# AS23049\_49703222\_lon\_V2\_RBC1\_BC6 \_rawlib.basecaller

Overall QC Status: PASS Sample QC Status: PASS Fusion QC Status: PASS Variations QC Status: PASS Job: 20230927 AS23049 [4787]

**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: 27-Sep-2023 4:32 Report Creator: mpvghtpe@gmail.com Report Date: 27-Sep-2023 18:50

## **Statistics**

#### **Molecular Barcode Statistics**

Total Fragments	Fragments with Complete Adapter	Number of Reads After Trimming Adapters
1,521,147	1,412,235	1,323,310

#### **Read Statistics**

Туре	Total Fragments (# / %)	Mapped (# / %)	Pass Alignment Filter (%)	On Target (%)
All Fragments	1,316,787 / 100.0	1,316,787 / 100.0	100.0	99.1
Unique Fragments	446,617 / 33.9	446,617 / 100.0	100.0	98.9

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

Туре	DNA Reads (# / %)	RNA Reads (# / %)	Ambiguous Reads (# / %)
All Fragments	100,742.0 / 7.7	941,172.0 / 72.1	262,836.0 / 20.1
Molecular Bins	41,810.0 / 9.5	282,650.0 / 64.0	117,430.0 / 26.6
Average Molecular Bins per GSP2	63.44	428.91	178.19
Unique Start Sites	16,441.0 / 27.0	38,576.0 / 63.3	14,454.0 / 23.7
Average Unique Start Sites per GSP2	25.63	78.68	25.49
Average Unique Start Sites per GSP2 Control	16.42	114.58	15.5





## **QC Statistics**

Avg. Unique DNA And Ambiguous Start Sites Per GSP2	Avg. Unique RNA Start Sites Per GSP2 Control
50.76	114.58

## **Miscellaneous Statistics**

On Target Deduplication Ratio
2.95:1

## **DNA/RNA Fragment Lengths**

DNA Median Fragment Length	DNA Mean Fragment Length	RNA Median Fragment Length	RNA Mean Fragment Length
137.0	146.6	133.0	140.3

# **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

% Intronic fusion

**↓** F Not enough unique start sites

O Transcriptional readthrough event

☐ Known ensembl paralogue

Fusion: COL1A1 → PDGFB			
Filters: ☑ ⑩	Reads: 3864 (83.76%)	<u>Segments</u>	
<b>GSP2:</b> PDGFB_chr22_39631845_24_+_A1_GSP 2	Start Sites: 244	chr17:48266156→48266103 COL1A1(-) NM_000088.3, exon:42	
Mutation Classification: Undefined		chr22:39631879→39631783	
Is Artifact: no		PDGFB(-) NM_002608.3, exon:2	

Fusion: COL1A1 → PDGFB			
Filters: 曼 ‡	Reads: 229 (6.06%)	<u>Segments</u>	
<b>GSP2:</b> PDGFB_chr22_39631845_24_+_A1_GSP 2	Start Sites: 93	chr17:48266156→48266111 COL1A1(-) NM_000088.3, exon:42	
Mutation Classification: Undefined		chr22:39639913→39639906	
Is Artifact: no		PDGFB(-) NM_002608.3, exon:1	