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# **snpFC Documentation**

***Release 0.0.1***

**Ram Krishna Shrestha**

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# CHAPTER 1

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## Introduction

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snpFC - a python tool for filtering and comparing SNPs from two or multiple VCF files.



## CHAPTER 2

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### How can I get it

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At the moment, latest snpFC tool can be obtained from github repo [link](#) . Just clone the repository or download as a compressed file.

I have a target to upload snpFC to python repository enabling to install using pip command.



## CHAPTER 3

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### Pre-requisites

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Assuming you have python3 already, install pyVCF

```
pip install pyvcf
```



## CHAPTER 4

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### Quick Start

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If you don't have patience, here is how to run snpFC

```
python scripts/filter_and_comparesnps.py --vcf testfiles/test1.vcf testfiles/test2.  
↪vcf --filter --compare --show --frequency 70 --pvalue 0.05 --genotype heterozygous -  
↪-quality 10 --rawreaddepth 5 --qualityreaddepth 5 --depthreference 5 --depthvariant_  
↪5 --outdir ./testfiles
```

The above command will output as following

```
Common and Unique SNPS in vcf File : testfiles/test1.vcf  
tig000000003 775 A G common  
tig000000003 776 T C common  
tig000000003 836 G C unique  
tig000000004 1166 G A common  
tig000000004 1190 G A unique  
Common and Unique SNPS in vcf File : testfiles/test2.vcf  
tig000000003 775 A G common  
tig000000003 776 T C common  
tig000000003 836 G T unique  
tig000000004 1166 G A common  
tig000000004 1190 G A,C unique  
The outputs are saved in these files : /home/shrestha/Sites/test1_snpanalysis.txt /  
↪home/shrestha/Sites/test2_snpanalysis.txt
```



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## Options

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Available options:

|                    |                                                           |
|--------------------|-----------------------------------------------------------|
| --vcf              | space separated two <b>or</b> more vcf files              |
| --filter           | filter snps                                               |
| --compare          | compare snps between vcf files                            |
| --frequency        | frequency threshold value to filter (default: 70)         |
| --pvalue           | pvalue threshold value to filter ( default: 0.05)         |
| --genotype         | genotype to filter (default: heterozygous)                |
| --quality          | genotype quality threshold to filter (default: 10)        |
| --rawreaddepth     | raw read depth threshold to filter (default: 5)           |
| --qualityreaddepth | quality read depth threshold to filter (default:5)        |
| --depthreference   | depth <b>in</b> reference threshold to filter (default:5) |
| --depthvariant     | depth <b>in</b> variant threshold to filter (default:5)   |
| --show             | show the compared snps on screen                          |
| --outdir           | output directory                                          |

To check the options:

```
python scripts/filter_and_comparesnps.py --help
```

## 5.1 Modules

### 5.1.1 filter\_and\_comparesnps

### 5.1.2 vcfFilter

## 5.2 Contributions

The project is contributed by Dr Dan Maclean and Dr Ram Krishna Shrestha

The project is supported by Gatesby Foundation.

## 5.3 Contact

For help, please contact [ram\\_krishna.shrestha@tsl.ac.uk](mailto:ram_krishna.shrestha@tsl.ac.uk)

## CHAPTER 6

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### Indices and tables

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