

FusionPlex[®] Sarcoma

This panel is no longer in use. This panel was replaced by ASARv3

Panel description

The Archer FusionPlex Sarcoma (ASARv1, # AB0003) panel targets 26 genes commonly mutated in sarcoma. The Anchored Multiplex PCR technology (RNA-NGS) identifies both known and novel fusion partners.

<i>Gene</i>	<i>RefSeq number</i>	<i>accession</i>	<i>Target Exons</i>	<i>Direction</i>	<i>Assay Type</i>
<i>ALK</i>	NM_004304		19, 20, 21, 22	5'	Fusion
<i>CAMTA1</i>	NM_015215		8, 9, 10	5'	Fusion
<i>CCNB3</i>	NM_033031		2, 3, 4, 5, 6	5'	Fusion
<i>CIC</i>	NM_015125		19, 20	3'	Fusion
<i>EPC1</i>	NM_025209		9, 10, 11	3'	Fusion
<i>EWSR1</i>	NM_005243		4,5,6,7,8,9,10,11,12,13	3'	Fusion
<i>FOXO1</i>	NM_002015		2,3	3'	Fusion
<i>FOXO1</i>	NM_002015		1,2	5'	Fusion
<i>FOXO1</i>	NM_002015		3	5'	Fusion
<i>FOXO1</i>	NM_002015		1	3'	Fusion
<i>FUS</i>	NM_004960		4, 5, 6, 7, 8, 9, 10, 11, 14	3'	Fusion
<i>GLI1</i>	NM_005269		4,5,6,7	5'	Fusion
<i>GLI1</i>	NM_005269		4,5,6,7	3'	Fusion
<i>HMGA2</i>	NM_003483		1,2,3,4,5	3'	Fusion
<i>JAZF1</i>	NM_175061		2,3,4	3'	Fusion
<i>MEAF6</i>	NM_001270875		4,5	3'	Fusion
<i>MKL2</i>	NM_014048		11,12,13	5'	Fusion
<i>NCOA2</i>	NM_006540		11, 12, 13, 14	5'	Fusion
<i>NTRK3</i>	NM_002530		13, 14, 15	3'	Fusion
<i>NTRK3</i>	NM_002530		13, 14, 15, 16	5'	Fusion
<i>PDGFB</i>	NM_002608		2,3	5'	Fusion
<i>PLAG1</i>	NM_002655		1,2,3,4	5'	Fusion
<i>ROS1</i>	NM_002944		31,32,33,34,35,36,37	5'	Fusion
<i>SS18</i>	NM_001007559		10, 11	5'	Fusion
<i>SS18</i>	NM_001007559		4,5,6,8,9,10	3'	Fusion
<i>STAT6</i>	NM_001178078		1,2,3,4,5,6,7,16,17,18,19	5'	Fusion
<i>TAF15</i>	NM_139215		6,7	5'	Fusion
<i>TAF15</i>	NM_139215		5,6,7	3'	Fusion

TCF12	NM_207036	4,5,6	3'	Fusion
TFE3	NM_006521	3,4,5,6	5'	Fusion
TFG	NM_006070	4,5,6,7	3'	Fusion
TFG	NM_006070	6	5'	Fusion
USP6	NM_004505	1,2,3,	5'	Fusion
YWHAE	NM_006761	5	3'	Fusion

Panel characteristics

General: Data analysis is performed in Archer Analysis software. The used “pipeline” version can be obtained on request. Unless otherwise stated only “ strong evidence” fusions are reported.

Detection of fusion, variants and exon skipping events can be detected depending on the variant type, sequence context, sample DNA quality and quantity.

For reliable analysis the following requirements have to be met:

- (1) QC score (Cq) should be < 30. If Cq 30-31, the fusionplex is less reliable, if Cq >31, quality is unsuitable.
- (2) The number of reads per sample should be > 1.5M
- (3) The percentage of RNA reads should be > 40%