For IN VITRO USE ONLY. Version 1, October 2024

UGT1A6 T181A (nt A541G) ToolSet[™] for LightCycler[™] Application on LightCycler 480

Lyophilized ToolSet for PCR using the LightCycler™ Instrument.

Order#: UGT1A6 181 - 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 Fast Start DNA Master HybProbes, 10 x conc. (Roche Cat.No.: 03003248001) and LightCycler® 480 Multiwell Plates 96, white (Roche Cat.No.: 04729692001)

1. ToolSet contents

Vial	Label	Content	Quantity
			UGT1A6 181 - 16
1, Red cap	OligoTool	lyophilized oligos for PCRcontains mutation detection	For 16 tests
		and anchor probe, primers	Dissolved:
			50 μL
2, Green cap	Control	- lyophilized DNA	Dissolved:
			20 μL
3, Blue cap	Solvent	- to dissolve OligoTool / Control	1000 μL of Solvent

Additional equipment and reagents required but not supplied: LightCycler® Fast Start DNA Master HybProbes, 10 x conc.Cat.No.: 03003248001, including 25mM MgCl₂; LightCycler® 480 instrument, LightCycler® 480 Multiwell Plates 96 white, DNA extraction materials

2. Introduction

2.1. Product overview

ToolSet description The ToolSet is specifically adapted for genotyping the amino acid T181A respectively

nucleotide A541G polymorphism in the UGT1A6 gene 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 times.

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

Primers? Probes?

You don't have to add primers. 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 UGT1A6 181–16, dissolved	2.8
Solvent UGT1A6 181-16	8.8
MgCl ₂ 25 mM	2.4
Fast Start DNA Master HybProbes,10x	2
Total Reaction Mix	16
+ Your DNA or Control UGT1A6 181-16, dissolved	4
Grand Total	20

Use Fast Start DNA Master Hybridization Probes 10x and MgCl, 25 mM from Roche LightCycler Fast Start DNA Master Hybridization Probes, 10 x conc. (Roche Cat.No.: 03003248001, 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 UGT1A6 181-16 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 UGT1A6 T181A ToolSet™ for LightCycler™ allows the detection of the A541G nucleotide exchange = T181A amino acid exchange in the UGT1A6 gene which is usually linked with the A552C nucleotide exchange = R184S amino acid exchange and associated with diminished activity of the UGT1A6 gene product – i.e. the glucuronidation enzyme UGT1A6. UGT1A6 metabolizes mainly planar phenols such as 4-naphtol, 4-methylphenol and planar arylamines like 1- and 2naphtylamine. Medical drugs such as aspirin and other NSAID's are also substrates for UGT1A6, and the protective effect of aspirin on colon adenoma risk was recently shown to be restricted to carriers of UGT1A6 181A mutant alleles (Cancer Res. 2001 61(9):3566-9; PubMedID: 11325819).

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



4. LightCycler 480 Settings and Experimental Protocol

For use with LC 480 Program Version 1.5 series. Detection: Dynamic Red 640 (498-640 nm)

Fast Start Enzyme Activation and DNA Denaturation

Cycle Program Data	Value	
Cycles	1	
Analysis Mode	None	
Temperature Targets	Segment 1	
Target Temperature (°C)	95	
Incubation time (s) Hold	600	
Temperature Transition Rate (°C/s) Ramp Rate	4.4	
Acquisitions (per °C)		
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	Quantification		
Temperature Targets	Segment 1	Segment 2	Segment 3
Target Temperature (°C)	95	53	72
Incubation time (s) Hold	5	12	18
Temperature Transition Rate (°/s) Ramp Rate	4.4	2.2	4.4
Acquisitions (per °C)			
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	Melting Curves	
Temperature Targets	Segment 1	Segment 2	Segment 3
Target Temperature (°C)	95	40	95
Incubation time (s) Hold	60	60	
Temperature Transition Rate (°/s) Ramp Rate	4.4	2.2	0.29
Acquisitions (per °C)			2
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) Hold	30
Temperature Transition Rate (°/s) Ramp Rate	2.2
Secondary Target Temperature (°C)	0
Step Size (°C)	0
Step Delay (Cycles)	0
Acquisition Mode	None

5. Typical results

Introduction

Use the Melting Curve Analysis 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 UGT1A6 T181A ToolSetTM for LightCyclerTM:

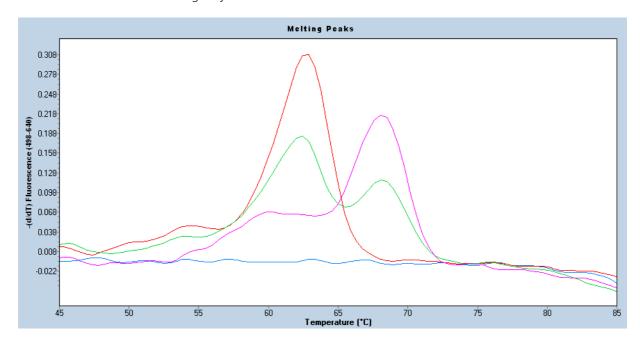


Figure 1: Melting curve analysis of possible genotypes of the UGT1A6 sequence.

RED: Homozygote wild type **TT181** DNA ($T_m = 62,6$ °C).

GREEN: The Heterozygote **T181A** Control DNA contained in the ToolSet (Green Cap tube).

PINK: Homozygote mutant **181AA** DNA ($T_m = 68.2$ °C).

Blue: NO DNA Control.

Conditions: LC 480 program version 1.5

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 +/- 1.0 °C. The UGT1A6 T181A 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 ratiogen.

6. Notices to Purchaser: Licenses and Trademarks, Prohibition of Resale

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