data availability statement All data relevant to the study are included in the article or uploaded as supplementary information.

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e-Table 1. Variants of *CFTR* identified in 103 Chinese children (100 families) with CF presented to Beijing Children's Hospital

Case No.	Sex	Age at Dx, y	Sweat conductivity, mmol/L	Region	Nucleotide change (HGVS nomenclature)	Amino acid change	Type of mutation	Genotype	Familial targeted mutation study	
									Father	Mother
1 ^[4]	M	11.58	101	Ex 13	c.1699G>T	p.Asp567Tyr	Missense	Compound	+/-	-/-
				Ex 24	c.3909C>G	p.Asn1303Lys	Missense	heterozygous	-/-	+/-
2 ^[4]	F	10.58	103	Ex 3	c.263T>G	p.Leu88X	Nonsense	Compound heterozygous	-/-	+/-
				In 13	c.1766+5G>T	-	Splicing		+/-	-/-
				3'UTR	c.*110C>G	-	Sequence variation		-/-	+/-
3 ^[4]	M	13.25	101	Ex 22	c.3700A>G	p.Ile1234Val	Missense	Compound	+/-	-/-
				Ex 8	c.960dupA	p.Ser321IlefsX43	Frameshift	heterozygous	-/-	-/-
4 ^[4]	F	13.67	99	Ex 3	c.263T>G	p.Leu88X	Nonsense	Compound	-/-	+/-
				Ex 18	c.2909G>A	p.Gly970Asp	Missense	heterozygous	+/-	-/-
5 ^[4]	M	7.17	106	Ex 4	c.326A>G	p.Tyr109Cys	Missense	Compound	-/-	+/-
				Ex 8	c.1000C>T	p.Arg334Trp	Missense	heterozygous	+/-	-/-

				Ex 12	c.1666A>G	p.Ile556Val	Missense		+/+	-/-
6 ^[4]	F	10.67	127	Ex 6	c.595C>T	p.His199Tyr	Missense	Heterozygous	ND	ND
7 ^[4]	F	7.75	118	Ex 3	c.223C>T	p.Arg75X	Nonsense	Compound heterozygous	ND	ND
				Ex 4	c.326A>G	p.Tyr109Cys	Missense		ND	ND
8 ^[4]	F	7.33	118	Ex 8	c.1000C>T	p.Arg334Trp	Missense	Heterozygous	+/-	-/-
9 ^[4]	F	10.17	105	Ex 3	c.263T>G	p.Leu88X	Nonsense	Homozygous	+/-	+/-
10 ^[4]	F	11.08	96	Ex 12	c.1666A>G	p.Ile556Val	Missense	Heterozygous	ND	ND
11 ^[4]	M	8.25	115	Ex 4	c.293A>G	p.Gln98Arg	Missense	Compound heterozygous	-/-	+/-
				Ex 5	c.558C>G	p.Asn186Lys	Missense		+/-	-/-
12 ^[4]	F	4.17	101	Ex 4	c.326A>G	p.Tyr109Cys	Missense	Compound heterozygous	ND	ND
				Ex 14	c.2374C>T	p.Arg792X	Nonsense		ND	ND
13 ^[4]	M	3.67	99	Ex 12	c.1666A>G	p.Ile556Val	Missense	Homozygous	+/-	+/-
14 ^[4]	F	12.67	122	Ex 4	c.293A>G	p.Gln98Arg	Missense	Heterozygous	-/-	+/-
15 ^[4]	M	11.00	ND	Ex 6	c.648G>A	p.Trp216X	Nonsense	Homozygous	+/-	+/-
				In 14	c.2491-126T>C	-	Sequence variation		+/-	+/-
16 ^[4]	F	10.33	ND	Ex 20	c.3196C>T	p.Arg1066Cys	Missense	Homozygous	ND	ND

17 ^[4]	M	11.17	ND	Ex 4	c.413_415dupTA C	p.Leu138dup	In-frame insertion	Homozygous	ND	ND
18 ^[4]	F	3.42	ND	Ex 8	c.1075C>T	p.Gln359X	Nonsense	Compound heterozygous	ND	ND
				Ex 20	c.3307delA	p.Ile1103X	Nonsense		ND	ND
19 ^[4]	F	14.00	115	Ex 18	c.2909G>A	p.Gly970Asp	Missense	Homozygous	+/-	+/-
20 ^[6]	F	0.42	130	Ex 3	c.223C>T	p.Arg75X	Nonsense	Compound heterozygous	-/-	+/-
				Ex 18	c.2909G>A	p.Gly970Asp	Missense		+/-	-/-
21 ^[6]	F	2.16	134	Ex 11	c.1521_1523delC TT	p.Phe508del	In-frame deletion	Homozygous	+/-	+/-
22 ^[6]	F	8.25	122	Ex 8	c.1000C>T	p.Arg334Trp	Missense	Compound heterozygous	+/-	-/-
				Ex 13	c.1733T>C	p.Leu578Pro	Missense		-/-	+/-
23 ^[6]	M	0.75	134	Ex 3	c.264_268delAT ATT	p.Leu88PhefsX21	Frameshift	Compound heterozygous	-/-	+/-
				Ex 23	c.3860delG	p.Gly1287GlufsX 2	Frameshift		+/-	-/-
24 ^[6]	F	0.42	105	Ex 3	c.263T>G	p.Leu88X	Nonsense	Compound heterozygous	-/-	+/-
				Ex 18	c.2909G>A	p.Gly970Asp	Missense		+/-	-/-

25 ^[6]	M	0.67	93	In 8	c.1116+1G>A	–	Splicing	Compound heterozygous	+/-	-/-
				Ex 18	c.2909G>A	p.Gly970Asp	Missense		-/-	+/-
26 ^[6]	M	0.25	ND	Ex 18	c.2909G>A	p.Gly970Asp	Missense	Compound heterozygous	+/-	-/-
				Ex 23	c.3718-?_3873+? del	–	Large deletion		-/-	+/-
27 ^[6]	F	0.92	115	Ex 14	c.2236_2246delG AGGCGATACTi nsAAAAATC	p.Glu746LysfsX8	Frameshift	Compound heterozygous	-/-	+/-
				Ex 25	c.3635delT	p.Val1212AlafsX 16	Frameshift		+/-	-/-
28 ^[6]	M	6.67	125	In 9	c.1210-3C>G	–	Splicing	Compound heterozygous	+/-	-/-
				In 24	c.3964-7A>G	–	Splicing		-/-	+/-
29 ^[6]	M	0.75	125	Ex 19	c.3068T>G	p.Ile1023Arg	Missense	Compound heterozygous	+/-	-/-
				Ex 6	c.595C>T	p.His199Tyr	Missense		-/-	+/-
30-1	F	7.67	151	Ex 5	c.532G>A	p.Gly178Arg	Missense	Homozygous	+/-	+/-
30-2 ^[6]	F	0.50	ND	Ex 5	c.532G>A	p.Gly178Arg	Missense	Homozygous	+/-	+/-

31-1	M	9.33	102	In 5	c.579+1_579+2in sACAT	–	Splicing	Compound heterozygous	+/-	-/-
				In 13	c.1766+5G>T	–	Splicing		-/-	+/-
31-2 ^[6]	F	8.83	141	In 5	c.579+1_579+2in sACAT	–	Splicing	Compound heterozygous	+/-	-/-
				In 13	c.1766+5G>T	–	Splicing		-/-	+/-
32	F	1.08	125	Ex 11	c.1521_1523delC TT	p.Phe508del	In-frame deletion	Compound heterozygous	+/-	-/-
				Ex 11	c.1429_1437delC CTTCAGAG	p.Pro477_Glu479 del	In-frame deletion		-/-	+/-
33	M	7.33	125	Ex 20	c.3209G>C	p.Arg1070Pro	Missense	Compound	+/-	-/-
				Ex 14	c.2328dupA	p.Val777SerfsX2	Frameshift	heterozygous	-/-	+/-
34	F	9.00	104	Ex 4	c.293A>G	p.Gln98Arg	Missense	Compound	+/-	-/-
				Ex 19	c.3068T>G	p.Ile1023Arg	Missense	heterozygous	-/-	+/-
35	F	5.67	109	Ex 11	c.1572C>A	p.Cys524X	Nonsense	Compound	+/-	-/-
				Ex 1	c.3G>A	p.Met1Ile	Missense	heterozygous	-/-	+/-
36	F	10.67	127	Ex 6	c.595C>T	p.His199Tyr	Missense		+/-	-/-

				Ex 14	c.2058_2061delT T	p.Phe687X	Nonsense	Compound heterozygous	-/-	+/-
37	F	14.33	140	Ex 11	c.1409T>A	p.Val470Glu	Missense	Compound heterozygous	+/-	-/-
				In 9	c.1210-3C>G	-	Splicing		-/-	+/-
38	M	13.25	109	Ex 10	c.1347_1350delA GAA	p.Arg450AspfsX 18	Frameshift	Compound heterozygous	ND	ND
				Ex 18	c.2909G>A	p.Gly970Asp	Missense		Compound ND	ND
39	M	10.50	83	In 23	c.3874-4522A>G	-	Splicing	Compound	+/-	-/-
				Ex 18	c.2936A>C	p.Asp979Ala	Missense	Heterozygous	-/-	+/-
40	M	9.83	141	Ex 8	c.1000C>T	p.Arg334Trp	Missense	Compound heterozygous	-/-	+/-
				Ex 3	c.264_268delAT ATT	p.Leu88PhefsX21	Frameshift		+/-	-/-
41	F	5.58	67	Ex 6	c.650A>G	p.Glu217Gly	Missense	Compound heterozygous	-/-	+/-
				Ex 10	c.1231A>G	p.Lys411Glu	Missense		+/-	-/-
42	M	7.00	130	In 13	c.1766+5G>T	-	Splicing	Compound heterozygous	-/-	+/-
				Ex 22	c.3484C>T	p.Arg1162X	Nonsense		+/-	-/-

43	M	3.83	ND	Ex 6	c.595C>T	p.His199Tyr	Missense	Compound heterozygous	+/-	-/-
				Ex 14	c.2290C>T	p.Arg764X	Nonsense		-/-	+/-
44	M	14.42	64	In 22	c.3718-2477C>T	-	Sequence variation	Compound heterozygous	+/-	-/-
				Ex 17	c.2834C>T	p.Ser945Leu	Missense		-/-	+/-
				Ex 4	c.374T>C	p.Ile125Thr	Missense		-/-	+/-
45	M	8.50	163	Ex 11	c.1521_1523delC TT	p.Phe508del	In-frame deletion	Compound heterozygous	+/-	-/-
				Ex 4	c.293A>G	p.Gln98Arg	Missense		-/-	+/-
46	M	10.17	60	Ex 4	c.374T>C	p.Ile125Thr	Missense	Compound heterozygous	-/-	+/-
				Ex 14	c.2042A>T	p.Glu681Val	Missense		+/-	-/-
47	M	13.08	99	Ex 17	c.2834C>T	p.Ser945Leu	Missense	Homozygous	+/-	+/-
48	F	11.25	118	Ex 14	c.2036G>A	p.Trp679X	Nonsense	Compound heterozygous	+/-	-/-
				In 2	c.164+2T >C	-	Splicing		-/-	+/-
49	F	4.67	85	Ex 4	c.325T>G	p.Tyr109Asp	Missense	Compound heterozygous	-/-	+/-
				Ex 20	c.3196C>T	p.Arg1066Cys	Missense		+/-	-/-
50	M	9.08	96	Ex 4	c.350G>A-7T	p.Arg117His-7T	Missense		-/-	+/-

				Ex 14	c.2036G>A	p.Trp679X	Nonsense	Compound heterozygous	+/-	-/-
51	F	11.00	96	Ex 18	c.2909G>A	p.Gly970Asp	Missense	Compound heterozygous	+/-	-/-
				Ex 8	c.1000C>T	p.Arg334Trp	Missense		-/-	+/-
52	M	14.50	114	Ex 3	c.223C>T	p.Arg75X	Nonsense	Compound heterozygous	+/-	-/-
				Ex 4	c.293A>G	p.Gln98Arg	Missense		-/-	+/-
53	F	3.92	122	Ex 21	c.3387delT	p.Gly1130ValfsX 4	Frameshift	Compound heterozygous	+/-	-/-
				Ex 11	c.1409T>A	p.Val470Glu	Missense		-/-	+/-
54	F	8.58	125	Ex 12	c.1657C>T	p.Arg553X	Nonsense	Compound heterozygous	+/-	-/-
				Ex 14	c.1810A>C	p.Thr604Pro	Missense		-/-	+/-
55	M	10.67	108	Ex 18	c.2909G>A	p.Gly970Asp	Missense	Compound heterozygous	+/-	-/-
				Ex 20	c.3196C>A	p.Arg1066Ser	Missense		-/-	+/-
56	M	2.08	111	In 13	c.1766+2T>C	-	Splicing	Compound heterozygous	+/-	-/-
				In 21	c.3469-2A>T	-	Splicing		-/-	+/-
57	F	9.42	108	In 23	c.3874-4522A>G	-	Sequence variation	Compound heterozygous	+/-	-/-

				Ex 10	c.1368delT	p.Ala457LeufsX 12	Frameshift		-/-	+/-
58	F	8.75	136	Ex 11	c.1523_1534delT TGGTGTTCCT	p.Phe508_Ser511 del	In-frame deletion	Compound heterozygous	+/-	-/-
				Ex 11	c.1393-?_1584+? del	-	Large deletion		-/-	+/-
59	F	7.75	97	Ex 18	c.2977G>T	p.Asp993Tyr	Missense	Compound heterozygous	ND	ND
				Ex 10	c.1265C>T	p.Ser422Phe	Missense		ND	ND
				In 19	c.3140-26A>G	-	Missense		ND	ND
60	M	1.75	144	Ex 24	c.3908delA	p.Asn1303ThrfsX 25	Frameshift	Compound heterozygous	+/-	-/-
				Ex 18	c.2909G>A	p.Gly970Asp	Missense		-/-	+/-
61	F	4.17	ND	Ex 15	c.2551C>T	p.Arg851X	Nonsense	Compound heterozygous	-/-	+/-
				Ex 10	c.1219G>T	p.Glu407X	Nonsense		+/-	-/-
				Promoter	c.-152G>C	-	Sequence variation		-/-	+/-

62	M	16.83	ND	Ex 3	c.223C>T	p.Arg75X	Nonsense	Compound heterozygous	ND	ND
				Ex 18	c.2909G>A	p.Gly970Asp	Missense		ND	ND
63	F	13.75	ND	Ex 14	c.1810A>C	p.Thr604Pro	Missense	Compound heterozygous	ND	ND
				Ex 8	c.940G>T	p.Gly314Trp	Missense		ND	ND
				3'UTR	c.*133delT	–	Sequence variation		ND	ND
64	F	12.50	168	Ex 4	c.374T>C	p.Ile125Thr	Missense	Compound heterozygous	+/-	-/-
				Ex 11	c.1521_1523delC TT	p.Phe508del	In-frame deletion		-/-	+/-
65	F	1.67	89	Ex 13	c.1703T>A	p.Leu568X	Nonsense	Homozygous	+/-	+/-
66	M	4.00	ND	Ex 12	c.1657C>T	p.Arg553X	Nonsense	Homozygous	+/-	+/-
67	F	12.00	77	Ex 23	c.3841C>T	p.Gln1281X	Nonsense	Compound heterozygous	+/-	-/-
				Ex 4	c.298C>T	p.Leu100Phe	Missense		-/-	+/-
68	M	11.00	161	In 14	c.2491-2A>G	–	Missense	Compound heterozygous	-/-	-/-
				Ex 20	c.3196C>T	p.Arg1066Cys	Missense		-/-	+/-
69	F	6.33	85	Ex 22	c.3472C>T	p.Arg1158X	Nonsense		+/-	-/-

				Ex 2-3	c.54-?_273+?del	–	Large deletion	Compound heterozygous	-/-	+/-
70	F	3.25	101	Ex 6	c.607A>T	p.Ile203Phe	Missense	Compound heterozygous	+/-	-/-
				Ex 18-20	c.2909-?_3367+? del	–	Large deletion		-/-	+/-
71	F	13.25	122	Ex 20	c.3140-?_3367+? del	–	Large deletion	Compound heterozygous	-/-	+/-
				Ex 3	c.233dupT	p.Trp79LeufsX32	Frameshift		+/-	-/-
72	F	7.33	118	Ex 8	c.1000C>T	p.Arg334Trp	Missense	Compound heterozygous	+/-	-/-
				Ex 11	c.1393-?_1584+? del	–	Large deletion		-/-	+/-
73	F	15.33	149	Ex 4	c.293A>G	p.Gln98Arg	Missense	Compound heterozygous	+/-	-/-
				Ex 11	c.1521_1523delC TT	p.Phe508del	In-frame deletion		-/-	+/-
74	M	5.08	130	Ex 12	c.1657C>T	p.Arg553X	Nonsense	Compound heterozygous	+/-	-/-
				Ex 19	c.3068T>G	p.Ile1023Arg	Missense		-/-	+/-

75	F	8.58	146	Ex 3	c.222delG	p.Arg75AspfsX1 5	Frameshift	Compound heterozygous	+/-	-/-
				Ex 3	c.263T>G	p.Leu88X	Nonsense		-/-	+/-
76-1	M	7.75	130	Ex 18-20	c.2909-?_3367+? del	-	Large deletion	Compound heterozygous	+/-	-/-
				Ex 20	c.3196C>T	p.Arg1066Cys	Missense		-/-	+/-
76-2	F	5.58	120	Ex 18-20	c.2909-?_3367+? del	-	Large deletion	Compound heterozygous	+/-	-/-
				Ex 20	c.3196C>T	p.Arg1066Cys	Missense		-/-	+/-
77	F	4.50	60	Ex 4	c.298C>T	p.Leu100Phe	Missense	Compound heterozygous	-/-	+/-
				Ex 6	c.650A>G	p.Glu217Gly	Missense		+/-	-/-
78	F	12.92	119	Ex 22	c.3476C>T	p.Ser1159Phe	Missense	Compound heterozygous	+/-	-/-
				In 21	c.3469-12T>G	-	Splicing		-/-	+/-
79	M	2.75	134	Ex 22	c.3484C>T	p.Arg1162X	Nonsense	Compound heterozygous	+/-	-/-
				Ex 10	c.1369G>C	p.Ala457Pro	Missense		-/-	+/-
80	F	10.67	126	Ex 18-21	c.2909-?_3468+? del	-	Large deletion	Compound heterozygous	+/-	-/-

				Ex 18	c.2909G>A	p.Gly970Asp	Missense		-/-	+/-
81	F	10.58	127	Ex 3	c.223C>T	p.Arg75X	Nonsense	Compound	+/-	-/-
				Ex 18	c.2909G>A	p.Gly970Asp	Missense	heterozygous	-/-	+/-
82	M	6.00	144	In 13	c.1766+5G>T	-	Splicing	Homozygous	+/-	+/-
83	F	3.17	87	Ex 3	c.264_268delATA TT	p.Leu88PhefsX21	Frameshift	Compound	+/-	-/-
				In 22	c.3718-2477C>T	-	Sequence variation	heterozygous	-/-	+/-
84	F	0.50	ND	Ex 2-3	c.54-?_273+?del	-	Large deletion	Compound	+/-	-/-
				Ex 18	c.2909G>A	p.Gly970Asp	Missense	heterozygous	-/-	+/-
85	F	11.00	130	Ex 3	c.233dupT	p.Trp79LeufsX32	Frameshift	Compound	+/-	-/-
				Ex 14	c.2489dupA	p.Glu831GlyfsX 5	Frameshift	heterozygous	-/-	+/-
86	F	11.33	115	Ex 18	c.2909G>A	p.Gly970Asp	Missense	Compound	+/-	-/-
				Ex 18-20	c.2909-?_3367+? del	-	Large deletion	heterozygous	-/-	+/-

87	M	12.50	124	Ex 3	c.263T>G	p.Leu88X	Nonsense	Compound heterozygous	+/-	-/-
				Ex 11	c.1514delA	p.Asn505IlefsX2 2	Frameshift		-/-	+/-
88	F	5.67	95	Ex 4	c.374T>C	p.Ile125Thr	Missense	Compound heterozygous	+/-	-/-
				Ex 22	c.3659C>T	p.Thr1220Ile	Missense		-/-	+/-
89	M	4.83	152	Ex 10	c.1388G>A	p.Gly463Asp	Missense	Compound heterozygous	+/-	-/-
				In 16	c.2657+5G>A	-	Splicing		-/-	+/-
90	F	12.00	ND	Ex 18	c.2909G>A	p.Gly970Asp	Missense	Compound heterozygous	+/-	-/-
				Ex 20	c.233dupT	p.Trp79LeufsX32	Frameshift		-/-	+/-
91	F	5.42	ND	Ex 6	c.595C>T	p.His199Tyr	Missense	Compound heterozygous	+/-	-/-
				Ex 20	c.3176T>G	p.Leu1059X	Nonsense		-/-	+/-
92	F	3.17	ND	Ex 4	c.319_326delGC TTCCTA	p.Ala107X	Nonsense	Compound heterozygous	-/-	+/-
				Ex 11	c.1456G>T	p.Gly486X	Nonsense		+/-	-/-
93	F	7.58	112	Ex 14	c.2374C>T	p.Arg792X	Nonsense	Compound heterozygous	+/-	-/-
				Ex 4-11	c.274-?_1584+?d el	-	Large deletion		-/-	+/-

94	M	7.17	81	Ex 14	c.1772T>C	p.Val591Ala	Missense	Compound heterozygous	+/-	-/-
				Ex 4-11	c.274-?_1584+?del	-	Large deletion		-/-	+/-
95	M	0.50	ND	Ex 17	c.2812G>T	p.Val938Leu	Missense	Compound heterozygous	+/-	-/-
				Ex 18	c.2909G>A	p.Gly970Asp	Missense		-/-	+/-
96	F	3.75	ND	Ex 18	c.2909G>A	p.Gly970Asp	Missense	Compound heterozygous	+/-	-/-
				Ex 4	c.464C>G	p.Ala155Gly	Missense		-/-	+/-
97	M	0.58	ND	In 13	c.1766+2T>C	-	Splicing	Homozygous	+/-	+/-
98	F	6.00	ND	Ex 3	c.263T>G	p.Leu88X	Nonsense	Compound heterozygous	+/-	-/-
				In 13	c.1766+2T>C	-	Splicing		-/-	+/-
99	F	13.50	68	Ex 18	c.2909G>A	p.Gly970Asp	Missense	Compound heterozygous	+/-	-/-
				In 15	c.2619+2T>A	-	Splicing		-/-	+/-
100	M	0.42	103	Ex 8	c.1064C>G			Compound heterozygous	+/-	-/-
				In 9	c.1210-3C>G	-	Splicing		-/-	+/-

Novel variants are formatted in bold.

e-Table 2. Spectrum of *CFTR* variants detected in 202 Chinese patients with CF

No.	Region	Mutations	Amino acid change	Pathogenic significance	Reported/Novel	No. of alleles	Frequency, %
		(HGVS nomenclature)					
1 ^[4, 6, 10, 12, 16, 20, 24, 25, 27, 29, 36, 51]	Ex 18	c.2909G>A	p.Gly970Asp	CF-causing	Reported	47	12.11
2 ^[4, 6, 10, 32, 34, 40, 42, 43, 46-48, 50]	In 13	c.1766+5G>T	–	CF-causing	Reported (Chinese/Thai)	21	5.41

3 ^[9, 10, 12, 13, 34, 42, 45, 48]	Ex 12	c.1657C>T	p.Arg553X	CF-causing	Reported	14	3.61
4 ^[4, 10, 12]	Ex 4	c.293A>G	p.Gln98Arg	CF-causing	Reported	12	3.09
5 ^[6, 10, 42, 47, 48]	Ex 19	c.3068T>G	p.Ile1023Arg	Unknown	Reported (Chinese only)	11	2.84
6 ^[4, 6, 30, 37]	Ex 8	c.1000C>T	p.Arg334Trp	CF-causing	Reported	10	2.58
7 ^[4, 6, 20, 21]	Ex 3	c.263T>G	p.Leu88X	CF-causing	Reported	10	2.58
8 ^[4, 6, 10, 19, 27]	Ex 6	c.595C>T	p.His199Tyr	CF-causing	Reported	9	2.32

9 ^[4, 10, 26, 38]	Ex 20	c.3196C>T	p.Arg1066Cys	CF-causing	Reported	8	2.06
10 ^[6, 20]	Ex 11	c.1521_1523delCTT	p.Phe508del	CF-causing	Reported	7	1.80
11 ^[18, 31, 38]	Ex 6	c.650A>G	p.Glu217Gly	Unknown	Reported	6	1.55
12 ^[4, 12, 28]	3Ex 12	c.1666A>G	p.Ile556Val	Unknown	Reported	6	1.55
13 ^[4, 6]	Ex 3	c.223C>T	p.Arg75X	CF-causing	Reported	5	1.29
14 ^[33]	Ex 4	c.374T>C	p.Ile125Thr	Unknown	Reported	5	1.29
15 ^[6, 16]	In 9	c.1210-3C>G	–	Unknown	Reported (Chinese only)	4	1.03
16 ^[37]	Ex 11	c.1409T>A	p.Val470Glu	Unknown	Reported (Chinese only)	4	1.03
17 ^[17, 39]	Ex 14	c.2036G>A	p.Trp679X	Suspected	Reported	4	1.03
18 ^[38]	Ex 17	c.2834C>T	p.Ser945Leu	CF-causing	Reported	4	1.03

19 ^[12]	Ex 18-20	c.2909-?_3367+?del	–	Suspected	Reported	4	1.03
20 ^[27, 29, 38]	Ex 25	c.4056G>C	p.Gln1352His	Unknown	Reported	4	1.03
21 ^[10]	Ex 2-3	c.54-?_273+?del	–	Suspected	Reported	3	0.77
22	Ex 3	c.233dupT	p.Trp79LeufsX32	CF-causing	Reported	3	0.77
23 ^[6]	Ex 3	c.264_268delATATT	p.Leu88PhefsX21	Suspected	Reported	3	0.77
24 ^[4]	Ex 4	c.326A>G	p.Tyr109Cys	Unknown	Reported	3	0.77
25 ^[6, 24, 29]	In 8	c.1116+1G>A	–	CF-causing	Reported	3	0.77
26 ^[10, 20]	Ex 14	c.2125C>T	p.Arg709X	CF-causing	Reported	3	0.77
27 ^[4, 20]	Ex 14	c.2374C>T	p.Arg792X	CF-causing	Reported	3	0.77
28 ^[6, 10, 20]	Ex 22	c.3635delT	p.Val1212AlafsX1 6	Suspected	Reported (Chinese only)	3	0.77
29 ^[40]	Promoter	c.-152G>C	–	Unknown	Reported	2	0.52
30 ^[26]	Ex 1	c.3G>A	p.Met1Ile	Unknown	Reported	2	0.52

31 ^[12, 33]	Ex 2	c.95T>C	p.Leu32Pro	Unknown	Reported	2	0.52
32	Ex 4-11	c.274-?_1584+?del	–	Suspected	Reported	2	0.52
33	Ex 4	c.298C>T	p.Leu100Phe	Unknown	Novel	2	0.52
34 ^[51]	Ex 4	c.319_326delGCTTCCTA	p.Ala107X	Suspected	Reported	2	0.52
35 ^[10]	Ex 4	c.325T>G	p.Tyr109Asp	Unknown	Reported (Chinese only)	2	0.52
36 ^[36]	Ex 4	c.380T>G	p.Leu127X	Suspected	Reported	2	0.52
37 ^[4]	Ex 4	c.413_415dupTAC	p.Leu138dup	CF-causing	Reported	2	0.52
38 ^[6]	Ex 5	c.532G>A	p.Gly178Arg	CF-causing	Reported	2	0.52
39 ^[4, 12]	Ex 5	c.558C>G	p.Asn186Lys	Unknown	Reported (Chinese only)	2	0.52
40 ^[10]	Ex 6	c.607A>T	p.Ile203Phe	Unknown	Reported (Chinese only)	2	0.52
41 ^[4]	Ex 6	c.648G>A	p.Trp216X	Suspected	Reported (Chinese only)	2	0.52

42 ^[4, 20]	Ex 8	c.960dupA	p.Ser321IlefsX43	Suspected	Reported (Chinese only)	2	0.52
43 ^[35]	Ex 10	c.1369G>C	p.Ala457Pro	Unknown	Reported	2	0.52
44 ^[10]	Ex 10	c.1388G>A	p.Gly463Asp	Unknown	Reported	2	0.52
45	Ex 11	c.1393-?_1584+?del	–	Suspected	Novel	2	0.52
46 ^[26]	Ex 11	c.1572C>A	p.Cys524X	CF-causing	Reported	2	0.52
47 ^[10, 12]	In 12	c.1679+2T>C	–	Unknown	Reported	2	0.52
48	Ex 13	c.1703T>A	p.Leu568X	Suspected	Reported	2	0.52
49	Ex 14	c.1810A>C	p.Thr604Pro	Unknown	Novel	2	0.52
50 ^[46, 43]	Ex 14	c.2083dupG	p.Glu695GlyfsX35	Suspected	Reported (Chinese only)	2	0.52
51 ^[19, 25]	Ex 14	c.2290C>T	p.Arg764X	CF-causing	Reported	2	0.52
52 ^[4]	In 14	c.2491-126T>C	–	Unknown	Reported (Chinese only)	2	0.52

53 ^[10, 38]	Ex 15	c.2547C>A	p.Tyr849X	CF-causing	Reported	2	0.52
54 ^[38]	Ex 15	c.2551C>T	p.Arg851X	CF-causing	Reported	2	0.52
55 ^[10, 12]	In 16	c.2658-1G>C	–	CF-causing	Reported	2	0.52
56 ^[46, 43]	Ex 17	c.2684G>A	p.Ser895Asn	Unknown	Reported (Chinese only)	2	0.52
57 ^[29, 38]	Ex 19	c.3062C>T	p.Pro1021Leu	Unknown	Reported (Chinese only)	2	0.52
58 ^[48]	In 19	c.3140-26A>G	–	CF-causing	Reported	2	0.52
59 ^[23]	Ex 20	c.3140- 454_c.3367+249del193ins 13	–	Suspected	Reported (Chinese only)	2	0.52
60 ^[10]	Ex 20	c.3196C>A	p.Arg1066Ser	Unknown	Reported	2	0.52
61	Ex 22	c.3484C>T	p.Arg1162X	CF-causing	Reported	2	0.52
62 ^[4, 20]	Ex 22	c.3700A>G	p.Ile1234Val	CF-causing	Reported	2	0.52

63	In 22	c.3717+10kbC>T	–	Unknown	Reported	2	0.52
64	In 22	c.3718-2477C>T	–	CF-causing	Reported	2	0.52
65	In 23	c.3874-4522A>G	–	Unknown	Reported	2	0.52
66 ^[36]	3'UTR	c.*133delT	–	Unknown	Reported	2	0.52
67 ^[44]	Ex 1	c.19G>T	p.Glu7X	Suspected	Reported (Chinese only)	1	0.26
68 ^[14]	Ex 2	c.54_164del30bp	–	Suspected	Reported (Chinese only)	1	0.26
69	In 2	c.164+2T>C	–	CF-causing	Reported	1	0.26
70 ^[18]	Ex 3	c.214G>A	p.Ala72Thr	Unknown	Reported	1	0.26
71	Ex 3	c.222delG	p.Arg75AspfsX16	Suspected	Novel	1	0.26
72 ^[22]	Ex 3	c.271G>A	p.Gly91Arg	CF-causing	Reported	1	0.26
73 ^[21]	Ex 4-6	c.274-?_743+?del	–	Suspected	Reported	1	0.26
74 ^[35]	Ex 4	c.320C>A	p.Ala107Asp	Unknown	Reported (Chinese only)	1	0.26
75	Ex 4	c.350G>A	p.Arg117His	CF-causing	Reported	1	0.26
76 ^[10]	Ex 4	c.405_406dupAC	p.Leu136HisfsX18	Suspected	Reported	1	0.26

77	Ex 4	c.464C>G	p.Ala155Gly	Unknown	Novel	1	0.26
78 ^[15]	Ex 5	c.567C>A	p.Asn189Lys	Unknown	Reported (Chinese only)	1	0.26
79 ^[6]	In 5	c.579+1_579+2insACAT	–	Unknown	Reported (Chinese only)	1	0.26
80 ^[30]	In 5	c.580-1G>T	–	CF-causing	Reported	1	0.26
81 ^[12]	Ex 7-11	c.744-?_1584+ ?del	–	Suspected	Reported (Chinese only)	1	0.26
82 ^[41]	Ex 7	c.753_754delAG	p.Arg251SerfsX6	Suspected	Reported (Chinese only)	1	0.26
83 ^[44]	Ex 7	c.860dupA	p.Asn287LysfsX21	Suspected	Reported (Chinese only)	1	0.26
84 ^[38]	Ex 7	c.861C>G	p.Asn287Lys	Unknown	Reported	1	0.26
85 ^[11]	Ex 7	c.865A>T	p.Arg289X	Suspected	Reported (Chinese only)	1	0.26
86 ^[48]	Ex 7	c.868C>T	p.Gln290X	CF-causing	Reported	1	0.26
87 ^[26]	In 7	c.870-1G>C	–	Unknown	Reported	1	0.26

88 ^[38]	Ex 8	c.884delT	p.Leu295ArgfsX8	Suspected	Reported (Chinese only)	1	0.26
89	Ex 8	c.940G>T	p.Gly314Trp	Unknown	Novel	1	0.26
90 ^[29]	Ex 8	c.1040G>A	p.Arg347His	CF-causing	Reported	1	0.26
91	Ex 8	c.1064C>G	p.Pro355Arg	Unknown	Novel	1	0.26
92 ^[4]	Ex 8	c.1075C>T	p.Gln359X	Suspected	Reported (Chinese only)	1	0.26
93 ^[27]	In 8	c.1117-1G>C	–	Unknown	Reported (Chinese only)	1	0.26
94	Ex 10	c.1219G>T	p.Glu407X	Suspected	Novel	1	0.26
95	Ex 10	c.1231A>G	p.Lys411Glu	Unknown	Reported	1	0.26
96^[41]	Ex 10	c.1240C>T	p.Gln414X	CF-causing	Reported	1	0.26
97	Ex 10	c.1265C>T	p.Ser422Phe	Unknown	Novel	1	0.26
98	Ex 10	c.1347_1350delAGAA	p.Arg450AspfsX1 8	Suspected	Novel	1	0.26
99 ^[38]	Ex 10	c.1352G>T	p.Gly451Val	Unknown	Reported	1	0.26
100	Ex 10	c.1368delT	p.Ala457LeufsX12	Suspected	Novel	1	0.26

101 ^[22]	Ex 10	c.1373G>A	p.Gly458Glu	Unknown	Reported (Chinese only)	1	0.26
102 ^[38]	In 10	c.1393-4C>A	–	Unknown	Reported (Chinese only)	1	0.26
103 ^[13]	Ex 11	c. 1423delC	p.Leu475TrpfsX52	Suspected	Reported (Chinese only)	1	0.26
104 ^[6]	Ex 11	c.1429_1437delCCTTCA GAG	p.Pro477_Glu479d el	Unknown	Reported (Chinese only)	1	0.26
105	Ex 11	c.1456G>T	p.Gly486X	Suspected	Reported	1	0.26
106	Ex 11	c.1514delA	p.Asn505IlefsX22	Suspected	Novel	1	0.26
107	Ex 11	c.1523_1534delTTGGTG TTTCCT	p.Phe508_Ser511d el	Suspected	Novel	1	0.26
108 ^[29]	Ex 11	c.1526G>C	p.Gly509Ala	Unknown	Reported (Chinese only)	1	0.26
109 ^[4]	Ex 13	c.1699G>T	p.Asp567Tyr	Unknown	Reported (Chinese only)	1	0.26
110 ^[10]	Ex 13	c.1716C>A	p.Asp572Glu	Unknown	Reported	1	0.26

					(Chinese only)		
111 ^[6]	Ex 13	c.1733T>C	p.Leu578Pro	Unknown	Reported (Chinese only)	1	0.26
112 ^[49]	In 13	c.1766+1G>T	–	CF-causing	Reported	1	0.26
113	In 13	c.1766+2T>C	–	Unknown	Reported	1	0.26
114	Ex 14	c.1772T>C	p.Val591Ala	Unknown	Novel	1	0.26
115 ^[20]	Ex 14	c.1997T>G	p.Leu666X	Suspected	Reported (Chinese only)	1	0.26
116	Ex 14	c.2042A>T	p.Glu681Val	Unknown	Novel	1	0.26
117 ^[38]	Ex 14	c.2052delA	p.Lys684AsnfsX38	CF-causing	Reported	1	0.26
118 ^[12]	Ex 14	c.2052dupA	p.Gln685ThrfsX4	CF-causing	Reported	1	0.26
119	Ex 14	c.2058_2061delTT	p.Phe687X	Suspected	Novel	1	0.26
120 ^[6]	Ex 14	c.2236_2246delGAGGCG ATACTinsAAAAATC	p.Glu746LysfsX8	Suspected	Reported (Chinese only)	1	0.26
121	Ex 14	c.2328dupA	p.Val777SerfsX2	Suspected	Novel	1	0.26
122 ^[10]	Ex 14	c.2353C>T	p.Arg785X	CF-causing	Reported	1	0.26
123	Ex 14	c.2475_2478dupCGAA	p.Glu827ArgfsX10	Suspected	Reported	1	0.26

					(Chinese only)		
124	Ex 14	c.2489dupA	p.Glu831GlyfsX5	Suspected	Novel	1	0.26
125	In 14	c.2491-2A>G	–	Unknown	Reported	1	0.26
126	In 15	c.2619+2T>A	–	Unknown	Reported	1	0.26
127	In 16	c.2657+5G>A	–	CF-causing	Reported	1	0.26
128 ^[32]	Ex 17	c.2805delA	p.Pro936HisfsX6	Suspected	Reported (Chinese only)	1	0.26
129	Ex 17	c.2812G>T	p.Val938Leu	Unknown	Reported	1	0.26
130 ^[20]	Ex 17	c.2907A>C	p.A969A	Unknown	Reported (Chinese only)	1	0.26
131	Ex 18-21	c.2909-?_3468+?del	–	Suspected	Novel	1	0.26
132	Ex 18	c.2936A>C	p.Asp979Ala	Unknown	Reported	1	0.26
133	Ex 18	c.2977G>T	p.Asp993Tyr	Unknown	Reported	1	0.26
134 ^[17]	In 18	c.2988+2T>C	–	Unknown	Reported (Chinese only)	1	0.26
135 ^[10]	Ex 19	c.2997_3000delAATT	p.Ile1000X	Suspected	Reported	1	0.26
136 ^[38]	Ex 19	c.3123dupA	p.Gln1042ThrfsX5	Suspected	Reported	1	0.26

137	Ex 20	c.3140-?_3367+?del	–	Suspected	Novel	1	0.26
138	Ex 20	c.3176T>G	p.Leu1059X	Suspected	Reported	1	0.26
139 ^[40]	Ex 20	c.3197G>A	p.Arg1066His	CF-causing	Reported	1	0.26
140	Ex 20	c.3209G>C	p.Arg1070Pro	Unknown	Reported	1	0.26
141 ^[4]	Ex 20	c.3307delA	p.Ile1103X	Suspected	Reported (Chinese only)	1	0.26
142	Ex 21	c.3387delT	p.Gly1130ValfsX4	Suspected	Reported	1	0.26
143 ^[18]	Ex 21	c.3406G>A	p.Ala1136Thr	Unknown	Reported	1	0.26
144	In 21	c.3469-12T>G	–	Unknown	Novel	1	0.26
145	In 21	c.3469-2A>T	–	Unknown	Novel	1	0.26
146	Ex 22	c.3472C>T	p.Arg1158X	CF-causing	Reported	1	0.26
147	Ex 22	c.3476C>T	p.Ser1159Phe	CF-causing	Reported	1	0.26
148 ^[11]	Ex 22	c.3653_3656dupAATA	p.Tyr1219X	Suspected	Reported (Chinese only)	1	0.26
149	Ex 22	c.3659C>T	p.Thr1220Ile	Unknown	Novel	1	0.26
150 ^[15]	Ex 22	c.3691delT	p.Ser1231ProfsX4	CF-causing	Reported	1	0.26
151 ^[6]	Ex 23	c.3718-?_3873+?del	–	Suspected	Reported	1	0.26

					(Chinese only)		
152	Ex 23	c.3841C>T	p.Gln1281X	Suspected	Reported	1	0.26
153 ^[6]	Ex 23	c.3860delG	p.Gly1287GlufsX2	Suspected	Reported (Chinese only)	1	0.26
154 ^[10]	Ex 24	c.3883_3886delATTT	p.Ile1295PhefsX32	CF-causing	Reported	1	0.26
155	Ex 24	c.3908delA	p.Asn1303ThrfsX2 5	CF-causing	Reported	1	0.26
156 ^[4]	Ex 24	c.3909C>G	p.Asn1303Lys	CF-causing	Reported	1	0.26
157 ^[6]	In 24	c.3964-7A>G	–	Unknown	Reported (Chinese only)	1	0.26
158 ^[4]	3'UTR	c.*110C>G	–	Unknown	Reported (Chinese only)	1	0.26

HGVS, Human Genome Variation Society.

Novel variants are formatted in bold.

e-Table 3. Demographic and clinical features of 24 Chinese CF patients with novel variants

Case No.	Sex	Age at Dx, y	Age of symptom onset, y	Weight at Dx, kg (centile)	Clinical presentation	Sputum pathogens	Fecal Sudan III stain	Sweat conductivity, mmol/L	Genotype	Distribution (Province)
33	M	7.33	7.00	18.0 (<3rd)	Bronchiectasis/Recurrent pneumonia/ABPA/Sinusitis/Steatorrhea/Liver disease/FTT/Finger clubbing	<i>P. aeruginosa/S. aureus</i>	+	125	c.2328dupA/c.3209G>C	Hebei
36	F	10.67	5.00	37.5 (50-75th)	Bronchiectasis/Recurrent pneumonia/Sinusitis/Liver disease/Diabetes/Finger clubbing	<i>H. influenzae/P. aeruginosa/S. aureus</i>	–	127	c.2058_2061delT T/c.595C>T	Shanxi

38	M	13.25	7.00	30.0 7 (<3rd)	Bronchiectasis/Recurrent pneumonia/ABPA/Asthma/Hemoptysis/Sinusitis/Nasal polyps/FTT/Pancreatitis/Finger clubbing	<i>P. aeruginosa/S. aureus/A. fumigatus/A. niger</i>	+	109	c.1347_1350delA GAA/c.2909G>A	Zhejiang
46	M	10.17	7.00	27.0 (10th)	Bronchiectasis/ Asthma/Sinusitis	–	–	60	c.2042A>T/c.374 T>C	Yunnan
54	F	8.58	5.00	27.0 (25- 50th)	Bronchiectasis/Sinusitis/Nasal polyps	<i>P. aeruginosa</i>	–	125	c.1810A>C/c.165 7C>T	Shanghai
56	M	2.08	1.00	9.5 (<3rd)	Bronchiectasis/Recurrent pneumonia/Sinusitis/Steatorrhea/FTT/Finger clubbing	<i>P. aeruginosa</i>	+	111	c.3469-2A>T/c.1766+2T >C	Henan
57	F	9.42	5.00	26.0 (10- 25th)	Bronchiectasis/Recurrent pneumonia/Sinusitis/Pancreatitis/Finger clubbing	<i>P. aeruginosa/A. fumigatus/A. flavus</i>	–	108	c.1368delT/c.387 4-4522A>G	Yunnan
58	F	8.75	2.00	19.0	Bronchiectasis/Recurrent	<i>P. aeruginosa/C.</i>	+	136	c.1393-?_1584+?	Henan

				(<3rd)	pneumonia/Sinusitis/Steatorrhea/Rectal prolapse/Liver disease/FTT/Finger clubbing	<i>Albicans</i>			del/c.1523_1534delTTGGTGTTT CCT	
59	F	7.75	0.75	21.0 (10-25th)	Bronchiectasis/Recurrent pneumonia/ABPA/Sinusitis/Finger clubbing	<i>P. aeruginosa/S. maltophilia</i>	-	97	c.1265C>T/c.2977G>T/c.3140-26A>G	Shandong
61	F	4.17	1.00	12.0 (<3rd)	Bronchiectasis/Recurrent pneumonia/Sinusitis/Meconium ileus/Steatorrhea/Rectal prolapse/Liver disease/FTT/Finger clubbing	<i>P. aeruginosa/S. aureus/C. lusitaniae</i>	+	ND	c.1219G>T/c.2551C>T/c.-152G>C	Zhejiang
63	F	13.75	5.00	29.0 (<3rd)	Bronchiectasis/Recurrent pneumonia/Sinusitis/Pancreatitis/Steatorrhea/FTT/Finger clubbing	<i>P. aeruginosa/A. flavus</i>	+	ND	c.1810A>C/c.940G>T/c.*133delT	Zhejiang

67	F	12.00	9.00	47.6 (75-90th)	Bronchiectasis/Sinusitis/ABPA/Wheeze	<i>P. aeruginosa/A. terreus</i>	-	77	c.298C>T/c.3841C>T	Anhui
71	F	13.25	8.00	35.0 (3rd-10th)	Bronchiectasis/Sinusitis/FTT/Finger clubbing	<i>P. aeruginosa/C. Albicans</i>	-	122	c.3140-?_3367+?del/c.233dupT	Shandong
72	F	7.33	1.00	19.0 (3rd-10th)	Bronchiectasis/Sinusitis/FTT	<i>K. pneumoniae/S. aureus</i>	-	118	c.1393-?_1584+?del/c.1000C>T	Shanxi
75	F	8.58	0.58	19.0 (<3rd)	Bronchiectasis/Recurrent pneumonia/ABPA/Sinusitis/Steatorrhea/Diabetes/FTT/Finger clubbing	<i>P. aeruginosa/A. fumigatus/C. Albicans</i>	+	146	c.222delG/c.263T>G	Jiangxi
77	F	4.50	4.00	19.0 (75th)	Atelectasis/Asthma/Sinusitis	-	-	60	c.298C>T/c.650A>G	Jiangsu
78	F	12.92	10.00	44.0 (50th)	Bronchiectasis/Hemoptysis/Sinusitis/Nasal polyps/Finger clubbing	<i>P. aeruginosa/M. abscessus/A. terreus/C. Albicans</i>	-	119	c.3469-12T>G/c.3476C>T	Liaoning

80	F	10.67	6.00	33.0 (25-50th)	Bronchiectasis/ABPA/Asthma/Sinusitis/Pancreatitis	<i>P. aeruginosa/S. aureus</i>	–	126	c.2909-?_3468+?del/c.2909G>A	Henan
85	F	11.00	10.00	30.5 (10-25th)	Bronchiectasis/Sinusitis/Appendicitis	<i>P. aeruginosa</i>	–	130	c.2489dupA/c.233dupT	Anhui
87	M	12.50	0.17	31.0 (3rd)	Bronchiectasis/Recurrent pneumonia/Sinusitis/Steatorrhea/Liver disease/FTT/Finger clubbing	<i>S. aureus/C. Albicans</i>	+	124	c.1514delA/c.263T>G	Sichuan
88	F	5.67	0.25	23.0 (90th)	Bronchiectasis/Asthma/Sinusitis	<i>S. pneumoniae/C. Albicans</i>	–	95	c.3659C>T/c.374T>C	Henan
94	M	7.17	6.00	19.0 (3rd)	Bronchiectasis/Sinusitis/FTT/Finger clubbing	<i>P. aeruginosa/C. Albicans</i>	–	ND	c.1772T>C/c.274-?_1584+?del)	Henan
96	F	3.75	3.00	13.0 (3rd-10th)	Bronchiectasis/ABPA/Sinusitis/FTT	–	–	ND	c.464C>G/c.2909G>A	Henan
102	M	0.42	0.08	8.0 (50th)	Pseudo-Bartter syndrome	–	–	103	c.1064C>G/c.1210-3C>G	Henan

Abbreviations: *A. fumigatus* = *Aspergillus fumigatus*; *A. flavus* = *Aspergillus flavus*; *A. niger* = *Aspergillus niger*; *A. terreus* = *Aspergillus terreus*; *C. Albicans* = *Candida Albicans*; *C. lusitaniae* = *Candida lusitaniae*; *H. influenzae* = *Haemophilus influenzae*; *K. pneumonia* = *Klebsiella pneumoniae*; *M. abscessus* = *Mycobacterium abscessus*; *MRSA* = *Methicillin-resistant Staphylococcus aureus*; *P. aeruginosa* = *Pseudomonas aeruginosa*; *S. aureus* = *Staphylococcus aureus*; *S. maltophilia* = *Stenotrophomonas maltophilia*; *S. pneumoniae* = *Streptococcus pneumoniae*; Dx = Diagnosis; FTT = Failure to thrive; ND = Not done; Y = Years.

Novel variants are formatted in bold.