CFTR mutations and IVS8-5T variant in newborns with hypertrypsinaemia and normal sweat test

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Abstract

Neonates positive for immunoreactive trypsinogen assay (IRT) and negative for sweat test have formerly been found to carry the major cystic fibrosis (CF) mutation, $\Delta F508$, much more frequently than the general population. Among the 716 IRT positive newborns detected by a three tier (IRT, mutation analysis plus meconium lactase assay, sweat test) CF screening programme in north eastern Italy during the period January 1993 to March 1996, we found 45 carriers, a number significantly higher than the expected 17 ($p < 0.001$). We speculated that some of these heterozygotes could actually be affected by a very mild form of CF, and carry on the other chromosome an undetected CFTR mutation or a DNA variant, such as the 5-thymidine allele in intron 8 of the CFTR gene (IVS8-5T). This hypothesis was tested in four samples: group A (the 45 carriers mentioned above), group B (51 non-carrier, IRT positive neonates), group C (50 IRT negative neonates), and group D (90 CF adult female carriers). Chromosomes with IVS8-5T were seven (7.78%) in group A, seven (6.86%) in group B, five (5%) in group C, and four in group D (2.22%). The 5T prevalence in group A was significantly higher ($p < 0.05$) compared to group D; similarly, a higher ($p < 0.05$) 5T frequency in group A compared to group C was detected by considering the chromosomes free from CFTR mutations. This study is consistent with previous papers in finding among neonates with high trypsin levels a CF carrier frequency significantly higher than that expected. It is also suggested that in at least some babies raised trypsin levels at birth could be a phenotypic expression of compound heterozygosity for a major CF mutation plus IVS8-5T.

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Measurement of immunoreactive trypsinogen concentration (IRT) in dried blood spots is now the most common technique for cystic fibrosis (CF) neonatal screening.¹ Since a considerable number of unaffected newborns show raised IRT levels, several laboratories have chosen to improve the screening specificity by

introducing mutation analysis for infants with hypertrypsinaemia. As a consequence of genetic testing, a few CF carriers are detected, and in the past unaffected IRT positive babies have been unexpectedly found to carry the major CF mutation, $\Delta F508$, much more frequently than the general population.^{2,3} It has therefore been postulated^{2,3} that IRT could indirectly detect some heterozygotes.

An alternative explanation could be that at least some of these babies are actually affected, and carry on the other chromosome a mild mutation or a DNA variant, associated with few symptoms and normal sweat chloride values. A possible candidate for this role is a DNA polymorphic sequence of five thymines in intron 8 of the cystic fibrosis transmembrane conductance regulator (CFTR) gene. The 5-thymidine allele (IVS8-5T), in contrast to the more common 7T and 9T variants, is responsible for low levels of normal CFTR mRNA, and has been very frequently found in men with a form of infertility called congenital bilateral absence of the vas deferens (CBAVD), a condition which has been suggested to be a primarily genital form of CF.⁴⁻⁶

We report here data obtained in the genetic screening of some frequent CFTR mutations plus the 5T allele in 96 IRT positive neonates.

Materials and methods

SCREENING STRATEGY

The results of CF neonatal screening in north eastern Italy from January 1993 to March 1996 were reviewed. The screening strategy used a three tier system, whose progressive steps were immunoreactive trypsinogen (IRT), mutation analysis with complementary meconium lactase determination, and sweat test. IRT levels were measured in dried blood spot specimens from neonates born in the Veneto and Trentino-Alto Adige regions. An immunoreactive trypsinogen concentration of 100 μg trypsin/l whole blood, lowered in May 1995 to 95 μg /l (equal to the 99.5th centile for an unselected neonatal population of 7223), was chosen as the cut off point. At or above this value, mutation analysis and meconium lactase determination were performed. Initially we tested for $\Delta F508$, R1162X, and N1303K, estimated by a cohort study⁷ to cover 61% of CF chromosomes in our area; from March 1995 10 other mutations were included (2183AAG, 3849+10KbCT, G542X, 1717-1GA, R553X, Q552X, G85E, 711+5GA, 3132delTG,

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2789+5GA), thus covering 85% of CF mutations.⁷

Diagnosis was immediately established in neonates, either homozygous or compound heterozygous, who were referred directly for clinical management of CF. If either one mutation tested positive or meconium lactase was above the cut off level of 0.5 U/l, a sweat test, which is regarded as the definitive test for CF, was immediately performed on the infant, thus discriminating between affected and unaffected subjects. In neonates with borderline chloride levels, sweat test was repeated in order to achieve either clearly positive or negative results. Sweat test was always associated with clinical evaluation, and frequently also with IRT retesting and fecal chymotrypsin determination.

The families of carriers detected incidentally were offered genetic counselling and gene analysis.

POPULATION UNDER STUDY

The CFTR gene variants of T tract length of intron 8 were determined in 236 subjects, all from the same geographical area, who were divided into four groups. Group A: 45 neonates who were IRT positive, carried one of the mutations screened for, but whose sweat test result was normal; group B: 51 neonates with high trypsin levels (>95 g/l) but no mutation detected; group C: 50 neonates with low trypsin levels, who were supposed to be representative of the general population polyT sequence distribution pattern; group D: 90 adults, who carried the same CFTR mutations as newborns in group A (every baby was matched to two adults with the same mutation), chosen from among first or second degree relatives of 90 unrelated CF patients. Only female relations were included in the last group, as the 5T allele is suggested to be significantly less frequent in the chromosomes of fathers of patients with CF, and therefore in male carriers, than in the general population.⁴

A plan for follow up of infants belonging to the first group and found to carry the 5T allele is being developed.

ANALYTICAL PROCEDURES

Immunoreactive trypsin was assayed by the Pharmacia standardised Delfia Neonatal IRT kit.⁸ Meconium lactase activity was determined as previously described.⁹ Dried blood spots on Guthrie cards were the sources of DNA. The procedure used to prepare DNA for PCR, based on those described by Singer-Sam *et al*¹⁰ and Walsh *et al*¹¹ for forensic material, is very simple and requires only boiling cells in a 5% suspension of chelating resin (Sigma, St Louis, MO).

During the period January 1993 to February 1995 mutations were analysed by restriction enzymes as previously described,¹² or by heteroduplex analysis (for $\Delta F508$) according to Rommens *et al*¹³; from March 1995 we switched to a reverse dot blot assay.¹⁴ The intron 8 polyT sequence was analysed with the nested PCR method according to Chillon *et al*.⁴ The confirmatory sweat electrolyte analysis

was performed at least twice by the classical method of Gibson and Cooke.¹⁵ Chloride levels above 60 mEq/kg were considered as positive, between 40 and 60 mEq/kg as borderline, and below 40 mEq/kg as negative, provided that the amount of sweat collected was at least 50 mg for a stimulation/collection skin area of 3.2×3.2 cm.

Comparison of the detected and expected carrier frequencies was performed by normal approximation to binomial distribution. The groups under study were analysed using Fisher's exact test.

Results

A total of 154 637 newborns were screened, of whom 716 were IRT positive (0.46%). Among these we found 58 affected subjects (median chloride 99 mEq/kg, range 61-121; observed CF incidence 1/2666) and 45 carriers (median chloride 20 mEq/kg, range 6-39). Two babies, both carrying a CFTR mutation but not the 5T variant, showed repeatedly borderline sweat chloride levels: they are being closely followed, but so far we cannot say whether they are affected or not and they have not been included in our figures.

The carrier frequency of 1/15 (29/429) observed when we tested for three mutations is significantly higher ($p < 0.001$) than the expected 1/44 for the same mutations. A significant difference ($p < 0.001$) was also found after we decided to screen for 13 mutations, when the observed carrier frequency was 1/14 (16/227) against the expected 1/32.⁷ By combining the data from the two periods, the expected number of carriers in a population of 656 (429+227) would be 17, a figure which is again much lower than the detected number of 45 ($p < 0.001$).

The 5T allele was found in 7/45 (15%) IRT positive neonates carrying a CF mutation, in 7/51 (13%) IRT positive neonates not carrying any of the sought CF mutations, in 5/50 (10%) IRT negative neonates, and in 4/90 (4%) adult carriers (table 1). The 5T incidence in groups C and D is similar to that previously reported in control populations from different geographical areas.^{4 16-18} The frequency of the 5T allele gradually decreased from group A to group D, with a significant difference between A and D ($p = 0.0459$). In the seven babies who carried both IVS8-5T and a CF mutation, the latter was always $\Delta F508$; in the four adults the mutations were $\Delta F508$ (twice), N1303K, and R1162X.

Table 2 shows IRT and meconium lactase, plus CFTR and polyT genotypes found in group A. Sweat chloride values and concomitant IRT and weight Z score are also included. In all but one (No 26) cases in which data were available, blood trypsinogen levels showed a time related decrease not consistent with CF. No definite failure to thrive or lung disease symptoms were found at the time of sweat test in any of the subjects who could be clinically evaluated, with the exception of subject 5, a severely premature baby. Both subjects 43 and 45 underwent surgery, the former because of a

Table 1 Frequencies of the polyT alleles in intron 8 of CFTR in the groups under study

	Description	5T allele (%)	7T allele (%)	9T allele (%)
Group A	Newborns IRT positive at birth 1 CFTR mutation detected	7/90 (7.78)*	44/90 (48.89)	39/90 (43.33)
Group B	Newborns IRT positive at birth No CFTR mutation detected	7/102 (6.86)	81/102 (79.41)	14/102 (13.73)
Group C	Newborns IRT negative at birth	5/100 (5)	85/100 (85)	10/100 (10)
Group D	Adults (female CF relatives) 1 CFTR mutation detected	4/180 (2.22)*	97/180 (53.89)	79/180 (43.89)

*A v D: p=0.0459.

congenital heart defect, the latter for Hirschsprung’s disease.

Discussion

The raised IRT levels in infants with CF are thought to be the result of “leakage” of trypsinogen into the bloodstream, derived from functioning, but ductally obstructed, pancre-

atic tissue.¹⁹ Both pancreatic insufficient and, to a lesser extent, pancreatic sufficient subjects with CF can be identified by the IRT assay.^{20 21} However, neonatal hypertrypsinaemia also occurs in many unaffected infants.^{1 9 22} High IRT levels in seemingly normal babies have not yet been fully explained: even though some pathological conditions such as perinatal asphyxia seem to cause false positive screening results,²³ many newborns have no real cause for their hypertrypsinaemia. This study is consistent with previous papers from other authors^{2 3} in finding among neonates with high trypsin levels a CF carrier frequency significantly higher than the expected one: testing for three mutations one newborn in every 15 was a carrier, instead of the anticipated 1/44. When we expanded the analysis to 13 mutations, again one newborn in every 14 was found to be a carrier, against the expected 1/32.⁷ Considering the two mutation analysis periods as a whole, there is again a significant difference between the 17 expected and the 45 carriers actually found. The explanation for such a peculiar phenomenon is a matter for conjecture. One possibility could be that neonatal IRT positivity is capable of indirectly detecting some heterozygotes who

Table 2 Screening and genotype data in group A. The IRT retest and weight Z score are reported when available, and refer to a mean age of 38 days (range 22-90). Normal values for IRT and IRT retest were considered respectively as below 100 (95 from May 1995 onwards) and 75 µg/l. Meconium lactase threshold was 0.5 U/g. Bold type is used for neonates carrying the 5T variant.

Subject	Sex	IRT (µg/l)	Meconium lactase (U/g)	Sweat chloride (mEq/kg)	CFTR mutation	PolyT genotype	IRT retest (µg/l)	Weight Z score
1	F	134	0	6	ΔF508	7/9	67	1.47
2	M	95	0	28	ΔF508	7/9		
3	F	408	0	7	2183AA→G	7/7	14	-0.29
4	M	150	0.72	16	N1303K	7/9	19	-0.47
5	F	106	Unknown	18	R1162X	7/7		-6.38
6	M	131	0	22	N1303K	9/9	27	-0.54
7	M	106	0	21	ΔF508	7/9	34	0.11
8	F	100	0	15	ΔF508	5/9	37	-0.01
9	F	105	0	25	ΔF508	5/9	51	-0.15
10	F	100	0	24	R1162X	7/7		
11	M	266	0	14	ΔF508	7/9	5	0.20
12	F	103	0	9	ΔF508	7/9		
13	F	105	0	32	ΔF508	7/9		
14	F	268	0	22	ΔF508	7/9	30	
15	M	110	Unknown	33	R1162X	7/7		
16	M	174	0	12	ΔF508	7/9	52	1.87
17	F	100	0	15	ΔF508	7/9		
18	M	140	0	9	ΔF508	7/9		
19	M	98	0	30	ΔF508	5/9		
20	M	110	1.2	10	ΔF508	7/9	11	0.26
21	F	102	0	20	G542X	7/9	45	0.77
22	F	111	0	16	N1303K	7/9		
23	F	100	0	16	ΔF508	7/9	54	-0.44
24	F	95	0	18	R553X	7/9		
25	F	285	0	16	ΔF508	7/9	20	0.28
26	M	117	0	23	ΔF508	7/9	101	
27	M	115	0	24	ΔF508	5/9	12	-0.4
28	F	236	0	8	ΔF508	7/9	21	0.26
29	M	192	0	19	N1303K	7/9	73	0.04
30	M	103	0	39	ΔF508	5/9	69	
31	M	133	0	10	ΔF508	7/9	65	1.17
32	M	144	0	30	R1162X	7/7	74	-0.83
33	F	123	0	20	ΔF508	7/9	66	-0.02
34	F	100	Unknown	30	ΔF508	5/9	65	
35	M	134	0	28	ΔF508	7/9		
36	M	294	0.8	12	R553X	7/7		
37	M	102	Unknown	32	2789+5G→A	7/7		
38	F	114	0	36	ΔF508	7/9		
39	M	123	0	9	R1162X	7/7	51	-0.36
40	F	118	0	33	ΔF508	5/9	16	0.18
41	M	134	0	10	ΔF508	7/9	15	1.36
42	M	97	0	9	ΔF508	7/9		
43	M	98	1.71	31	ΔF508	7/9		
44	F	122	Unknown	23	ΔF508	7/9		
45	F	101	0	23	ΔF508	9/9		

show mild biological abnormalities as early as the first days of life. However, carriers have always been considered to be clinically and biologically asymptomatic, and it is difficult to explain why some of them show this feature while others do not.

Alternatively, either the carrier frequency in the general population could be more than the calculated one, or the relative frequency of the tested mutations could be higher. Both options are unlikely, the former because such a huge number of carriers would be consistent with an incidence of CF much greater than that currently observed, the latter because we have previously shown the distribution of CF mutations in our population by characterising more than 90% of the mutated alleles from a cohort of 225 CF chromosomes.⁷

Finally, one could postulate that at least some of these newborns carry an undetectable mild mutation or a DNA variant on the other chromosome, associated with normal sweat chloride values, but capable of raising trypsin levels.

A polymorphic repeat which could play this role is the 5-thymidine allele in intron 8. The proportion of the exon 9 transcript is inversely related to the length of the poly-thymidine tract in the sequence of the acceptor splice site of intron 8, the 5T allele causing a high proportion of an abnormal, alternatively spliced CFTR mRNA.²⁴ Subjects with one CF mutation on one chromosome and the 5T allele on the other have little normal CFTR, and their phenotypes are quite diverse, including mild CF, disseminated bronchiectasis, CBAVD, and the absence of fertility problems.^{4 25 26} We speculate that neonatal hypertrypsinaemia could also be included in this wide clinical spectrum. In fact, the prevalence of the 5T allele was only slightly higher in group A compared to B, and in B compared to C.

The differences in the IVS8-5T frequencies would become much higher (group A *v* C, $p < 0.05$) if we considered two chromosomes per subject in groups B (7/102) and C (5/100) and only one chromosome per subject in group A (7/45), according to the hypothesis that the 5T variant is always in *trans* with $\Delta F508$. This assumption is supported by previous data,¹⁶ showing that $\Delta F508$ was linked to the 9T allele in 85 $\Delta F508$ chromosomes. To the best of our knowledge, there are no published reports of a CFTR gene carrying both this mutation and the 5T allele.

The 5T phase relative to $\Delta F508$ was studied in four IRT positive babies whose parents agreed to be tested. $\Delta F508$ was never associated in *cis* with the 5T allele, thus supporting our hypothesis. However, using different numbers of chromosomes per person in the populations under comparison could be questionable. A more recognised approach to the study of empirical gene frequencies for a particular allele consists in calculating a ratio between the number of chromosomes positive for the allele and the total number of tested chromosomes, whether or not the status of the

chromosomes regarding other mutations is known.

To follow this convention and at the same time overcome the difficulties connected with contrasting genetically heterogeneous populations like A, B, and C, we chose to compare two samples of carriers, the IRT positive and CF heterozygous babies in group A and a control population of female CF carriers (group D). The significantly higher ($p < 0.05$) prevalence of the IVS8-5T variant which we found in the former substantiates the hypothesis that some of these newborns are actually affected by a very mild form of CF, thus explaining the high IRT levels. Similarly, it could be speculated that other CFTR variants or mild mutations were actually responsible for the hypertrypsinaemia in the remaining neonates in group A and, when affecting both chromosomes, in the ones in group B. A thorough study of the CFTR gene in these subjects should clarify the issue.

In conclusion, the incidence of CF carriers in infants with a raised IRT is higher than the one expected in the general population. In at least some babies raised trypsin levels at birth could be a phenotypic expression of compound heterozygosity for a major CF mutation plus the 5T polymorphism. Considering the wide range of manifestations connected with this genotype, which can range from completely healthy to CBAVD or even mild CF, it is extremely difficult to predict the clinical outcome of these newborns, and to provide satisfactory genetic counselling for the family. Close clinical follow up should help in clarifying the extent of the disease in these subjects, with special regard to the three males, who could possibly turn out to be affected by CBAVD.

Newborn CF carriers with hypertrypsinaemia should perhaps deserve a closer look before being dismissed as "unaffected".

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