

2013-12-01 Web release

ClinVar released a few modifications to the web site December 1, 2013. These descriptions do not include changes in content.

Modifications to the tabular display

Added a filter to support restriction to records for which clinical significance was reviewed by an expert panel. This is visible right now only for CFTR ([http://www.ncbi.nlm.nih.gov/clinvar/?term=cfr\[gene\]](http://www.ncbi.nlm.nih.gov/clinvar/?term=cfr[gene])). The order of the filters was also modified so that Clinical Significance and Review Status are closer to the top.

When multiple genes are included in a structural variation, only the first few are listed, with the link [..See more](#) provided to show all genes on the full display page.

Modifications to the full display

Description of the variation

If there are more than 5 HGVS expressions for a variation, only the first 5 are shown by default. The rest can be displayed by clicking on [...more](#).

Description of the phenotype

Up to three alternate terms for the same phenotype are reported on the ClinVar full display, with a link to MedGen if more are available.

Clinical Assertions tab

If there are citations in PubMed listed in the Clinical Assertions tab, a link is now provided to find all records in ClinVar citing the same publication. This link is provided to make it easier to determine whether there are multiple records based on the same publication.