

Supplementary Table 4 Summary of genetic diagnosis obtained in a cohort of 166 patients affected with syndromic craniosynostosis (only patients with positive results of genetic testing were calculated)

	<i>FGFR1</i>		<i>FGFR2</i>		<i>FGFR3</i>		<i>TWIST1</i>		<i>Total</i>		<i>Another</i>		<i>Total</i>	
	Patients	Frequency (%)	Patients	Frequency (%)	Patients	Frequency (%)	Patients	Frequency (%)	Patients	Frequency (%)	Patients	Frequency (%)	Patients	Frequency (%)
Sex														
Female	2	3	13	19	10	14	9	13	34	48	10	14	44	62
Male	1	1	6	9	3	4	8	12	18	25	9	13	27	38
Total	3	4	19	27	13	18	17	24	52	73	19	27	71	100
Presence of additional clinical features														
Isolated	0	0	4	8	8	16	8	16	20	39	5	10	25	49
Syndromic	0	0	6	12	1	2	5	10	12	24	14	27	26	51
Total	0	0	10	20	9	18	13	25	32	63	19	37	51	100
Number of affected sutures														
Multiple	0	0	8	16	5	10	7	14	20	41	12	24	32	65
Single	0	0	1	2	4	8	5	10	10	20	7	14	17	35
Total	0	0	9	18	9	18	12	24	30	61	19	39	49	100
Occurrence														
Familial	3	4	2	3	3	4	9	13	17	24	2	3	19	27
Sporadic	0	0	17	24	10	14	8	11	35	49	17	24	52	73
Total	3	4	19	27	13	18	17	24	52	73	19	27	71	100