# 20141002_data_release_notes

## Overview of changes in the release of October 2, 2014

### Overview of submissions:  2014

| Date | Total Submissions |
|---|---|
| Jan 01, 2014 | 68204 |
| Feb 01, 2014 | 73492 |
| Mar 01, 2014 | 83343 |
| Apr 01, 2014 | 111501 |
| May 01, 2014 | 112349 |
| Jun 01, 2014 | 117209 |
| Jul 01, 2014 | 127132 |
| Aug 01, 2014 | 127557 |
| Sep 1, 2014 | 143114 |
| Oct 1, 2014 | 143601 |

### Content

| Brief | Explanation |
|---|---|
| Clinical significance for variants submitted by OMIM | In August, ClinVar updated the clinical significance reported on behalf of OMIM for somatic variants, and for variants reported to cause named subtypes of familial hypercholesterolemia, but did not update the date the clinical significance was last evaluated. These dates were corrected the second week of September, 2014, to reflect the date of the change in August. |
| Submission from Blueprint Genetics | Blueprint Genetics added about 120 records based on their clinical testing results. |
| GTR test ids | GTR test identifiers from the Genetic Testing Registry are now searchable in ClinVar. Some submitters provide GTR test ids as part of the method used to detect the variant, e.g. http://www.ncbi.nlm.nih.gov/clinvar/?term=GTR000501759 |
| SCRP submission | A new BRCA1/BRCA2 submission from the Sharing Clinical Reports Project includes 85 assertion updates and 99 novel SCRP assertions. |
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# Overview of Submissions: 2013

| 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 |