# AL24001\_47913451\_lon\_V2\_RBC2\_BC3

Overall QC Status: PASS Sample QC Status: PASS Fusion QC Status: PASS Variations QC Status: PASS Job: 20240131\_AL24001 [4791]

**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: 30-Jan-2024 22:32 Report Creator: mpvghtpe@gmail.com

Report Date: 30-Jan-2024 23:12



#### **Statistics**

#### **Molecular Barcode Statistics**

Total Fragments	Fragments with Complete Adapter	Number of Reads After Trimming Adapters
3,500,000	3,317,345	3,154,055

#### **Read Statistics**

Туре	Total Fragments (# / %)	Mapped (# / %)	Pass Alignment Filter (%)	On Target (%)
All Fragments	3,143,100 / 100.0	3,143,100 / 100.0	100.0	96.9
Unique Fragments	194,055 / 6.2	194,055 / 100.0	100.0	95.7

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

Туре	DNA Reads (# / %)	RNA Reads (# / %)	Ambiguous Reads (# / %)
All Fragments	162,749.0 / 5.3	2,533,171.0 / 83.2	349,304.0 / 11.5
Molecular Bins	12,129.0 / 6.5	137,262.0 / 73.9	36,389.0 / 19.6
Average Molecular Bins per GSP2	74.41	842.1	223.25
Unique Start Sites	5,454.0 / 25.4	14,029.0 / 65.4	5,498.0 / 25.6
Average Unique Start Sites per GSP2	34.24	113.72	35.69
Average Unique Start Sites per GSP2 Control	41.75	205.62	88.5

## **QC Statistics**

Avg. Unique DNA And Ambiguous Start Sites Per GSP2	Avg. Unique RNA Start Sites Per GSP2 Control	
69.28	205.62	

## **Miscellaneous Statistics**

On	Target Deduplication Ratio
	16.39:1

## **DNA/RNA Fragment Lengths**

DNA Median Fragment Length	DNA Mean Fragment Length	RNA Median Fragment Length	RNA Mean Fragment Length
134.0	143.0	143.0	149.0

# **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

△ Fusion expression imbalance

1 Low confidence

 $\triangle$  User-annotated true positive

% Intronic fusion

1. Not enough unique start sites

O Transcriptional readthrough event

☐ Known ensembl paralogue

Fusion: CCDC6 $ ightarrow$ RET			
Filters: ☑ ⊚	Reads: 979 (86.18%)	<u>Segments</u>	
<b>GSP2</b> : RET_chr10_43612037_23A1_GSP2	Start Sites: 150	chr10:61666101→61665880 CCDC6(-) NM_005436.4, exon:1	
Mutation Classification: Undefined  Is Artifact: no		chr10:43612032→43612179 RET(+) NM_020630.4, exon:12	