

Sequence Alignment/Map Optional Fields Specification

The SAM/BAM Format Specification Working Group

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The master version of this document can be found at <https://github.com/samtools/hts-specs>.
This printing is version a5b2693-dirty from that repository, last modified on the date shown above.

This document is a companion to the *Sequence Alignment/Map Format Specification* that defines the SAM and BAM formats, and to the *CRAM Format Specification* that defines the CRAM format.¹ Alignment records in each of these formats may contain a number of optional fields, each labelled with a *tag* identifying that field's data. This document describes each of the predefined standard tags, and discusses conventions around creating new tags.

1 Standard tags

Predefined standard tags are listed in the following table and described in greater detail in later subsections. Optional fields are usually displayed as **TAG:TYPE:VALUE**; the *type* may be one of **A** (character), **B** (general array), **f** (real number), **H** (hexadecimal array), **i** (integer), or **Z** (string).

Tag	Type	Description
AM	i	The smallest template-independent mapping quality of segments in the rest
AS	i	Alignment score generated by aligner
BC	Z	Barcode sequence identifying the sample
BQ	Z	Offset to base alignment quality (BAQ)
BZ	Z	Phred quality of the unique molecular barcode bases in the OX tag
CC	Z	Reference name of the next hit
CG	B,I	BAM only: CIGAR in BAM's binary encoding if (and only if) it consists of >65535 operators
CM	i	Edit distance between the color sequence and the color reference (see also NM)
C0	Z	Free-text comments
CP	i	Leftmost coordinate of the next hit
CQ	Z	Color read base qualities
CS	Z	Color read sequence
CT	Z	Complete read annotation tag, used for consensus annotation dummy features
E2	Z	The 2nd most likely base calls
FI	i	The index of segment in the template
FS	Z	Segment suffix
FZ	B,S	Flow signal intensities
GC	?	Reserved for backwards compatibility reasons
GQ	?	Reserved for backwards compatibility reasons
GS	?	Reserved for backwards compatibility reasons
H0	i	Number of perfect hits
H1	i	Number of 1-difference hits (see also NM)
H2	i	Number of 2-difference hits
HI	i	Query hit index
IH	i	Number of stored alignments in SAM that contains the query in the current record

¹See [SAMv1.pdf](#) and [CRAMv3.pdf](#) at <https://github.com/samtools/hts-specs>.

LB	Z	Library
MC	Z	CIGAR string for mate/next segment
MD	Z	String for mismatching positions
MF	?	Reserved for backwards compatibility reasons
MI	Z	Molecular identifier; a string that uniquely identifies the molecule from which the record was derived
MQ	i	Mapping quality of the mate/next segment
NH	i	Number of reported alignments that contains the query in the current record
NM	i	Edit distance to the reference
OC	Z	Original CIGAR
OP	i	Original mapping position
OQ	Z	Original base quality
OX	Z	Original unique molecular barcode bases
PG	Z	Program
PQ	i	Phred likelihood of the template
PT	Z	Read annotations for parts of the padded read sequence
PU	Z	Platform unit
Q2	Z	Phred quality of the mate/next segment sequence in the R2 tag
QT	Z	Phred quality of the sample-barcode sequence in the BC (or RT) tag
QX	Z	Quality score of the unique molecular identifier in the RX tag
R2	Z	Sequence of the mate/next segment in the template
RG	Z	Read group
RT	Z	Barcode sequence (deprecated; use BC instead)
RX	Z	Sequence bases of the (possibly corrected) unique molecular identifier
SA	Z	Other canonical alignments in a chimeric alignment
SM	i	Template-independent mapping quality
SQ	?	Reserved for backwards compatibility reasons
S2	?	Reserved for backwards compatibility reasons
TC	i	The number of segments in the template
U2	Z	Phred probability of the 2nd call being wrong conditional on the best being wrong
UQ	i	Phred likelihood of the segment, conditional on the mapping being correct
X?	?	Reserved for end users
Y?	?	Reserved for end users
Z?	?	Reserved for end users

1.1 Additional Template and Mapping data

AM:i:int The smallest template-independent mapping quality of segments in the rest.

AS:i:score Alignment score generated by aligner.

BQ:Z:qualities Offset to base alignment quality (BAQ), of the same length as the read sequence. At the i -th read base, $BAQ_i = Q_i - (BQ_i - 64)$ where Q_i is the i -th base quality.

CC:Z:rname Reference name of the next hit; '=' for the same chromosome.

CG:B:I,encodedCigar Real CIGAR in its binary form if (and only if) it contains >65535 operations. This is a BAM file only tag as a workaround of BAM's incapability to store long CIGARs in the standard way. SAM and CRAM files created with updated tools aware of the workaround are not expected to contain this tag. See also the footnote in Section 4.2 of the SAM spec for details.

CP:i:pos Leftmost coordinate of the next hit.

E2:Z:bases The 2nd most likely base calls. Same encoding and same length as SEQ. See also U2 for associated quality values.

FI:i:int The index of segment in the template.

FS:Z:str Segment suffix.

H0:i:count Number of perfect hits.

H1:i:count Number of 1-difference hits (see also NM).

H2:i:count Number of 2-difference hits.

HI:i:i Query hit index, indicating the alignment record is the *i*-th one stored in SAM.

IH:i:count Number of stored alignments in SAM that contains the query in the current record.

MC:Z:cigar CIGAR string for mate/next segment.

MD:Z:[0-9]+((([A-Z]|\^[A-Z])+)[0-9]+)* String for mismatching positions.

The MD field aims to achieve SNP/indel calling without looking at the reference. For example, a string '10A5^AC6' means from the leftmost reference base in the alignment, there are 10 matches followed by an A on the reference which is different from the aligned read base; the next 5 reference bases are matches followed by a 2bp deletion from the reference; the deleted sequence is AC; the last 6 bases are matches. The MD field ought to match the CIGAR string.

MQ:i Mapping quality of the mate/next segment.

NH:i Number of reported alignments that contains the query in the current record.

NM:i Edit distance to the reference, including ambiguous bases but excluding clipping.

PQ:i Phred likelihood of the template, conditional on both the mapping being correct.

Q2:Z:qualities Phred quality of the mate/next segment sequence in the R2 tag. Same encoding as QUAL.

R2:Z:bases Sequence of the mate/next segment in the template. See also Q2 for any associated quality values.

SA:Z:(*rname*,*pos*,*strand*,*CIGAR*,*mapQ*,*NM*;)+ Other canonical alignments in a chimeric alignment, formatted as a semicolon-delimited list. Each element in the list represents a part of the chimeric alignment. Conventionally, at a supplementary line, the first element points to the primary line.

SM:i Template-independent mapping quality.

TC:i The number of segments in the template.

U2:Z Phred probability of the 2nd call being wrong conditional on the best being wrong. The same encoding and length as QUAL. See also E2 for associated base calls.

UQ:i Phred likelihood of the segment, conditional on the mapping being correct.

1.2 Metadata

RG:Z:readgroup The read group to which the read belongs. If @RG headers are present, then *readgroup* must match the RG-ID field of one of the headers.

LB:Z:library The library from which the read has been sequenced. If @RG headers are present, then *library* must match the RG-LB field of one of the headers.

PG:Z Program. Value matches the header PG-ID tag if @PG is present.

PU:Z:platformunit The platform unit in which the read was sequenced. If @RG headers are present, then *platformunit* must match the RG-PU field of one of the headers.

CO:Z:text Free-text comments.

1.3 Barcodes

DNA Barcodes can be used to identify the provenance of the underlying the reads. There are currently several usages of barcodes that may co-exist: Sample Barcode, Cell Barcode, and Molecular-barcode.

- The sample barcode actually identifies the Library so that multiple libraries can be combined and sequenced together. After sequencing, each barcode identifies the "readgroup" which is the assigned to the correct library and sample/individual.
- The cell barcode is similar to the sample barcode but the expectation is that every cell will be randomly assigned a barcode (whose content could be predetermined or random.) The cell barcode helps identify when reads came from different cell in a "single-cell" sequencing experiment. Unlike in Sample Barcode there is no control over which cell "gets" which barcode.
- The Molecular barcode can be used to identify the "original" molecule that is being replicated, so that duplicate marking isn't overzealous, or to identify the long molecule form which linked reads come from.

BC:Z:sequence Barcode sequence (Identifying the sample/library), with any quality scores (optionally) stored in the **QT** tag. The **BC** tag should match the **QT** tag in length. In the case of multiple unique molecular identifiers (e.g., one on each end of the template) the recommended implementation concatenates all the barcodes and places a hyphen ('-') between the barcodes from the same template.

QT:Z:qualities Phred quality of the sample-barcode sequence in the **BC** (or **RT**) tag. Same encoding as **QUAL**, i.e., Phred score + 33. In the case of multiple unique molecular identifiers (e.g., one on each end of the template) the recommended implementation concatenates all the quality strings with spaces (' ') between the different strings from the same template.

RX:Z:sequence+ Sequence bases from the unique molecular identifier. These could be either corrected or uncorrected. Unlike **MI**, the value may be non-unique in the file. Should be comprised of a sequence of bases. In the case of multiple unique molecular identifiers (e.g., one on each end of the template) the recommended implementation concatenates all the barcodes with a hyphen ('-') between the different barcodes.

If the bases represent corrected bases, the original sequence can be stored in **OX** (similar to **OQ** storing the original qualities of bases.)

QX:Z:qualities+ Phred quality of the unique molecular identifier sequence in the **RX** tag. Same encoding as **QUAL**, i.e., Phred score + 33. The qualities here may have been corrected (Raw bases and qualities can be stored in **OX** and **BZ** respectively.) The lengths of the **QX** and the **RX** tags must match. In the case of multiple unique molecular identifiers (e.g., one on each end of the template) the recommended implementation concatenates all the quality strings with a space (' ') between the different strings.

MI:Z:str Molecular Identifier. A unique ID within the SAM file for the source molecule from which this read is derived. All reads with the same **MI** tag represent the group of reads derived from the same source molecule.

OX:Z:sequence+ Raw (uncorrected) unique molecular identifier bases, with any quality scores (optionally) stored in the **BZ** tag. In the case of multiple unique molecular identifiers (e.g., one on each end of the template) the recommended implementation concatenates all the barcodes with a hyphen ('-') between the different barcodes.

BZ:Z:qualities+ Phred quality of the (uncorrected) unique molecular identifier sequence in the **OX** tag. Same encoding as **QUAL**, i.e., Phred score + 33. The **OX** tags should match the **BZ** tag in length. In the case of multiple unique molecular identifiers (e.g., one on each end of the template) the recommended implementation concatenates all the quality strings with a space (' ') between the different strings.

RT:Z:sequence Deprecated alternative to **BC** tag originally used at Sanger.

1.4 Original data

OC:Z:cigar Original CIGAR, usually before realignment.

OP:i:pos Original mapping position, usually before realignment.

OQ:Z:qualities Original base quality, usually before recalibration. Same encoding as QUAL.

1.5 Annotation and Padding

CT:Z:strand;type(;key(=value))* Complete read annotation tag, used for consensus annotation dummy features.

The CT tag is intended primarily for annotation dummy reads, and consists of a *strand*, *type* and zero or more *key=value* pairs, each separated with semicolons. The *strand* field has four values as in GFF3, and supplements FLAG bit 0x10 to allow unstranded ('.'), and stranded but unknown strand ('?') annotation. For these and annotation on the forward strand (*strand* set to '+'), do not set FLAG bit 0x10. For annotation on the reverse strand, set the *strand* to '-' and set FLAG bit 0x10.

The *type* and any *keys* and their optional *values* are all percent encoded according to RFC3986 to escape meta-characters '=', '%', ';', '|', or non-printable characters not matched by the `isprint()` macro (with the C locale). For example a percent sign becomes '%25'.

PT:Z:start;end;strand;type(;key(=value))*(\|start;end;strand;type(;key(=value)))* Read annotations for parts of the padded read sequence.

The PT tag value has the format of a series of tags separated by '|', each annotating a sub-region of the read. Each tag consists of *start*, *end*, *strand*, *type* and zero or more *key=value* pairs, each separated with semicolons. *Start* and *end* are 1-based positions between one and the sum of the M/I/D/P/S/=X CIGAR operators, i.e. SEQ length plus any pads. Note any editing of the CIGAR string may require updating the 'PT' tag coordinates, or even invalidate them. As in GFF3, *strand* is one of '+' for forward strand tags, '-' for reverse strand, '.' for unstranded or '?' for stranded but unknown strand. The *type* and any *keys* and their optional *values* are all percent encoded as in the CT tag.

1.6 Technology-specific data

FZ:B,S:intensities Flow signal intensities on the original strand of the read, stored as `(uint16_t) round(value * 100.0)`.

1.6.1 Color space

CM:i:distance Edit distance between the color sequence and the color reference (see also NM).

CS:Z:sequence Color read sequence on the original strand of the read. The primer base must be included.

CQ:Z:qualities Color read quality on the original strand of the read. Same encoding as QUAL; same length as CS.

2 Locally-defined tags

You can freely add new tags. Note that tags starting with 'X', 'Y', or 'Z' and tags containing lowercase letters in either position are reserved for local use and will not be formally defined in any future version of this specification.

If a new tag may be of general interest, it may be useful to have it added to this specification. Additions can be proposed by opening a new issue at <https://github.com/samtools/hts-specs/issues> and/or by sending email to samtools-devel@lists.sourceforge.net.

Appendix A SAM Tags History

This lists the date of each tagged SAM version along with changes that have been made while that version was current.

1.5: 23 May 2013 to current

- Add UMI-related tags (RX, QX, OX, BZ, MI) and clarified usage of sample barcode tag BC. (August 2017)
- SAMtags.txt (this file) created with tags from SAMv1