

Catalog Number	Kit Size
TBS42020-100	100 assays
TBS42020-200	200 assays

### DESCRIPTION

The Universal Aspergillus qPCR Kit is designed for identifying aspergillus genus species in a one PCR reaction using real-time quantitative polymerase chain reaction (qPCR) and probe fluorescence label. The detection of target DNA confirms ingredient authenticity or prevents food fraud, ethical issues, or health concerns.

### PRINCIPLE

Authenticating ingredients using real-time PCR is based on the amplification of a specific region of the relevant target genome. The amplified product is detected using target-specific fluorescent probes that bind to the amplified product. As the PCR product accumulates, there is an increased fluorescent signal from the bound probes. Monitoring the fluorescence intensities during the PCR run allows the detection of the accumulating PCR product in real time.

The Universal Aspergillus qPCR Kit includes aspergillus target positive and negative Controls, and PCR internal controls, qPCR super mix, primer-probe mix, in which the probe is labeled with FAM for aspergillus species, and Hex is labeled for PCR internal control. These aids in the straightforward interpretation of the results.

### KEY FEATURES

- ❖ Highly sensitivity and specificity for aspergillus detection.
- ❖ High efficiency: the optimal systemic conditions for PCR amplifications.
- ❖ Streamlined protocol: Just add DNA Template, and water.
- ❖ No cross reactivity with other species.

### APPLICATIONS

Detect aspergillus-derived DNA in plant, cannabis, cannabis ingredients, grain, food, herbals, and animal feed.

### KIT CONTENTS

Name	100RXN	200RXN
qPCR Super Mix (AG1)	0.8 mL	1.6 mL
Primer-probe Mix (AG2)	0.6 mL	1.2 mL
Positive Control DNA (AG <sup>+</sup> )	60 µL	100 µL
Negative Control DNA (AG <sup>-</sup> )	60 µL	100 µL

The Aspergillus probe is labeled with **FAM**, and PCR internal control is labeled with **Hex**.

### STORAGE CONDITION

The kit is shipped on ice and stored at -20°C for long-term storage. Shelf life of 12 months after receipt.

### PCR PROTOCOL

1. Set up PCR reaction for each sample in 20 µL

Reaction Component	Volume (µL)
qPCR Super Mix (AG1)	7.0
Primer-probe Mix (AG2)	5.0
Nuclease-free Water	3.0
DNA sample	5.0
Final Volume	20 µL

Internal control should be included as below: Positive Control (5 µL DNA /reaction) Negative Control (5 µL DNA/reaction)

2. Suggested PCR conditions

Step	Amplification	PCR	
	HOLD	CYCLE (40 cycles)	
		Denature	Anneal/ Extend
<b>Temperature</b>	95 °C	95 °C	60 °C
<b>Time</b>	2 min	15 sec	60 sec

### DATA ANALYSIS

Positive Reaction: Sample Ct < or = 37, and Positive, Negative and Blank controls are normal.

Negative Reaction: Sample Ct ≥ 38, and Positive, Negative and Blank controls are normal.

PCR internal control is positive in all samples, positive and negative controls. The positive response indicates a normal PCR amplification. Otherwise, the PCR reaction may be inhibited.

Repeat Reaction: If one of the control reactions is not normal, PCR reaction is failed, and should be repeated.

### RELATIVE PRODUCTS

- TBS6025: Microbial DNA Magnetic Extraction
- TBS42021: Aspergillus Flavus qPCR
- TBS42022: Aspergillus Fumigatus qPCR
- TBS42023: Aspergillus Niger qPCR
- TBS42024: Aspergillus Terreus qPCR
- TBS42025: 4-In-1 Aspergillus qPCR
- TBS42026: O157H7 E. Coli qPCR
- TBS42027: STEC qPCR
- TBS42028: Salmonella qPCR
- TBS42029: STEC and Salmonella Multiple qPCR
- TBS42030: Mycoplasma Detection qPCR
- TBS42031: Listeria Monocytogen qPCR
- TBS42032: Listeria Genus qPCR
- TBS42033: Bacillus Cereus qPCR

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