Knowledge Output:

- chr– Chrosome id
- pos Position
- ref-Reference allel
- alt- Reference allele as used to get consequences
- **gene** the gene symbol
- hgvs_protein_change the HGVS protein sequence name
- hgvs_nucleotide_change- the HGVS coding sequence name
- pathogenicity Clinical significance in Knowledge database
- **disease** related disease about this variant in Knowledge database
- pubmed_id related literature about this variant in Knowledge database
- dbsnp dbSNP id
- from variant source