# AS22007\_lon\_V2\_RBC2\_BC34\_rawlib.basecaller

Overall QC Status: PASS Sample QC Status: PASS Fusion QC Status: PASS Variations QC Status: PASS Job: 20220317 AS22007 [4722]

**Type:** RNA FusionRNA SNP/InDel Targeted Mutations: Archer Comprehensive Targets NIH v1.3.1 2 Include Non-Targeted

Variants: No

Software Version: Suite\_Analysis\_v6.2.7

Analysis Date: 17-Mar-2022 13:45
Report Creator: mpvghtpe@gmail.com

Report Date: 19-Jun-2023 3:23



#### **Statistics**

#### **Molecular Barcode Statistics**

Total Fragments	Fragments with Complete Adapter	Number of Reads After Trimming Adapters
3,432,132	3,068,970	2,835,181

#### **Read Statistics**

Туре	Total Fragments (# / %)	Mapped (# / %)	Pass Alignment Filter (%)	On Target (%)
All Fragments	2,819,318 / 100.0	2,819,318 / 100.0	100.0	97.9
Unique Fragments	594,143 / 21.1	594,143 / 100.0	100.0	98.3

#### **DNA/RNA Statistics**

Туре	DNA Reads (# / %)	RNA Reads (# / %)	Ambiguous Reads (# / %)
All Fragments	154,438.0 / 5.6	2,252,650.0 / 81.6	352,746.0 / 12.8
Molecular Bins	46,310.0 / 7.9	417,898.0 / 71.6	119,761.0 / 20.5
Average Molecular Bins per GSP2	78.89	711.92	204.02
Unique Start Sites	18,374.0 / 29.8	37,690.0 / 61.2	16,621.0 / 27.0
Average Unique Start Sites per GSP2	32.37	92.17	32.93
Average Unique Start Sites per GSP2 Control	37.0	200.83	46.58

#### **QC Statistics**

Avg. Unique DNA And Ambiguous Start Sites Per GSP2	Avg. Unique RNA Start Sites Per GSP2 Control
64.62	200.83

#### **Miscellaneous Statistics**

On Target Deduplication Ratio
4.73:1

#### **DNA/RNA Fragment Lengths**

DNA Median Fragment Length	DNA Mean Fragment Length	RNA Median Fragment Length	RNA Mean Fragment Length
120.0	133.7	126.0	132.4

## **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

**↓** F Not enough unique start sites

O Transcriptional readthrough event

☐ Known ensembl paralogue

Fusion: EWSR1 → FLI1			
Filters: ☑ ⊚	Reads: 998 (19.70%)	<u>Segments</u>	
<b>GSP2:</b> EWSR1_chr22_29683087_28_+_A1_GSP 2	Start Sites: 161	chr22:29682912→29683123 EWSR1(+) NM_005243.3, exon:7	
Mutation Classification: Undefined		chr11:128651853→128651918	
Is Artifact: no		FLI1(+) NM_002017.4, exon:5	

Fusion: EWSR1 → FLI1			
Filters: ∰ ‡	Reads: 17 (0.37%)	<u>Segments</u>	
<b>GSP2:</b> EWSR1_chr22_29683087_28_+_A1_GSP 2	Start Sites: 14	chr22:29683088→29683122 EWSR1(+) NM_005243.3, exon:7	
Mutation Classification: Undefined		chr11:128651853→128651918	
Is Artifact: no		FLI1(+) NM_002017.4, exon:5	