20140306_data_release_notes

Summary of changes in this data release

Number of submissions

Date	Total Submissions
Apr 05, 2013	30333
May 01, 2013	30386
Jun 01, 2013	39047
Jul 01, 2013	39170
Aug 01, 2013	45901
Sep 01, 2013	50263
Oct 01, 2013	52047
Nov 01, 2013	64750
Dec 01, 2013	64881
Jan 01, 2014	68204
Feb 01, 2014	73492
Mar 01, 2014	83343

Content

Brief	Explanation
clinical variants from dbSNP	Some clinically relevant variants were submitted to dbSNP prior to the release of ClinVar. Some of those datasets have been submitted to ClinVar.
xml corrections	We have added some consistency checking so that the content of the HGVS and XRef elements are more consistent. We appreciate your feedback; we know we are not finished yet.

Expected modifications in the few months

Brief	Explanation
Representation on GRCh38	ClinVar plans to report information on both GRCh37 and GRCh38 for the indefinite future. We will start reporting SequenceLocation for GRCh38 in our next release.