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& Empirical bioscience	PIS-048 AT-HF-RB-Units	Version: 004 Effective Date: 07/06/20 Author: Beth Lowe CO#: 062620-1

Print only pages 2 for Customers



Product Information Sheet

Product Name: AlloyTaq™ Polymerase, High Fidelity with 10X Reaction Buffer*

Concentration: AlloyTaq Polymerase, High Fidelity: 5U/µI, AlloyTaq Reaction Buffer: 10X

Storage and Handling: Upon arrival store at -20°C for provided expiration date

Ordering Information:

Item Number	Units	Number of Tubes and Volumes
AT-HF-RB-500	500	AlloyTaq Polymerase, High Fidelity 500 Units: 1x100μL, 10X AlloyTaq Reaction Buffer: 2x1mL
AT-HF-RB-1000	1000	AlloyTaq Polymerase, High Fidelity 500 Units: 2x100μL, 10X AlloyTaq Reaction Buffer: 4x1mL

Product Description:

PIS-048 Version 004

AlloyTaq™ Polymerase is an optimized blend of Taq and Integrity High Fidelity Polymerases™ from species pyrococcus GBD. With the 3'-5' proofreading ability of Integrity and the robust amplification of Tag, AlloyTaq™ is not only great for routine PCR, but also high fidelity and longer and more difficult amplicons. This composition increases fidelity of Taq by greater than two-fold. AlloyTaq™ Polymerase will produce blunt ended overhang products at the 3' end.

10X AlloyTaq™ Reaction Buffer:

Specially optimized reaction buffer specifically manufactured for AlloyTaq™ Polymerase, High Fidelity. This Buffer is supplied in a 10X concentration and should be diluted for use.

Protocol: Minimize Freeze thaw of master mix to avoid loss of performance. The following reaction set up and general cycling conditions are recommended but can vary depending on the template and primers being used. Reaction set-up for a 50uL Reaction:

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Component	Volume	Final Concentration			
10X AlloyTaq™ Reaction Buffer	5 µl	1X			
Upstream Primer, 10 μM	0.5-5.0 µl	0.1-1.0μM			
Downstream Primer, 10 μM	0.5-5.0 µl	0.1-1.0μM			
dNTP, 10mM	1-2.5 µl	200-500μM			
DNA Template	XμI	> 1ng			
AlloyTaq™ Polymerase	0.25-1uL	1.25U-5U			
Nuclease Free Water	to 50 µl	N.A.			

Thermal cycling conditions: The following general cycling conditions are recommended but can vary depending on the template and primers being used.

Cycling Step	Temperature	Holding Time	Cycles
Initial Denaturation	95°C	2min	1
Denaturation	98°C	10sec	00.00
Annealing#	68°C	60sec	20-30
Extension	70°C	(1min/kb)	
Final Extension	72°C	10min	1

^{*}Annealing will depend on primer length and composition. Generally, begin 5°C below primer T_m.

*This product is intended for Research Use Only. This product is manufactured under ISO13485:2016 Quality System Requirements and is available for use as a Raw Material for use in IVD applications. Please contact Empirical Bioscience for further details.

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