

DATA SHEET

Version: 3 Revision date: 09/04/2024 Canvax Reagents, S.L.U. Luis de Mercado Street, 19 Boecillo Technological Park 47151, Boecillo Valladolid, Spain.

Tlf: +34 983 54 85 63 info@canvaxbiotech.com

www.canvaxbiotech.com

1. Identification

Product name Horse-Power™ Red DNA Polymerase

Master Mix 2.5X

100 rxn

Cat. No P0027-S

2. Description

Horse-Power[™] Red DNA Polymerase is a ready-to-use 2,5x master mix that contains all PCR reaction components: TruePure[™] dNTPs, PCR buffer, Mg2+ and Horse-Power Taq DNA polymerase. Only primers and template need to be added.

The mix also contains an agarose loading buffer including a red dye for visual tracking of DNA migration and a dense compound to facilitate the drop-down of the samples into the well agarose gels.

3. Composition

Item	Quantity
Horse-Power™ Red DNA Polymerase Master Mix 2.5X	0.8 mL

Concentration: (2,5X (Buffer Red 2,5X; dNTPs 0,5 mM each; HorsePower Taq DNA polymerase 0,250 U/µL, Glycerol 30%).

4. Features

- Ready-to-use.
- Adds extra nucleotides (preferentially adenine) without template at 3 ends leaving 3 overhangs PCR fragments. This fact allows the popular TA-cloning or GC cloning.
- > Both save times in the PCR process and in agarose loading samples.

Quality:

- Functionally tested in PCR.
- Undetected bacterial DNA (by PCR).
- Undetectable nucleases activity (endo-, exo- and ribo-).

5. Storage specifications

Store at -20° C

6. Applications

- > Design for medium or high throughput applications (ex. colony screening).
- ➤ PCR fragments amplification for TA or GC cloning. (preferably use a proofreading polymerase for cloning purpose and a blunt cloning vector) (see pSpark® DNA Cloning System Cat. No: C0001)

7. Assay conditions

Enzyme activity is assayed in the following mixture: 25mM Tris-HCl pH 9.0 at 25°C, 50mM KCl, 2mM MgCl2, 0.1mg/mL gelatine, 200 μ M de dATP, dGTP, dTTP, 100 μ M[α 32-P]dCTP (0.05 μ Ci/nmol) and 12.5 μ g activated salmon sperm DNA.





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8. Recomended PCR Assay (20µL assay)

Red DNA Polymerase Master mix 2.5X	8µl (1X)
Forward Primer (15µM)	1μl (0.75 pmol/μL)
Reverse Primer (15µM)	1μl (0.75 pmol/μL)
Template DNA	plasmide: 30-75ng; gDNA: 100-500ng
PCR grade H20	up to 20 μL

Cycling instructions: 94°C 5:00, 25-30x (94°C 0:35, Tm 0:35, 72°C 1'/kb), 72°C 7:00, 4°C ∞)

Red dye Agarose Mobility			
Agarose Gel Concentration (%)	Effective separation of: (bp)	Migrati on Rate (bp)	
0,7	800-12000	3000	
1,0	400-8000	1500	
1,5	200-3000	900	
2,0	100-2000	300	
3,0	25-1000	> 100	

^{*} in TAE Buffer

9. Further information

Product Use Limitation This product is developed, designed, and sold exclusively only for research purposes use. The product was not tested for use in diagnostics or for drug development, nor is it suitable for administration to humans or animals.

Disclaimer

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