

# DEMO | wf-human-variation CNV report

Results generated through the wf-human-variation nextflow workflow provided by Oxford Nanopore Technologies.

Research use only 2024-05-22 2.2.0

## Introduction

This report contains CNVs detected using QDNAseq, as part of the wf-human-variation workflow.

## Chromosomal Sex

Undetermined

## Median Read Length

8244bp

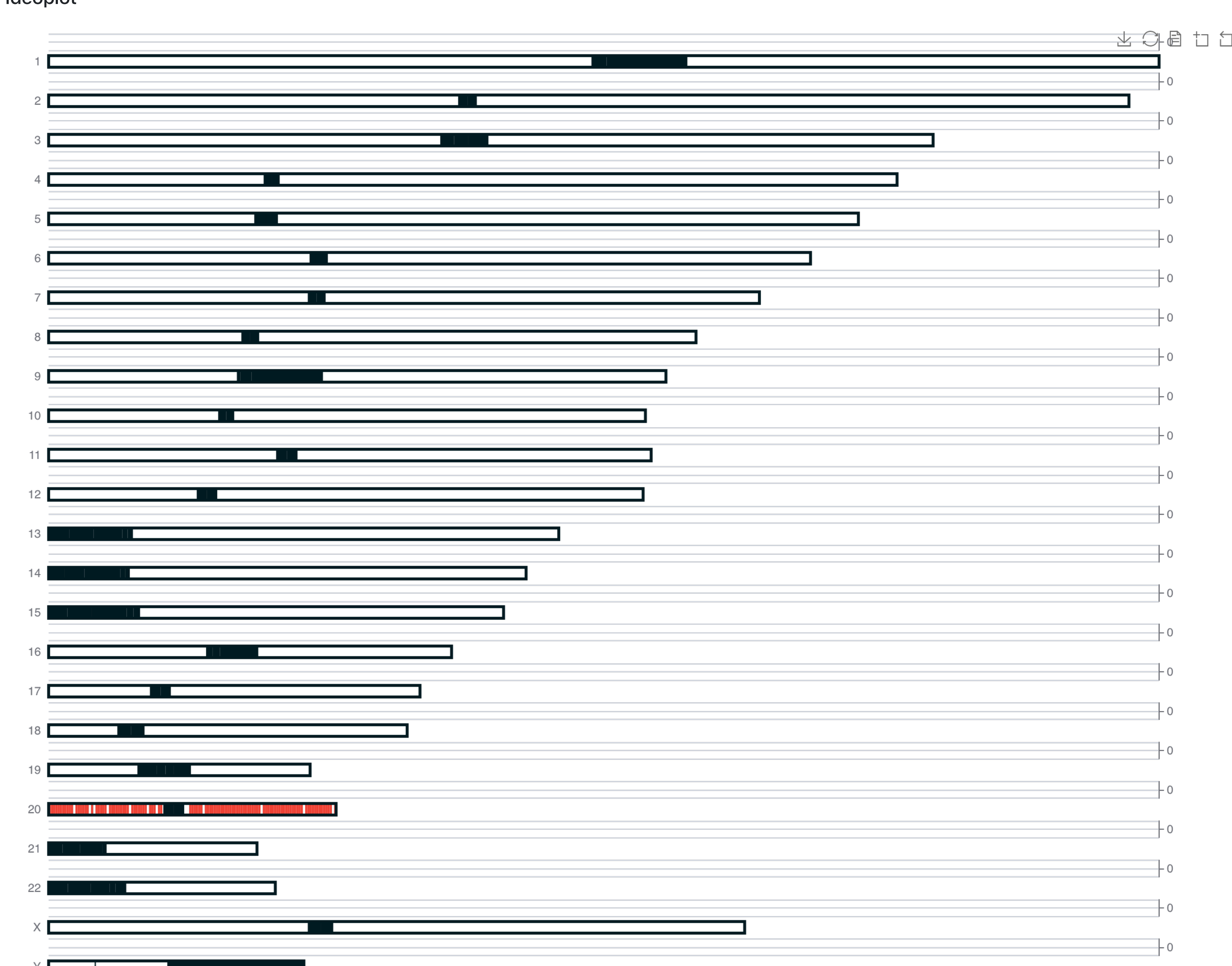
## Total Reads

414,758

## Chromosome Copy Summary

0 copies	1 copies	2 copies	3 copies	4 copies
-	-	1 10 11 12 13 14 15 16 17 18 19 2 21 22 3 4 5 6 7 8 9 X Y	-	20

## Ideoplot

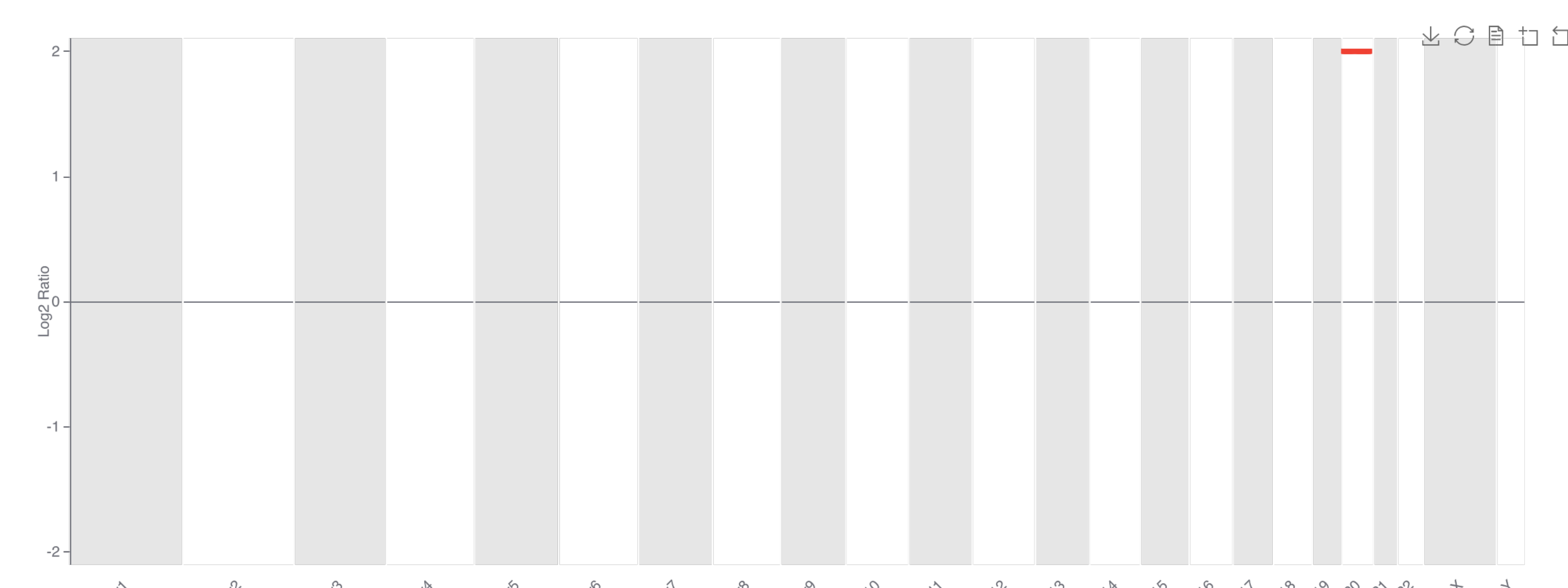


Red: Increased copy number detected  
 Blue: Decreased copy number detected  
 Yellow: QDNAseq no call  
 Black: 'acent', 'gvar', or 'stalk'. Read counts of <-2 converted to 2

## Detailed CNV Results

QDNAseq Smoothed Read Counts

Bin size 500Kbp, No call not displayed



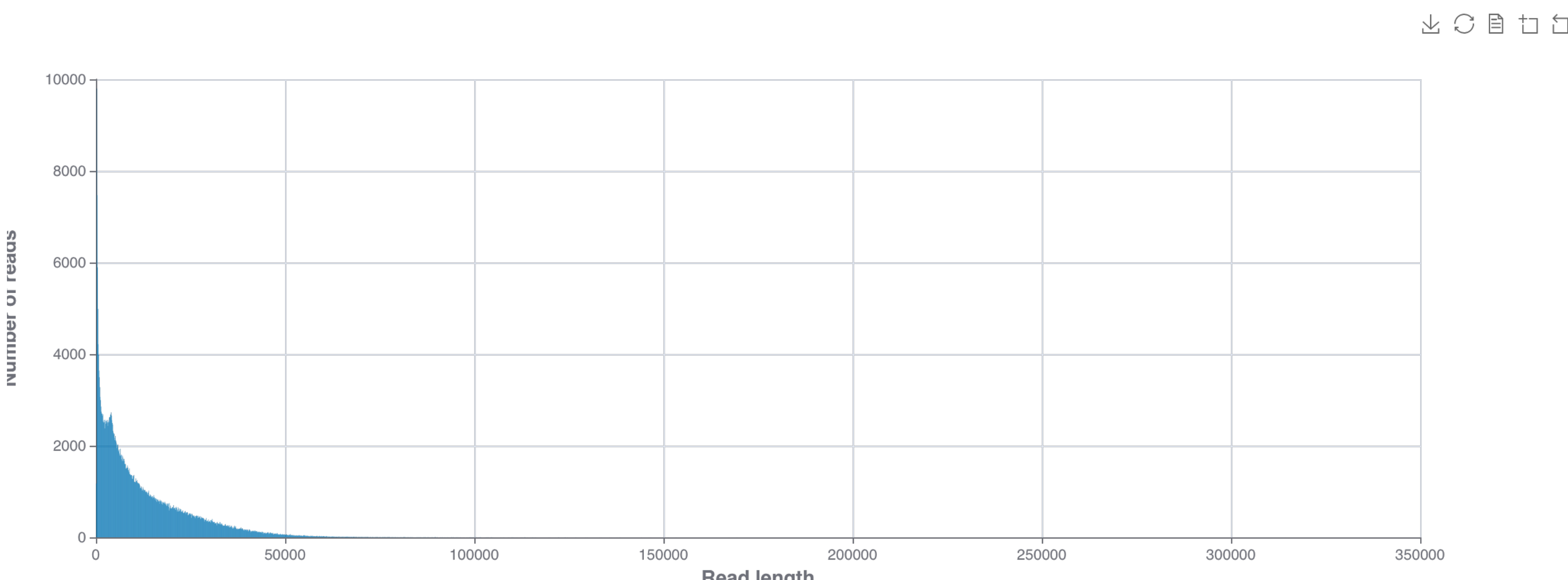
10 entries per page

chr	start	end	genes	read count	call
chr20	0	500000	<a href="#">Genes</a>	11.904	2
chr20	500000	1000000	<a href="#">Genes</a>	11.725	2
chr20	1000000	1500000	<a href="#">Genes</a>	11.809	2
chr20	1500000	2000000	<a href="#">Genes</a>	11.686	2
chr20	2000000	2500000	<a href="#">Genes</a>	11.775	2
chr20	2500000	3000000	<a href="#">Genes</a>	11.756	2
chr20	3000000	3500000	<a href="#">Genes</a>	11.737	2
chr20	3500000	4000000	<a href="#">Genes</a>	11.741	2
chr20	4000000	4500000	<a href="#">Genes</a>	11.771	2
chr20	4500000	5000000	<a href="#">Genes</a>	11.782	2

Showing 1 to 10 of 106 entries

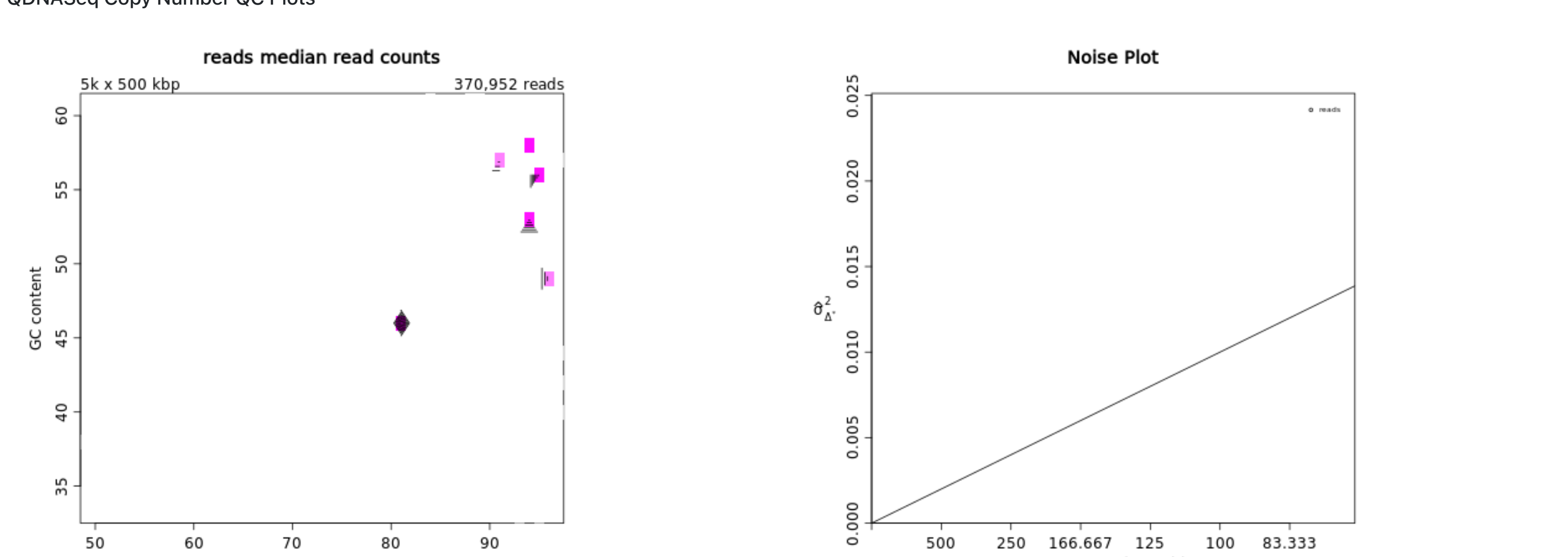
## Quality Control Data

### Histogram of Read Lengths



N.B. The read length affects the results of QDNAseq analysis. In future versions we will provide preset parameters based on detected read length

### QDNAseq Copy Number QC Plots



Median read counts per genomic bin shown as a function of GC content and mappability Read count relationship between sequence depth and noise

## Software versions

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Name	Version
pysam	0.21.0
R	4.2.2
QDNAseq	1.34.0
samtools	1.17

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## Workflow parameters

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Key	Value
help	False
version	False
disable_ping	False
threads	40
aws_image_prefix	None
aws_queue	None
out_dir	/project/ctb-noncodo/VnanoOperations/wf-human-variation-test/QDNAseq
snp	True
sv	True
mod	False

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## About this report

This report was produced using the [epi2me-labs/wf-human-variation](#) nextflow workflow (2.2.0).

Oxford Nanopore Technologies products are not intended for use for health assessment or to diagnose, treat, mitigate, cure or prevent any disease or condition.