



VERSION 2

APR 09, 2023

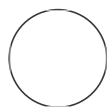
OpenVent Eco DNA Polymerase Production V.2

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Beneficial Bio

Low-cost, high-quality ...



Jenny Molloy

OPEN ACCESS

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<https://protocols.io/view/openvent-eco-dna-polymerase-production-ciudues6> Version created by [Jenny Molloy](#)

MANUSCRIPT CITATION:

Bhadra, Sanchita, et al. "Preparation and Use of Cellular Reagents: A Low-resource Molecular Biology Reagent Platform." *Current Protocols* 2.3 (2022): e387.

Bhadra, Sanchita, et al. "Producing molecular biology reagents without purification." *PLoS One* 16.6 (2021): e0252507.

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Protocol status: Working
We use this collection and it's working

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ABSTRACT

COLLECTION integer ID:
72293

This collection contains the protocols for production, quality control, packaging and use of Beneficial Bio's dehydrated OpenVent DNA Polymerase.

Keywords: PCR, Reagent
Manufacturing

Format: 32 tubes (4x8-tube PCR strips) 20ul of Enzyme each

OpenVent DNA polymerase is an extremely high thermostable and high fidelity enzyme suitable for routine PCR applications and amplification of GC-rich or looped sequences. The enzyme is compatible with a wide range of templates and its robustness guarantees reliable amplification results in almost all PCR applications. The Eco and room temperature stable format - PCR reaction tube strips containing the dehydrated enzyme (10 reactions per tube), offers affordability and flexibility.

FEATURES

- Dehydrated formulation enables better stability at room temperature
- Ideal for a wide range of applications including GC-rich or looped sequences amplification
- Amplification of low copy DNA targets
- Eco formulation and flexible format

DOCUMENTS

[Product Manual](#)

[Product Specification Sheet \(PSS\)](#)

[Certificate of Analysis \(CoA\)](#)

PROTOCOL GUIDANCE

Select one of the following options for enzyme production based on your lab infrastructure:

- IPTG-based induction requires a shaking incubator, large benchtop centrifuge and IPTG
- Plate-based autoinduction requires none of the above

After producing the enzymes, perform the three QC protocols in any order:

- SDS-PAGE to determine expression has been successful
- Functionality testing in PCR
- Nuclease contamination testing

IMAGE ATTRIBUTION

Beneficial Bio Ltd

FILES

Protocol



NAME

Plate Protein Expression on Autoinduction media

VERSION 3

CREATED BY

Stephane Fadanka

[OPEN](#) →

Protocol



NAME

Production of cellular reagents using IPTG

VERSION 1

CREATED BY

Nadine Mowoh

[OPEN](#) →

Protocol



NAME

Functionality test (OpenVent polymerase, PCR Master Mixes)

VERSION 2

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Protocol



NAME

Nuclease Test (OpenVent polymerase, PCR Master Mix, DNA loading dye)

VERSION 1

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Nadine Mowoh

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Protocol



NAME

Assessing protein purity using SDS PAGE

VERSION 1

CREATED BY

Nadine Mowoh

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