



# GeneAce SYBR<sup>®</sup> qPCR Mix $\alpha$

## I. Description

GeneAce SYBR<sup>®</sup> qPCR Mix  $\alpha$  is a convenient 2X premix for real-time qPCR using SYBR<sup>®</sup> Green I dye. This product was optimized for instruments that require a high ROX reference signal for fluorescence normalization (Compatible instruments: ABI PRISM<sup>®</sup> 7000/7700/7900HT, ABI 7300, and ABI GeneAmp<sup>®</sup> 5700, etc.).

This product provides high specificity and reproducibility and allows for reducing non-specific amplification by using optimized buffer components and Hot-Start Gene Taq NT, a chemically modified Taq DNA Polymerase.

## II. Storage

-20°C, protected from light.

This product should be protected from light. After thawing, the master mix may be stored at 4°C for up to one month or returned to -20°C for long term storage.

## III. Components

	60 rxns	300 rxns
2 x GeneAce SYBR qPCR Mix $\alpha$ <sup>(*)</sup>	1.5 ml x 1	1.5 ml x 5

<sup>(\*)</sup> It contains Hot-Start Gene Taq NT, dNTP Mixture, Mg<sup>2+</sup>, SYBR<sup>®</sup> Green I, ROX reference dye, stabilizer.

## IV. Notes

- **The PCR must start with an initial 10 minutes incubation at 95°C to activate the chemically modified hot-start Taq DNA polymerase.**
- **We recommend two-step cycling protocols. (See V. Typical PCR protocol)**
- **This product cannot be used for runs in "Fast"<sup>(\*)2</sup> mode on ABI real time instruments.**

<sup>(\*)2</sup> 95°C 20 sec → (95°C 3 sec, 60°C 30 sec) x cycle numbers

- Always ensure that the product has been fully thawed and mixed before use. Gently mix the mixtures without creating bubbles.
- Uracil-N-Glycosylase treatment is unusable on this product.

## V. Typical PCR Protocol

Choose an appropriate total reaction volume, depending on the instrument used. (e.g., 50  $\mu$ l or 25  $\mu$ l)

Component	Volume	Volume
2 x GeneAce SYBR qPCR Mix $\alpha$	25.0 $\mu$ l	12.5 $\mu$ l
25 $\mu$ M each Primers	1.0 $\mu$ l	0.5 $\mu$ l
Template	5.0 $\mu$ l	2.5 $\mu$ l
d.d.H <sub>2</sub> O	up to	50.0 $\mu$ l
		25.0 $\mu$ l

Recommended cycler conditions

**95°C 10 min. Enzyme activation <sup>(\*)3</sup>**

**95°C 30 sec.**  
**60°C 1 min. ) 45 cycles**



Melting curve analysis

Optimum PCR condition depends on primer sequence and template DNA and so on.

<sup>(\*)3</sup> Ensure that the cycling program includes the DNA polymerase activation step. Initial 10 minutes incubation at 95°C is required.

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# GeneAce SYBR<sup>®</sup> qPCR Mix $\alpha$

## I. 製品説明

GeneAce SYBR<sup>®</sup> qPCR Mix  $\alpha$ は、ROXを用いてウェル間の蛍光シグナルの補正を行うリアルタイム PCR 装置 (ABI PRISM<sup>®</sup> 7000 / 7700 / 7900HT、ABI 7300、ABI GeneAmp<sup>®</sup> 5700 等)に対応したインターカレーター法用のリアルタイム PCR 用試薬です。2x 濃度に予め調製されたマスターミックスが小分け分注済みです。

本製品は、ホットスタートPCR用酵素 Hot-Start Gene Taq NT と最適化されたバッファーにより、非特異的増幅を抑制し、高い特異性と再現性を実現しています。

## II. 保存

-20°C (遮光)

4°C保存(遮光)も可能ですが、その場合は1ヶ月以内にご使用下さい。

## III. 製品内容

試薬	60 反応用	300 反応用
2 x GeneAce SYBR qPCR Mix $\alpha$ *1)	1.5 ml x 1 本	1.5 ml x 5 本

\*1) Hot-Start Gene Taq NT, dNTP Mixture, Mg<sup>2+</sup>, SYBR<sup>®</sup> Green I, ROX Reference Dye, stabilizer を含んでいます。

## IV. 注意

- 化学修飾を施したホットスタート PCR 用酵素を用いているため、酵素活性化ステップ(95°C 10min.)を必ず実施して下さい。
- 推奨 PCR サイクル条件にてご使用下さい。(V.使用例 参照)
- ABI リアルタイム PCR 装置のランモード「Fast」\*2)には対応しておりません。
  - \*2) 95°C 20 sec - (95°C 3 sec, 60°C 30 sec) x サイクル数
- 使用時は、泡立てないように穏やかに転倒混和し、試薬を十分均一にしてからご使用下さい。
- Uracil-N-Glycosylaseによるキャリアオーバー処理は出来ません。

## V. 使用例

ご使用の装置に対応した反応量量でご使用下さい。

<反応液(例)>	[ 50 $\mu$ l 系 ]	[ 25 $\mu$ l 系 ]
2 x GeneAce SYBR qPCR Mix $\alpha$	25.0 $\mu$ l	12.5 $\mu$ l
25 $\mu$ M each Primers	1.0 $\mu$ l	0.5 $\mu$ l
Template	5.0 $\mu$ l	2.5 $\mu$ l
d.d.H <sub>2</sub> O	up to 50.0 $\mu$ l	25.0 $\mu$ l

<推奨 PCR サイクル条件>\*3)

95°C 10 min.  
95°C 30 sec.  
60°C 1 min. ) 45 cycles



融解曲線解析

プライマーの設計や鋳型 DNA 等により反応の至適条件が変わることがあります。

\*3) 酵素の活性化ステップは、95°C10 分間必ず行って下さい。

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