

NGS based identification of mutational hotspots for targeted therapy in anaplastic thyroid carcinoma

Supplementary Materials

Supplementary Table 1: Mutations in the analysed genes 118 ATC primary tumour samples and the relative prevalence of each mutation in %. See Supplementary_Table_1

Supplementary Table 2: Panel of analyzed genes and exons

Gene	Exon
<i>ALK</i>	20–25
<i>BRAF</i>	11, 15
<i>CDKN2A</i>	2
<i>EGFR</i>	18–21
<i>ERBB2</i>	20, 23
<i>HRAS</i>	2–4
<i>KRAS</i>	2–4
<i>KIT</i>	9–11, 13, 17, 18
<i>MET</i>	14
<i>mTOR</i>	4, 5
<i>NRAS</i>	2–4
<i>PDGFRA</i>	12, 14, 18
<i>PIK3CA</i>	10, 21
<i>RB1</i>	18
<i>RET</i>	10, 11, 13–16
<i>TSC2</i>	30
<i>p53</i>	4–9