

**Supplementary Table 1 Somatic mutations**

<b>Sample ID</b>	<b>Gene</b>	<b>Chromosome</b>	<b>Chromosome position</b>	<b>Exon</b>	<b>Coverage</b>	<b>Mutation Frequency (%)</b>	<b>AA change</b>	<b>Co</b>	
<b>MTC29</b>	RET	10	43617416	16	85	21,18	c.2753T>C	p.Met918Thr	CC
<b>MTC31</b>	RET	10	43617416	16	1751	64,13	c.2753T>C	p.Met918Thr	CC
<b>MTC33</b>	RET	10	43617416	16	606	64,13	c.2753T>C	p.Met918Thr	CC
<b>MTC34</b>	EGFR	7		21	2093	51,84	c.2543C>T	p.Pro848Leu	CC
<b>MTC34</b>	RET	10	43617416	16	647	50,85	c.2753T>C	p.Met918Thr	CC
<b>MTC34</b>	RET	10	43609101	10			c.1858T>C	p.Cys620Arg	CC

The reference sequences used were NM\_020975.4 for RET and NM\_005228 for EGFR