Online DNA variant interpretation

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https://mobidetails.iurc.montp.inserm.fr/MD/
Purpose: easy access to data required for DNA variant interpretation

- Direct links to relevant Pubmed IDs
- Full HGVS nomenclature + defgen export
- Population Frequencies + LOVD matches + UCSC links
- Missense + splicing predictions
- Interver direct link + ACMG classification
Simple architecture based on modern web technologies
Create your variant in a few clicks and interpret the data in a single web page...
Many other possibilities...

On any device