Variation Annotation with SnpEff

SnpEff is a variant annotation and effect prediction tool. It annotates and predicts the effects of genetic variants (such as amino acid changes).

A typical SnpEff use case would be:

-Input: The inputs are predicted variants (SNPs, insertions, deletions and MNPs). The input file is usually obtained as a result of a sequencing experiment, and it is usually in variant call format (VCF).

-Output: SnpEff analyzes the input variants. It annotates the variants and calculates the effects they produce on known genes (e.g. amino acid changes).



How to Use This Sample

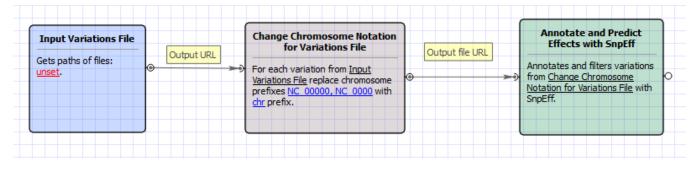
If you haven't used the workflow samples in UGENE before, look at the "How to Use Sample Workflows" section of the documentation.

Workflow Sample Location

The workflow sample "Variation Annotation with SnpEff" can be found in the "NGS" section of the Workflow Designer samples.

Workflow Image

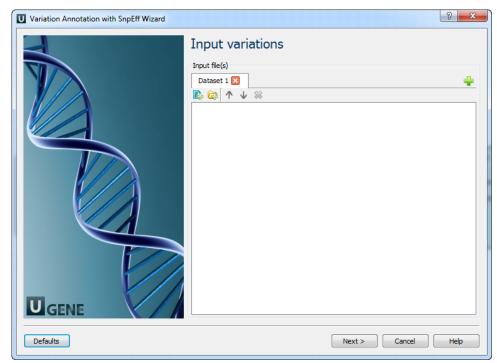
The opened workflow looks as follows:



Workflow Wizard

The wizard has 3 pages.

1. Input Variations: On this page you must input variations file(s).



2. Change chromosome notation for variations: On this page you can change the chromosome notation for variations.

U Variation Annotation with SnpEff Wizard			? ×
	Change chromosome notation for variations		
	Parameters Replace prefixes Replace by	NC_00000;NC_0000	
	Replace by	chr	
Defaults		< Back Next > Cancel	Help

The following parameters are available:

Replace prefixes	Input the list of chromosome prefixes that you would like to replace. For example "NC_000". Separate different prefixes by semicolons.
Replace by	Input the prefix that should be set instead, for example "chr".

3. <u>SnpEff Parameters:</u> On this page you can modify SnpEff parameters.

U Variation Annotation with SnpEff Wizard		<u>ନ୍</u>	
	SnpEff parameters Parameters Genome	Homo sapiens (hg19) 🔻	
	Additional Hide additional parameters Cannonical transcripts	ralse v	
	HGVS nomenclature Annotate Loss of function	False	
	Annotate TFBSs motifs Upstream/downstream length	False No upstream/downstream interval (0 bases)	
	1		
Defaults	< Ba	dk Next > Cancel Help	

The following parameters are available:

Genome	Select the target genome. Genome data will be downloaded if it is not found.	
Cannonical transcripts	Use only cannonical transcripts	

HGVS nomenclature	Annotate using HGVS nomenclature
Annotate Loss of function	Annotate Loss of function (LOF) and Nonsense mediated decay (NMD)
Annotate TFBSs motifs	Annotate transcription factor binding site motifs (only available for latest GRCh37)
Upstream/downstream length	Upstream and downstream interval size. Eliminate any upstream and downstream effect by using 0 length

4. Output: On this page you need input output parameters.

Variation Annotation with SnpEff Wizard				? ×
	Output Output folder Output directory Custom directory	Workflow		•
Defaults	< 8	ack Apply	Cancel Run	Help