

CATS
(Cancer Genomic Test Standardized)
Format

Document for details

By Section of Genomic Data Management,
C-CAT

v1.6.0

2026/05/11

Contents

1. Introduction.....	4
1-1. Objectives.....	4
1-2. Terms	4
1-3. About the condition field	5
1-4. Format information	5
2. Matters specific to C-CAT	6
2-1. Sending format of files	6
2-2. Scope of inputs	6
2-3. Requests	6
2-4. Notes	7
3. metaData tag	8
3-1. schemaVersion key	8
3-2. referenceGenome tag	8
3-2-1. Tags within referenceGenome tag	8
3-2-2. Examples of referenceGenome tag.....	9
3-3. configOptions tag.....	9
3-3-1. Tags within configOptions tag.....	10
3-3-2. Example of configOptions tag.....	14
3-4. comments tag	15
3-4-1. Tags within comments tag	15
3-4-2. Example of comments tag	15
4. testInfo tag.....	17
4-1. Tags within testInfo tag	17
4-2. Example of testInfo tag.....	17
5. sequencingSamples tag.....	19
5-1. Tags within sequencingSamples tag.....	19
5-2. Example of sequencingSamples tag.....	22
6. variants tag	25
6-1. shortVariants tag	25
6-1-1. Tags within shortVariants tag	25
6-1-2. Example of shortVariants tag	32
6-2. copyNumberAlterations tag	38
6-2-1. Tags within copyNumberAlterations tag	38
6-2-2. Example of copyNumberAlterations tag	41
6-3. structuralVariations tag.....	43
6-3-1. Tags within structuralVariations tag.....	43

6-3-2. Example of structuralVariations tag	50
7. otherBiomarkers tag	57
7-1. Tags within otherBiomarkers tag	57
7-2. Example of otherBiomarkers tag	58
8. expressions tag.....	60
8-1. Tags within expressions tag	60
8-2. Example of expressions tag.....	62
9. alterationNotes tag	64
9-1. Tag within alterationNotes tag.....	64
9-2. Example of alterationNotes tag	65
10. armLevelChanges tag	67
10-1. Tags within armLevelChanges tag	67
10-2. Example of armLevelChanges tag	68
11. nonHumanContents tag	70
11-1. Tags within nonHumanContents tag	70
11-2. Example of nonHumanContents tag	70
12. wildTypes tag	72
12-1. Tags within wildTypes tag	72
12-2. Example of wildTypes tag.....	72
13. druggability tag.....	74
13-1. Tags within druggability tag.....	74
13-2. Example of druggability tag.....	75
14. pathogenicity tag	78
14-1. Tags within pathogenicity tag	78
14-2. Example of pathogenicity tag	79
15. Other notes.....	81
15-1. itemId	81
15-1-1. Example of itemId description	81
15-2. matePieceLocation	82
15-2-1. Example of matePieceLocation description	82
16. For inquires	84

1. Introduction

1-1. Objectives

Differences in the formats for results in comprehensive genomic profiling tests of cancer makes it difficult for a third party to annotate alterations with candidate drugs and clinical trials using the same software under a simple framework. To efficiently promote uniformity and homogeneity in the interpretation of cancer genomic test results, it is necessary to define a standardized format to present alteration data in cancer comprehensive genome profiling tests.

This document describes the CATS (cancer genomic test standardized) format, a standardized format for presenting alteration data in cancer comprehensive genomic profiling tests. The data schema of the CATS format is defined by the JSON definition file "schema.json", the specifications of which are explained in this document.

This format is used as follows: A laboratory that performs cancer comprehensive genome profiling tests sends alteration data in the CATS format to a test annotation organization such as C-CAT. The organization annotates alteration data in the CATS format with candidate drugs and clinical trials, taking into account the clinical data of patients. This format only applies to alteration data, not clinical data, because clinical data are stored in electronic medical records in hospitals and are unlikely to be accessible to testing companies.

1-2. Terms

Term	Description
Test report	A report summarizing the results of comprehensive cancer genome profiling tests issued by testing companies.
Test annotation organization	An organization that receives alteration data in cancer comprehensive genome profiling tests from testing companies and clinical data from hospitals and uses the cancer knowledge base to annotate each alteration with candidate drugs and clinical trials. For example, C-CAT.
Test annotation document	A document that associates individual alterations in a cancer comprehensive genomic profiling test with candidate drugs and clinical trials for each patient. An example is the C-CAT Findings document.
Cancer knowledge base	A database that associates alterations in cancer with candidate drugs and clinical trials. Examples include C-CAT CKDB (<u>c</u> ancer <u>k</u> nowledge <u>d</u> atab <u>a</u> se) and OncoKB (Chakravarty et al, 2017, JCO Precision Oncology).

1-3. About the condition field

- Required: Required field when the JSON parent tag exists. When the parent tag is optional, the required field is described as "[required]" because it is assumed that the parent tag exists.
- Optional: Recommended field that may be associated with drug and clinical trial information in test annotation documents or that increases the accuracy of test annotation documents according to test reports. If it is strongly recommended, it is described as "optional (recommended)".

1-4. Format information

- Character code: UTF8
- Type: JSON
- Extension: json

2. Matters specific to C-CAT

2-1. Sending format of files

Alteration data in cancer comprehensive genomic profiling tests should be sent to C-CAT from the testing company in CATS (cancer genomic test sandardized) format.

2-2. Scope of inputs

Quality-assured data on alterations are in the scope of inputs in CATS format. Please do not input alterations that are determined as to be false positives. You can choose whether or not to show alterations and their associated knowledge base annotations in C-CAT Findings by the tag ("reported" and "grade") described later.

Note: In the following sections, the terms "Molecular Alterations" "Genetic Alterations" and "Annotation Of Molecular Alterations" as well as the items included in each category, are shown in the table below.

Term	Items
Molecular Alterations	shortVariants, copyNumberAlterations, structuralVariations, otherBiomarkers, expressions, armLevelChanges, nonHumanContents, wildTypes
Genetic Alterations	shortVariants, copyNumberAlterations, structuralVariations, wildTypes
Annotation Of Molecular Alterations	druggability, pathogenicity, alterationNotes

With these above attentions, please be sure to input data on alterations in test report or equivalent report. Please be sure to output data on alterations that are approved by the concerned authorities in C-CAT Findings.

2-3. Requests

- It is recommended to provide inputs for as many optional fields as possible. As a result, more information on drugs or clinical trials may be added to C-CAT Findings, and C-CAT Findings based on test reports will be more accurate. Additionally, more information linked to these fields may be added in future versions, even if such information does not appear in the current version of C-CAT Findings. C-CAT may separately contact the testing company for inputs in optional fields as necessary.

- Please try your best to input quality-assured data on all alterations (including those with the tag of "reported": false, as explained below) in CATS format. Otherwise, in case of any changes in the format or specifications of test reports, C-CAT may send inquiries to the testing company and the production of C-CAT Findings may be delayed.

2-4. Notes

Detailed precautions specific to C-CAT are indicated by ‘*’ in the Description below.

3. metaData tag

This tag is used to define the metadata.

It contains 4 keys: schemaVersion, referenceGenome, configOptions, and comments.

Key	Condition	Data type	Description
metaData	required	object	Aggregation tag for metadata.

3-1. schemaVersion key

Key	Condition	Data type	Description
schemaVersion	required	string regex: ^[0-9\.\-]+	Schema version of this format.

3-2. referenceGenome tag

Key	Condition	Data type	Description
referenceGenome	required	object	For information on a reference genome sequence.

3-2-1. Tags within referenceGenome tag

Key	Condition	Data type	Description
name	optional	string regex: ^.+	Name of a reference genome sequence used in your test.
grcRelease	required	string regex: ^GRC.+	GRC (Genome Reference Consortium) release ID of a reference genome sequence.
descriptions	optional	array (length: 0-N, string regex: ^.+)	Description of the reference genome sequence in the name tag. See the contents tag within the comments tag for usable languages and new lines.

3-2-2. Examples of referenceGenome tag

(Example1. for NCBI)

```
"referenceGenome": {  
  "name": "GRCh38.p13",  
  "grcRelease": "GRCh38.p13",  
  "descriptions": [  
    "Homo sapiens (human) genome assembly GRCh38 (hg38) from the Genome Reference Consortium."  
  ]  
}
```

(Example2. for UCSC)

```
"referenceGenome": {  
  "name": "hg38Patch11",  
  "grcRelease": "GRCh38.p11",  
  "descriptions": [  
    "GRCh38 Genome Reference Consortium Human Reference 38 (GCA_000001405.22))"  
  ]  
}
```

(Example3. for GDC)

```
"referenceGenome": {  
  "name": "GRCh38.d1.vd1",  
  "grcRelease": "GRCh38",  
  "descriptions": [  
    "Homo sapiens (human) genome assembly GRCh38 (hg38) from GDC, GRCh38.d1.vd1"  
  ]  
}
```

3-3. configOptions tag

This tag controls matching to cancer knowledge bases, such as C-CAT CKDB.

Key	Condition	Data type	Description
configOptions	optional	object	Aggregation tag that controls matching to cancer knowledge bases.

3-3-1. Tags within configOptions tag

Key	Condition	Data type	Description
typeLabelsInterpretedAsKbAmplification	optional	array (length: 1-4, string) [choice]	<p>The testing company's labels for genetic alterations, which are interpreted as copyNumberAlteration's "amplification" (copy number amplification) in cancer knowledge bases.</p> <p>Choose from the following options (must not be duplicated in an array).</p> <ul style="list-style-type: none"> • "copyNumberAlterationType: amplification" • "copyNumberAlterationType: gain" • "copyNumberAlterationType: duplication" • "structuralVariationType: duplication" <p>(Default: "copyNumberAlterationType: amplification", "copyNumberAlterationType: gain", "copyNumberAlterationType: duplication")</p>
typeLabelsInterpretedAsKbLoss	optional	array (length: 1-5, string) [choice]	<p>The testing company's labels for genetic alterations, which are interpreted as copyNumberAlteration's "loss" (copy number loss) in cancer knowledge bases.</p> <p>Choose from the following options (must not be duplicated in an array).</p> <ul style="list-style-type: none"> • "copyNumberAlterationType: loss" • "copyNumberAlterationType: deletion" • "copyNumberAlterationType: homozygous deletion" • "structuralVariationType: deletion" • "structuralVariationType: exon skipping" <p>(Default: "copyNumberAlterationType: loss", "copyNumberAlterationType: deletion", "copyNumberAlterationType: homozygous deletion")</p>

typeLabelsInterpretedAsKbGeneFusion	optional	array (length: 1-4, string) [choice]	<p>The testing company's labels for genetic alterations, which are interpreted as structuralVariation's "geneFusion" (gene fusion) in cancer knowledge bases.</p> <p>Choose from the following options (must not be duplicated in an array).</p> <ul style="list-style-type: none"> • "structuralVariationType: gene fusion" • "structuralVariationType: gene fusion and frameshift variant" • "structuralVariationType: bidirectional gene fusion" • "structuralVariationType: other" <p>(Default: "structuralVariationType: gene fusion", "structuralVariationType: gene fusion and frameshift variant", "structuralVariationType: bidirectional gene fusion")</p>
typeLabelsInterpretedAsKbInversion	optional	array (length: 1-3, string) [choice]	<p>The testing company's labels for genetic alterations, which are interpreted as structuralVariation's "inversion" (gene inversion) in cancer knowledge bases.</p> <p>Choose from the following options (must not be duplicated in an array).</p> <ul style="list-style-type: none"> • "structuralVariationType: inversion" • "structuralVariationType: truncation" • "structuralVariationType: other" <p>(Default: "structuralVariationType: inversion")</p>
typeLabelsInterpretedAsKbDeletion	optional	array (length: 1-8, string) [choice]	<p>The testing company's labels for genetic alterations, which are interpreted as structuralVariation's "deletion" (gene deletion) in cancer knowledge bases.</p> <p>Choose from the following options (must not be duplicated in an array).</p> <ul style="list-style-type: none"> • "structuralVariationType: deletion" • "structuralVariationType: truncation" • "structuralVariationType: splice variant" • "structuralVariationType: exon skipping" • "structuralVariationType: other" • "copyNumberAlterationType: deletion"

			<ul style="list-style-type: none"> • "copyNumberAlterationType: homozygous deletion" • "copyNumberAlterationType: loss" (Default: "structuralVariationType: deletion")
typeLabelsInterpretedAsKbDuplication	optional	array (length: 1-6, string) [choice]	<p>The testing company's labels for genetic alterations, which are interpreted as structuralVariation's "duplication" (gene duplication) in cancer knowledge bases.</p> <p>Choose from the following options (must not be duplicated in an array).</p> <ul style="list-style-type: none"> • "structuralVariationType: duplication" • "structuralVariationType: tandem duplication" • "structuralVariationType: other" • "copyNumberAlterationType: duplication" • "copyNumberAlterationType: amplification" • "copyNumberAlterationType: gain" (Default: "structuralVariationType: duplication", "structuralVariationType: tandem duplication")
typeLabelsInterpretedAsKbTruncation	optional	array (length: 1-7, string) [choice]	<p>The testing company's labels for genetic alterations, which are interpreted as structuralVariation's "truncation" (gene truncation) in cancer knowledge bases.</p> <p>Choose from the following options (must not be duplicated in an array).</p> <ul style="list-style-type: none"> • "structuralVariationType: truncation" • "structuralVariationType: deletion" • "structuralVariationType: inversion" • "structuralVariationType: other" • "copyNumberAlterationType: loss" • "copyNumberAlterationType: deletion" • "copyNumberAlterationType: homozygous deletion" (Default: "structuralVariationType: truncation")
typeLabelsInterpretedAsKbExonSkipping	optional	array (length: 1-4, string) [choice]	<p>The testing company's labels for genetic alterations, which are interpreted as structuralVariation's "exon skipping" (exon skipping) in cancer knowledge bases.</p> <p>Choose from the following options (must not be duplicated in an array).</p>

			<ul style="list-style-type: none"> • "structuralVariationType: exon skipping" • "structuralVariationType: splice variant" • "structuralVariationType: deletion" • "structuralVariationType: other" (Default: "structuralVariationType: exon skipping")
typeLabelsInterpretedAsTranslocation	optional	array (length: 1-2, string) [choice]	<p>The testing company's labels for genetic alterations, which are interpreted as structuralVariation's "translocation" (translocation) in cancer knowledge bases.</p> <p>Choose from the following options (must not be duplicated in an array).</p> <ul style="list-style-type: none"> • "structuralVariationType: translocation" • "structuralVariationType: other" (Default: "structuralVariationType: translocation")
typeLabelsInterpretedAsKbStructuralVariation	optional	array (length: 1-18, string) [choice]	<p>The testing company's labels for genetic alterations, which are interpreted as structuralVariation's "structuralVariation" (gene structuralVariation) in cancer knowledge bases.</p> <p>Choose from the following options (must not be duplicated in an array).</p> <ul style="list-style-type: none"> • "structuralVariationType: gene fusion" • "structuralVariationType: gene fusion and frameshift variant" • "structuralVariationType: bidirectional gene fusion" • "structuralVariationType: duplication" • "structuralVariationType: tandem duplication" • "structuralVariationType: deletion" • "structuralVariationType: inversion" • "structuralVariationType: truncation" • "structuralVariationType: splice variant" • "structuralVariationType: exon skipping" • "structuralVariationType: translocation" • "structuralVariationType: other" • "copyNumberAlterationType: amplification" • "copyNumberAlterationType: gain" • "copyNumberAlterationType: duplication" • "copyNumberAlterationType: loss" • "copyNumberAlterationType: deletion"

			<ul style="list-style-type: none"> • "copyNumberAlterationType: homozygous deletion" (Default: "structuralVariationType: other")
--	--	--	-------------------------------------------------------------------------------------------------------------------------------------------------

3-3-2. Example of configOptions tag

(Example)

```
"configOptions": {
  "typeLabelsInterpretedAsKbAmplification": [
    "copyNumberAlterationType: amplification",
    "copyNumberAlterationType: gain",
    "copyNumberAlterationType: duplication"
  ],
  "typeLabelsInterpretedAsKbLoss": [
    "copyNumberAlterationType: loss",
    "copyNumberAlterationType: deletion",
    "structuralVariationType: deletion"
  ],
  "typeLabelsInterpretedAsKbGeneFusion": [
    "structuralVariationType: gene fusion",
    "structuralVariationType: gene fusion and frameshift variant",
    "structuralVariationType: bidirectional gene fusion"
  ],
  "typeLabelsInterpretedAsKbInversion": [
    "structuralVariationType: inversion"
  ],
  "typeLabelsInterpretedAsKbDeletion": [
    "structuralVariationType: deletion"
  ],
  "typeLabelsInterpretedAsKbDuplicationDeletion": [
    "structuralVariationType: duplication",
    "structuralVariationType: tandem duplication"
  ],
  "typeLabelsInterpretedAsKbTruncation": [
    "structuralVariationType: truncation"
  ]
}
```

3-4. comments tag

You can comment on molecular alterations, annotation of molecular alterations, and information on sequencing samples (sequencingSamples). This tag contains the itemIds and contents tags.

Key	Condition	Data type	Description
comments	optional	array (length: 1-N, object)	Aggregation tag for comment information. Each object in the array must be unique.

3-4-1. Tags within comments tag

Key	Condition	Data type	Description
itemIds	required	array (length: 0-N, string regex: ^.+)\$)	The itemIds (multiple unique itemIds possible) to indicate alterations and information on sequencing samples (sequencingSamples). Possible to comment on a test overall if you set the length of this key to be zero. * The content will not be shown in C-CAT Findings, if itemId is specified.
contents	required	array (length: 1-N, string regex: ^.+)\$)	The content of the comment for itemId. The description can be in English or Japanese. Please use array elements if you make new lines, because we ignore line feed codes in this tag. * The total number of characters for each element is within 4,000 characters including the line feed code.

3-4-2. Example of comments tag

(Example)

```
"comments": [  
  {
```

```
    "itemIds": [],
```

Note: If the length of the itemIds array is zero, it represents a comment on this test overall.

```
    "contents": [  
      "Amplification of the FGFR1 gene is observed in 5 to 20% of squamous cell
```

```
      carcinomas, and it has been reported that FGFR1 is sensitive to FGFR inhibitors in  
      vitro.",
```

```
      "FGFR2 and FGFR3 gene activating mutations and FGFR3 gene fusions have been  
      reported one after another, and their frequency is low at around 3%, but therapeutic  
      effects with FGFR inhibitors are expected."
```

Note: If you want to make new lines in test annotation documents, sentences or phrases should be separated as elements of an array.

```
]
},
{
  "itemIds": [
    "variant-1"
```

Note: The itemId of the mutation should be included to comment on a specific mutation.

```
],
  "contents": [
    "TSC1 functions independently of TSC2 and mTORC1."
  ]
},
{
  "itemIds": [
    "variant-1",
    "variant-5"
```

Note: You can comment on multiple mutations altogether by listing multiple itemIds.

```
],
  "contents": [
    "Although CD4 T cell percentage in Tsc1-/- mice was not strongly affected by Bim deficiency in vivo, TCR-mediated apoptosis of Tsc1-/- Bcl2l11-/- double knockout CD4 T cells was less pronounced compared with that of Tsc1-/- cells. (Kai Yang et al.)"
  ]
}
]
```


4. testInfo tag

This tag is used to provide test information.

Key	Condition	Data type	Description
testInfo	required	object	Aggregation tag for test information.

4-1. Tags within testInfo tag

Key	Condition	Data type	Description
testId	required	string regex: ^.+	Any ID used by the testing company.
testType	required	string [choice]	The combination of specimens used in the test. <ul style="list-style-type: none">• "tumor-only": test using tumor samples only• "tumor and matched-normal": test using tumor and matched normal samples• "tumor-only (cell-free)": test using cell-free tumor samples only• "tumor (cell-free) and matched-normal": test using cell-free tumor samples and normal samples
targetRegion Version	optional	string regex: ^.+	Target area version defined in the bed file.
softwareName	optional	string regex: ^.+	Name of gene analysis software.
softwareVersion	optional	string regex: ^.+	Version of gene analysis software.
panelName	required	string regex: ^.+	Name of your genomic profiling (gene panel) test. * Please inform C-CAT beforehand if you want to use a genomic test NOT approved to use under National Health Insurance.
panelVersion	required	string regex: ^.+	Version of your test.

4-2. Example of testInfo tag

(Example)

```
"testInfo": {  
  "testId": "123456789012319000001",  
  "testType": "tumor and matched-normal",  
  "targetRegionVersion": "target region A",  
  "softwareName": "variant caller A",  
  "softwareVersion": "ver.1.2",  
  "panelName": "Multi-gene Panel A",  
  "panelVersion": "ver.1.03-00"  
}
```

5. sequencingSamples tag

This tag is used to provide the information on sequencing results in the NGS run.

Key	Condition	Data type	Description
sequencingSamples	required	array (length: 1-N, object)	Aggregation tag for information on sequencing results. Each object in the array must be unique.

The maximum length of this array is 4, since the tumorOrNormal tag and the nucleicAcid tag can each take two values "tumor" or "normal" and "DNA" or "RNA", respectively.

5-1. Tags within sequencingSamples tag

Key	Condition	Data type	Description
itemId	required	string regex: ^.+ \$	The ID of the sample. It must be unique within this case report.
tumorOrNormal	required	string [choice]	Whether the sequencing results are of tumor specimen or normal specimen. • "tumor" • "normal"
nucleicAcid	required	string [choice]	Whether the sequencing results are derived from DNA or RNA samples. • "DNA": DNA sample • "RNA": RNA sample
resultQuality	optional	object	The information of the sequencing results determined by the testing company.
state	[required]	string [choice]	The quality of the sequencing results. • "pass": Pass • "reference": Reference purpose only • "fail": Fail
reason	optional	string regex: ^.+ \$	The reason and details for the state. e.g.) Possibility of contamination. Low read depth.
specimenType	optional	string regex: ^.+ \$	The name of the specimen type written in the test report. e.g.) FFPE Intact

medianReadDepth	optional (recommended)	number	Median read depth.
meanReadDepth	optional (recommended)	number	Mean read depth.
readCountMetrics	optional (recommended)	array (length: 1-N, object)	Information on the number of reads. Each object in the array must be unique.
value	[required]	number	Measured value.e.g.) 76442556
labelName	[required]	string [choice]	Label for measured value. <ul style="list-style-type: none"> • "allReadCount": the number of all reads including unmapped reads • "mappedReadCount": the number of mapped reads * Please inform C-CAT beforehand if you use options other than the above.
mappingType	[required]	string [choice]	Type of mapping. <ul style="list-style-type: none"> • "onTarget": target • "totalMapped": target and off target • "all": all reads including unmapped * Please inform C-CAT beforehand if you use options other than the above.
duplicateType	[required]	string [choice]	Type of duplication. <ul style="list-style-type: none"> • "unique": unique read • "duplicate": duplicate read • "all": all read * Please inform C-CAT beforehand if you use options other than the above. * If mappingType is "all", then duplicateType is "all".
reported	[required]	boolean	Whether or not the number of reads is reported in test report or similar documents.

readPercentageMetrics	optional (recommended)	array (length: 1-N, object)	Information on the percentage of reads. Each object in the array must be unique.
value	[required]	number	Measured value. e.g.) 91.52
labelName	[required]	string [choice]	Label for measured value. • "duplicateReadPercentage": percentage of duplicate reads • "mappedReadPercentage": percentage of mapped reads * Please inform C-CAT beforehand if you use options other than the above.
numerator	[required]	object	Numerator information.
value	optional	number	Refer to the description in the readCountMetrics tag.
mappingType	required	string [choice]	Refer to the description in the readCountMetrics tag.
duplicateType	required	string [choice]	Refer to the description in the readCountMetrics tag.
denominator	[required]	object	Denominator information.
value	optional	number	Refer to the description in the readCountMetrics tag.
mappingType	required	string [choice]	Refer to the description in the readCountMetrics tag.
duplicateType	required	string [choice]	Refer to the description in the readCountMetrics tag.

reported	[required]	boolean	Whether or not the percentage of reads is reported in test report or similar documents.
tumorContent	optional	array (length: 1-N, object)	The value of the tumor content written in the test report.
value	[required]	number	Tumor content. The unit is %.
type	[required]	string [choice]	Type of tumor content. <ul style="list-style-type: none"> • "pathological": pathologically measured. • "estimate": estimated from sequence data. • "unclear": unclear.
descriptions	optional	array (length: 1-N, string regex: ^.+)\$)	Description of sequencing sample. Refer to the contents tag within the comments tag for usable languages and new lines.

5-2. Example of sequencingSamples tag

(Example)

```
"sequencingSamples": [
{
  "itemId": "sequence-1-tumor-dna",
  "tumorOrNormal": "tumor",
  "nucleicAcid": "DNA",
  "resultQuality": {
    "state": "pass"
  },
  "specimenType": "FFPE",
  "medianReadDepth": 247,
  "meanReadDepth": 238,
  "readPercentageMetrics": [
    {
      "value": 91.52,
      "labelName": "duplicateReadPercentage",
      "numerator": {
        "mappingType": "totalMapped",
        "duplicateType": "duplicate"
      },
      "denominator": {
```

```

    "mappingType": "totalMapped",
    "duplicateType": "all"
  },
  "reported": true
},
{
  "value": 87.31,
  "labelName": "mappedReadPercentage",
  "numerator": {
    "mappingType": "onTarget",
    "duplicateType": "unique"
  },
  "denominator": {
    "mappingType": "totalMapped",
    "duplicateType": "unique"
  },
  "reported": true
}
],
},
{
  "itemId": "sequence-2-tumor-rna",
  "tumorOrNormal": "tumor",
  "nucleicAcid": "RNA",
  "resultQuality": {
    "state": "reference",
    "reason": "degraded"
  },
  "readCountMetrics": [
    {
      "value": 11928428,
      "labelName": "allReadCount",
      "mappingType": "all",
      "duplicateType": "all",
      "reported": true
    },
    {
      "value": 9542742,
      "labelName": "mappedReadCount",
      "mappingType": "totalMapped",
      "duplicateType": "all",
      "reported": true
    }
  ],
  "tumorContent": [
    {
      "value": 50,
      "type": "pathological"
    }
  ]
},

```

```
{
  "itemId": "sequence-3-normal-dna",
  "tumorOrNormal": "normal",
  "nucleicAcid": "DNA"
}
]
```


6. variants tag

This tag is used to provide information on detected genetic alterations. It contains shortVariants tag, copyNumberAlterations tag, and structuralVariations tag.

Key	Condition	Data type	Description
variants	optional	object	Aggregation tag for information on genetic alterations.

6-1. shortVariants tag

This tag is used to define information on SNV (single nucleotide variation), insertion, deletion, delins (simultaneous insertion and deletion), indel (insertion and deletion), and MNV (multiple-nucleotide variation).

Key	Condition	Data type	Description
shortVariants	optional	array (length: 1-N, object)	Aggregation tag for information on single nucleotide variation (SNV), insertion (insertion), deletion (deletion), deletion and insertion (delins), insertion and deletion (indel), and multiple-nucleotide variant (MNV) of nucleotides. Each object in the array must be unique.

6-1-1. Tags within shortVariants tag

key	Condition	Data type	Description
itemId	required	string regex: ^.+	An ID assigned to a genetic alteration. It must be unique within this case report.
chromosome	required	string regex: ^[a-zA-Z0-9_#-]+\$	Chromosome number.
position	required	integer	Physical position in a chromosome. Please use the 1-based coordinate system and describe according to VCF v4.5 (e.g., as stated on page 21 of VCF v4.5, when the reference base is atCga and

			the variant base is at-ga, and C position in the reference base is 3, denote the physical position as "position": 2, "referenceAllele": "TC", and "alternateAllele": "T". As described below, "referenceAllele" represents a reference base and "alternateAllele" represents a variant base.)
cytoband	optional	string regex: ^.+	Cytoband where the variant exists.
referenceAllele	required	string regex: ^[ACGTN]+\$	Reference base. Please describe according to VCF v4.5 (see the example above).
alternateAllele	required	string regex: ^[ACGTN¥*]+\$	Variant base. Please describe according to VCF v4.5 (see the example above). Multi-alleles – tri-alleles and more – should be listed as different elements in the shortVariants tag. In that case, indicate the itemIds and that the variants are multi-alleles in the comments tag.
alternateAlleleFrequency	required	number	Variant allele frequency (ranging from 0 to 1).
totalReadDepth	optional (recommend)	integer	Total read depth (minimum value 0).
alternateAlleleReadDepth	optional (recommend)	integer	Variant allele read depth (minimum value 0).

variantType	optional	string [choice]	<p>Short variant type written in the test report. Select one from:</p> <ul style="list-style-type: none"> • "SNV" • "insertion" • "deletion" • "delins" • "indel" • "MNV" <p>* Please inform C-CAT beforehand if you use other types.</p>
transcripts	required	array (length: 1-N, object)	<p>Information on the representative transcript. Each object in the array must be unique.</p> <p>* When creating C-CAT Findings, the sequence is scanned from the beginning and the first transcript information for each gene is used.</p>
transcriptId	[required]	string regex: ^[^\$s]+\$	<p>Transcription product ID (e.g., NM_000368.<u>4</u>). It is strongly recommended to include sub-numbers (underlined in red above) for accurate computation. If no transcriptId is present due to a mutation in an intergenic region, then you can define "transcriptId": null.</p>
transcriptDatabaseName	[required]	string [choice]	<p>Database of the transcript ID.</p> <p>Select from the following two options.</p> <ul style="list-style-type: none"> • "RefSeq" • "Ensembl"

			If the transcriptId is null, then you can define "transcriptDatabaseName": null.
transcriptDatabaseVersion	optional	string regex: ^.+	Version of the database. * If this is not defined, the version determined by C-CAT will be used.
regionStructure	optional	array (length: 1-N, object)	Information of the variant's structure.
type	[required]	string [choice]	Additional information on gene position, as written in the test report. Select one from: <ul style="list-style-type: none"> • "promoter" • "upstream" • "downstream" • "intergenic" • "5'UTR" • "3'UTR" • "exon" • "intron" * Please consult C-CAT if you want to use other types.
number	optional	integer	If "exon" or "intron" is selected for type, entering the exon number or intron number is recommended.
totalNumber	optional	integer	If "exon" or "intron" is selected for type, entering the exon number or intron number is recommended.
geneSymbol	[required]	string regex: ^[^\$]+	Name of the gene, as written in the test report. Enter "geneSymbol": null if a gene does not exist. Enter an associated gene such as "TERT promoter", "TERT" should be listed.

			The "promoter" is described in the "regionType" tag.
geneSymbolSource	optional (recommend)	string regex: ^.+	The source for the geneSymbol written in the test report. e.g. HGNC. Ensembl
geneSymbolId	optional (recommend)	string regex: ^.+	ID for the geneSymbol written in the test report. e.g. if "geneSymbol": "BRAF" "geneSymbolSource": "HGNC" then "geneSymbolId": "1097"
regionName	optional	string regex: ^.+	Name of specific region where the alteration exists, which is written in the test report. e.g. D4Z4 repeat
strand	optional	string [choice]	Direction of transcription. If the orientation is the same as that of the reference genome sequence, it is "+"; if the orientation is opposite, it is "-". If the transcriptId is null, then you can define "strand": null.
cdsChange	[required]	string regex: ^.+	Enter changes at the DNA level, as written in the test report. In principle, the cdsChange should conform to HGVS. When an RNA is not transcribed as in an intergenic region, you can input null (in addition to the notation of non-coding regions such as n.*).

aminoAcidsChange	[required]	string regex: ^.+	<p>Enter changes at the protein level, as written in the test report.</p> <p>Amino acids should be written with a single letter. In principle, other amino acids should confirm to HGVS.</p> <p>If no change is observed in amino acids as in an untranslated region, you can input null.</p>
shortVariantName	optional	string regex: ^.+	<p>Names specially assigned given by testing company to single-nucleotide variants, insertions or deletions, insertion-deletions.</p> <p>e.g. EGFR Exon 20 Insertions</p>
calculatedEffects	optional	array (length: 1-N, string regex: ^.+)	<p>Note the effects of alterations on transcripts, such as "splicing_variant", using Sequence Ontology terms.</p> <p>This field corresponds to "Effect (Sequence Ontology)" of the snpEff tool and VCF "Func.refGene" of the annovar tool. For annovar, this is explained at Output file 1 (refSeq gene annotation) on Gene-based Annotation in User Guide, whereby terms provided by annovar can be converted to Sequence Ontology terms.</p>

			One term should be assigned to each element of an array. Each string in the array must be unique.
skippedExonRanges	optional (recommend)	array (length: 1-N, object)	Exon with skipping potential.
transcriptId	[required]	string regex: ^[^¥s]+\$	Refer to the description in the transcripts tag.
transcriptDatabaseName	[required]	string [choice]	Refer to the description in the transcripts tag.
transcriptDatabaseVersion	optional	string regex: ^\.+\$	Refer to the description in the transcripts tag.
geneSymbol	[required]	string regex: ^[^¥s]+\$	Refer to the description in the transcripts tag.
strand	optional	string [choice]	Refer to the description in the transcripts tag.
exonRange	[required]	array (length: 2, integer)	Range of skipped exon numbers. Start and end exon numbers are included in the range.
totalExonNumber	optional	integer	Total exon number.
sampleItemId	required	string regex: ^\.+\$	Described the itemId of sequencing sample information (sequencing Samples). Represents sample information in which the mutation was detected.
variantOrigin	optional (recommend)	string [choice]	Somatic or germline origin. * C-CAT uses a different cancer knowledge base, according to whether alterations are somatic or germline. If no input is

			<p>provided, the knowledge base for somatic alterations is used.</p> <ul style="list-style-type: none"> • "somatic": derived from somatic cells • "germline": derived from germline cells • "likely somatic": typically in a tumor-only test, the alteration is likely to be somatic, and the knowledge base for somatic alterations is used. • "likely germline": typically in a tumor-only test, the alteration is likely to be germline, and the knowledge base for germline alterations is used.
tumorNormalPairingStatus	optional	string [choice]	<p>Combination of samples used for detection.</p> <ul style="list-style-type: none"> • "tumor-only" • "tumor and matched-normal" • "normal-only"

6-1-2. Example of shortVariants tag

```
(Example1. for SNV)
{
  "itemId": "variant-1",
  Note: The itemId is a character string at the discretion of the testing company for the detected genetic alteration.
  "chromosome": "9",
  "position": 135781005,
  "cytoband": "q34.13",
  "referenceAllele": "C",
  "alternateAllele": "G",
```


Note: Describe "position" and "referenceAllele", "alternateAllele" according to the rules of VCF v4.5.

```
"alternateAlleleFrequency": 0.54,
"alternateAlleleReadDepth": 108,
"totalReadDepth": 200,
"variantType": "SNV",
"transcripts": [
  {
    "transcriptId": "NM_000368.4",
    "transcriptDatabaseName": "RefSeq",
    "transcriptDatabaseVersion": "Release 99",
    "regionStructure": [
      {
        "type": "exon",
        "number": 15,
        "totalNumber": 23
      }
    ],
    "geneSymbol": "TSC1",
    "cdsChange": "c.1960C>G",
    "aminoAcidsChange": "p.Q654E",
    "calculatedEffects": [
      "missense_variant"
    ]
  }
],
"sampleItemId": "sequence-1-tumor-dna",
"variantOrigin": "somatic",
"tumorNormalPairingStatus": "tumor and matched-normal"
}
```

(Example2. for insertion)

```
{
  "itemId": "variant-2",
  "chromosome": "7",
  "position": 55249008,
  "cytoband": "p11.2",
  "referenceAllele": "T",
  "alternateAllele": "TGGACAACCC",
  "alternateAlleleFrequency": 0.4953,
  "alternateAlleleReadDepth": 368,
  "totalReadDepth": 743,
  "variantType": "insertion",
  "transcripts": [
    {
      "transcriptId": "NM_005228.3",
      "transcriptDatabaseName": "RefSeq",
      "transcriptDatabaseVersion": "Release 99",
      "regionStructure": [
        {
```

```

    "type": "exon",
    "number": 20,
    "totalNumber": 28
  }
],
"geneSymbol": "EGFR",
"cdsChange": "c.2307_2315dupGGACAACCC",
"aminoAcidsChange": "p.P772_H773insDNP",
"shortVariantName": "EGFR Exon 20 Insertions"
}
],
"sampleItemId": "sequence-1-tumor-dna",
"variantOrigin": "somatic",
"tumorNormalPairingStatus": "tumor and matched-normal"
}

```

(Example3. for deletion)

```

{
  "itemId": "variant-3",
  "chromosome": "1",
  "position": 27097751,
  "cytoband": "p.36.11",
  "referenceAllele": "TC",
  "alternateAllele": "T",
  "alternateAlleleFrequency": 0.1203,
  "alternateAlleleReadDepth": 32,
  "totalReadDepth": 266,
  "variantType": "deletion",
  "transcripts": [
    {
      "transcriptId": "ENST00000324856.13",
      "transcriptDatabaseName": "Ensembl",
      "transcriptDatabaseVersion": "v99",
      "regionStructure": [
        {
          "type": "exon",
          "number": 12,
          "totalNumber": 20
        }
      ],
      "geneSymbol": "ARID1A",
      "cdsChange": "c.3340delC",
      "aminoAcidsChange": "p.P1115fs*46",
      "calculatedEffects": [
        "frameshift_variant"
      ]
    }
  ],
  "sampleItemId": "sequence-3-normal-dna",
  "variantOrigin": "somatic",
  "tumorNormalPairingStatus": "tumor and matched-normal"
}

```

```
}
```

(Example4. for delins)

```
{
  "itemId": "variant-4",
  "chromosome": "1",
  "position": 26696982,
  "cytoband": "p36.11",
  "referenceAllele": "GC",
  "alternateAllele": "TT",
  "alternateAlleleFrequency": 0.0993,
  "alternateAlleleReadDepth": 52,
  "totalReadDepth": 524,
  "variantType": "delins",
  "transcripts": [
    {
      "transcriptId": "NM_006015.6",
      "transcriptDatabaseName": "RefSeq",
      "transcriptDatabaseVersion": "Release 99",
      "regionStructure": [
        {
          "type": "exon",
          "number": 1,
          "totalNumber": 20
        }
      ],
      "geneSymbol": "ARID1A",
      "cdsChange": "c.579_580delinsTT",
      "aminoAcidsChange": "p.E193_P194delinsDS"
    }
  ],
  "sampleItemId": "sequence-1-tumor-dna",
  "variantOrigin": "somatic",
  "tumorNormalPairingStatus": "tumor and matched-normal"
}
```

(Example5. for "TERT promoter")

```
{
  "itemId": "variant-5",
  "chromosome": "5",
  "position": 1295228,
  "cytoband": "p15.33",
  "referenceAllele": "G",
  "alternateAllele": "A",
  "alternateAlleleFrequency": 0.163,
  "alternateAlleleReadDepth": 15,
  "totalReadDepth": 92,
  "variantType": "SNV",
  "transcripts": [
    {
```

```

"transcriptId": "NM_198253.3",
"transcriptDatabaseName": "RefSeq",
"transcriptDatabaseVersion": "Release 99",
"regionStructure": [
  {
    "type": "promoter"
  }
],
"geneSymbol": "TERT",
"regionName": "TERT promoter",
"cdsChange": "c.-124C>T",
"aminoAcidsChange": null,
"shortVariantName": "TERT promoter c.-124C>T",
"calculatedEffects": [
  "TF_binding_site_variant"
]
}
],
"sampleItemId": "sequence-1-tumor-dna",
"variantOrigin": "somatic",
"tumorNormalPairingStatus": "tumor and matched-normal"
}

```

(Example6. for "Exon skipping")

```

{
  "itemId": "variant-6",
  "chromosome": "7",
  "position": 116411892,
  "cytoband": "p11.2",
  "referenceAllele": "TCTGTTTAAAGATCTGGGCAGTGAATTAGTTCG",
  "alternateAllele": "T",
  "alternateAlleleFrequency": 0.023,
  "alternateAlleleReadDepth": 23,
  "totalReadDepth": 1000,
  "variantType": "SNV",
  "transcripts": [
    {
      "transcriptId": "NM_000245.2",
      "transcriptDatabaseName": "RefSeq",
      "transcriptDatabaseVersion": "Release 99",
      "geneSymbol": "MET",
      "cdsChange": "c.2888-10_2909delCTGTTTAAAGATCTGGGCAGTGAATTAGTTCG",
      "aminoAcidsChange": null,
      "shortVariantName": "MET Exon 14 skipping"
    }
  ],
  "skippedExonRanges": [
    {
      "transcriptId": "NM_000245.2",
      "transcriptDatabaseName": "RefSeq",
      "transcriptDatabaseVersion": "Release 99",

```

```
    "geneSymbol": "MET",  
    "exonRange": [14, 14],  
    "totalExonNumber": 21  
  }  
],  
  "sampleItemId": "sequence-1-tumor-dna",  
  "variantOrigin": "somatic",  
  "tumorNormalPairingStatus": "tumor and matched-normal"  
}
```

6-2. copyNumberAlterations tag

This tag is used to provide information on copy number alterations (CNAs).

Key	Condition	Data type	Description
copyNumberAlterations	optional	array (length: 1-N, object)	Aggregation tag for information on copy number alterations (CNAs). Each object in the array must be unique.

6-2-1. Tags within copyNumberAlterations tag

Key	Condition	Data type	Description
itemId	required	string regex: ^.+	An ID assigned to a genetic alteration. It must be unique within this case report.
chromosome	optional (recommend)	string regex: ^[a-zA-Z0-9_]+	Chromosome number.
startPosition	optional (recommend)	integer	Physical starting position in a chromosome. Please use the 1-based coordinate system.
startCytoband	optional	string regex: ^.+	Cytoband at the starting position.
endPosition	optional (recommend)	integer	Physical ending position in a chromosome. Please use the 1-based coordinate system.
endCytoband	optional	string regex: ^.+	Cytoband at the ending position.
copyNumberMetrics	optional (recommend)	array (length: 1-N, object)	Measurements and units of the copy number alteration. Array of objects composed of two keys, value and unit. If there are two or more values in different units, register them

			as an array with the length of 2 or more. Each object in the array must be unique.
value	[required]	number	Aberrated copy number measurement.
unit	[required]	string [choice]	Unit for measured value. Select one from: <ul style="list-style-type: none"> • "copy number": Copy number • "fold-change": Ratio of (standardized) reading depth of tumor samples to normal samples • "log2 fold-change": log2 transformation of "fold-change" • "fraction-of-gene": fraction of a CNA region to the gene region of interest * Please consult C-CAT if you want to use other units.
copyNumberAlterationType	required	string [choice]	CNA type written in the test report or equivalent report. Select one from: <ul style="list-style-type: none"> • "amplification" • "gain" • "duplication" • "loss" • "deletion" • "homozygous deletion" • "neutral" * Please consult C-CAT if you want to use other types.
transcripts	required	array (length: 1-N, object)	Refer to the description in the shortVariants tag.
transcriptId	optional	string regex: $^{\wedge}[\p{s}]^{+}$	Refer to the description in the shortVariants tag.
transcriptDatabaseName	optional	string [choice]	Refer to the description in the shortVariants tag. If the transcriptId is entered, this key is recommended to input as well.

transcriptDatabaseVersion	optional	string regex: ^.+	Refer to the description in the shortVariants tag.
regionStructure	optional	array (length: 1-N, object)	Refer to the description in the shortVariants tag.
type	[required]	string [choice]	Refer to the description in the shortVariants tag.
number	optional	integer	Refer to the description in the shortVariants tag.
totalNumber	optional	integer	Refer to the description in the shortVariants tag.
geneSymbol	[required]	string regex: ^[^s]+	Refer to the description in the shortVariants tag.
geneSymbolSource	optional (recommend)	string regex: ^.+	Refer to the description in the shortVariants tag.
geneSymbolId	optional (recommend)	string regex: ^.+	Refer to the description in the shortVariants tag.
regionName	optional	string regex: ^.+	Refer to the description in the shortVariants tag.
strand	optional	string [choice]	Refer to the description in the shortVariants tag.
cdsChange	optional	string regex: ^.+	Refer to the description in the shortVariants tag.
aminoAcidsChange	optional	string regex: ^.+	Refer to the description in the shortVariants tag.
copyNumberAlterationName	optional	string regex: ^.+	Names for specially assigned copy number alterations given by the testing company.

calculatedEffects	optional	array (length: 0-N, string regex: ^.+)\$	Refer to the description in the shortVariants tag.
skippedExonRanges	optional (recommend)	array (length: 1-N, object)	Refer to the description in the shortVariants tag.
transcriptId	[required]	string regex: ^[^¥s]+\$	Refer to the description in the shortVariants tag.
transcriptDatabaseName	[required]	string [choice]	Refer to the description in the shortVariants tag.
transcriptDatabaseVersion	optional	string regex: ^.+)\$	Refer to the description in the shortVariants tag.
geneSymbol	[required]	string regex: ^[^¥s]+\$	Refer to the description in the shortVariants tag.
strand	optional	string [choice]	Refer to the description in the shortVariants tag.
exonRange	[required]	array (length: 2, integer)	Refer to the description in the shortVariants tag.
totalExonNumber	optional	integer	Refer to the description in the shortVariants tag.
sampleItemId	required	string regex: ^.+)\$	Refer to the description in the shortVariants tag.
variantOrigin	optional (recommend)	string [choice]	Refer to the description in the shortVariants tag.
tumorNormalPairingStatus	optional	string [choice]	Refer to the description in the shortVariants tag.

6-2-2. Example of copyNumberAlterations tag

(Example)

```
{
  "itemId": "variant-9",
  "chromosome": "1",
  "startPosition": 8921059,
  "startCytoband": "p36.23",
  "endPosition": 8939151,
```

```
"endCytoband": "p36.23",
"copyNumberMetrics": [
  {
    "value": 0.2309,
    "unit": "fold-change"
  },
  {
    "value": -2.1147,
    "unit": "log2 fold-change"
  }
],
"copyNumberAlterationType": "loss",
"transcripts": [
  {
    "transcriptDatabaseName": "RefSeq",
    "transcriptDatabaseVersion": "Release 99",
    "geneSymbol": "ENO1"
  }
],
"sampleItemId": "sequence-1-tumor-dna",
"variantOrigin": "somatic",
"tumorNormalPairingStatus": "tumor and matched-normal"
}
```

6-3. structuralVariations tag

This tag is used to provide information on structuralVariations such as fusions, duplications, large deletions, and inversions.

Key	Condition	Data type	Description
structuralVariations	optional	array (length: 1-N, object)	Aggregation tag for information on structuralVariations, such as fusions, duplications, large deletions, and inversions. Each object in the array must be unique.

6-3-1. Tags within structuralVariations tag

Key	Condition	Data type	Description
itemId	required	string regex: ^.+	An ID assigned to a genetic alteration. It must be unique within this case report.
breakends	required	array (length: 2, object)	Two breakends of the structuralVariation. Each object in the array must be unique.
chromosome	required	string regex: ^[a-zA-Z0-9_#-]+\$	Chromosome number.
startPosition	required	integer	Physical starting position in a chromosome. Please use the 1-based coordinate system.
startCytoband	optional	string regex: ^.+	Cytoband at the starting position.
endPosition	required	integer	Physical ending position in a chromosome. Please use the 1-based coordinate system.

endCytoband	optional	string regex: ^.+	Cytoband at the ending position.
referenceAllele	optional (recommend)	string regex: ^[ACGTN]+	Reference base. Please describe according to VCF v4.5. See "XV-2. matePieceLocation" below for a detailed explanation.
alternateAllele	optional (recommend)	string regex: ^[ACGTN][0-9]+	Variant base. Please describe according to VCF v4.5. See "XV-2. matePieceLocation" below for a detailed explanation.
matePieceLocation	optional (recommend)	string [choice]	If the sequence of interest is bound to another sequence on the upstream (or downstream) of this sequence of interest along the reference genome sequence, it is input as "upstream" (or "downstream"). It is strongly recommended to input this key for accurately identifying the genomic changes. See "XV-2. matePieceLocation" below for a detailed explanation.
supportingReadCount	optional	integer	The number of reads that support the breakends, or the numerator of alternateAlleleFrequency.
totalReadCount	optional	integer	The total number of reads that support and do not support breakends, or the denominator of alternateAlleleFrequency.
alternateAlleleFrequency	optional (recommend)	number	Variant allele frequency of a breakend (ranging from 0 to 1). Please input a variant allele frequency calculated for each breakend.
transcripts	required	array (length:	Refer to the description in the shortVariants tag.

		1-N, object)	
transcriptId	optional	string regex: ^[^¥s]+\$	Refer to the description in the shortVariants tag.
transcriptDatabaseName	optional	string [choice]	Refer to the description in the shortVariants tag. If the transcriptId is entered, this key is recommended to input as well.
transcriptDatabaseVersion	optional	string regex: ^.+ \$	Refer to the description in the shortVariants tag.
readingFrameStatus	optional	string regex: ^.+ \$	Frame prediction for transcripts (e.g., inFrame, outOfFrame).
regionStructure	optional (recommend)	array (length: 1-N, object)	Refer to the description in the shortVariants tag.
type	[required]	string [choice]	Refer to the description in the shortVariants tag.
number	optional (recommend)	integer	Refer to the description in the shortVariants tag.
totalNumber	optional	integer	Refer to the description in the shortVariants tag.
geneSymbol	[required]	string regex: ^[^¥s]+\$	Refer to the description in the shortVariants tag.
geneSymbolSource	optional (recommend)	string regex: ^.+ \$	Refer to the description in the shortVariants tag.
geneSymbolId	optional (recommend)	string regex: ^.+ \$	Refer to the description in the shortVariants tag.

regionName	optional (recommend)	string regex: ^.+ \$	Refer to the description in the shortVariants tag.
strand	optional	string [choice]	Refer to the description in the shortVariants tag.
cdsChange	optional	string regex: ^.+ \$	Refer to the description in the shortVariants tag.
aminoAcidsChange	optional	string regex: ^.+ \$	Refer to the description in the shortVariants tag.
calculatedEffects	optional	array (length: 1-N, string regex: ^.+ \$)	Refer to the description in the shortVariants tag.
orderedGenePairs	optional (recommend)	array (length: 1-N, array)	<p>An array of an ordered pair(s) of gene names that traverses a breakpoint of a structuralVariation, where the pair(s) is written in the transcriptional order.</p> <p>This array has, as a child element(s), an array(s) in length 2 that stores a pair of geneSymbols of transcripts. The geneSymbols in each pair should be aligned in the transcriptionally forward direction and should be unique. For example, for geneSymbols A and B, if A is in the upstream of transcription and B is the downstream, it is represented as</p> <p>"orderedGenePairs": [["A", "B"]].</p> <p>* This directional information is taken into account in the cancer knowledge base search. For example, in the case of</p>

			<p>[["A", "B"]], ["A", "B"] alone is searched. In the case of [["A", "B"], ["B", "A"]], both ["A", "B"] and ["B", "A"] are respectively searched. You can control the search in more detail, such as by [["A", "B"], ["B", "A"], ["C", "B"]].</p> <p><u>* Please specify such an appropriate input in this tag if you want to control how gene pairs are searched.</u></p> <p>* If this tag is omitted, the pair of geneSymbols under the breakends tag is treated as a pair without information on the transcriptional direction, and both direction of the pair will be respectively searched in the knowledge base.</p>
(Child element of orderedGenePairs)	required	array (length: 2, string regex: ^.+)\$	An array of strings of length 2 with the geneSymbol described in transcripts aligned in the forward direction of transcription. e.g.) ["EML4", "ALK"]
affectedGenes	optional (recommend)	array (length: 1-N, string regex: ^.+)\$	Genes affected by gene structuralVariation. If multiple genes are deleted due to long deletion, this key is recommended to input.
skippedExonRanges	optional (recommend)	array (length: 1-N, object)	Exon information skipped by structuralVariation. If "exon skipping" is selected for structuralVariationType, this key is recommended to input as well.
transcriptId	[required]	string regex: ^[^\s]+\$	Refer to the description in the shortVariants tag.
transcriptDatabaseName	[required]	string [choice]	Refer to the description in the shortVariants tag.

			If the transcriptId is entered, this key is recommended to input as well.
transcriptDatabaseVersion	optional	string regex: ^.+	Refer to the description in the shortVariants tag.
geneSymbol	[required]	string regex: ^[^\$]+	Refer to the description in the shortVariants tag.
strand	optional	string [choice]	Refer to the description in the shortVariants tag.
exonRange	[required]	array (length: 2, integer)	Range of skipped exon numbers. Start and end exon numbers are included in the range.
totalExonNumber	optional	integer	Refer to the description in the shortVariants tag.
insertedSequence	optional	string regex: ^[ACGTN]+\$	Sequence inserted between the two breakends of the genome sequence. If an inserted sequence does not exist, input null.
supportingReadCount	optional	integer	The number of reads that support structuralVariation, or the numerator of alternateAlleleFrequency.
totalReadCount	optional	integer	The total number of reads that support and do not support structuralVariation, or the denominator of alternateAlleleFrequency.
alternateAlleleFrequency	optional (recommend)	number	Variant allele frequency of a structuralVariation (ranging from 0 to 1). Please input a value calculated for a structuralVariation.
expressionLevelMetrics	optional	array (length: 1-N, object)	Information on expression levels in RNA-seq. Each object in the array must be unique.
value	[required]	number	Value of an expression level
unit	[required]	string	Unit of an expression level.

		[choice]	<p>Select one from the following options:</p> <ul style="list-style-type: none"> • "TPM" • "FPKM" • "FPM" • "RPKM" • "RPM" <p>* Please consult C-CAT if you want to use other units.</p>
type	optional	string [choice]	<p>Type of summary statistics. Select one from the following options:</p> <ul style="list-style-type: none"> • "mean" • "median" • "standard deviation" <p>(Default: null)</p> <p>If the expression level represents a measured value (not a summary statistic), you can input null.</p> <p>* Please consult C-CAT if you want to use other choices.</p>
sampleSize	optional	integer	<p>Sample size when calculating summary statistics.</p> <p>* If a non-null value is inputted in type, it is recommended to input this item as well.</p> <p>(Default: null)</p>
isControl	optional	boolean	<p>Whether the expression level was taken from a control (normal) sample (true) or from a tumor sample (false).</p> <p>(Default: false)</p>
structuralVariationNames	optional (recommend)	array (length: 1-N, string regex: ^.+)\$	<p>Names for specially assigned structuralVariations name given by the testing company. e.g. IGH (IGHJ4) CRLF2 upstream translocation</p> <p>Each string in the array must be unique.</p>
structuralVariationType	required	string [choice]	<p>StructuralVariation type written in the test report. Select one from:</p>

			<ul style="list-style-type: none"> • "gene fusion" • "gene fusion and frameshift variant" • "bidirectional gene fusion" • "duplication" • "tandem duplication" • "deletion" • "inversion" • "truncation" • "splice variant" • "exon skipping" • "translocation" • "other" <p>* Please consult C-CAT if you want to use other types.</p>
sampleItemId	required	string regex: ^.+	Refer to the description in the shortVariants tag.
variantOrigin	optional (recommend)	string [choice]	Refer to the description in the shortVariants tag.
tumorNormalPairingStatus	optional	string [choice]	Refer to the description in the shortVariants tag.

6-3-2. Example of structuralVariations tag

(Example1. In cases where orderedGenePairs is present)

```
{
  "itemId": "variant-13",
  "breakends": [
    {
      "chromosome": "2",
      "startPosition": 42510050,
      "startCytoband": "p21",
      "endPosition": 42510050,
      "endCytoband": "p21",
      "referenceAllele": "A",
      "alternateAllele": "A[1:29445240[",
      "matePieceLocation": "downstream",
      "transcripts": [
        {
          "transcriptId": "NM_001145076.3",
          "transcriptDatabaseName": "RefSeq",
          "transcriptDatabaseVersion": "Release 99",
          "regionStructure": [
            {
```

```

        "type": "exon",
        "number": 7,
        "totalNumber": 22
      }
    ],
    "geneSymbol": "EML4"
  }
],
{
  "chromosome": "2",
  "startPosition": 29445240,
  "startCytoband": "p23.2",
  "endPosition": 29445240,
  "endCytoband": "p23.2",
  "referenceAllele": "A",
  "alternateAllele": "]2:42510050]A",
  "matePieceLocation": "upstream",
  "transcripts": [
    {
      "transcriptId": "NM_001353765.2",
      "transcriptDatabaseName": "RefSeq",
      "transcriptDatabaseVersion": "Release 99",
      "regionStructure": [
        {
          "type": "exon",
          "number": 3,
          "totalNumber": 10
        }
      ],
      "geneSymbol": "ALK"
    }
  ]
},
"orderedGenePairs": [
  ["EML4", "ALK"]
],
  "supportingReadCount": 30,
  "totalReadCount": 430,
  "alternateAlleleFrequency": 0.07,
  "structuralVariationType": "gene fusion",
  "sampleItemId": "sequence-1-tumor-dna",
  "variantOrigin": "somatic",
  "tumorNormalPairingStatus": "tumor and matched-normal"
}

```

(Example2. When describing the totalReadCount and alternateAlleleFrequency for each
breakend)

```

{

```

```

"itemId": "variant-14",
"breakends": [
{
  "chromosome": "2",
  "startPosition": 42510050,
  "startCytoband": "p21",
  "endPosition": 42510050,
  "endCytoband": "p21",
  "matePieceLocation": "downstream",
  "totalReadCount": 330,
  "alternateAlleleFrequency": 0.06,
  "transcripts": [
    {
      "transcriptId": "NM_001145076.3",
      "transcriptDatabaseName": "RefSeq",
      "transcriptDatabaseVersion": "Release 99",
      "regionStructure": [
        {
          "type": "exon",
          "number": 7,
          "totalNumber": 22
        }
      ],
      "geneSymbol": "EML4"
    }
  ],
},
{
  "chromosome": "2",
  "startPosition": 29445240,
  "startCytoband": "p23.2",
  "endPosition": 29445240,
  "endCytoband": "p23.2",
  "matePieceLocation": "upstream",
  "totalReadCount": 570,
  "alternateAlleleFrequency": 0.07,
  "transcripts": [
    {
      "transcriptId": "NM_001353765.2",
      "transcriptDatabaseName": "RefSeq",
      "transcriptDatabaseVersion": "Release 99",
      "regionStructure": [
        {
          "type": "exon",
          "number": 3,
          "totalNumber": 10
        }
      ],
      "geneSymbol": "ALK"
    }
  ]
}
]

```

```

    }
  ],
  "orderedGenePairs": [
    ["EML4", "ALK"]
  ],
  "supportingReadCount": 30,
  "structuralVariationType": "gene fusion",
  "sampleItemId": "sequence-1-tumor-dna",
  "variantOrigin": "somatic",
  "tumorNormalPairingStatus": "tumor and matched-normal"
}

```

(Example3. In cases where insertedSequence is present)

```

{
  "itemId": "variant-15",
  "breakends": [
    {
      "chromosome": "14",
      "startPosition": 234567,
      "startCytoband": "p13",
      "endPosition": 234567,
      "endCytoband": "p13",
      "matePieceLocation": "downstream",
      "transcripts": [
        {
          "transcriptId": null,
          "transcriptDatabaseName": "RefSeq",
          "transcriptDatabaseVersion": "Release 99",
          "geneSymbol": null
        }
      ]
    },
    {
      "chromosome": "2",
      "startPosition": 321672,
      "startCytoband": "p25.3",
      "endPosition": 321672,
      "endCytoband": "p25.3",
      "matePieceLocation": "upstream",
      "transcripts": [
        {
          "transcriptId": null,
          "transcriptDatabaseName": "RefSeq",
          "transcriptDatabaseVersion": "Release 99",
          "geneSymbol": "LINC01865"
        }
      ]
    }
  ],
  "insertedSequence": "GTNNNNNCAT",
  "supportingReadCount": 30,

```

```

"alternateAlleleFrequency": 0.07,
"structuralVariationType": "other",
"sampleItemId": "sequence-1-tumor-dna",
"variantOrigin": "somatic",
"tumorNormalPairingStatus": "tumor and matched-normal"
}

```

(Example4. In cases of exon skipping)

```

{
  "itemId": "variant-16",
  "breakends": [
    {
      "chromosome": "7",
      "startPosition": 116771654,
      "startCytoband": "p22.1",
      "endPosition": 116771654,
      "endCytoband": "p22.1",
      "transcripts": [
        {
          "transcriptId": "NM_000245",
          "transcriptDatabaseName": "RefSeq",
          "transcriptDatabaseVersion": "Release 99",
          "regionStructure": [
            {
              "type": "exon",
              "number": 13,
              "totalNumber": 21
            }
          ],
          "geneSymbol": "MET"
        }
      ]
    },
    {
      "chromosome": "7",
      "startPosition": 116774881,
      "startCytoband": "p22.1",
      "endPosition": 116774881,
      "endCytoband": "p22.1",
      "transcripts": [
        {
          "transcriptId": "NM_000245",
          "transcriptDatabaseName": "RefSeq",
          "transcriptDatabaseVersion": "Release 99",
          "regionStructure": [
            {
              "type": "exon",
              "number": 15,
              "totalNumber": 21
            }
          ]
        }
      ]
    }
  ]
}

```

```

    "geneSymbol": "MET"
  }
]
},
"skippedExonRanges": [
{
  "transcriptId": "NM_000245",
  "transcriptDatabaseName": "RefSeq",
  "transcriptDatabaseVersion": "Release 99",
  "geneSymbol": "MET",
  "exonRange": [14, 14],
  "totalExonNumber": 21
}
],
"supportingReadCount": 30,
"alternateAlleleFrequency": 0.07,
"structuralVariationType": "exon skipping",
"sampleItemId": "sequence-2-tumor-rna",
"variantOrigin": "somatic",
"tumorNormalPairingStatus": "tumor-only"
}

```

(Example5. In cases of translocation)

```

{
  "itemId": "variant-17",
  "breakends": [
    {
      "chromosome": "14",
      "startPosition": 105857793,
      "startCytoband": "p11.2",
      "endPosition": 105857793,
      "endCytoband": "p11.2",
      "matePieceLocation": "downstream",
      "transcripts": [
        {
          "transcriptId": null,
          "transcriptDatabaseName": null,
          "geneSymbol": "IGH",
          "regionName": "IGH"
        }
      ]
    },
    {
      "chromosome": "8",
      "startPosition": 127735729,
      "startCytoband": "p11.23",
      "endPosition": 127735729,
      "endCytoband": "p11.23",
      "matePieceLocation": "upstream",
      "transcripts": [

```

```
{
  "transcriptId": null,
  "transcriptDatabaseName": null,
  "regionStructure": [
    {
      "type": "upstream"
    }
  ],
  "geneSymbol": "MYC"
}
],
"supportingReadCount": 120,
"totalReadCount": 435,
"alternateAlleleFrequency": 0.31,
"structuralVariationType": "translocation",
"sampleItemId": "sequence-1-tumor-dna",
"variantOrigin": "somatic",
"tumorNormalPairingStatus": "tumor-only"
}
```


7. otherBiomarkers tag

This tag is used to provide information on biomarkers other than genetic alterations defined in the variants tag. Currently, Micro-Satellite Instability (MSI), Tumor Mutation Burden (TMB), HRD and ctDNA Tumor Fraction can be noted.

Key	Condition	Data type	Description
otherBiomarkers	optional	array (length: 1-N, object)	Aggregation tag for information on biomarkers. Each object in the array must be unique.

7-1. Tags within otherBiomarkers tag

Key	Condition	Data type	Description
itemId	required	string regex: ^.+\$\$	An ID assigned to a biomarker. It must be unique within this case report.
biomarkerType	required	string [choice]	Type of biomarkers. Select one from: <ul style="list-style-type: none"> • "TMB": Tumor Mutation Burden • "MSI": Micro-Satellite Instability • "HRD": Homologous Recombination Deficiency • "ctDNA Tumor Fraction": ctDNA Tumor Fraction * Inform C-CAT beforehand if you want other biomarkers.
biomarkerMetrics	optional (recommend)	array (length: 1-N, object)	Inspection value information. Each object in the array must be unique. * When creating C-CAT Findings, the first object in the array is used first.
value	[required]	number	Measured value. e.g.) 5.15
unit	[required]	string regex: ^.+\$\$	Unit for the measured value. Units may vary depending on test types. e.g.) % If the value is unitless, you can input null.
type	[required]	string regex: ^.+\$\$	Type of inspection value. e.g.) Mutations per megabase MSIsensor score percentage of MSI sites

state	optional (recommend)	string [choice]	<p>Select the biomarker status from the following options.</p> <ul style="list-style-type: none"> • "high" • "low" • "intermediate" • "stable" • "not high" • "positive" • "negative" • "cannot be determined" <p>* Inform C-CAT beforehand if you want other options.</p>
descriptions	optional	array (length: 1-N, string regex: ^.+)\$	<p>Descriptions such as how to obtain the testing value and the meaning of the value. Refer to the contents tag within the comments tag for usable languages and new lines.</p>
biomarkerOrigin	optional (recommend)	string [choice]	Refer to the description in the variantOrigin tag in shortVariants tag.
sampleItemId	required	string regex: ^.+\$\$	Refer to the description in the shortVariants tag.
tumorNormalPairingStatus	optional	string [choice]	Refer to the description in the shortVariants tag.

7-2. Example of otherBiomarkers tag

```
(Example)
"otherBiomarkers": [
{
  "itemId": "biomarker-1",
  "biomarkerType": "MSI",
  "biomarkerMetrics": [
    {
      "value": 5.15,
      "unit": "%",
      "type": "percentage of MSI sites"
    },
    {
      "value": 2,
      "unit": null,
      "type": "MSI sensor score"
    }
  ]
}
```

```
],
```

Note: If one inspection item has values in multiple units, use array notation in the "biomarkerMetrics" tag.

```
  "state": "stable",
  "descriptions": [
    "MSI sensor score 10 points or more was MSI-H, 3 points or more and less than 10
    points was indeterminate (MSI-I), and less than 3 points was microsatellite stable
    (MSS).",
    "https://www.gi-cancer.net/gi/ronbun/archives/201901-01.html"
  ],
  "sampleItemId": "sequence-1-tumor-dna",
  "tumorNormalPairingStatus": "tumor and matched-normal" },
{
  "itemId": "biomarker-2",
  "biomarkerType": "TMB",
  "biomarkerMetrics": [
    {
      "value": 34.5680122,
      "unit": "Muts/Mb",
      "type": "Mutations per megabase"
    }
  ],
  "state": "high",
  "sampleItemId": "sequence-1-tumor-dna",
  "reported": true
},
{
  "itemId": "biomarker-3",
  "biomarkerType": "HRD",
  "biomarkerMetrics": [
    {
      "value": 0.94,
      "unit": null,
      "type": "HRD"
    }
  ],
  "state": "positive",
  "sampleItemId": "sequence-1-tumor-dna"
}
]
```

8. expressions tag

This tag is used to provide the information on gene expression.

Key	Condition	Data type	Description
expressions	optional	array (length: 1-N, object)	Aggregation tag for information on gene expressions. Each object in the array must be unique.

8-1. Tags within expressions tag

Key	Condition	Data type	Description
itemId	required	string regex: ^.+ \$	An ID assigned to an expression. It must be unique within this case report.
readCount	optional (recommend)	integer	Number of reads.
status	optional	string [choice]	Select the expression level status from the following options. <ul style="list-style-type: none"> • "up-regulated" • "down-regulated" • "not differentially expressed" * Please consult C-CAT if you want to use other types.
transcripts	required	array (length: 1-N, object)	Refer to the description in the shortVariants tag.
transcriptId	optional	string regex: ^[^¥s]+ \$	Refer to the description in the shortVariants tag.
transcriptDatabaseName	optional	string [choice]	Refer to the description in the shortVariants tag. If the transcriptId is entered, this key is recommended to input as well.
transcriptDatabaseVersion	optional	string regex: ^.+ \$	Refer to the description in the shortVariants tag.
regionStructure	optional	array (length: 1-N, object)	Refer to the description in the shortVariants tag.

type	[required]	string [choice]	Refer to the description in the shortVariants tag.
number	optional	integer	Refer to the description in the shortVariants tag.
totalNumber	optional	integer	Refer to the description in the shortVariants tag.
geneSymbol	[required]	string regex: ^[^¥s]+\$	Refer to the description in the shortVariants tag.
geneSymbolSource	optional (recommend)	string regex: ^.+ \$	Refer to the description in the shortVariants tag.
geneSymbolId	optional (recommend)	string regex: ^.+ \$	Refer to the description in the shortVariants tag.
regionName	optional	string regex: ^.+ \$	Refer to the description in the shortVariants tag.
strand	optional	string [choice]	Refer to the description in the shortVariants tag.
cdsChange	optional	string regex: ^.+ \$	Refer to the description in the shortVariants tag.
aminoAcidsChange	optional	string regex: ^.+ \$	Refer to the description in the shortVariants tag.
calculatedEffects	optional	array (length: 1-N, string regex: ^.+ \$)	Refer to the description in the shortVariants tag.
expressionLevelMetrics	required	array (length: 1-N, object)	Refer to the description in the structuralVariations tag.
value	[required]	number	Refer to the description in the structuralVariations tag.

unit	[required]	string [choice]	Refer to the description in the structuralVariations tag.
type	optional (recommend)	string [choice]	Refer to the description in the structuralVariations tag.
sampleSize	optional	integer	Refer to the description in the structuralVariations tag.
isControl	optional	boolean	Refer to the description in the structuralVariations tag.
sampleItemId	required	string regex: ^.+\$\$	Refer to the description in the shortVariants tag.

8-2. Example of expressions tag

(Example)

```
"expressions": [
{
  "itemId": "expression-1",
  "readCount": 9928,
  "transcripts": [
    {
      "transcriptDatabaseName": "RefSeq",
      "transcriptDatabaseVersion": "Release 99",
      "geneSymbol": "MET"
    }
  ],
  "expressionLevelMetrics": [
    {
      "value": 28.24,
      "unit": "TPM",
      "isControl": false
    },
    {
      "value": 142.7,
      "unit": "TPM",
      "type": "mean",
      "sampleSize": 20,
      "isControl": true
    },
    {
      "value": 134.2,
      "unit": "TPM",
      "type": "standard deviation",
      "sampleSize": 20,
      "isControl": true
    }
  ]
}
```

```
    ],  
    "sampleItemId": "sequence-4-normal-rna"  
  }  
]
```

9. alterationNotes tag

This tag is used to provide supplementary information about the molecular alterations, druggability and pathogenicity as documented in the test report or similar documents.

Key	Condition	Data type	Description
alterationNotes	required	array(length: 0~N, object)	Aggregation tag for information on alteration notes. Each object in the array must be unique. * This tag should be described for all molecular alterations, druggability, and pathogenicity.

9-1. Tag within alterationNotes tag

Key	Condition	Data type	Description
itemId	required	string regex: ^.+\$	The itemId to indicate a molecular alterations, druggability, and pathogenicity.
reported	required	ブール型	Whether or not the alteration is reported in test report or similar documents. * In case of true, the variant is outputted to C-CAT finding document.
reportSections	optional	array(length: 1~N, string regex: ^.+\$)	The name of the section or page in the test report or similar documents where the alteration is described.
approved	optional	Boolean	Whether the variant is approved (true) or not (false) in the test report or equivalent report. (Default: true)
grade	optional	string [choice]	The variant grade treated in the test report or the other related report. Select one from: <ul style="list-style-type: none"> • "clear": unambiguous detection • "equivocal": equivocal detection, such as low confidence • "suppl": supplementary information related to variants regardless of the accuracy of detection. (Default: "clear")

			* In the case of clear or equivocal, the variant is annotated with cancer knowledge base. Please consult C-CAT beforehand if you use other types.
--	--	--	---------------------------------------------------------------------------------------------------------------------------------------------------

9-2. Example of alterationNotes tag

```
(Example)
"alterationNotes": [
  {
    "itemId": "shortVariant-1",
    "reported": true,
    "reportSections": [
      "Somatic variant"
    ],
    "approved": true,
    "grade": "clear"
  },
  {
    "itemId": "copyNumberAlteration-1",
    "reported": true,
    "reportSections": [
      "Variant of Uncertain Significance"
    ],
    "approved": false,
    "grade": "equivocal"
  },
  {
    "itemId": "structuralVariation-1",
    "reported": true,
    "reportSections": [
      "Somatic variant"
    ],
    "approved": false,
    "grade": "suppl"
  },
  {
    "itemId": "otherBiomarker-1",
    "reported": true,
    "reportSections": [
      "Appendix"
    ],
    "approved": true,
    "grade": "clear"
  },
  {
    "itemId": "expression-1",
    "reported": true,
    "reportSections": [
      "Appendix"
    ]
  }
]
```

```

    ],
    "approved": false,
    "grade": "clear"
  },
  {
    "itemId": "armLevelChanges-1",
    "reported": true,
    "reportSections": [
      "Appendix"
    ],
    "approved": false,
    "grade": "clear"
  },
  {
    "itemId": "nonHumanContents-1",
    "reported": true,
    "reportSections": [
      "Appendix"
    ],
    "approved": false,
    "grade": "suppl"
  },
  {
    "itemId": "wildType-1",
    "reported": true,
    "reportSections": [
      "Appendix"
    ]
  },
  {
    "itemId": "druggability-1",
    "reported": true,
    "reportSections": [
      "Drug"
    ],
    "approved": false
  },
  {
    "itemId": "pathogenicity-1",
    "reported": true,
    "reportSections": [
      "Appendix"
    ],
    "approved": false
  }
]

```

10. armLevelChanges tag

This tag is used to provide the information about chromosome – chromosome arm change.

Key	Condition	Data type	Description
armLevelChanges	optional	array (length: 1-N, object)	Aggregation tag for armLevelChange information. Each object in the array must be unique.

10-1. Tags within armLevelChanges tag

Key	Condition	Data type	Description
itemId	required	string regex: ^\.+\$	An ID assigned to chromosome – chromosome arm change. It must be unique within this case report.
armLevelChangeType	required	string [choice]	Type of the change for chromosome or chromosome arm level change. Select one from: For more information, see International System for Human Cytogenomic Nomenclature (ISCN) 2020 for the details of these abbreviations. <ul style="list-style-type: none"> • "amp": amplification • "del": deletion • "t": translocation • "inv": inversion • "dup": duplication • "cx": complex reconstruction * Please consult C-CAT if you want to use other types.
cytobands	required	array (length: 1-N, object)	Cytoband information.
chromosome	[required]	string regex: ^[a-zA-Z0-9_¥-]+\$	Chromosome number.
arm	optional (recommend)	string [choice]	Short or long arm. Select from the following two options. <ul style="list-style-type: none"> • "p": short arm • "q": long arm

region	optional (recommend)	string regex: ^.+	Cytoband region
armLevelChangeNames	optional (recommend)	array (length: 1-N, string regex: ^.+)	Name of chromosome – chromosome arm change written in the test report. Each string in the array must be unique.
sampleItemId	required	string regex: ^.+	Refer to the description in the shortVariants tag.

10-2. Example of armLevelChanges tag

(Example)

```
"armLevelChanges": [
{
  "itemId": "arm-1",
  "armLevelChangeType": "amp",
  "cytobands": [
    {
      "chromosome": "7",
      "arm": "q"
    }
  ],
  "sampleItemId": "sequence-1-tumor-dna",
  "reported": true
},
{
  "itemId": "arm-2",
  "armLevelChangeType": "del",
  "cytobands": [
    {
      "chromosome": "17",
      "arm": "p"
    }
  ],
  "sampleItemId": "sequence-1-tumor-dna",
  "reported": true
},
{
  "itemId": "arm-3",
  "armLevelChangeType": "t",
  "cytobands": [
    {
      "chromosome": "X",
      "arm": "q",
```

```

    "region": "28"
  },
  {
    "chromosome": "14",
    "arm": "q",
    "region": "11.2"
  }
],
"armLevelChangeNames": [
  "t(X;14)(q28;q11.2)"
],
"sampleItemId": "sequence-1-tumor-dna",
"reported": true
},
{
  "itemId": "arm-4",
  "armLevelChangeType": "amp",
  "cytobands": [
    {
      "chromosome": "21"
    }
  ],
  "armLevelChangeNames": [
    "iAMP21"
  ],
  "sampleItemId": "sequence-1-tumor-dna",
  "reported": true
}
]

```

11. nonHumanContents tag

This tag is used to provide the information on non-human content.

Key	Condition	Data type	Description
nonHumanContents	optional	array (length: 1-N, object)	Aggregation tag for information on non-human contents. Each object in the array must be unique.

11-1. Tags within nonHumanContents tag

Key	Condition	Data type	Description
itemId	required	string regex: ^.+ \$	The ID of the item. It must be unique within this case report.
organism	required	string regex: ^.+ \$	The name of the species, as listed in the test report.
contentMetrics	optional	array (length: 1-N, object)	Refer to the description in the biomarkerMetrics tag in otherBiomarkers tag.
value	[required]	number	Refer to the description in the biomarkerMetrics tag in otherBiomarkers tag.
unit	[required]	string regex: ^.+ \$	Refer to the description in the biomarkerMetrics tag in otherBiomarkers tag.
type	[required]	string regex: ^.+ \$	Refer to the description in the biomarkerMetrics tag in otherBiomarkers tag.
descriptions	optional	array (length: 1-N, string regex: ^.+ \$)	Description on nonHumanContent. Refer to the contents tag within the comments tag for usable languages and new lines.
sampleItemId	required	string regex: ^.+ \$	Refer to the description in the shortVariants tag.

11-2. Example of nonHumanContents tag

(Example)

```
"nonHumanContents": [  
  {  
    "itemId": "nonHuman-1",  
    "organism": "HBV",  
    "contentMetrics": [  
      {  
        "value": 65,  
        "unit": "reads-per-million",  
        "type": "virus-derived read sequences"  
      }  
    ],  
    "descriptions": [  
      "Hepatitis B virus."  
    ],  
    "sampleItemId": "sequence-1-tumor-dna",  
    "reported": false  
  }  
]
```

12. wildTypes tag

This tag provides information on wild type.

Key	Condition	Data type	Description
wildTypes	optional	array (length: 1-N, object)	Aggregation tag for information on wild type.

12-1. Tags within wildTypes tag

Key	Condition	Data type	Description
itemId	[required]	string regex: ^\.+\$	An ID assigned to a wild type. It must be unique within this case report.
geneSymbol	[required]	string regex: ^[^¥\$]+	Refer to the description in the shortVariants tag.
geneSymbolSource	optional (recommend)	string regex: ^\.+\$	Refer to the description in the shortVariants tag.
geneSymbolId	optional (recommend)	string regex: ^\.+\$	Refer to the description in the shortVariants tag.
transcriptId	optional	string regex: ^[^¥\$]+	Refer to the description in the shortVariants tag.
origin	optional	string [choice]	Refer to the description in the variantOrigin tag in shortVariants tag.
wildTypeName	optional	string regex: ^\.+\$	The name of the wild type written in the test report.
descriptions	optional	array (length: 1-N, string regex: ^\.+\$)	Comment. Refer to the contents tag in the comments tag for usable languages and new lines.

12-2. Example of wildTypes tag

```
(Example)
"wildTypes": [
{
```



```
"itemId": "wildType-1",
"geneSymbol": "BRAF",
"geneSymbolSource": "HGNC",
"geneSymbolId": "1097",
"transcriptId": "ENST00000288602.6",
"origin": "somatic"
},
{
  "itemId": "wildType-2",
  "geneSymbol": "KRAS",
  "geneSymbolSource": "HGNC",
  "geneSymbolId": "6407"
},
{
  "itemId": "wildType-3",
  "geneSymbol": "NRAS",
  "geneSymbolSource": "HGNC",
  "geneSymbolId": "7989"
}
]
```

13. druggability tag

This tag provides information on druggability.

Key	Condition	Data type	Description
druggability	optional	object	Aggregation tag for information on druggability.

13-1. Tags within druggability tag

Key	Condition	Data type	Description
companionDiagnostics	optional (recommend)	array (length: 1-N, object)	Information on companion diagnostics. Each object in the array must be unique.
itemId	[required]	string regex: ^.+ \$	An ID assigned to a druggability. It must be unique within this case report.
alterationItemIds	[required]	array (length: 1-N, string regex: ^.+ \$)	Target molecular alterations(itemId). If a drug corresponds to a combination of multiple molecular alterations, multiple itemIds should be specified.
drugs	[required]	array (length: 1-N, object)	Information on drugs. ※For a single agent, specify only one array element, and for combinations, specify multiple array elements.
drugName	required	string regex: ^.+ \$	The name of the drug written in the test report.
innDrugName	required	string regex: ^.+ \$	International Nonproprietary Name.
descriptions	optional	array (length: 1-N, string regex: ^.+ \$)	Description on druggability. Refer to the contents tag within the comments tag for usable languages and new lines.
comprehensiveGenomicProfiling	optional (recommend)	array (length: 1-N, object)	Information on comprehensive genomic profiling. Each object in the array must be unique.

itemId	[required]	string regex: ^.+ \$	An ID assigned to a druggability. It must be unique within this case report.
alterationItemIds	[required]	array (length: 1- N, string regex: ^.+ \$)	Target molecular alterations(itemId). If a drug corresponds to a combination of multiple molecular alterations, multiple itemIds should be specified.
drugs	[required]	array (length: 1- N, object)	Information on drugs. ※For treatment options using a single agent, specify only one array element, and for treatment options using combinations, specify multiple array elements.
drugName	required	string regex: ^.+ \$	The name of the drug written in the test report.
innDrugName	required	string regex: ^.+ \$	International Nonproprietary Name.
knowledgeBase	optional	array (length: 1- N, string regex: ^.+ \$)	The name of the knowledge base.
evidenceLevel	optional (recommend)	string regex: ^.+ \$	Evidence level.
cancerCode	optional	string regex: ^.+ \$	Cancer code.
cancerName	optional	string regex: ^.+ \$	Cancer name.
descriptions	optional	array (length: 1- N, string regex: ^.+ \$)	Description on druggability. Refer to the contents tag within the comments tag for usable languages and new lines.

13-2. Example of druggability tag

(Example)

```

"druggability": {
  "companionDiagnostics": [
    {
      "itemId": "druggability-CDx-1",
      "alterationItemIds": [
        "variant-1"
      ],
      "drugs": [
        {
          "drugName": "PERJETA®",
          "innDrugName": "pertuzumab"
        },
        {
          "drugName": "HERCEPTIN®",
          "innDrugName": "trastuzumab"
        }
      ]
    },
    {
      "itemId": "druggability-CDx-2",
      "alterationItemIds": [
        "variant-2"
      ],
      "drugs": [
        {
          "drugName": "PHESGO®",
          "innDrugName": "pertuzumab, trastuzumab, and vorhyaluronidase alfa"
        }
      ]
    }
  ],
  "comprehensiveGenomicProfiling": [
    {
      "itemId": "druggability-CGP-1",
      "alterationItemIds": [
        "variant-2"
      ],
      "drugs": [
        {
          "drugName": "alectinib",
          "innDrugName": "alectinib",
          "knowledgeBase": [
            "OncoKB"
          ],
          "evidenceLevel": "A",
          "cancerCode": "NSCLC",
          "cancerName": "Non-Small Cell Lung Cancer"
        }
      ]
    }
  ]
}
]

```

}

14. pathogenicity tag

This tag provides information on pathogenicity.

Key	Condition	Data type	Description
pathogenicity	optional	array (length: 1-N, object)	Aggregation tag for information on pathogenicity.

14-1. Tags within pathogenicity tag

Key	Condition	Data type	Description
itemId	[required]	string regex: ^.+ \$	An ID assigned to a pathogenicity. It must be unique within this case report.
alterationItemIds	[required]	array (length: 1-N, string regex: ^.+ \$)	Target molecular alterations(itemId). Basically, a single itemId is described. However, if pathogenicity is composed of multiple itemIds, all relevant itemIds should be described.
knowledgeBase	[required]	string regex: ^.+ \$	Database name. e.g.) COSMIC, ClinVar
knowledgeBaseVersion	optional	string regex: ^.+ \$	Database version.
knowledgeBaseRecords	[required]	array (length: 1-N, object)	Database record information. Each object in the array must be unique.
recordId	optional (recommend)	string regex: ^.+ \$	Record ID.
field	required	array (length: 1-N, object)	Record information.
name	required	string regex: ^.+ \$	Field name. e.g.) CLNSIG
value	required	string regex:	Value for field name. e.g.) Pathogenic

		^.+ \$	
descriptions	optional	array (length: 1- N, string regex: ^.+ \$)	Comment. Refer to the contents tag in the comments tag for usable languages and new lines.

14-2. Example of pathogenicity tag

```
(Example)
"pathogenicity": [
{
  "itemId": "pathogenicity-001",
  "alterationItemIds": [
    "TEST001"
  ],
  "knowledgeBase": "COSMIC",
  "knowledgeBaseVersion": "vXX(20240123)",
  "knowledgeBaseRecords": [
    {
      "recordId": "COSM000",
      "field": [
        {
          "name": "MUTATION_SIGNIFICANCE_TIER",
          "value": "3"
        },
        {
          "name": "COSMIC_SAMPLE_MUTATED",
          "value": "1"
        }
      ]
    }
  ],
  "descriptions": [
    "MUTATION_SIGNIFICANCE_TIER: 3",
    "COSMIC_SAMPLE_MUTATED: 1"
  ]
},
{
  "itemId": "pathogenicity-002",
  "alterationItemIds": [
    "TEST002",
    "TEST003"
  ],
  "knowledgeBase": "ClinVar",
  "knowledgeBaseVersion": "20240321",
  "knowledgeBaseRecords": [
    {
      "recordId": "12345",
      "field": [
```

```
[
  {
    "name": "CLNREVSTAT",
    "value": "criteria_provided,_multiple_submitters,_no_conflicts"
  },
  {
    "name": "CLNSIG",
    "value": "Pathogenic"
  }
]
},
"descriptions": [
  "CLNREVSTAT: criteria_provided,_multiple_submitters,_no_conflicts",
  "CLNSIG: Pathogenic"
]
}
```


15. Other notes

Precautions.

15-1. itemId

The value of itemId must be unique within the file. The value can be any string.

15-1-1. Example of itemId description

Examples of itemId values for each tag are as follows.

- For variants tag

"itemId": "variant-1"

"itemId": "variant-2"

"itemId": "variant-3"

- For otherBiomarkers tag

"itemId": "biomarker-1"

"itemId": "biomarker-2"

"itemId": "biomarker-3"

- For sequencingSamples tag

"itemId": "sequence-1-tumor-dna"

"itemId": "sequence-2-tumor-rna"

"itemId": "sequence-3-normal-dna"

- For druggability tag

"itemId": "druggability-1"

"itemId": "druggability-2"

"itemId": "druggability-3"

15-2. matePieceLocation

Here, we provide an explanation for matePieceLocation in the breakends tag. We assume that "upstream" and "downstream" correspond to the direction of decreasing and increasing positional coordinates along a chromosome in a reference genome sequence, respectively.

Note: These "upstream" and "downstream" are different from those in the transcriptional direction.

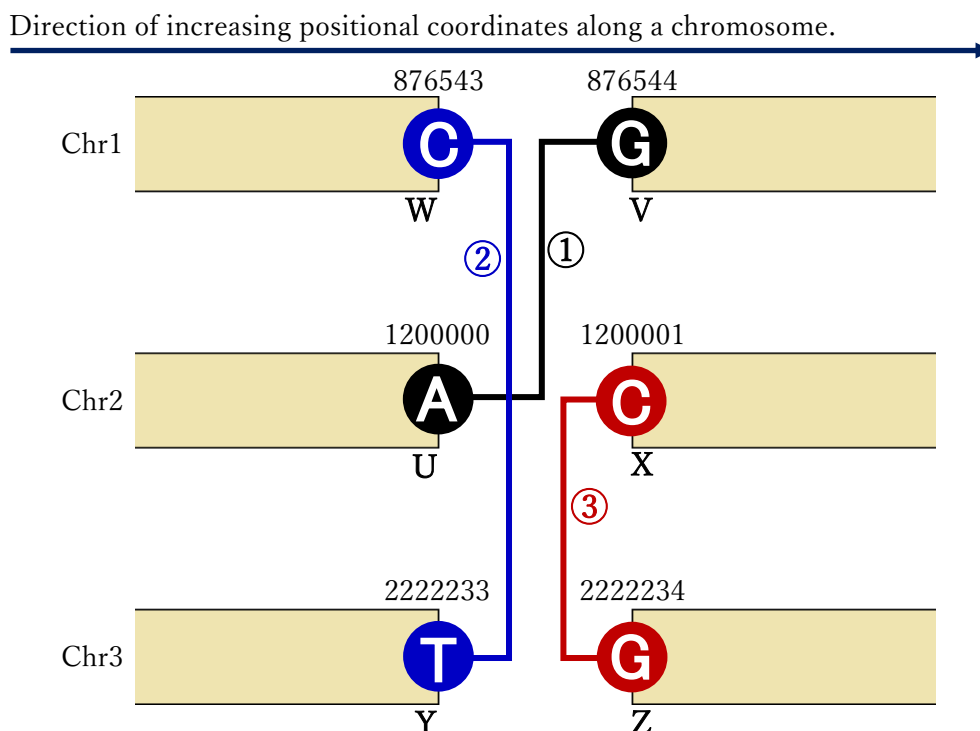
15-2-1. Example of matePieceLocation description

We illustrate how to input matePieceLocation and present representations in VCF format (v4.3). The examples below are modified from the figure in the following documents:

The Variant Call Format Specification VCF v4.5 and BCF v2.2

<https://samtools.github.io/hts-specs/VCFv4.5.pdf>

Example of structuralVariations:



Descriptions in the standardized format:

• Example①

For the chromosome 2 junction point, the downstream sequence of the junction point is replaced with another sequence; therefore, the value of matePieceLocation is "downstream".

In contrast, for the chromosome 1 junction point, the upstream sequence of the junction point is replaced with another sequence; therefore, the value of matePieceLocation is "upstream".

(Example①)

```
"breakends": [  
  {  
    "chromosome": "2",  
    "startPosition": 1200000,  
    "endPosition": 1200000,  
    "matePieceLocation": "downstream"  
  },  
  {  
    "chromosome": "1",  
    "startPosition": 876654,  
    "endPosition": 876654,  
    "matePieceLocation": "upstream"  
  }  
]
```

• Example②

For the chromosome 1 junction point, the downstream sequence of the junction point is replaced with another sequence; therefore, the value of matePieceLocation is "downstream".
The same is true for chromosome 3; thus, the value of matePieceLocation is "downstream".

(Example②)

```
"breakends": [  
  {  
    "chromosome": "1",  
    "startPosition": 876543,  
    "endPosition": 876543,  
    "matePieceLocation": "downstream"  
  },  
  {  
    "chromosome": "3",  
    "startPosition": 2222233,  
    "endPosition": 2222233,  
    "matePieceLocation": "downstream"  
  }  
]
```

• Example③

For the chromosome 2 junction point, the upstream sequence of the junction point is replaced with another sequence; therefore, the value of matePieceLocation is "upstream".
The same is true for the chromosome 3 junction point; thus, the value of matePieceLocation is "upstream".

(Example③)

```
"breakends": [  
  {  
    "chromosome": "2",  
    "startPosition": 1200001,  
    "endPosition": 1200001,  
    "matePieceLocation": "upstream"  
  },  
  {  
    "chromosome": "3",  
    "startPosition": 2222234,  
    "endPosition": 2222234,  
    "matePieceLocation": "upstream"  
  }  
]
```

Description in VCF format:

The following representation for the above figure is based on VCF v4.5: The numbers in the leftmost column in the table below correspond to the numbers in the figure above.

	#CHROM	POS	ID	REF	ALT	QUAL	FILTER	INFO
①	1	876544	bnd_V	G]2:1200000]G	6	PASS	SVTYPE=BN
	2	1200000	bnd_U	A	A[1:876544[6	PASS	SVTYPE=BN
②	1	876543	bnd_W	C	C]3:2222233]	6	PASS	SVTYPE=BN
	3	2222233	bnd_Y	T	T]1:876543]	6	PASS	SVTYPE=BN
③	2	1200001	bnd_X	C	[3:2222234[C	6	PASS	SVTYPE=BN
	3	2222234	bnd_Z	G	[2:1200001[G	6	PASS	SVTYPE=BN

16. For inquiries

Please contact the C-CAT Help Desk.

E-Mail: helpdesk_c-cat@ml.res.ncc.go.jp