

Thermo Scientific ABsolute Blue QPCR SYBR Green Mix Plus ROX Vial

Description

ABsoluteTM Blue QPCR SYBR[®] Green Mix Plus ROX Vial has been developed to quantify DNA and cDNA*. With the exception of primers and template, this 2X mix contains all the components required to perform a rapid, sensitive and reproducible QPCR reaction:

- <u>Thermo-Start</u>TM DNA Polymerase, a chemically modified hot-start version of Thermoprime Plus DNA Polymerase, which prevents non-specific amplification during the reaction set-up. This enzyme requires an activation step at 95°C for 15 minutes.
- <u>Proprietary reaction buffer</u> which provides highly sensitive, specific and consistent fluorescence readings for real-time and end-point analysis. This buffer has been optimized for MgCl₂ and enhancers to improve amplification across a wide range of templates including plant DNA and GC rich fragments. It contains an inert blue dye to assist in the visualization of the ABsolute Blue QPCR SYBR Green Mix after aliquoting into the reaction well.
- dNTP's, including <u>dTTP</u> to improve reaction sensitivity and efficiency compared to dUTP.
- <u>SYBR[®] Green I</u>, a dye which fluoresces after binding of the double-stranded DNA. The overall fluorescence increases proportionally to the double-stranded DNA concentration.

ROX, passive reference dye for normalization of data (separate vial).

Kit Contents		
Vial	Pack Size (cap color)	
	А	В
ABsolute Blue QPCR SYBR Green (2X)	2 x 1.25ml (green)	16 x 1.25ml (green)
$MgCl_2$ (1 M)	100µ1 (clear)	100µ1 (clear)
ROX Reference Dye (1 mM)	25µl (brown)	25µl (brown)

Kit Contents

* For RNA template, use VersoTM SYBR[®] Green 1-Step QRT-PCR Kit Plus ROX Vial (AB-4104)

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INFORMATION

Thermo-StartTM DNA Polymerase

The enzyme requires an activation step at 95°C for 15 minutes.

Thermo-StartTM has 5' to 3' polymerization and exonuclease activity but lacks 3' to 5' exonuclease activity (proofreading).

Blue Dye

This proprietary inert blue dye allows quick and easy visualization of the amount of the mix in the well, minimizing aliquoting errors. It does not interfere with the QPCR reaction and is only available in master mix format.

ROX Dye

ROX is an internal passive reference dye used to normalize the fluorescent reporter signal generated in QPCR. A separate vial of ROX is included in this kit for optional addition to the ABsolute Blue QPCR SYBR Green Mix. The final concentration will vary depending on the real time cycler. For example, for a concentration of 100 nM ROX in a <u>final</u> 1X QPCR reaction mix, dilute ROX (1 mM) 40 times i.e. 5 μ l ROX Reference Dye + 195 μ l PCR grade water and add 10 μ l of the diluted ROX solution to each 1.25 ml vial of ABsolute Blue QPCR SYBR Green Mix or 40 μ l to each 5 ml vial of ABsolute Blue QPCR SYBR Green Mix.

MgCl₂

The initial concentration of $MgCl_2$ in the ABsolute Blue QPCR SYBR Green Mix corresponds to 3 mM in the <u>final</u> 1X reaction. This concentration is effective over a broad range of templates. Some assays may be improved further with $MgCl_2$ optimization. A separate vial of 1 M $MgCl_2$ is therefore supplied with each kit.

MgCl₂ concentration can be increased as follows: each 2.5 μ l or 10 μ l addition of MgCl₂ to the 1.25 ml or 5 ml undiluted ABsolute Blue QPCR SYBR Green Mix respectively corresponds to an increase of 1 mM in the <u>final</u> 1X reaction. Scale up or down accordingly. Mix thoroughly by inverting the vial ten to twenty times. **Do not vortex.**





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Cycler Compatibility

ABsoluteTM Blue QPCR SYBR[®] Green Mix Plus ROX Vial is compatible with all blockbased QPCR instruments and the Rotor-GeneTM.

Storage Conditions

Store at -20°C until ready for use. ABsolute[™] Blue QPCR SYBR[®] Green Mix Plus ROX Vial is stable for a minimum of 12 months. The reagents can be stored at 4°C for up to 1 month. Avoid repeated freeze thawing. The ROX and SYBR[®] Green dyes are light sensitive; exposure should be minimized. Shipped on ice within the UK and on dry ice for international and within the US.

Additional Info

- The use of disposable gloves, DNase and RNase free filter tips and plastics is recommended.
- For optimal results, the recommended amplicon length is in the range of 60 to 300 bp.
- As best performance is achieved with dTTP, the ABsolute Blue QPCR SYBR Green Mix contains a nucleotide mix with dTTP instead of dUTP.

Tips before use

Thaw the reagents on ice, mix the solutions and spin down before use to recover the maximum amount. Do not vortex the ABsolute Blue QPCR SYBR Green Vial. Briefly centrifuge to avoid bubbles within the wells, as these will interfere with the fluorescence. Always include a no template control (NTC).



PROTOCOL

Example of Reaction Mix preparation for a 25 µl final reaction:

		Volume	Final Concentration
	ABsolute Blue QPCR SYBR Green	12.5 µl	1X
Reaction	(2X)		
Mix	Forward primer (1 μ M) ^a	1.75 µl	70 nM
IVIIX	Reverse primer $(1 \ \mu M)^{a}$	1.75 μl	70 nM
	Water (PCR grade) ^b	variable	
	Template (DNA or cDNA) ^c	1 - 5 µl	<250 ng/reaction
	Total volume	25 µl	

Example of a **QPCR thermal cycling program**:

	Temp.	Time	Number of cycle
Enzyme activation	95°C	15 min	1 cycle
Denaturation	95°C	15 sec	
Annealing ^d	50-60°C	30 sec	40 cycles
Extension ^e	72°C	30 sec	

It is recommended to perform a melt curve to confirm the specificity of the reaction. Example of a **melt curve program** ^f:

Denaturation	95°C	30 sec	1 cycle
Starting temp.	60°C	30 sec	1 cycle
Melting step ^g	60°C	10 sec	80 cycles

Notes

- a For optimization, a primer titration should be performed from 50 nM to 300 nM final concentration. Scale up or down the volume and concentration as appropriate.
- b The volume of the total reaction should be completed up to 25 μ l with water.
- c The volume of template to add to the QPCR reaction can be adjusted as required. For standard templates only 1 µl should be added to reduce the carryover of any PCR inhibitor. This volume can be increased up to 5 µl for low copy number templates.
- d Annealing temperature dependent on primer sequence.
- a Amerianing temperature dependent on primer sequence.
 b Time of extension depends on the length of the amplicon. If the amplicon exceeds 300 bp amplification time should be adapted (Thermo-Start[™] DNA Polymerase extends approximately at 1000 bp/min).
 f Melt curve program may vary depending on instrument manufacturer and software.
- g Increase set point temperature by 0.5°C per cycle.



Quality control

ABsolute Blue QPCR SYBR Green Mix is tested functionally using QPCR. The product must demonstrate linearity of amplification over a specified serial dilution of human genomic DNA.

Ordering Information

AB-4166/A	ABsolute [™] Blue QPCR SYBR [®] Green Mix Plus ROX Vial	200 x 25 µl rxns
AB-4166/B	ABsolute [™] Blue QPCR SYBR [®] Green Mix Plus ROX Vial	1,600 x 25 µl rxns
AB-4167/A	ABsolute [™] Blue QPCR SYBR [®] Green Mix Plus ROX Vial	400 x 25 µl rxns
AB-4167/B	ABsolute [™] Blue QPCR SYBR [®] Green Mix Plus ROX Vial	4,000 x 25 µl rxns

All formats are supplied with an additional vial of 1 M MgCl₂.

Related Products

Cat. No.	Description	Quantity
AB-0600/W	Thermo-Fast TM 96 Non-Skirted, white *	25 plates
AB-0800/W	Thermo-Fast [™] 96 Skirted PCR Plate, white *	25 plates
AB-0900/W	Thermo-Fast [™] 96 Semi-Skirted PCR Plate, white *	25 plates
AB-1170	ABsolute [™] QPCR Seal (adhesive seal)	50 sheets
AB-0812	Clear Seal Diamond (heat seal)	100 sheets
AB-0866	Ultra Clear Cap Strips (8 caps)	120 strips

* For Cycler compatibility and other color choices, see our latest catalogue or visit www.abgene.com

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Troubleshooting:	Email	Phone
North America (US, Canada, Central/South America)	Techservice.genomics @thermofisher.com	+1 (800) 235-9880
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