



FastGene® BAC-free HS Taq is the optimal solution for microbial studies

The FastGene[®] BAC-free HS Tag DNA Polymerase is based on the single-subunit, wild-type Tag DNA polymerase of the thermophilic bacterium *Thermus aquaticus*. It is however not a bacterial recombinant protein but purified from an eukaryotic expression system.

NEW

Applications

- High throughput PCR of bacterial genomes
- Amplification of low copy DNA templates
- Multiplex PCR
- Specific amplification of complex templates
- RT-PCR





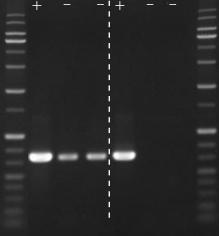


Fig. 1: Amplification of a non-ribosomal gene using E. coli DNA (+) or no template control. No template control (-) were amplified with standardly produced Taq vs FastGene® BAC-free HS Taq. The conventional Taq produced taq variasticette Die being a non-template control while there was no product in the FastGene® **BAC**free HS Taq. This indicates a bacterial genomic DNA contamination of the conventional Taq polymerase

Eukaryotic Expression System - No more false positive Performing PCR with bacterial templates could lead to a false positive, when using Taq enzymes purified from E. coli expression systems due to a contamination of the Tag enzyme with prokaryotic genomes. The FastGene® BAC-free HotStart Tag DNA Polymerase is produced using eukaryotic cells. Hence, no bacterial genome is present.

Ordering Information

Cat. No.	Product	Content
LS33	FastGene® BAC-free HS TAQ Polymerase	500 Units
LSO5	FastGene® DNA Releasy	300µl, 10 Rxn
LSO6	FastGene® DNA Releasy	1.5 ml, 50 Rxn



