

TNF β A329G ToolSet™ for LightCycler™

Lyophilized ToolSet for PCR using the LightCycler™ Instrument. Licensed by Roche Diagnostics GmbH

Order#: TNF β 329 - 16

1 ToolSet for 16 reactions

Store at 4°C, protected from light.
Exposure to light may especially damage
the OligoTool™ tube (vial with red cap).

For use with LightCycler-DNA Master Hybridization Probes, 10 x conc. (Roche Cat.No.: 2 015 102)

1. ToolSet contents

| Vial | Label | Content | Quantity |
|---------------------|------------------|--|--|
| | | | TNF β 329 - 16 |
| 1, Red cap | OligoTool | - lyophilized oligos for PCR - contains mutation detection and anchor probe, primers | For 16 tests Dissolved: 50 μ L |
| 2, Green cap | Control | - lyophilized heterozygous DNA | Dissolved: 20 μ L |
| 3, Blue cap | Solvent | - to dissolve OligoTool / Control | 1000 μ L of Solvent |

Additional equipment and reagents required but not supplied :
LightCycler-DNA Master Hybridization Probes, 10 x conc.Cat.No.: 2 015 102, including 25mM MgCl₂; LightCycler
instrument, LightCycler capillaries, DNA extraction materials

2. Introduction

2.1. Product overview

ToolSet description The ToolSet is specifically adapted for genotyping TNF β nucleotide position A329G by LightCycler PCR with Melting Curve Analysis. Fluorescent detection and anchor probes and the primer pair have been optimized for specific amplification of targets and optimal genotype discrimination.

Control material Heterozygous control DNA, lyophilized.

Storage of ToolSet and Solutions Store at +4°C when lyophilized, protected from light.
The unopened lyophilized ToolSet is stable at +4°C for 12 months from date of manufacture if protected from light. When dissolved store at +4°C for a maximum of 4 weeks, or at -20°C for longer periods (months), protected from light. Avoid freezing and thawing.

3. Preparation for LightCycler PCR

Toolset preparation Dissolve the content of the OligoTool tube (Red Cap) with 50 µl of Solvent.
Dissolve the content of the Control tube (Green Cap) with 20 µl of Solvent.

1. Before opening tubes, centrifuge them quickly.
2. Add Solvent into OligoTool tube and Control tube as above.
3. Recap tubes, vortex gently.
4. Before opening tubes, centrifuge them quickly.
5. Proceed to Reaction Mix preparation.

Primers ? You don't have to add primers.

Probes ? You don't have to add probes.

Reaction Mix Preparation For 1 (One) reaction, prepare the Reaction Mix as shown in the following table :

| Reagent | µL |
|-----------------------------------|------|
| OligoTool TNF β A329G dissolved | 2.8 |
| Solvent TNF β A329G | 10.4 |
| MgCl ₂ 25 mM | 0.8 |
| Master Hybridization Probes 10x | 2 |
| Total Reaction Mix | 16 |
| + Your DNA or Control TNF β A329G | 4 |
| Grand Total | 20 |

Use Master Hybridization Probes 10x and MgCl₂ 25 mM from Roche LightCycler-DNA Master Hybridization Probes, 10 x conc.

(Roche Cat.No.: 2 015 102, including 25mM MgCl₂).

For multiple reactions, multiply the indicated volumes appropriately.

Positive Control Always run a positive control with the samples.
Use the dissolved heterozygous Control TNF β A329G DNA (Green Cap).

Negative control Always run a negative control with the samples. To prepare a negative control, replace the template DNA with Solvent (Blue Cap).

Extraction of genomic DNA You can use different Kits for DNA isolation, either with a manual method or with an automated system. The elution buffers should be salt-free. Example : Roche High Pure PCR Template Preparation Kit (Cat.No. 1 796 828)

Application The TNF β A329G ToolSet™ for LightCycler™ allows the detection of the single point mutation at position 329 which has been associated with altered transcription of the TNF α gene, altered generation of the gene product – i.e. the cytokine TNF α - and with a variety of disease states.

More information is available in the *Genes* section at www.Genes-4U.com.

Note : This ToolSet was developed for use in life science research only.

4. LightCycler Settings and Experimental Protocol

Denaturation

| Cycle Program Data | Value |
|-----------------------------------|------------------|
| Cycles | 1 |
| Analysis Mode | None |
| Temperature Targets | Segment 1 |
| Target Temperature (°C) | 95 |
| Incubation time (s) | 120 |
| Temperature Transition Rate (°/s) | 20.0 |
| Secondary Target Temperature (°C) | 0 |
| Step Size (°C) | 0 |
| Step Delay (Cycles) | 0 |
| Acquisition Mode | None |

Amplification

| Cycle Program Data | Value | | |
|-----------------------------------|------------------|------------------|------------------|
| Cycles | 50 | | |
| Analysis Mode | None | | |
| Temperature Targets | Segment 1 | Segment 2 | Segment 3 |
| Target Temperature (°C) | 95 | 60 | 72 |
| Incubation time (s) | 1 | 10 | 13 |
| Temperature Transition Rate (°/s) | 20.0 | 20.0 | 5.0 |
| Secondary Target Temperature (°C) | 0 | 0 | 0 |
| Step Size (°C) | 0 | 0 | 0 |
| Step Delay (Cycles) | 0 | 0 | 0 |
| Acquisition Mode | None | Single | None |

Melting Curve Analysis

| Cycle Program Data | Value | | |
|-----------------------------------|------------------|------------------|------------------|
| Cycles | 1 | | |
| Analysis Mode | Melting Curves | | |
| Temperature Targets | Segment 1 | Segment 2 | Segment 3 |
| Target Temperature (°C) | 95 | 45 | 80 |
| Incubation time (s) | 30 | 120 | 0 |
| Temperature Transition Rate (°/s) | 20.0 | 20.0 | 0.1 |
| Secondary Target Temperature (°C) | 0 | 0 | 0 |
| Step Size (°C) | 0 | 0 | 0 |
| Step Delay (Cycles) | 0 | 0 | 0 |
| Acquisition Mode | None | None | Continuous |

Cooling

| Cycle Program Data | Value |
|-----------------------------------|------------------|
| Cycles | 1 |
| Analysis Mode | None |
| Temperature Targets | Segment 1 |
| Target Temperature (°C) | 40 |
| Incubation time (s) | 30 |
| Temperature Transition Rate (°/s) | 20.0 |
| Secondary Target Temperature (°C) | 0 |
| Step Size (°C) | 0 |
| Step Delay (Cycles) | 0 |
| Acquisition Mode | None |

Fluorescence display mode

Use F2/F1 or preferably F2 with colour compensation. For LC Program Versions 3.3 or lower : gains F1=1; F2=15. For LC Program Versions 3.5 and higher : use automatic gain control.

5. Typical results

Introduction

Use the Melting Curve program to genotype the human genomic DNA research samples. The melting peaks allow discrimination between the homozygous (wild type or mutant) and the heterozygous samples. Figure 1 shows a typical result obtained with the TNF β A329G ToolSet™ for LightCycler™ :

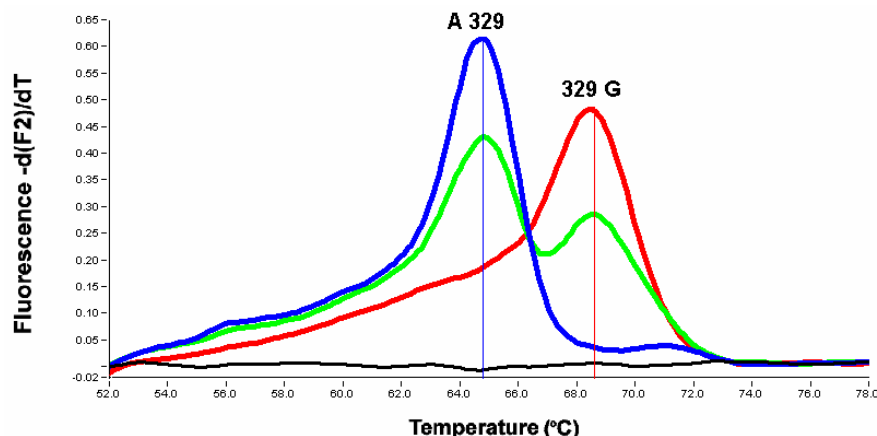


Figure 1 : Melting curve analysis of the three possible genotypes of the TNF β sequence at nt 329 .

BLUE : Homozygote for A329 (**wild type**), **RED** : Homozygous for 329G,
GREEN : The A329G Heterozygote Control contained in the ToolSet, Control TNF β A329G -16.
Conditions : LC Program 3.3, Color compensation and Digital Filter enabled,
Calculation Method : Polynomial, Degrees to Average : 6.
Red Cursor : T_m = 68.6 °C, Blue Cursor : T_m = 64.7 °C

Note : The values for the respective melting temperatures may vary for +/- 2.5 °C between different experiments. The Delta T between the melting peaks for different genotypes may vary +/- 0.5 °C. The TNF β A329G ToolSet™ has been developed for and validated with the LightCycler™ and its original accessory materials and reagents. Performance of the ToolSet with other instruments, accessories and reagents has not been validated by Genes-4U.

7. Notices to Purchaser

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