Genetic Report for Sample HG00512

Sample HG00512 likely has Disease Z due to the identified structural varia
chr1 at position 90082. The identified structural variant in the genome of th
individual is pathogenic based on previous research. More information ab
this genetic variant can be found in the ClinVar database using the ClinVar
reference ID 91011.

Genetic Report for Sample HG02055

Sample HG02055 likely has Disease X due to the identified structural varia
chr1 at position 10746. The identified structural variant in the genome of th
individual is pathogenic based on previous research. More information ab
this genetic variant can be found in the ClinVar database using the ClinVar
reference ID 1234.

Genetic Report for Sample HG03540

Sample HG03540 likely has Disease Y due to the identified structural varia
chr1 at position 10626. The identified structural variant in the genome of th
individual is likely pathogenic based on previous research. More informati
about this genetic variant can be found in the ClinVar database using the
ClinVar reference ID 5678.

Genetic Report for Sample PGP1

Sample PGP1 likely has Disease Z due to the identified structural variant c
chr1 at position 90082. The identified structural variant in the genome of th
individual is pathogenic based on previous research. More information ab
this genetic variant can be found in the ClinVar database using the ClinVar
reference ID 91011.