
vcfpy Documentation

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Installation Getting Started

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VCFPy is a Python 3 library with good support for both reading and writing VCF files. The documentation is split into three parts (accessible through the navigation on the left):

Installation & Getting Started Instructions for the installation of the module and some examples to get you started.

API Documentation This section contains the API documentation for the module

Project Info More information on the project, including the changelog, list of contributing authors, and contribution instructions.

CHAPTER 1

Quick Example

```
#!/usr/bin/env python
# -*- coding: utf-8 -*-
import vcfpy

# Open input, add FILTER header, and open output file
reader = vcfpy.Reader.from_path('input.vcf')
reader.header.add_filter_line(vcfpy.OrderedDict([
    ('ID', 'DP10'), ('Description', 'total DP < 10')]))
writer = vcfpy.Writer.from_path('/dev/stdout', reader.header)

# Add "DP10" filter to records having less than 10 reads
for record in reader:
    ad = sum(c.data.get('DP', 0) for c in record.calls)
    if ad < 10:
        record.add_filter('DP10')
    writer.write_record(record)
```


CHAPTER 2

Features

- Support for reading and writing VCF v4.3
- Interface to `INFO` and `FORMAT` fields is based on `OrderedDict` allows for easier modification than `PyVCF` (also I find this more pythonic)
- Read (and jump in) and write BGZF files just using `vcfpy`

Frequently Asked Questions

Why another Python library for VCF? I've been using PyVCF with quite some success in the past. However, the main bottleneck of PyVCF is when you want to modify the per-sample genotype information. There are some issues in the tracker of PyVCF but none of them can really be considered solved. I tried several hours to solve these problems within PyVCF but this never got far or towards a complete rewrite...

For this reason, VCFPy was born and here it is!

Why Python 3 only? As I'm only using Python 3 code, I see no advantage in carrying around support for legacy Python 2 and maintaining it. At a later point when VCFPy is known to be stable, Python 2 support might be added if someone contributes a pull request.

What's the state? VCFPy is the result of two full days of development plus some maintenance work later now (right now). I'm using it in several projects but it is not as battle-tested as PyVCF.

What's the difference to PyVCF? The main difference is technical. Instead of using `collections.namedtuple` for storing the call annotation, VCFPy uses `collections.OrderedDict`. This has the advantage that (1) access to optional settings is much more pythonic using `.get(KEY, DEFAULT)` instead of `getattr()`. Further, (2) adding call annotations (FORMAT) fields is able without any performance penalty where for PyVCF, `copy.deepcopy` has to be used at some point which is very slow. There has not been any movement in supporting modifying FORMAT fields in PyVCF and here is a library that does this well.

What's the aim? The aim of the project is to provide simple yet efficient read and write access to VCF files. Eventually, PySAM will probably be a better choice once it has a Python wrapper for the VCF part of `htslib`. However, as this is still missing, VCFPy is a good solution for the time being.

3.1 Installation

3.1.1 Stable release

To install vcfpy, run this command in your terminal:

```
$ pip install vcfpy
```

This is the preferred method to install VCFPy, as it will always install the most recent stable release.

If you don't have `pip` installed, this [Python installation guide](#) can guide you through the process.

3.1.2 From sources

The sources for vcfpy can be downloaded from the [Github repo](#).

You can either clone the public repository:

```
$ git clone git://github.com/bihealth/vcfpy
```

Or download the [tarball](#):

```
$ curl -OL https://github.com/bihealth/vcfpy/tarball/master
```

Once you have a copy of the source, you can install it with:

```
$ python setup.py install
```

3.2 Getting Started

After installation, you can use VCFPy in your project simply by importing the module.

```
import vcfpy
```

That's all, continue and look at the list of examples.

3.3 Examples

This chapter contains several examples for the most important use cases of VCFPy.

3.3.1 Reading VCF Files

The following is an example for reading VCF files and writing out a TSV file with the genotype calls of all SNVs. You can find the example Python and VCF file in the sources below the directory `examples/vcf_to_tsv`.

```
#!/usr/bin/env python
# -*- coding: utf-8 -*-
import vcfpy

# Open file, this will read in the header
reader = vcfpy.Reader.from_path('input.vcf')

# Build and print header
header = ['#CHROM', 'POS', 'REF', 'ALT'] + reader.header.samples.names
print('\t'.join(header))

for record in reader:
    if not record.is_snv():
        continue
    line = [record.CHROM, record.POS, record.REF]
```

```

line += [alt.value for alt in record.ALT]
line += [call.data.get('GT') or './.' for call in record.calls]
print('\t'.join(map(str, line)))

```

The program call looks as follows.

```

$ ./vcf_to_tsv.py
↪ #CHROM      POS      REF      ALT      BLANK      NA12878      NA12891      NA12892
chr22      42522392      G      A      0/0      0/1      0/1      0/
↪ 0      0/0      0/0      0/0
chr22      42522597      C      T      0/1      0/0      0/0      0/
↪ 0      0/0      0/0      0/0
chr22      42522613      G      C      0/1      0/1      0/0      0/
↪ 1      0/1      0/1      0/1
chr22      42523003      A      G      0/1      1/1      0/1      0/
↪ 1      0/1      0/1      0/1
chr22      42523209      T      C      0/1      1/1      0/1      0/
↪ 1      0/1      0/1      0/1
chr22      42523211      T      C      0/0      0/1      0/1      0/
↪ 0      0/0      0/0      0/0
chr22      42523409      G      T      0/1      0/1      0/0      0/
↪ 1      0/1      0/1      0/1
chr22      42523491      C      T      0/1      0/0      0/0      0/
↪ 0      0/0      0/0      0/0
chr22      42523507      A      G      0/1      0/0      0/0      0/
↪ 0      0/0      0/0      0/0
chr22      42523805      C      T      0/0      0/0      0/1      0/
↪ 0      0/0      0/0      0/0
chr22      42523943      A      G      0/1      1/1      0/1      0/
↪ 1      0/1      0/1      0/1
chr22      42524435      T      A      0/1      0/0      0/0      0/
↪ 0      0/0      0/1      0/1
[...]
```

3.3.2 Writing VCF Files

The following shows how to add values to the FILTER column to records of an existing VCF file. Adding to existing records is simpler than constructing them from scratch, of course.

```

#!/usr/bin/env python
# -*- coding: utf-8 -*-
import vcfpy

# Open input, add FILTER header, and open output file
reader = vcfpy.Reader.from_path('input.vcf')
reader.header.add_filter_line(vcfpy.OrderedDict([
    ('ID', 'DP10'), ('Description', 'total DP < 10')]))
writer = vcfpy.Writer.from_path('/dev/stdout', reader.header)

# Add "DP10" filter to records having less than 10 reads
for record in reader:
    ad = sum(c.data.get('DP', 0) for c in record.calls)
    if ad < 10:
        record.add_filter('DP10')
    writer.write_record(record)

```

The program call looks as follows.

```
##fileformat=VCFv4.3
##contig=<ID=20,length=62435964>
##INFO=<ID=NS,Number=1,Type=Integer,Description="Number of Samples With Data">
##INFO=<ID=DP,Number=1,Type=Integer,Description="Total Depth">
##INFO=<ID=AF,Number=A,Type=Float,Description="Allele Frequency">
##INFO=<ID=AA,Number=1,Type=String,Description="Ancestral Allele">
##INFO=<ID=DB,Number=0,Type=Flag,Description="dbSNP membership, build 129">
##INFO=<ID=H2,Number=0,Type=Flag,Description="HapMap2 membership">
##FILTER=<ID=q10,Description="Quality below 10">
##FILTER=<ID=s50,Description="Less than 50% of samples have data">
##FORMAT=<ID=GT,Number=1,Type=String,Description="Genotype">
##FORMAT=<ID=GQ,Number=1,Type=Integer,Description="Genotype Quality">
##FORMAT=<ID=DP,Number=1,Type=Integer,Description="Read Depth">
##FORMAT=<ID=HQ,Number=2,Type=Integer,Description="Haplotype Quality">
##FILTER=<ID=DP10,Description="total DP < 10">
```

CHROM	POS	ID	REF	ALT	QUAL	FILTER	INFO	FORMAT
20	14370	rs6054257	G	A	29	PASS	NS=3;	GT:GQ:DP:HQ
								0 0:48:1:51,51
								1 0:48:8:51,
								51
								1/1:43:5:.,.
20	17330	.	T	A	3	q10	NS=3;DP=11;AF=0.	GT:GQ:DP:HQ
								0 0:49:3:58,50
								0 1:3:5:65,3
								0/0:41:3:40,
								30
20	1110696	rs6040355	A	G,				GT:GQ:DP:HQ
								0 0:54:7:56,60
								0 0:48:4:51,51
								0/
								0:61:2:40,30
20	1230237	.	T	.	47	PASS	NS=3;DP=13;	GT:GQ:DP:HQ
								0 0:54:7:56,60
								0 0:48:4:51,51
								0/
								0:61:2:40,30
20	1234567	microsat1	GTC	G,GTCT	50	PASS;		GT:GQ:DP
								0/1:35:4
								0/2:17:2
								1/
								1:40:3

3.3.3 Jumping in Tabix-indexed Files

The following shows a small program that extracts a genomic region from the input VCF file and writes it to stdout.

```
#!/usr/bin/env python
# -*- coding: utf-8 -*-
import vcfpy

# Open input, add FILTER header, and open output file
reader = vcfpy.Reader.from_path('input.vcf.gz')
writer = vcfpy.Writer.from_path('/dev/stdout', reader.header)

# Fetch region 20:1,110,694-1,230,237. Note that the coordinates
# in the API call are zero-based and describe half-open intervals.
for record in reader.fetch('20', 1110695, 1230237):
    writer.write_record(record)
```

The program call looks as follows.

```
##fileformat=VCFv4.3
##fileDate=20090805
##source=myImputationProgramV3.1
```

```
##reference=file:///seq/references/1000GenomesPilot-NCBI36.fasta
##contig=<ID=1,length=249250621>
##contig=<ID=2,length=243199373>
##contig=<ID=20,length=62435964>
##phasing=partial
##INFO=<ID=NS,Number=1,Type=Integer,Description="Number of Samples With Data">
##INFO=<ID=DP,Number=1,Type=Integer,Description="Total Depth">
##INFO=<ID=AF,Number=A,Type=Float,Description="Allele Frequency">
##INFO=<ID=AA,Number=1,Type=String,Description="Ancestral Allele">
##INFO=<ID=DB,Number=0,Type=Flag,Description="dbSNP membership, build 129">
##INFO=<ID=H2,Number=0,Type=Flag,Description="HapMap2 membership">
##FILTER=<ID=q10,Description="Quality below 10">
##FILTER=<ID=s50,Description="Less than 50% of samples have data">
##FORMAT=<ID=GT,Number=1,Type=String,Description="Genotype">
##FORMAT=<ID=GQ,Number=1,Type=Integer,Description="Genotype Quality">
##FORMAT=<ID=DP,Number=1,Type=Integer,Description="Read Depth">
##FORMAT=<ID=HQ,Number=2,Type=Integer,Description="Haplotype Quality">

#CHROM      POS      ID      REF      ALT      QUAL      FILTER      INFO      FOR
20          1110696      rs6040355      A      G,
#T          67      PASS      NS=2;DP=10;AF=0.333,0.667;AA=T;
#DB          GT:GQ:DP:HQ|2:21:6:23,27      2|1:2:0:18,2      2/2:35:4:40,30
20          1230237      .      T      .      47      PASS      NS=3;DP=13;
#AA=T      GT:GQ:DP:HQ      0|0:54:7:56,60      0|0:48:4:51,51      0/
#0:61:2:40,30
```

3.4 Best Practice

While not strictly part of the documentation of VCFPy, we include some notes on hints that we consider best practice when building VCF processing applications.

3.4.1 Keep Input Verbatim Where Possible

Try to keep the input verbatim if there is no strong reason for adjusting it. Strong reasons include fixing `Type` or `Number` in header lines describing arrays of strings, for example.

Whenever possible, keep the header order intact. VCFPy does this automatically for you (in contrast to PyVCF).

3.4.2 Prefer Soft-Filters over Hard-Filters

Soft-filters mean annotating your VCF records in the `FILTER` column whereas **Hard**-filters mean removing records from VCF file. In many situations, it is useful to keep around all VCF records and just annotate why they are to be dropped. Then, in the last step, only the interesting ones are kept.

This makes tracing back easier when and why a record was removed.

3.5 Header

Contents

- *Header*
 - *vcfpy.OrderedDict*
 - *vcfpy.Header*
 - *vcfpy.HeaderLine*
 - *vcfpy.header_without_lines*
 - *vcfpy.SimpleHeaderLine*
 - *vcfpy.AltAlleleHeaderLine*
 - *vcfpy.MetaHeaderLine*
 - *vcfpy.PedigreeHeaderLine*
 - *vcfpy.SampleHeaderLine*
 - *vcfpy.ContigHeaderLine*
 - *vcfpy.FilterHeaderLine*
 - *vcfpy.CompoundHeaderLine*
 - *vcfpy.InfoHeaderLine*
 - *vcfpy.FormatHeaderLine*
 - *vcfpy.FieldInfo*
 - *vcfpy.SamplesInfos*

3.5.1 vcfpy.OrderedDict

Convenience export of `OrderedDict`. When available, the `cyorderreddict`, a Cython-reimplementation of `OrderedDict` is used for Python before 3.5 (from 3.5, Python ships with a fast, C implementation of `OrderedDict`).

class `vcfpy.OrderedDict`

Dictionary that remembers insertion order

clear () → None. Remove all items from od.

copy () → a shallow copy of od

fromkeys (*S* [, *v*]) → New ordered dictionary with keys from *S*.
If not specified, the value defaults to None.

move_to_end ()
Move an existing element to the end (or beginning if `last=False`).

Raises `KeyError` if the element does not exist. When `last=True`, acts like a fast version of `self[key]=self.pop(key)`.

pop (*k* [, *d*]) → *v*, remove specified key and return the corresponding value. If key is not found, *d* is returned if given, otherwise `KeyError` is raised.

popitem () → (*k*, *v*), return and remove a (key, value) pair.
Pairs are returned in LIFO order if `last` is true or FIFO order if false.

setdefault (k, d) $\rightarrow$ `od.get(k,d)`, also set `od[k]=d` if `k` not in `od`

3.5.2 vcfpy.Header

class `vcfpy.Header` (*lines=None, samples=None*)

Represent header of VCF file

While this class allows mutating records, it should not be changed once it has been assigned to a writer. Use **`:py:method:`~Header.copy``** to create a copy that can be modified without problems.

This class provides function for adding lines to a header and updating the supporting index data structures. There is no explicit API for removing header lines, the best way is to reconstruct a new `Header` instance with a filtered list of header lines.

add_contig_line (*mapping*)

Add “contig” header line constructed from the given mapping

Parameters **mapping** – `OrderedDict` with mapping to add. It is recommended to use `OrderedDict` over `dict` as this makes the result reproducible

Returns `False` on conflicting line and `True` otherwise

add_filter_line (*mapping*)

Add FILTER header line constructed from the given mapping

Parameters **mapping** – `OrderedDict` with mapping to add. It is recommended to use `OrderedDict` over `dict` as this makes the result reproducible

Returns `False` on conflicting line and `True` otherwise

add_format_line (*mapping*)

Add FORMAT header line constructed from the given mapping

Parameters **mapping** – `OrderedDict` with mapping to add. It is recommended to use `OrderedDict` over `dict` as this makes the result reproducible

Returns `False` on conflicting line and `True` otherwise

add_info_line (*mapping*)

Add INFO header line constructed from the given mapping

Parameters **mapping** – `OrderedDict` with mapping to add. It is recommended to use `OrderedDict` over `dict` as this makes the result reproducible

Returns `False` on conflicting line and `True` otherwise

add_line (*header_line*)

Add header line, updating any necessary support indices

Returns `False` on conflicting line and `True` otherwise

copy ()

Return a copy of this header

filter_ids ()

Return list of all filter IDs

format_ids ()

Return list of all format IDs

get_format_field_info (*key*)

Return *FieldInfo* for the given INFO field

get_info_field_info(*key*)

Return *FieldInfo* for the given INFO field

get_lines(*key*)

Return header lines having the given *key* as their type

has_header_line(*key*, *id_*)

Return whether there is a header line with the given ID of the type given by *key*

Parameters

- **key** – The VCF header key/line type.
- **id** – The ID value to compare fore

Returns *True* if there is a header line starting with `##${key}=` in the VCF file having the mapping entry ID set to *id_*.

info_ids()

Return list of all info IDs

lines = *None*

list of `:py:HeaderLine` objects

samples = *None*

SamplesInfo object

3.5.3 vcfpy.HeaderLine

class `vcfpy.HeaderLine`(*key*, *value*)

Base class for VCF header lines

copy()

Return a copy

key = *None*

str with key of header line

serialize()

Return VCF-serialized version of this header line

3.5.4 vcfpy.header_without_lines

`vcfpy.header_without_lines`(*header*, *remove*)

Return *Header* without lines given in *remove*

remove is an iterable of pairs *key*/*ID* with the VCF header key and ID of entry to remove. In the case that a line does not have a mapping entry, you can give the full value to remove.

```
# header is a vcfpy.Header, e.g., as read earlier from file
new_header = vcfpy.without_header_lines(
    header, [('assembly', None), ('FILTER', 'PASS')])
# now, the header lines starting with "##assembly=" and the "PASS"
# filter line will be missing from new_header
```

3.5.5 vcfpy.SimpleHeaderLine

```
class vcfpy.SimpleHeaderLine (key, value, mapping)
    Base class for simple header lines, currently contig and filter header lines

    Don't use this class directly but rather the sub classes.

    Raises vcfpy.exceptions.InvalidHeaderException in the case of missing key "ID"

    copy ()
        Return a copy

    mapping = None
        collections.OrderedDict with key/value mapping of the attributes
```

3.5.6 vcfpy.AltAlleleHeaderLine

```
class vcfpy.AltAlleleHeaderLine (key, value, mapping)
    Alternative allele header line

    Mostly used for defining symbolic alleles for structural variants and IUPAC ambiguity codes

    classmethod from_mapping (mapping)
        Construct from mapping, not requiring the string value

    id = None
        name of the alternative allele
```

3.5.7 vcfpy.MetaHeaderLine

```
class vcfpy.MetaHeaderLine (key, value, mapping)
    Alternative allele header line

    Used for defining set of valid values for samples keys

    classmethod from_mapping (mapping)
        Construct from mapping, not requiring the string value

    id = None
        name of the alternative allele
```

3.5.8 vcfpy.PedigreeHeaderLine

```
class vcfpy.PedigreeHeaderLine (key, value, mapping)
    Header line for defining a pedigree entry

    classmethod from_mapping (mapping)
        Construct from mapping, not requiring the string value

    id = None
        name of the alternative allele
```

3.5.9 vcfpy.SampleHeaderLine

```
class vcfpy.SampleHeaderLine (key, value, mapping)
    Header line for defining a SAMPLE entry
```

```
classmethod from_mapping (mapping)  
    Construct from mapping, not requiring the string value  
  
id = None  
    name of the alternative allele
```

3.5.10 vcfpy.ContigHeaderLine

```
class vcfpy.ContigHeaderLine (key, value, mapping)  
    Contig header line  
  
    Most importantly, parses the 'length' key into an integer  
  
classmethod from_mapping (mapping)  
    Construct from mapping, not requiring the string value  
  
id = None  
    name of the contig  
  
length = None  
    length of the contig, None if missing
```

3.5.11 vcfpy.FilterHeaderLine

```
class vcfpy.FilterHeaderLine (key, value, mapping)  
    FILTER header line  
  
    description = None  
        description for the filter, None if missing  
  
classmethod from_mapping (mapping)  
    Construct from mapping, not requiring the string value  
  
id = None  
    token for the filter
```

3.5.12 vcfpy.CompoundHeaderLine

```
class vcfpy.CompoundHeaderLine (key, value, mapping)  
    Base class for compound header lines, currently format and header lines  
  
    Compound header lines describe fields that can have more than one entry.  
  
    Don't use this class directly but rather the sub classes.  
  
    copy ()  
        Return a copy  
  
    mapping = None  
        OrderedDict with key/value mapping
```

3.5.13 vcfpy.InfoHeaderLine

```
class vcfpy.InfoHeaderLine (key, value, mapping)  
    Header line for INFO fields
```

Note that the `Number` field will be parsed into an `int` if possible. Otherwise, the constants `HEADER_NUMBER_*` will be used.

description = None
description, should be given, None if not given

classmethod from_mapping(mapping)
Construct from mapping, not requiring the string value

id = None
key in the INFO field

source = None
source of INFO field, None if not given

type = None
value type

version = None
version of INFO field, None if not given

3.5.14 vcfpy.FormatHeaderLine

class vcfpy.FormatHeaderLine(key, value, mapping)
Header line for FORMAT fields

description = None
description, should be given, None if not given

classmethod from_mapping(mapping)
Construct from mapping, not requiring the string value

id = None
key in the INFO field

source = None
source of INFO field, None if not given

type = None
value type

version = None
version of INFO field, None if not given

3.5.15 vcfpy.FieldInfo

class vcfpy.FieldInfo(type_, number, description=None, id_=None)
Core information for describing field type and number

description = None
Description for the header field, optional

id = None
The id of the field, optional.

number = None
Number description, either an int or constant

type = None
The type, one of `INFO_TYPES` or `FORMAT_TYPES`

3.5.16 vcfpy.SamplesInfos

class vcfpy.SamplesInfos (*sample_names*, *parsed_samples=None*)

Helper class for handling the samples in VCF files

The purpose of this class is to decouple the sample name list somewhat from *Header*. This encapsulates subsetting samples for which the genotype should be parsed and reordering samples into output files.

Note that when subsetting is used and the records are to be written out again then the `FORMAT` field must not be touched.

copy ()

Return a copy of the object

is_parsed (*name*)

Return whether the sample name is parsed

name_to_idx = None

mapping from sample name to index

names = None

list of sample that are read from/written to the VCF file at hand in the given order

parsed_samples = None

set with the samples for which the genotype call fields should be read; can be used for partial parsing (speedup) and defaults to the full list of samples, None if all are parsed

3.6 Input/Output

Contents

- *Input/Output*
 - *vcfpy.Reader*
 - *vcfpy.Writer*

3.6.1 vcfpy.Reader

class vcfpy.Reader (*stream*, *path=None*, *tabix_path=None*, *record_checks=None*,
parsed_samples=None)

Class for parsing of files from file-like objects

Instead of using the constructor, use the class methods *from_stream()* and *from_path()*.

On construction, the header will be read from the file which can cause problems. After construction, *Reader* can be used as an iterable of *Record*.

Raises `InvalidHeaderException` in the case of problems reading the header

Note: It is important to note that the `header` member is used during the parsing of the file. **If you need a modified version then create a copy, e.g., using `:py:method:~vcfpy.header.Header.copy`.**

Note: If you use the `parsed_samples` feature and you write out records then you must not change the `FORMAT` of the record.

close()

Close underlying stream

fetch(*chrom_or_region*, *begin=None*, *end=None*)

Jump to the start position of the given chromosomal position and limit iteration to the end position

Parameters

- **chrom_or_region** (*str*) – name of the chromosome to jump to if *begin* and *end* are given and a samtools region string otherwise (e.g. “chr1:123,456-123,900”).
- **begin** (*int*) – 0-based begin position (inclusive)
- **end** (*int*) – 0-based end position (exclusive)

classmethod from_path(*path*, *tabix_path=None*, *record_checks=None*, *parsed_samples=None*)

Create new [Reader](#) from path

Note: If you use the `parsed_samples` feature and you write out records then you must not change the `FORMAT` of the record.

Parameters

- **path** – the path to load from (converted to `str` for compatibility with `path.py`)
- **tabix_path** – optional string with path to TBI index, automatic inferral from `path` will be tried on the fly if not given
- **record_checks** (*list*) – record checks to perform, can contain ‘INFO’ and ‘FORMAT’

classmethod from_stream(*stream*, *path=None*, *tabix_path=None*, *record_checks=None*, *parsed_samples=None*)

Create new [Reader](#) from file

Note: If you use the `parsed_samples` feature and you write out records then you must not change the `FORMAT` of the record.

Parameters

- **stream** – file-like object to read from
- **path** – optional string with path to store (for display only)
- **record_checks** (*list*) – record checks to perform, can contain ‘INFO’ and ‘FORMAT’
- **parsed_samples** (*list*) – list of `str` values with names of samples to parse call information for (for speedup); leave to `None` for ignoring

header = None

the Header

parsed_samples = None
if set, list of samples to parse for

parser = None
the parser to use

path = None
optional `str` with the path to the stream

record_checks = None
checks to perform on records, can contain 'FORMAT' and 'INFO'

stream = None
stream (file-like object) to read from

tabix_file = None
the `pysam.TabixFile` used for reading from index bgzip-ed VCF; constructed on the fly

tabix_path = None
optional `str` with path to tabix file

3.6.2 vcfpy.Writer

class `vcfpy.Writer` (*stream*, *header*, *path=None*)
Class for writing VCF files to file-like objects

Instead of using the constructor, use the class methods `from_stream()` and `from_path()`.

The writer has to be constructed with a `Header` object and the full VCF header will be written immediately on construction. This, of course, implies that modifying the header after construction is illegal.

close()
Close underlying stream

classmethod `from_path` (*path*, *header*)
Create new `Writer` from path

Parameters

- **path** – the path to load from (converted to `str` for compatibility with `path.py`)
- **header** – VCF header to use, lines and samples are deep-copied

classmethod `from_stream` (*stream*, *header*, *path=None*, *use_bgzf=None*)
Create new `Writer` from file

Note that for getting bgzf support, you have to pass in a stream opened in binary mode. Further, you either have to provide a `path` ending in `".gz"` or set `use_bgzf=True`. Otherwise, you will get the notorious “TypeError: ‘str’ does not support the buffer interface”.

Parameters

- **stream** – file-like object to write to
- **header** – VCF header to use, lines and samples are deep-copied
- **path** – optional string with path to store (for display only)
- **use_bgzf** – indicator whether to write bgzf to `stream` if `True`, prevent if `False`, interpret `path` if `None`

header = None
the `:py:class:`~vcfpy.header.Header` to write out, will be deep-copied into the Writer on initialization`

path = **None**
 optional `str` with the path to the stream

stream = **None**
 stream (`file`-like object) to read from

write_record(*record*)
 Write out the given `vcfpy.record.Record` to this `Writer`

3.7 Exceptions

Contents

- *Exceptions*
 - *vcfpy.VCFPyException*
 - *vcfpy.InvalidHeaderException*
 - *vcfpy.InvalidRecordException*
 - *vcfpy.IncorrectVCFFormat*
 - *vcfpy.HeaderNotFound*

3.7.1 vcfpy.VCFPyException

exception `vcfpy.VCFPyException`
 Base class for module's exception

3.7.2 vcfpy.InvalidHeaderException

exception `vcfpy.InvalidHeaderException`
 Raised in the case of invalid header formatting

3.7.3 vcfpy.InvalidRecordException

exception `vcfpy.InvalidRecordException`
 Raised in the case of invalid record formatting

3.7.4 vcfpy.IncorrectVCFFormat

exception `vcfpy.IncorrectVCFFormat`
 Raised on problems parsing VCF

3.7.5 vcfpy.HeaderNotFound

exception `vcfpy.HeaderNotFound`
 Raised when a VCF header could not be found

3.8 Records

Contents

- *Records*
 - *Record-Related Constants*
 - *vcfpy.Record*
 - *vcfpy.Call*
 - *vcfpy.AltRecord*
 - *vcfpy.Substitution*
 - *vcfpy.SV*
 - *vcfpy.BreakEnd*
 - *vcfpy.SingleBreakEnd*
 - *vcfpy.SymbolicAllele*

3.8.1 Record-Related Constants

The following constants are also available as `vcfpy.CONSTANT`.

```
vcfpy.record.HOM_REF = 0
    Code for homozygous reference

vcfpy.record.HOM_ALT = 2
    Code for homozygous alternative

vcfpy.record.FIVE_PRIME = '5'
    code for five prime orientation BreakEnd

vcfpy.record.THREE_PRIME = '3'
    code for three prime orientation BreakEnd

vcfpy.record.FORWARD = '+'
    code for forward orientation

vcfpy.record.REVERSE = '-'
    code for reverse orientation
```

3.8.2 vcfpy.Record

```
class vcfpy.Record(CHROM, POS, ID, REF, ALT, QUAL, FILTER, INFO, FORMAT, calls)
    Represent one record from the VCF file

    Record objects are iterators of their calls

    ALT = None
        A list of alternative allele records of type AltRecord

    CHROM = None
        A str with the chromosome name
```

FILTER = None

A list of strings for the FILTER column

FORMAT = None

A list of strings for the FORMAT column

ID = None

A list of the semicolon-separated values of the ID column

INFO = None

An OrderedDict giving the values of the INFO column, flags are mapped to True

POS = None

An int with a 1-based begin position

QUAL = None

The quality value, can be None

REF = None

A str with the REF value

add_filter (*label*)

Add label to FILTER if not set yet, removing PASS entry if present

add_format (*key*, *value=None*)

Add an entry to format

The record's calls `data[key]` will be set to `value` if not yet set and `value` is not None. If `key` is already in `FORMAT` then nothing is done.

affected_end

Return affected start position in 0-based coordinates

For SNVs, MNVs, and deletions, the behaviour is based on the start position and the length of the REF. In the case of insertions, the position behind the insert position is returned, yielding a 0-length interval together with `affected_start()`

affected_start

Return affected start position in 0-based coordinates

For SNVs, MNVs, and deletions, the behaviour is the start position. In the case of insertions, the position behind the insert position is returned, yielding a 0-length interval together with `affected_end()`

begin = None

An int with a 0-based begin position

call_for_sample = None

A mapping from sample name to entry in `self.calls`

calls = None

A list of genotype *Call* objects

end = None

An int with a 0-based end position

is_snv()

Return True if it is a SNV

3.8.3 vcfpy.Call

class vcfpy.Call (*sample*, *data*, *site=None*)

The information for a genotype callable

By VCF, this should always include the genotype information and can contain an arbitrary number of further annotation, e.g., the coverage at the variant position.

called = None

whether or not the variant is fully called

data = None

an OrderedDict with the key/value pair information from the call's data

gt_alleles = None

the allele numbers (0, 1, ...) in this calls or None for no-call

gt_bases

Return the actual genotype bases, e.g. if VCF genotype is 0/1, could return ('A', 'T')

gt_phase_char

Return character to use for phasing

gt_type

The type of genotype, returns one of HOM_REF, HOM_ALT, and HET.

is_filtered (*require=None, ignore=None*)

Return True for filtered calls

Parameters

- **ignore** (*iterable*) – if set, the filters to ignore, make sure to include 'PASS', when setting, default is ['PASS']
- **require** (*iterable*) – if set, the filters to require for returning True

is_het

Return True for heterozygous calls

is_phased

Return boolean indicating whether this call is phased

is_variant

Return True for non-hom-ref calls

plodity = None

the number of alleles in this sample's call

sample = None

the name of the sample for which the call was made

site = None

the *Record* of this *Call*

3.8.4 vcfpy.AltRecord

class vcfpy.AltRecord (*type_=None*)

An alternative allele Record

Currently, can be a substitution, an SV placeholder, or breakend

serialize()

Return str with representation for VCF file

type = None

String describing the type of the variant, could be one of SNV, MNV, could be any of teh types described in the ALT header lines, such as DUP, DEL, INS, ...

3.8.5 vcfpy.Substitution

class vcfpy.**Substitution** (*type_, value*)
A basic alternative allele record describing a REF->AltRecord substitution

Note that this subsumes MNVs, insertions, and deletions.

value = None
The alternative base sequence to use in the substitution

3.8.6 vcfpy.SV

vcfpy.SV

3.8.7 vcfpy.BreakEnd

class vcfpy.**BreakEnd** (*mate_chrom, mate_pos, orientation, mate_orientation, sequence, within_main_assembly*)
A placeholder for a breakend

mate_chrom = None
chromosome of the mate breakend

mate_orientation = None
orientation breakend's mate

mate_pos = None
position of the mate breakend

orientation = None
orientation of this breakend

sequence = None
breakpoint's connecting sequence

serialize()
Return string representation for VCF

within_main_assembly = None
bool specifying if the breakend mate is within the assembly (True) or in an ancillary assembly (False)

3.8.8 vcfpy.SingleBreakEnd

class vcfpy.**SingleBreakEnd** (*orientation, sequence*)
A placeholder for a single breakend

3.8.9 vcfpy.SymbolicAllele

class vcfpy.**SymbolicAllele** (*value*)
A placeholder for a symbolic allele

The allele symbol must be defined in the header using an ALT header before being parsed. Usually, this is used for succinct descriptions of structural variants or IUPAC parameters.

value = None
The symbolic value, e.g. 'DUP'

3.9 Contributing

Contributions are welcome, and they are greatly appreciated! Every little bit helps, and credit will always be given.

You can contribute in many ways:

3.9.1 Types of Contributions

Report Bugs

Report bugs at <https://github.com/bihealth/vcfpy/issues>.

If you are reporting a bug, please include:

- Your operating system name and version.
- Any details about your local setup that might be helpful in troubleshooting.
- Detailed steps to reproduce the bug.

Fix Bugs

Look through the GitHub issues for bugs. Anything tagged with “bug” and “help wanted” is open to whoever wants to implement it.

Implement Features

Look through the GitHub issues for features. Anything tagged with “enhancement” and “help wanted” is open to whoever wants to implement it.

Write Documentation

vcfpy could always use more documentation, whether as part of the official vcfpy docs, in docstrings, or even on the web in blog posts, articles, and such.

Submit Feedback

The best way to send feedback is to file an issue at <https://github.com/bihealth/vcfpy/issues>.

If you are proposing a feature:

- Explain in detail how it would work.
- Keep the scope as narrow as possible, to make it easier to implement.
- Remember that this is a volunteer-driven project, and that contributions are welcome :)

3.9.2 Get Started!

Ready to contribute? Here’s how to set up *vcfpy* for local development.

1. Fork the *vcfpy* repo on GitHub.
2. Clone your fork locally:

```
$ git clone git@github.com:your_name_here/vcfpy.git
```

3. Install your local copy into a virtualenv. Assuming you have virtualenvwrapper installed, this is how you set up your fork for local development:

```
$ mkvirtualenv vcfpy
$ cd vcfpy/
$ python setup.py develop
```

4. Create a branch for local development:

```
$ git checkout -b name-of-your-bugfix-or-feature
```

Now you can make your changes locally.

5. When you're done making changes, check that your changes pass flake8 and the tests, including testing other Python versions with tox:

```
$ flake8 vcfpy tests
$ python setup.py test or py.test
$ tox
```

To get flake8 and tox, just pip install them into your virtualenv.

6. Commit your changes and push your branch to GitHub:

```
$ git add .
$ git commit -m "Your detailed description of your changes."
$ git push origin name-of-your-bugfix-or-feature
```

7. Submit a pull request through the GitHub website.

3.9.3 Pull Request Guidelines

Before you submit a pull request, check that it meets these guidelines:

1. The pull request should include tests.
2. If the pull request adds functionality, the docs should be updated. Put your new functionality into a function with a docstring, and add the feature to the list in README.rst.
3. The pull request should work for Python 3.3, 3.4 and 3.5. Check https://travis-ci.org/bihealth/vcfpy/pull_requests and make sure that the tests pass for all supported Python versions.

3.9.4 Tips

To run a subset of tests:

```
$ py.test tests.test_vcfpy
```

3.10 Credits

3.10.1 Development Lead

- Manuel Holtgrewe <manuel.holtgrewe@bihealth.de>

3.10.2 Contributors

None yet. Why not be the first?

3.11 History

3.11.1 v0.11.2 (2018-04-16)

- Removing `pip` module from `setup.py` which is not recommended anyway.

3.11.2 v0.11.1 (2018-03-06)

- Working around problem in HTSJDK output with incomplete `FORMAT` fields (#127). Writing out `.` instead of keeping trailing empty records empty.

3.11.3 v0.11.0 (2017-11-22)

- The field `FORMAT/FT` is now expected to be a semicolon-separated string. Internally, we will handle it as a list.
- Switching from warning helper utility code to Python `warnings` module.
- Return `str` in case of problems with parsing value.

3.11.4 v0.10.0 (2017-02-27)

- Extending API to allow for reading subsets of records. (Writing for sample subsets or reordered samples is possible through using the appropriate `names` list in the `SamplesInfos` for the `Writer`).
- Deep-copying header lines and samples infos on `Writer` construction
- Using `samples` attribute from `Header` in `Reader` and `Writer` instead of passing explicitly

3.11.5 0.9.0 (2017-02-26)

- Restructuring of `requirements.txt` files
- Fixing parsing of no-call `GT` fields

3.11.6 0.8.1 (2017-02-08)

- PEP8 style adjustments
- Using versioneer for versioning
- Using `requirements*.txt` files now from `setup.py`
- Fixing dependency on `cyordereddict` to be for Python <3.6 instead of <3.5
- Jumping by samtools coordinate string now also allowed

3.11.7 0.8.0 (2016-10-31)

- Adding `Header.has_header_line` for querying existence of header line
- `Header.add*_line` return a `bool` no indicating any conflicts
- Construction of `Writer` uses samples within header and no extra parameter (breaks API)

3.11.8 0.7.0 (2016-09-25)

- Smaller improvements and fixes to documentation
- Adding Codacy coverage and static code analysis results to README
- Various smaller code cleanup triggered by Codacy results
- Adding `__eq__`, `__neq__` and `__hash__` to data types (where applicable)

3.11.9 0.6.0 (2016-09-25)

- Refining implementation for breakend and symbolic allele class
- Removing `record.SV_CODES`
- Refactoring parser module a bit to make the code cleaner
- Fixing small typos and problems in documentation

3.11.10 0.5.0 (2016-09-24)

- Deactivating warnings on record parsing by default because of performance
- Adding validation for `INFO` and `FORMAT` fields on reading (#8)
- Adding predefined `INFO` and `FORMAT` fields to `pyvcf.header` (#32)

3.11.11 0.4.1 (2016-09-22)

- Initially enabling codeclimate

3.11.12 0.4.0 (2016-09-22)

- Exporting constants for encoding variant types
- Exporting genotype constants HOM_REF, HOM_ALT, HET
- Implementing `Call.is_phased`, `Call.is_het`, `Call.is_variant`, `Call.is_phased`, `Call.is_hom_ref`, `Call.is_hom_alt`
- Removing `Call.phased` (breaks API, next release is 0.4.0)
- Adding tests, fixing bugs for methods of `Call`

3.11.13 0.3.1 (2016-09-21)

- Work around `FORMAT/FT` being a string; this is done so in the Delly output

3.11.14 0.3.0 (2016-09-21)

- `Reader` and `Writer` can now be used as context manager (with `with`)
- Including license in documentation, including Biopython license
- Adding support for writing bgzf files (taken from Biopython)
- Adding support for parsing arrays in header lines
- Removing `example-4.1-bnd.vcf` example file because v4.1 tumor derival lacks ID field
- Adding `AltAlleleHeaderLine`, `MetaHeaderLine`, `PedigreeHeaderLine`, and `SampleHeaderLine`
- Renaming `SimpleHeaderFile` to `SimpleHeaderLine`
- Warn on missing `FILTER` entries on parsing
- Reordered parameters in `from_stream` and `from_file` (#18)
- Renamed `from_file` to `from_stream` (#18)
- Renamed `Reader.jump_to` to `Reader.fetch`
- Adding `header_without_lines` function
- Generally extending API to make it esier to use
- Upgrading dependencies, enabling pyup-bot
- Greatly extending documentation

3.11.15 0.2.1 (2016-09-19)

- First release on PyPI

3.12 License

3.12.1 VCFPy License

You can find the License of VCFPy below.

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3.12.2 Biopython License Agreement

The bgzf writing code is taken from the Biopython project. You can find a copy of the license below.

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