



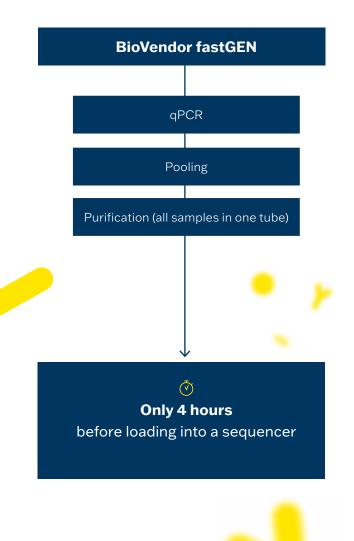
fastGEN

A breakthrough system for a one-step NGS library preparation



lt's faster. Way faster!

The fastGEN technology is based on a rapid and easy-handling workflow that significantly lowers the probability of mistakes.



4 reasons why fastGEN worths attention

BioVendor Group developed user-friendly diagnostic NGS kits with excellent analytical parameters and high-speed processing.

Unique

Simple, ultra-sensitive, specific and effective technology

Reliable

Significantly lower probability of mistakes, uniform coverage for different amplicons and different samples

Fast

Excellent speed (<30 min hands-on time)

Robust

Robustness achieved by using short amplicons obtained by a single polymerase chain reaction with special tagged hybrid primers



Cloud-based database bioinformatic and interpretation system

Key features

- User-friendliness
- Automatized bioinformatic analysis
- Advanced quality control of sequencing data
- Easy variants filtration
- Local clinical variants database
- Visualization of NGS data
- Internal interpretation of variants and sharing between clinics
- Clinical report generation
- Custom baseline creation

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Find out more about the software on our <u>website.</u>



Quick and easy detection of mutations in genes

NRAS | KRAS | BRAF



Contact our NGS team



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