

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

Icd 9 Codes For Metabolic Panel

Arturo Cuomo

A red circular graphic with a gradient, appearing as a semi-circle or a partial circle, located to the right of the author's name.

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