



## Drowning under NGS data?

Juggling with difficult and complex software to browse through huge amounts of data, searching for clinically relevant variants?

You want:

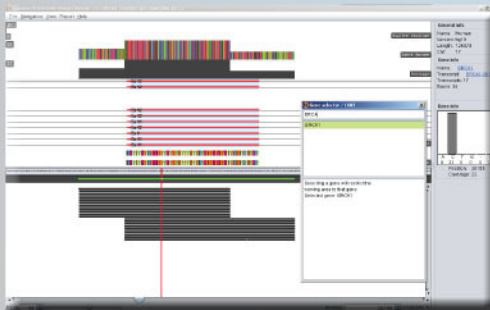
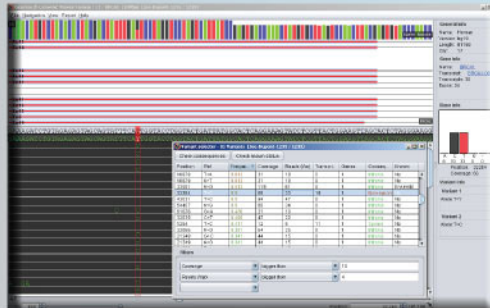
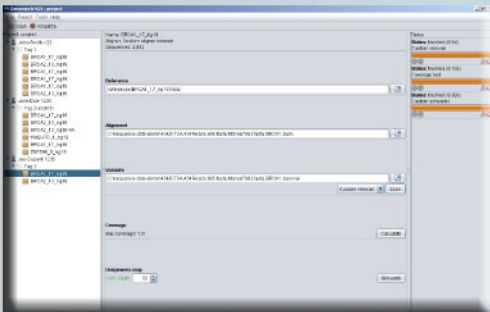
User friendly software capable of managing patient centric data?

Intelligent variant detection and filtering system?

Variant consequence prediction support?

Manage your own variant database, integrated in a DNA sequence analysis software?

To look up various public database such as LOVD, UMD, Ensembl?



PhenoSystems SA is partner in:



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