AS24009_50351729_lon_V2_RBC1_BC18_rawlib.basecaller

Overall QC Status: PASS Sample QC Status: PASS Fusion QC Status: PASS Variations QC Status: PASS Job: 20240530 AS24009 [3536]

Type: RNA FusionRNA SNP/InDel Targeted Mutations: Archer Comprehensive Targets NIH v1.3.1 Include Non-Targeted

Variants: No

Software Version: Suite_Analysis_v6.2.7

Analysis Date: 30-May-2024 2:59
Report Creator: mpvghtpe@gmail.com
Report Date: 30-May-2024 18:13



Statistics

Molecular Barcode Statistics

Total Fragments	Fragments with Complete Adapter	Number of Reads After Trimming Adapters
3,500,000	3,324,509	3,195,072

Read Statistics

Туре	Total Fragments (# / %)	Mapped (# / %)	Pass Alignment Filter (%)	On Target (%)
All Fragments	3,183,314 / 100.0	3,183,314 / 100.0	100.0	99.2
Unique Fragments	166,410 / 5.2	166,410 / 100.0	100.0	98.3

DNA/RNA Statistics

Туре	DNA Reads (# / %)	RNA Reads (# / %)	Ambiguous Reads (# / %)
All Fragments	366,736.0 / 11.6	1,877,389.0 / 59.4	914,772.0 / 29.0
Molecular Bins	21,590.0 / 13.2	85,403.0 / 52.2	56,649.0 / 34.6
Average Molecular Bins per GSP2	19.88	78.64	52.16
Unique Start Sites	10,829.0 / 24.2	26,587.0 / 59.5	11,201.0 / 25.1
Average Unique Start Sites per GSP2	10.22	27.79	12.1
Average Unique Start Sites per GSP2 Control	12.42	59.92	5.83

QC Statistics

Avg. Unique DNA And Ambiguous Start Sites Per GSP2	Avg. Unique RNA Start Sites Per GSP2 Control	
22.25	59.92	

Miscellaneous Statistics

On Target Deduplication Ratio
19.30:1

DNA/RNA Fragment Lengths

DNA Median Fragment Length	DNA Mean Fragment Length	RNA Median Fragment Length	RNA Mean Fragment Length
113.0	127.9	120.0	129.6

Reportable Variants

None Found

Reportable Isoforms

☑ Passed all strong-evidence filters

♣ Likely off-target mispriming event

Exact breakpoint known

☎ Cross contamination

✗ User-annotated false positive

■ Known fusion partners in Archer Quiver™

☐ Percent GSP2 reads below threshold

 $\Delta \hat{}$ Fusion expression imbalance

1 Low confidence

 $\ensuremath{\bigtriangleup}$ User-annotated true positive

% Intronic fusion

1. Not enough unique start sites

O Transcriptional readthrough event

☐ Known ensembl paralogue

Fusion: PAX3 → FOXO1		
Filters: ☑ ⊚	Reads: 172 (36.06%)	<u>Segments</u>
GSP2: PAX3_chr2_223084865_21A1_GSP2 Mutation Classification: Undefined	Start Sites: 97	chr2:223085053→223084859 PAX3(-) NM_181457.3, exon:7 chr13:41134997→41134805
Is Artifact: no		FOXO1(-) NM_002015.3, exon:2