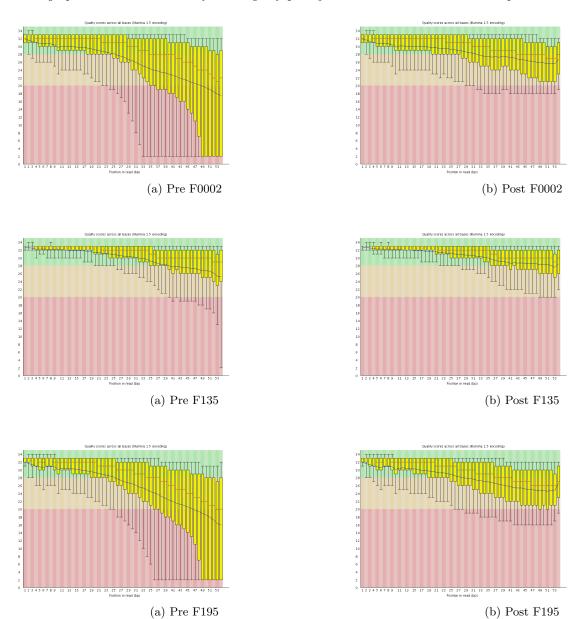
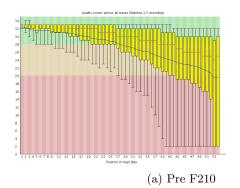
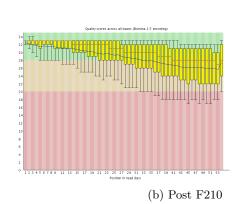
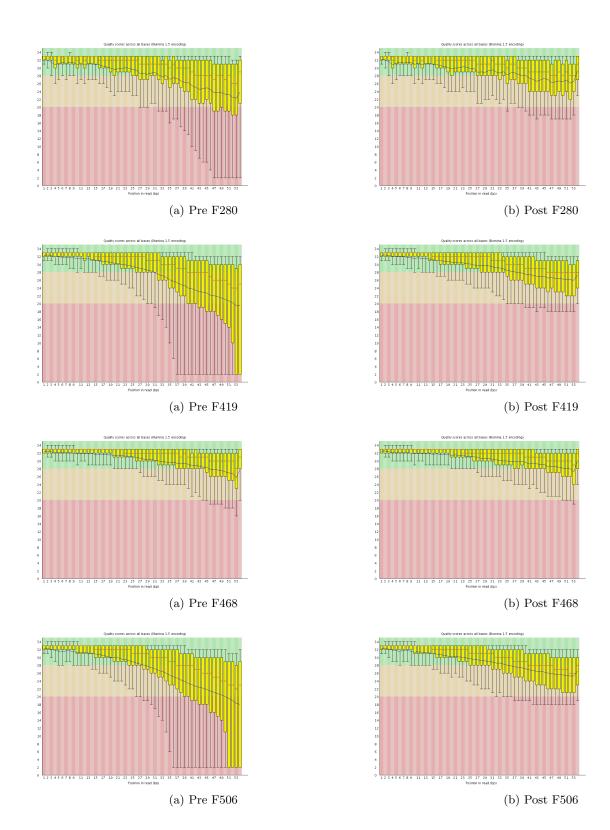
1 Base Quality Graph

This graph shows an overview of the range of quality values across all bases at each position in the FastQ file.



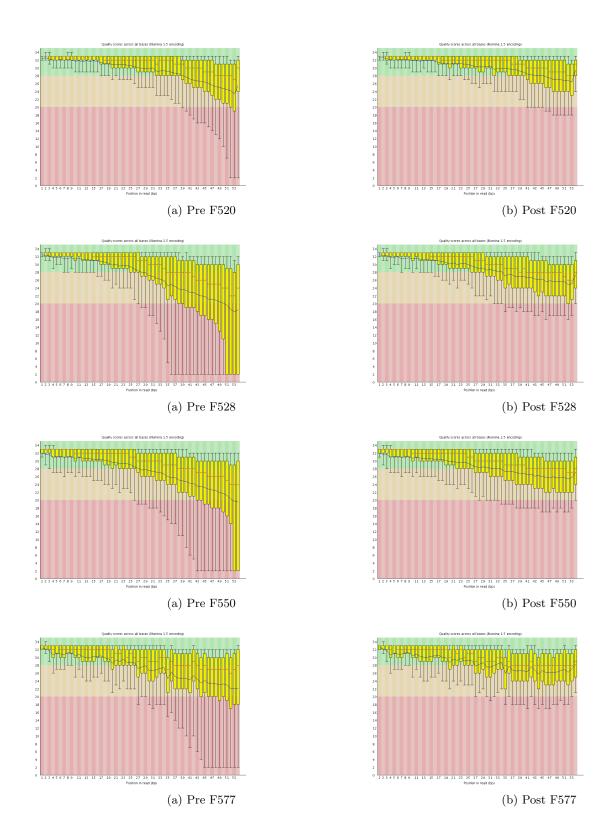






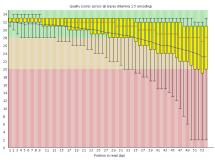
2 Per Base Sequence Content Graph

This graph plots out the proportion of each base position in a file for which each of the four normal DNA bases has been called.

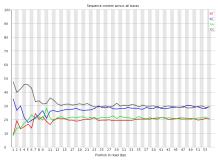


3 Data Summary

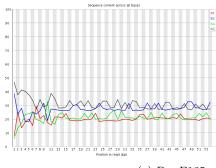
 $This\ section\ contains\ post\ trimming\ FastQC\ summary\ results.$



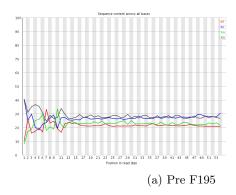
(a) Pre F_2129



(a) Pre F0002



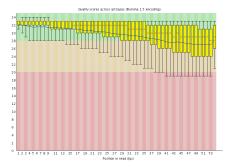
(a) Pre F135



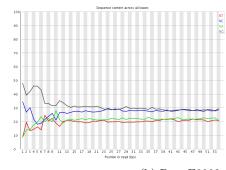
3.1 F0002.fastq.gz

Basic Statistics:

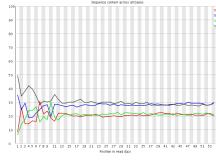
Total Sequences 16995797



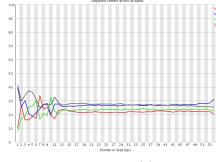
(b) Post F_2129



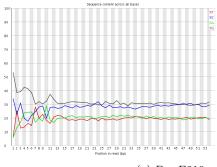
(b) Post F0002



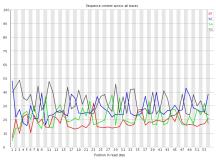
(b) Post F135



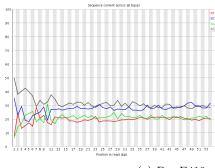
(b) Post F195



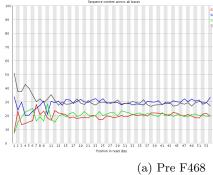
(a) Pre F210

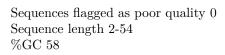


(a) Pre F280

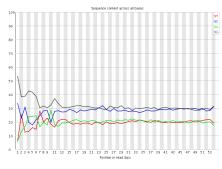


(a) Pre F419

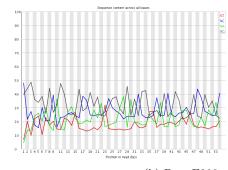




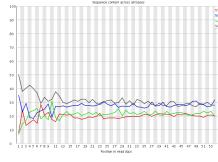
Other Statistics Summary:



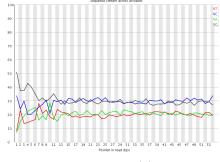
(b) Post F210



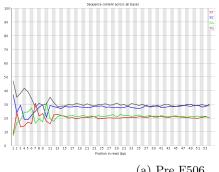
(b) Post F280

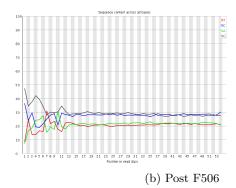


(b) Post F419

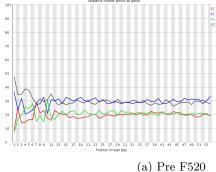


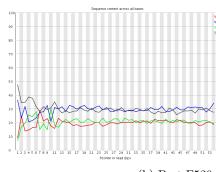
(b) Post F468











(b) Post F520

Metrics ResultBasic Statistics PASS Per base sequence quality PASS Per tile sequence quality FAIL Per sequence quality scores PASS Per base sequence content FAIL This could've failed due to fragmentation bias in the first 13bp. Checking... Only fragment bias in the first 13bp found. Per sequence GC content PASS Per base N content PASS Sequence Length Distribution WARN Sequence Duplication Levels WARN

Overrepresented sequences WARN Adapter Content PASS

Kmer Content FAIL

3.2 F135.fastq.gz

Basic Statistics:

Total Sequences 7877229 Sequences flagged as poor quality 0 Sequence length 2-54 %GC58

Other Statistics Summary:

Metrics ResultBasic Statistics PASS Per base sequence quality PASS Per tile sequence quality PASS

Per sequence quality scores PASS
Per base sequence content FAIL
This could've failed due to fragmentation bias in the first 13bp. Checking...
Only fragment bias in the first 13bp found.Per sequence GC content PASS
Per base N content PASS
Sequence Length Distribution WARN
Sequence Duplication Levels FAIL
Overrepresented sequences WARN
Adapter Content PASS
Kmer Content FAIL

3.3 F195.fastq.gz

Basic Statistics:

Total Sequences 11278843 Sequences flagged as poor quality 0 Sequence length 2-54 %GC 55

Other Statistics Summary:

Metrics ResultBasic Statistics PASS Per base sequence quality PASS

Per tile sequence quality PASS

Per sequence quality scores PASS

Per base sequence content FAIL

This could've failed due to fragmentation bias in the first 13bp. Checking... Only fragment bias in the first 13bp found.Per sequence GC content PASS

Per base N content PASS

Sequence Length Distribution WARN

Sequence Duplication Levels PASS

Overrepresented sequences WARN

Adapter Content PASS

Kmer Content FAIL

3.4 F210.fastq.gz

Basic Statistics:

Total Sequences 9924625 Sequences flagged as poor quality 0 Sequence length 2-54 %GC 60

Other Statistics Summary:

Metrics ResultBasic Statistics PASS Per base sequence quality PASS Per tile sequence quality PASS Per sequence quality scores PASS Per base sequence content FAIL This could've failed due to fragmentation bias in the first 13bp. Checking... Only fragment bias in the first 13bp found.Per sequence GC content WARN Per base N content PASS Sequence Length Distribution WARN Sequence Duplication Levels WARN Overrepresented sequences WARN Adapter Content PASS Kmer Content FAIL

3.5 F280.fastq.gz

Basic Statistics:

Total Sequences 12461046 Sequences flagged as poor quality 0 Sequence length 2-54 %GC 60

Other Statistics Summary:

Metrics ResultBasic Statistics PASS

Per base sequence quality PASS

Per tile sequence quality FAIL

Per sequence quality scores PASS

Per base sequence content FAIL

This could've failed due to fragmentation bias in the first 13bp. Checking...

The error isn't just due to fragment bias. Please check the reads to remove possible adapter contamination

Per sequence GC content FAIL

Per base N content PASS

Sequence Length Distribution WARN

Sequence Duplication Levels FAIL

Overrepresented sequences FAIL

Adapter Content FAIL

Kmer Content FAIL

3.6 F419.fastq.gz

Basic Statistics:

Total Sequences 10936459 Sequences flagged as poor quality 0 Sequence length 2-54 %GC 58

Other Statistics Summary:

Metrics ResultBasic Statistics PASS

Per base sequence quality PASS

Per tile sequence quality FAIL

Per sequence quality scores PASS

Per base sequence content FAIL

This could've failed due to fragmentation bias in the first 13bp. Checking...

Only fragment bias in the first 13bp found.Per sequence GC content WARN Per base N content PASS
Sequence Length Distribution WARN
Sequence Duplication Levels FAIL
Overrepresented sequences FAIL
Adapter Content PASS
Kmer Content FAIL

3.7 F468.fastq.gz

Basic Statistics:

Total Sequences 12369854 Sequences flagged as poor quality 0 Sequence length 2-54 %GC 59

Other Statistics Summary:

Metrics ResultBasic Statistics PASS Per base sequence quality PASS Per tile sequence quality PASS Per sequence quality scores PASS

Per base sequence content FAIL

This could've failed due to fragmentation bias in the first 13bp. Checking... Only fragment bias in the first 13bp found.Per sequence GC content FAIL

Per base N content PASS

Sequence Length Distribution WARN

Sequence Duplication Levels FAIL

Overrepresented sequences FAIL

Adapter Content PASS

Kmer Content FAIL

3.8 F506.fastq.gz

Basic Statistics:

Total Sequences 12663958 Sequences flagged as poor quality 0 Sequence length 2-54 %GC 58

Other Statistics Summary:

Metrics ResultBasic Statistics PASS

Per base sequence quality PASS

Per tile sequence quality FAIL

Per sequence quality scores PASS

Per base sequence content FAIL

This could've failed due to fragmentation bias in the first 13bp. Checking...

Only fragment bias in the first 13bp found.Per sequence GC content PASS

Per base N content PASS

Sequence Length Distribution WARN Sequence Duplication Levels WARN Overrepresented sequences WARN Adapter Content PASS Kmer Content FAIL

$3.9 \quad F520.fastq.gz$

Basic Statistics:

Total Sequences 8590866Sequences flagged as poor quality 0 Sequence length 2-54%GC 59

Other Statistics Summary:

Metrics ResultBasic Statistics PASS

Per base sequence quality PASS

Per tile sequence quality WARN

Per sequence quality scores PASS

Per base sequence content FAIL

This could've failed due to fragmentation bias in the first 13bp. Checking...

Only fragment bias in the first 13bp found.Per sequence GC content FAIL

Per base N content PASS

Sequence Length Distribution WARN

Sequence Duplication Levels FAIL

Overrepresented sequences FAIL

Adapter Content PASS

Kmer Content FAIL

3.10 F528.fastq.gz

Basic Statistics:

Total Sequences 17323169 Sequences flagged as poor quality 0 Sequence length 2-54 %GC 62

Other Statistics Summary:

Metrics ResultBasic Statistics PASS

Per base sequence quality PASS

Per tile sequence quality FAIL

Per sequence quality scores PASS

Per base sequence content FAIL

This could've failed due to fragmentation bias in the first 13bp. Checking...

The error isn't just due to fragment bias. Please check the reads to remove possible adapter contamination

Per sequence GC content FAIL

Per base N content PASS

Sequence Length Distribution WARN

Sequence Duplication Levels FAIL Overrepresented sequences FAIL Adapter Content FAIL Kmer Content FAIL

3.11 F550.fastq.gz

Basic Statistics:

Total Sequences 11367271 Sequences flagged as poor quality 0 Sequence length 2-54 %GC 58

Other Statistics Summary:

Metrics ResultBasic Statistics PASS

Per base sequence quality PASS

Per tile sequence quality PASS

Per sequence quality scores PASS

Per base sequence content FAIL

This could've failed due to fragmentation bias in the first 13bp. Checking...

The error isn't just due to fragment bias. Please check the reads to remove possible adapter contamination

Per sequence GC content WARN

Per base N content PASS

Sequence Length Distribution WARN

Sequence Duplication Levels FAIL

Overrepresented sequences FAIL

Adapter Content FAIL

Kmer Content FAIL

3.12 F577.fastq.gz

Basic Statistics:

Total Sequences 15228352Sequences flagged as poor quality 0 Sequence length 2-54 %GC 59

Other Statistics Summary:

No. 1. D. 1. D. 1. C. 1. 1.

Metrics ResultBasic Statistics PASS

Per base sequence quality PASS

Per tile sequence quality FAIL

Per sequence quality scores PASS

Per base sequence content FAIL

This could've failed due to fragmentation bias in the first 13bp. Checking...

The error isn't just due to fragment bias. Please check the reads to remove possible adapter contamination

Per sequence GC content FAIL

Per base N content PASS

Sequence Length Distribution WARN

Sequence Duplication Levels FAIL Overrepresented sequences FAIL Adapter Content FAIL Kmer Content FAIL

3.13 F_2129.fastq.gz

Basic Statistics:

Total Sequences 11206374 Sequences flagged as poor quality 0 Sequence length 2-54 %GC 57

Other Statistics Summary:

Metrics ResultBasic Statistics PASS
Per base sequence quality PASS
Per tile sequence quality FAIL
Per sequence quality scores PASS
Per base sequence content FAIL
This could've failed due to fragmentation bias in the first 13bp. Checking...
Only fragment bias in the first 13bp found.Per sequence GC content WARN
Per base N content PASS
Sequence Length Distribution WARN
Sequence Duplication Levels FAIL
Overrepresented sequences WARN
Adapter Content PASS

4 Post Mapping Data

4.1 F0002.fastq.gz

Post Mapping Metrics

Kmer Content FAIL

4.2 F135.fastq.gz

Post Mapping Metrics

4.3 F195.fastq.gz

Post Mapping Metrics

4.4 F210.fastq.gz

Post Mapping Metrics

4.5 F280.fastq.gz

Post Mapping Metrics

#======#All numbers are READ count

Total records: 12462040

QC failed: 0

Optical/PCR duplicate: 0

Non primary hits 0

Unmapped reads: 3411636

mapq < mapq_cut (non-unique): 8704839

mapq >= mapq cut (unique): 345565

Read-1: 0 Read-2: 0

Reads map to '+': 175027 Reads map to '-': 170538 Non-splice reads: 345565

Splice reads: 0

Reads mapped in proper pairs: 0

Proper-paired reads map to different chrom:0

4.6 F419.fastq.gz

Post Mapping Metrics

#All numbers are READ count

Total records: 10939668

QC failed: 0

Optical/PCR duplicate: 0 Non primary hits 0

Unmapped reads: 2716427

mapq < mapq_cut (non-unique): 7511502

mapq >= mapq_cut (unique): 711739

Read-1: 0 Read-2: 0

Reads map to '+': 334514 Reads map to '-': 377225 Non-splice reads: 711739

Splice reads: 0

Reads mapped in proper pairs: 0

Proper-paired reads map to different chrom:0 $\,$

4.7 F468.fastq.gz

Post Mapping Metrics

Total records: 12371766

 QC failed: 0

Optical/PCR duplicate: 0 Non primary hits 0

Unmapped reads: 1355496

mapq < mapq_cut (non-unique): 10361834

mapq >= mapq cut (unique): 654436

Read-1: 0 Read-2: 0

Reads map to '+': 325237 Reads map to '-': 329199 Non-splice reads: 654436

Splice reads: 0

Reads mapped in proper pairs: 0

Proper-paired reads map to different chrom:0

4.8 F506.fastq.gz

Post Mapping Metrics

4.9 F520.fastq.gz

Post Mapping Metrics

Total records: 8592246

QC failed: 0

Optical/PCR duplicate: 0 Non primary hits 0

Unmapped reads: 1746473

mapq < mapq_cut (non-unique): 6290950

mapq >= mapq cut (unique): 554823

Read-1: 0 Read-2: 0

Reads map to '+': 263588 Reads map to '-': 291235 Non-splice reads: 554823

Splice reads: 0

Reads mapped in proper pairs: 0

4.10 F528.fastq.gz

Post Mapping Metrics

#All numbers are READ count

Total records: 17324676

 QC failed: 0

Optical/PCR duplicate: 0 Non primary hits 0

Unmapped reads: 4465909

 $mapq < mapq_cut$ (non-unique): 12673017

 $mapq >= mapq_cut$ (unique): 185750

Read-1: 0Read-2: 0

Reads map to '+': 89052 Reads map to '-': 96698 Non-splice reads: 185750

Splice reads: 0

Reads mapped in proper pairs: 0

Proper-paired reads map to different chrom:0

4.11 F550.fastq.gz

Post Mapping Metrics

#-----

#All numbers are READ count

Total records: 11368382

QC failed: 0

Optical/PCR duplicate: 0

Non primary hits 0

Unmapped reads: 3642734

mapq < mapq_cut (non-unique): 7160646

 $mapq >= mapq_cut$ (unique): 565002

Read-1: 0 Read-2: 0

Reads map to '+': 279923 Reads map to '-': 285079 Non-splice reads: 565002

Splice reads: 0

Reads mapped in proper pairs: 0

Proper-paired reads map to different chrom:0

4.12 F577.fastq.gz

Post Mapping Metrics

Total records: 15229162

QC failed: 0

Optical/PCR duplicate: 0 Non primary hits 0

Unmapped reads: 8717524

mapq < mapq_cut (non-unique): 5899753

mapq >= mapq cut (unique): 611885

Read-1: 0 Read-2: 0

Reads map to '+': 294881 Reads map to '-': 317004 Non-splice reads: 611885

Splice reads: 0

Reads mapped in proper pairs: 0

Proper-paired reads map to different chrom:0

$4.13 \quad F_2129.fastq.gz$

Post Mapping Metrics

Total records: 11207762

QC failed: 0

Optical/PCR duplicate: 0 Non primary hits 0 Unmapped reads: 2867241

mapq < mapq_cut (non-unique): 7313329

 $mapq >= mapq_cut$ (unique): 1027192

Read-1: 0 Read-2: 0

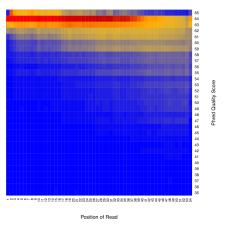
Reads map to '+': 521179 Reads map to '-': 506013 Non-splice reads: 1027192

Splice reads: 0

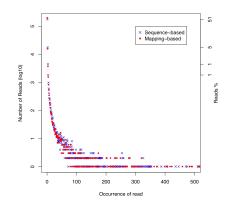
Reads mapped in proper pairs: 0

Proper-paired reads map to different chrom:0

5 Post Mapping Visual Data

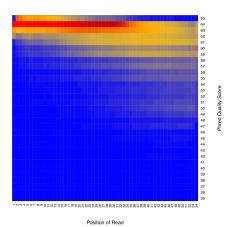


(a) Alignment Quality

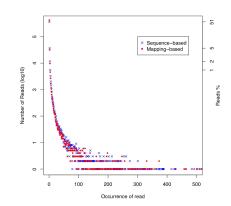


(b) Duplication Rate

Figure 27: F280.fastq.gz

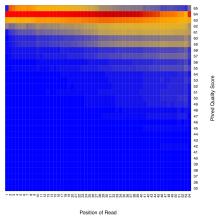


(a) Alignment Quality

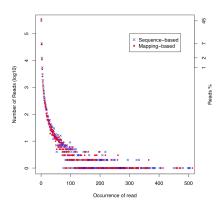


(b) Duplication Rate

Figure~28:~F419.fastq.gz

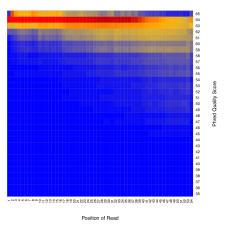


(a) Alignment Quality

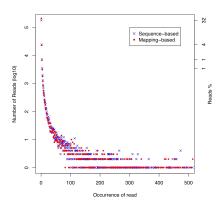


(b) Duplication Rate

Figure 29: F468.fastq.gz

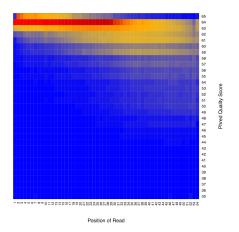


(a) Alignment Quality

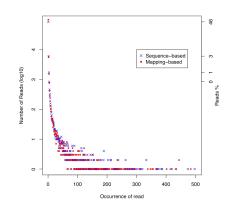


(b) Duplication Rate

Figure 30: F520.fastq.gz

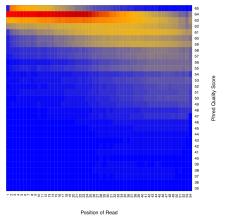


(a) Alignment Quality

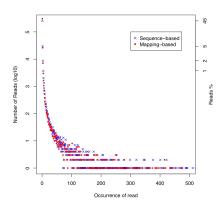


(b) Duplication Rate

Figure~31:~F528.fastq.gz

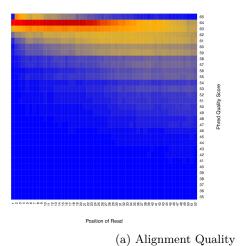


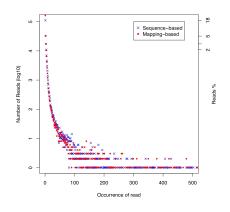
(a) Alignment Quality



(b) Duplication Rate

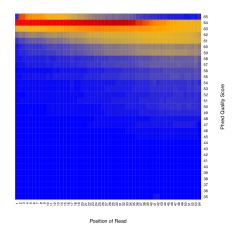
Figure 32: F550.fastq.gz



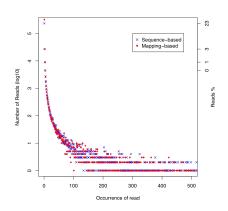


(b) Duplication Rate

Figure 33: F577.fastq.gz



(a) Alignment Quality



(b) Duplication Rate

Figure 34: $F_2129.fastq.gz$