



Submitting SNPs using a Reference Sequence Position

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I have to submit SNPs discovered by my group, but do not have any of the sequence data for the variations — just the position on a reference sequence.

There are three ways you can submit:

1. If you do not have the sequence data, but have a GenBank accession and a position as reference, you can send this information to dbSNP submissions, and we can use this information to get the sequence from GenBank.
2. You can now submit batches of SNPs yourself using dbSNP's [Human Variation: Annotate and Submit Batch Data](#) site, although you must be able to describe the variations using [HGVS nomenclature](#) in order to submit them. Click on the “submission help” link located at the upper right corner of the page for help getting started.
3. Finally, if you want to submit you variations one at a time, you can use the [Human Variation: Search, Annotate, Submit](#) site. Here is an example of how this resource might be used for a submission: if you query using NM_000212.2:c.176T>C as your search term, you will find that [rs5918](#) has already been assigned to that location. You can then add value that SNP by providing the URL to your database.

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