





- eVaiutilities: Data management utilities for eVai
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DOI: 10.xxxxx/draft

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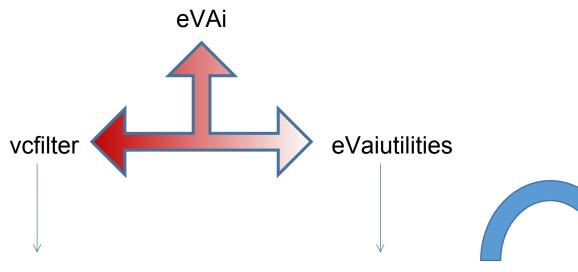
Funded by ABM project: Development of a universal fast-response platform, based on RNA technology, ensuring the national drug and epidemiological safety. 2021/ABM/05/00004-00



Key Features:

- *** RUST, Conda ,Asynchronous, SQLITE
- **···>** Population scale analysis
- **...** ▶ Variant summary for easy annotation.
- **···>** Annotate variant

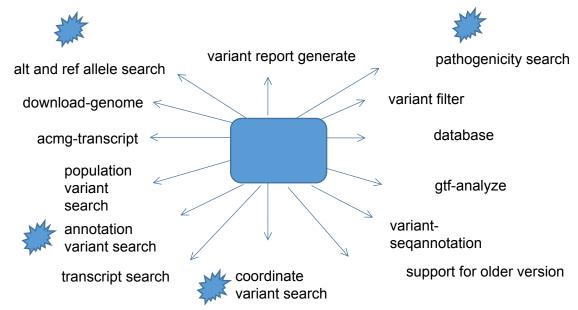




For population search across variants with specific coverage, quality and zygosity

For population search across ACMG standards

- eVaiutilities and vcffilter: analyzer for variant annotation.
- https://github.com/IBCHgenomic/eVaiutilities
- https://github.com/IBCHgenomic/vcfilter





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asynchronous vcf filter for human genomics.
      *************
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     ***********
Usage: vcfilter <COMMAND>
Commands:
 default-vcf-filter
                            vcf filter according to quality, coverage and zygosity
 quality-vcf-filter
                            vcf filter according to quality
 coverage-vcf-filter
                            vcf filter according to coverage
                            vcf filter according to zygosity
 zygosity-vcf-filter
 variant-default-vcf-filter vcf filter default quality40, coverage10 and zygosity multi
 variant-quality-vcf-filter vcf filter according to quality and variant
 variant-coverage-vcf-filter vcf filter according to coverage and variant
 variant-zygosity-vcf-filter vcf filter according to zygosity and variant
 help
                            Print this message or the help of the given subcommand(s)
Options:
  -h, --help
               Print help
  -V. --version Print version
```

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User interface of vcfilter
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annotating and analyzing eVai results.
      **************
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     Funded through Development of a universal fast-response platform,
     based on RNA technology, ensuring the national drug and epidemiological safety.
     2021/ABM/05/00004-00 to
     Prof. Luiza Handschuh: luizahan@ibch.poznan.pl.
     ************
Usage: eVaiutilities <COMMAND>
Commands:
  variant-analyzer
                                 analyzer for the variants
  variant-filter
                                filter the variants
  variant-database
                                create variant database
  gtf-analyze
                                analyze the corresponding gtf
  variant-seq
                                prepare the variant seg annotation
                                download the human genome
  download-genome
                                variant-transcriptids
  acmg-transcript
  sequence-profile
                                sequence profiling
  population-variant-search
                                search for the variant across population
  coordinate-search
                                search according to coordinates
                                search according to coordinates and variant
  coordinate-search-variant
  annotation-search
                                 search according to annotation
                                pathogenicity filter
  pathogenicity-filter
  population-variant-searcholder search for the variant across population older version
  coordinate-searcholder
                                 search according to coordinates older version
  coordinate-searc-variantholder search according to coordinates and variant older version
  annotation-searcholder
                                search according to annotation older version
  pathogenicity-filterolder
                                pathogenicity filter older version
  transcript-search
                                search by the specific transcript
  transcript-searcholder
                                search by the specific transcript older
  alt-allele
                                search for the ref allele
  alt-ref-allele
                                search according to ref allele and alt allele
  alt-allele-older
                                search for the ref allele older version
  alt-ref-allele-older
                                 search according to ref allele and alt allele older version
                                Print this message or the help of the given subcommand(s)
  help
Options:
  -h. —help
                Print help
 -V, -version Print version
```

User interface of eVai-analyzer