

# Briefly: Bioinformatics File Formats

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# Overview

- **ASCII Text**
  - **Sequence**
    - **Fasta, Fastq**
  - **~Annotation**
    - **TSV, CSV, BED, GFF, GTF, VCF, SAM**
- **Binary (Data, Compressed, Executable)**
  - **Data**
    - **HDF5**
    - **BAM / CRAM**
    - **2bit**
  - **Compressed**
    - **gzip, bzip2, bgzip**
  - **Executable**

TEXT

# Fasta

>m54050R1\_180210\_051102/4194473/0\_1421

CCCGGCTGCCCCGCCCCGCTCGAAGCGATGACTTGCCGGCGGGCCGACGCGATTAGCTGCCGCGCATGCGATGCGGCCGCGGGCGGGCGTGCTGACCTGGCTGGCGGTG  
TTGAGCTGCTATACATCCGGCAACAACGCTGCCCAACGACTGACCTGACCGGCCGCCTCGATCCTGGCGGCCGCGGGCCTGGCCTGCGCTTTTCCTTCTTCTCTTTC  
CTTC

>m54050R1\_180210\_051102/4194473/1497\_4602

GGCGCGCTGATCGGCAAAACGGCTGGGGCGGCCGGAACACCTTTCAACCGTCGCCAACCGCGATCGCCGCGCGCACCCCGCCTTCCGCGCCGCTGTGGCGTTCTCTCG  
CCCGTCTACTCTACTGGCATCCGTCTCATTTCTCCCGCTCTTCCCTCCACCCCTTCCCTGCTCACCGCTTCCGTCTTTTTGTCAACCTCTCCTCTGGGCCGACGAC  
GTCGCCGCTACTGCGACAAAAACGAGGTCGACAAGGCCCGCCGTTACGACCGTCACCCCGAATTCCATCCGGCTGCTCCGCGGT

>m54050R1\_180210\_051102/4194551/0\_17688

ACCGGACGTACCGCGGGCGGGGGCCTCCCCCGGGTGGCTCGGGTGCAGCGCAAATCCTTTCTTTGCTGACCCACCTGCGCAGCGAGTGTGAATCTGTGCGGATCG  
AGAAAACAAGAAACCCGGCGGGGCCCTGCCTGACGCGCGCCCGTCCCGCCGCGCCCCCTTCCGCTTGGCGACGTGAGTTTTTGACGGGAGGTTTGTGCTTTCGACAGA  
CGGGTCCGCCAGCACCCCTCGTCGAGTCCCGTTAACTCAGGAAGAACTCCAGTTGGCCCGGGCATCTGCCAACGCCTCCGGGG

>m54050R1\_180210\_051102/4194551/17752\_17812

AAACATATTATTTTTTATTACTCAAATAATTATTATATTCACCTAATTTTCTTTATTATT

>m54050R1\_180210\_051102/4194552/0\_89

CAGATCGGGGCCAGCATGGCCACCCGTCCTGCACGTCTACGCGACTTCGCCGGTGGGGATCGGCAGCGGGAACGGCTCGCGGGCTGG

>m54050R1\_180210\_051102/4194552/162\_490

GCCGCACCCGAGCCGTTCCCGCTGCCGATCCACACCGTCGACGTGCGCGTCGACGTGCAGCCGGCGTCCATGCTTGCCCCGATCTTGGGCTAACAAGCCGCTGCTGA  
CACCGACGGACGCCACCGCCCGCGACCACTGGCCCGGGCCTCGGTGATGGCGCTGTCTACCGTCGCGCATTCGCGGCTCGGCATCTATCAGCCTCGGTGCCGCA  
CGGTCATCGACGATGGCGAAACCGTCACTGCACGTTTTTCATGACGCGGGGCAGGCAGCGAACCGGGGCACATCGGGCATCTACGCT

# Fasta

```
>m54050R1_180210_051102/4194473/0_1421
```

```
CCCGGCTGCCCCGCCCCGCTCGAAGCGATGACTTGCCGGCGGGCCCGACGCGATTAGCTGCCGCGCATGCGATGCGGCCGCGGGCGGGCGTGCTGACCTGGCTGGCGGTG  
TTGAGCTGCTATACATCCGGCAACAACGCTGCCCAACGACTGACCTGACCGGCCGCCTCGATCCTGGCGGCCGCCGGCCTGGCCTGCGCTTTTCCTTCTTCTTTTC  
CTTC
```

Header symbol “>” also redirects stuff into files, so be careful using > in bash commands!

Header text (sequence ID) has formats particular to different organizations and different software, but really has no consistent rules that you can rely on.

Sequence can contain: newline characters (“\n”), ACGT, N, acgt, n, x, . or - (gaps), IUPAC ambiguity codes BDHV etc., alternates like [A/T], amino acid single letter codes (protein fasta; sometimes file name is ‘sequence.fna’ for fasta nucleic acid, or ‘sequence.faa’ for fasta amino acid)

# Fastq ... “fasta + qualities”

```
@SN638:981:HK7HWBCXX:2:1101:14799:2762 1:N:0:TTAGGC
TGGCGCAACTGCCGATCACCATCGACACCAACGGGTATCTGGTCGCCAAC
+
GGGGGIIIIIIIIIIIIIIIGIIIIIIIIIIIIIIIIIIIIIIIG
@SN638:981:HK7HWBCXX:2:1101:14784:2782 1:N:0:TTAGGC
CATCATCGAGGACAGCGCCGGTGACCTGGCGGCCCGCATCGGTGCCCCC
+
GGGGGIIIIIIIIIIIIIIIIIIIIIIIGIIIIIIIIIIIIIGIIII
@SN638:981:HK7HWBCXX:2:1101:14983:2799 1:N:0:TTAGGC
CGGCGCCGTTGCTGCTGCTGCCGGTGCTGCTTTCGGCGCTGATCGTGCGG
+
GGGGGIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIGIIII
@SN638:981:HK7HWBCXX:2:1101:14763:2901 1:N:0:TTAGGC
CCTGACGACGGCACGAAGGACCTCTTCGTCCACTACTCCGAGATCCAGGG
+
GAGGGIGIGGGGGGGGIA.<GGGIGGAGGGGIIIGIIIGGIIIG<GA.<<GA
```

@Header1  
Sequence  
+Header2  
Qualities

Blocks of four lines for each sequence (sequences shouldn't occupy more than one line, as they can in fasta). Second header line (starting with “+”) is mandatory, sometimes contains the same header as the first line (that starts with “@”). Why??

The  $n^{\text{th}}$  quality character applies to the  $n^{\text{th}}$  nucleotide, and is a number that is *encoded in a single character from the ASCII table*.

# Fastq ... “fasta + qualities”

@SN638:981:HK7HWBCXX:2:1101:14799:2762 1:N:0:TTAGGC  
TGGCGCAACTGCCGATCACCATCGACACCAACGGGTATCTGGTCGCCAAC  
+  
GGGGGIIIIIIIIIIIIIIIGIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIG

@SN638:981:HK7HWBCXX:2:1101:14784:2782 1:N:0:TTAGGC  
CATCATCGAGGACAGCGCCGGTGACCTGGCGGCCCGCATCGGTGCCCCC  
+  
GGGGGIIIIIIIIIIIIIIIIIIIIIIIGIIIIIIIIIIIIIGIIII

@SN638:981:HK7HWBCXX:2:1101:14983:2799 1:N:0:TTAGGC  
CGGCGCCGTTGCTGCTGCTGCCGGTGCTGCTTTCGGCGCTGATCGTGCGG  
+  
GGGGGIIIGIII  
@SN638:981:HK7HWBCXX:2:1101:14763:2901 1:N:0:TTAGGC  
CCTGACGACGGCACGAAGGACCTCTTCGTCCACTACTCCGAGATCCAGGG  
+  
GAGGGIGIGGGGGGGGIA.<GGGIGGAGGGGIIGIIGGGIIIG<GA.<<GA

The “I” for base 16 (“C”) means that that base has a quality of (I ’s decimal value: 73) - 33 = 40 (sometimes referred to as “Q40”). Why 33? Because there are 32 non-printable “characters” at the beginning of the ASCII table! (type ‘man ascii’)

Q40 means that the probability of error (that C is actually the wrong basecall) is:

$$p_e = 10^{(-40 / 10)} = 0.0001, \text{ or } 1 \text{ in } 10,000$$

see also: [https://en.wikipedia.org/wiki/FASTQ\\_format](https://en.wikipedia.org/wiki/FASTQ_format)

# CSV and TSV - comma/tab-separated values

	B01	B02	B03	B04
PDCD1	0	0	0	0
GAL3ST2	0	0	0	0
D2HGDH	55	71	89	101
ING5	1	1	1	1
DTYMK	2	5	7	12
ATG4B	0	0	0	0
THAP4	136	158	85	161
BOK	0	0	0	0
STK25	145	175	195	141

For example, abundances of mRNAs from genes (count data).

(First tab character - “\t” - in column names sometimes omitted for ease of reading by R scripts).



# BED - tsv with defined column meanings

chr7	127471196	127472363	Pos1	0	+	column	meaning
chr7	127472363	127473530	Pos2	0	+	1	chromosome name
chr7	127473530	127474697	Pos3	0	+	2	feature start coordinate (0-based...?)
chr7	127474697	127475864	Pos4	0	+	3	feature stop coordinate (0-based...?)
chr7	127475864	127477031	Neg1	0	-	4	feature name
chr7	127477031	127478198	Neg2	0	-	5	score (1-1000)
chr7	127478198	127479365	Neg3	0	-	6	strand ('+' or '-' or '.' for unknown or not applicable)
chr7	127479365	127480532	Pos5	0	+	...	...
chr7	127480532	127481699	Neg4	0	-		

Number of columns used shouldn't vary within a particular file.

see also:

<https://genome.ucsc.edu/FAQ/FAQformat.html#format1>

# GFF / GTF - tsv with defined column meanings

```
chr22  TeleGene  enhancer  10000000  10001000  500  +  .  touch1
chr22  TeleGene  promoter  10010000  10010100  900  +  .  touch1
chr22  TeleGene  promoter  10020000  10025000  800  -  .  touch2
```

column	meaning
--------	---------

- |   |  |
|---|--|
| 1 | chromosome / scaffold name                                     |
| 2 | source (e.g. software that generated this feature / gene call) |
| 3 | feature name (e.g. “exon1”, “enhance”r, “3’-UTR”)              |
| 4 | feature start coordinate (1-based)                             |
| 5 | feature stop coordinate (1-based)                              |
| 6 | score (1-1000)   |
| 7 | strand (‘+’ or ‘-’ or ‘.’ for unknown or not applicable)       |
| 8 | reading frame (0, 1, 2, or “.” if N/A)                         |
| 9 | group (allows grouping features together)                      |

GTF is newer, and shares the first eight (8) columns. Column 9 has additional restrictions in format (gene\_id, transcript\_id, etc.)

see also: <https://genome.ucsc.edu/FAQ/FAQformat.html#format3>

# VCF - tsv with defined column meanings

```
##fileformat=VCFv4.2
##fileDate=20090805
##source=myImputationProgramV3.1
##reference=file:///seq/references/1000GenomesPilot-NCBI36.fasta
##contig=<ID=20,length=62435964,assembly=B36,md5=f126cdf8a6e0c7f379d618ff66beb2da,species="Homo sapiens",taxonomy=x>
##phasing=partial
##INFO=<ID=NS,Number=1,Type=Integer,Description="Number of Samples With Data">
##INFO=<ID=DP,Number=1,Type=Integer,Description="Total Depth">
##INFO=<ID=AF,Number=A,Type=Float,Description="Allele Frequency">
##INFO=<ID=AA,Number=1,Type=String,Description="Ancestral Allele">
##INFO=<ID=DB,Number=0,Type=Flag,Description="dbSNP membership, build 129">
##INFO=<ID=H2,Number=0,Type=Flag,Description="HapMap2 membership">
##FILTER=<ID=q10,Description="Quality below 10">
##FILTER=<ID=s50,Description="Less than 50% of samples have data">
##FORMAT=<ID=GT,Number=1,Type=String,Description="Genotype">
##FORMAT=<ID=GQ,Number=1,Type=Integer,Description="Genotype Quality">
##FORMAT=<ID=DP,Number=1,Type=Integer,Description="Read Depth">
##FORMAT=<ID=HQ,Number=2,Type=Integer,Description="Haplotype Quality">
#CHROM POS ID REF ALT QUAL FILTER INFO FORMAT sample07 ...
20 14370 rs6054257 G A 29 PASS NS=3;DP=14;AF=0.5;DB;H2 GT:GQ:DP:HQ 0|0:48:1:51,51
20 17330 . T A 3 q10 NS=3;DP=11;AF=0.017 GT:GQ:DP:HQ 0|0:49:3:58,50
20 111069 rs6040355 A G,T 67 PASS NS=2;DP=10;AF=0.333,0.667;AA=T;DB GT:GQ:DP:HQ 1|2:21:6:23,27
20 123027 . T . 47 PASS NS=3;DP=13;AA=T GT:GQ:DP:HQ 0|0:54:7:56,60
20 123457 microsat1 GTC G,GTCT 50 PASS NS=3;DP=9;AA=G GT:GQ:DP 0/1:35:4
```

# SAM - tsv with defined column meanings

<http://www.htslib.org/>

See also samtools man page: <http://samtools.sourceforge.net/>

SAM spec grew out of 1000 Genomes Project (see Li et al. 2009 *Bioinformatics* 25:2078)

SAM is plain text; BAM is binary, compressed version of SAM; CRAM is further compressed but not widely used / recognizable by many tools.

# SAM - tsv with defined column meanings

[illegible]

# SAM - tsv with defined column meanings

Col	Field	Type	Regex/Range	Brief description
1	QNAME	String	[!-?A-~]{1,254}	Query template NAME
2	FLAG	Int	[0,2 <sup>16</sup> -1]	bitwise FLAG
3	RNAME	String	\*  [!-( )+-<>-~] [!-~]*	Reference sequence NAME
4	POS	Int	[0,2 <sup>31</sup> -1]	1-based leftmost mapping POSition
5	MAPQ	Int	[0,2 <sup>8</sup> -1]	MAPping Quality
6	CIGAR	String	\*  ([0-9]+[MIDNSHPX=])+	CIGAR string
7	RNEXT	String	\* =  [!-( )+-<>-~] [!-~]*	Ref. name of the mate/next read
8	PNEXT	Int	[0,2 <sup>31</sup> -1]	Position of the mate/next read
9	TLEN	Int	[-2 <sup>31</sup> +1,2 <sup>31</sup> -1]	observed Template LENgth
10	SEQ	String	\*  [A-Za-z=.]+	segment SEQUENCE
11	QUAL	String	[!-~]+	ASCII of Phred-scaled base QUALity+33

# BINARY

# HDF5

- “Hierarchical Data Format” used across many industries
- PacBio read data *no longer* comes in bas.h5 / bax.h5 files (instead, you get BAM files) ... so let's forget about HDF5!



# BAM / CRAM - compressed SAM

- **\* Don't dump binary formats to your terminal / shell ...**
- **Indexing both BAM and CRAM allow *rapid* random read access to any coordinate range, without uncompressing whole file first**
- **CRAM restricts sequence alphabet, so compression ratio can be greater**
- **CRAM does *lossy* compression of base qualities, also helps compression ratio**

# 2bit

- **Old format used for sequence in UCSC Genome Browser**
- **Can only store 4 bases per position:**
  - **00 = A**
  - **01 = C**
  - **10 = G**
  - **11 = T**
  - **... N? Lower case acgt for soft masking? Nope ...**

*Questions ... comments ... confusion?*