

Product Datasheet

HiDi® DNA polymerase 9001

Product Name HiDi® DNA polymerase

Catalog Number #9001

Description HiDi® stands for **H**igh **D**iscrimination of mismatches at the 3'-terminus of primers in PCR. This enzyme family is optimized for this feature and is the first choice for applications that rely on this property such as allele-specific PCR (asPCR), also known as allele-specific amplification (ASA).

Comparison studies with competitor products show that the HiDi® DNA polymerase family is the first choice for highly selective PCRs, such as genotyping by allele-specific PCR, HLA genotyping, analysis of single CpG methylation sites or the detection of mutations in a high background of wild-type sequences.

By using HiDi® DNA polymerase, less than 10 copies of a mutation can be detected in a background of >10,000 wild-type copies without any other tedious assay optimization.

Applications:

- Allele-specific PCR (asPCR), allele-specific amplification (ASA)
- HLA genotyping
- Analysis of single CpG methylation sites by PCR (methylation specific PCR, MSP)
- Mutation detection by PCR even in a high background of wild-type sequences
- Genotyping e.g., in CRISPR/Cas and TALEN approaches

This polymerase is also available as a **full-length Taq DNA polymerase** with a nuclease domain, featuring 100% compatibility with hydrolysis probes (TaqMan® probes etc.).

Several independently conducted studies show that HiDi® DNA polymerase is ideally suited for use in asPCR in numerous research areas ranging from mutation detection to genome editing. Please see "References" below.

For research use and further manufacturing. Designed and manufactured under ISO13485

Tested Applications End-Point, Real-Time

Brand myPOLS Biotec

Storage -20°C

References

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Primordial Germ Cell Migration and Histological and Molecular Characterization of Gonadal Differentiation in Pachón Cavefish *Astyanax mexicanus*

Imarazene B, Beille S, Jouanno E, Branthone A, Thermes V, Thomas M, Herpin A, Rétaux S, Guiguen Y Sex Dev. 2021 Mar 10;1-18.

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<https://doi.org/10.1016/j.stemcr.2019.11.008>

A New Protocol for the Detection of Sterigmatocystin-producing Aspergillus Section Versicolores Using a High Discrimination Polymerase.

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Kikuchi Y, Hara-Kudo Y, Terajima J, Sugita-Konishi Y.
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Algal Research, 2019 May, 101469

Link to publication:

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Genome Res. 2018 Feb;28(2):223-230

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diatom *Phaeodactylum tricornutum* by DNA-free genome editing.

Serif M, Dubois G, Finoux AL, Teste MA, Jallet D, Daboussi F.
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Content	S pack: 250 U, 5 U/ μ l, 1 x 50 μ l HiDi® DNA polymerase; 1 x 1.25 ml 10x HiDi reaction buffer
	M pack: 250 U, 5 U/ μ l, 1 x 200 μ l HiDi® DNA polymerase; 2 x 1.25 ml 10x HiDi reaction buffer