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The Genetic Database Website provides access to curated information regarding all de novo mutations observed and reported in scientific literature. Using the database, researchers can check whether a de novo mutation in a gene or a more specific locus has been reported previously. Those investigating rare diseases will be able to eliminate some findings quickly as non-causally related if that mutation is observed across other, more common diseases in the database.

TECHNOLOGY SUMMARY
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FEATURES AND BENEFITS
- Increases clinician and researcher access to genetic disease information.
- Facilitates rapid identification of diseases related to specific de novo mutations.

INVENTOR PROFILE
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