

Source (code type)	Code	Description
GP records (Read code)	1JB0	Suspected Downs syndrome
	677C4	Carrier of fragile X gene mutation
	C301	Phenylketonuria
	EMISNQCA42	Cause of learning disabilities: down's syndrome
	Eu842	[X]Rett's syndrome
	F1y0	Fragile X associated tremor ataxia syndrome
	PJ0	Down's syndrome - trisomy 21
	PJ00	Trisomy 21, meiotic nondisjunction
	PJ01	Trisomy 21, mosaicism
	PJ02	Partial trisomy 21 in Down's syndrome
	PJ0-2	trisomy 21
	PJ0-98	Down's syndrome
	PJ0z	Down's syndrome NOS
	PJ1..	Patau's syndrome - trisomy 13
	PJ10.	Trisomy 13, meiotic nondisjunction
	PJ11.	Trisomy 13, mosaicism
	PJ12.	Trisomy 13, translocation
	PJ1z.	Patau's syndrome NOS
	PJ2	Edward's syndrome - trisomy 18
	PJ20	Trisomy 18, meiotic nondisjunction
	PJ21	Trisomy 18, mosaicism
	PJ22	Partial trisomy 18 in Edward's syndrome
	PJ2z	Edward's syndrome NOS
	PJ30.	Deletion of long arm of chromosome 21

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