Table 1. RET mutations in MEN2A syndrome.

Exon	Codon No	Amino acid substitution
10	609	Cys to Arg
		Cys to Tyr
	611	Cys to Trp
		Cys to Tyr
	618	Cys to Phe
		Cys to Ser
		Cys to Gly
		Cys to Arg
		Cys to Tyr
		Cys to Stop
	620	Cys to Arg
		Cys to Tyr
		Cys to Phe
		Cys to Ser
		Cys to Gly
11	630	Cys to Phe
	634	Cys to Arg
		Cys to Tyr
		Cys to Phe
		Cys to Ser
		Cys to Gly
		Cys to Tyr
	666	Lys to Glu
13	768	Gly to Asp
	790	Leu to Phe
	791	Tyr to Phe
14	804	Val to Met
		Val to Leu
15	891	Ser to Ala