TAKE A QUANTUM LEAP

Accelerated answers with 90-min hybridization



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SureSelect QXT FOR CLINICAL RESEARCH

- Complete and accurate variant calling from only 50 ng of sample
- 3.5x faster workflow, same day sample to sequencing

Fastest Workflow

- Same day sample to sequencing from only 50 ng of gDNA
- Quick 90 minute hybridization and only 3.5 hr total hands-on time

Superior Performance

- High sensitivity, specificity and coverage of regions of interest
- Complete and accurate variant profiling with less sequencing

Complete and Flexible Solution

- Solutions for library prep, capture, QC, automation and analysis
- · Enables analysis of genomes, exomes or custom targets

LEARN MORE ABOUT SureSelect www.agilent.com/genomics/QXT

Easy customization, faster analysis

- Discover SureDesign: Choose a Disease Research Panel or create a custom panel design in minutes
- · SureCall Software: NGS data analysis software for non-bioinformaticians

Visit our NGS Resource Center for more information: www.agilent.com/genomics/NGSresource

DELIVERING FASTER ANSWERS IS IN OUR GENES

The superior coverage and efficient workflow provided by Agilent's NGS target enrichment portfolio gets you better answers, faster.

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