



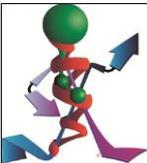
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A Geno Technology, Inc. (USA) brand name

# *Toothpick™-PCR*

(Cat. #786-410)



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## INTRODUCTION

*Toothpick*<sup>™</sup>-PCR allows for the rapid analysis of plasmid DNA by the polymerase chain reaction (PCR) from a bacterial colony itself. There is no requirement for growing bacteria, performing “minipreps” or purifying the plasmid DNA. Simply pick a bacterial colony and screen with *Toothpick*<sup>™</sup>-PCR to see if you have the right construct. This kit has enough reagents for the screening of 300 colonies.

The 6X Glow BromoBlue<sup>™</sup> Dye is Ficoll based and there is no need to add ethidium bromide to the running buffer as the dye contains ethidium bromide. Glow BromoBlue<sup>™</sup> Dye provides intense DNA bands with little background or band distortion and can be used with any type of agarose or acrylamide gels.

## ITEM(S) SUPPLIED (Cat# 786-410)

|   |       |
|---|-------|
| Plasmid Screening <i>Toothpick</i> <sup>™</sup> | 4.5ml |
| 6X Glow BromoBlue <sup>™</sup> Dye              | 1.5ml |

## STORAGE CONDITIONS

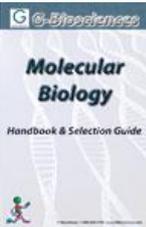
The kit is shipped at ambient temp. Store it at 4°C, upon arrival. The kit is stable for one year, if stored and used properly.

## PROTOCOL

1. Pick a freshly plated bacterial colony (1-2 mm in diameter) with a sterile toothpick or pipette tip and suspend transfer to the bottom of a 0.6ml tube or microtiter plate well. Remember to clearly label the colonies picked for easy identification of positive colonies later.
2. Add 15µl of the Plasmid Screening *Toothpick*<sup>™</sup> solution to the tube or well and vigorously vortex to resuspend the bacteria. Incubate at room temperature for 1-2 minutes and mix again.
3. Heat the sample at 100°C in a boiling water bath for 5 minutes.
4. Vortex and chill on ice for 2 minutes.
5. Transfer 1µl sample to a 50µl PCR reaction mix (Primers, deoxynucleotides, polymerase and polymerase buffer).
6. Perform the PCR reaction as normal.
7. After the PCR reaction is complete, transfer 10µl PCR reaction mix to a fresh tube and add 2µl of the 6X Glow BromoBlue<sup>™</sup> Dye to the sample. Mix and load into the wells of an agarose gel. No additional ethidium bromide is required.
8. Once electrophoresis is complete visualize PCR products under UV.

## RELATED PRODUCTS

Download our Molecular Biology Handbook.



<http://info2.gbiosciences.com/complete-molecular-biology-handbook>

For other related products, visit our website at [www.GBiosciences.com](http://www.GBiosciences.com) or contact us.

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