# bioticol

# qPCR KITS



## What is different in biotical qPCR KITS?

- All our gPCR kits are produced in the UE (SPAIN), are CE marked and has been validated under the new EU regulatory mark of sanitary products. that will be implemented in 2022.
- The products are LYOPHILIZATED, it is an important value comparing with other manufacturers:
  - NO COLD CHAIN REQUIRED. The product is stable at 25°C.
  - DOUBLE OF EXPIRY DATE (24 Months) comparing with other competitors.
- DEDICATED SUPORT TO THE DISTRIBUTOR: We have created manuals for use our kits in the most common thermocyclers of the market. In case of a new machine, we support you for a perfect adjustment of the machine.
- Specialized catalogue with high added value products.
- Customization options in controls, strips measures and highs.

### **Customization options**

- 96 Strips or 48 Strips option in all references.
- Low Strip, high Strip or Tube.
- 3 types of controls available:

#### **INTERNAL CONTROL**

A

-Q

It is a **parallel reaction** ran in the same reaction well with the same PCR reagents except primer and probe, the reaction is followed in different fluorescence channel (HEX or Cy5).

#### IC serves to verify the PCR assay:

- The reaction well has properly rehydrated
- The reagents are in proper condition
- The PCR system has executed the thermal cycle and the optical system is working

#### **EXTRACTION CONTROL**

EC (green vial) is a non-infectious nucleic acid lyophilized with the sequence of target (amplicon).

EC HEX or Cv5 channel.

Extraction Control can be used to monitor nucleic acid isolation and/or as PCR inhibition control.

#### **ENDOGENOUS CONTROL**

It is a **parallel reaction** ran in the same reaction well with the same PCR reagents except primer and probe.

Housekeeping gene (human RNase P gene).

Human housekeeping genes are involved in basic cell maintenance and, therefore, are expected to be present in all nucleated human cells and maintain relatively constant expression levels.



MENINGITIS	• N. meningitis + H
	• E. coli + S. Agala
TRANSPLANTATION	• Herpes virus 1 + 2
IRANSPLANTATION	
	• Herpes virus 6 +
	• Poliomavirus BK
RESPIRATORY VIRUS	• FLU (H1N1, H5N1,
	• FLU A + FLU B +
	• SARS-COV 2 qPG
RESPIRATORY BACTERIA	• S. pneumoniae +
	Bordetella pertuss
GASTROINTESTINAL BACTERIA	• Eschericia coli (E
	Shigella / Entero
	• Yersinia enteroco
	Helicobacter pylo
ANTIBIOTIC RESISTANCE	Clarithromycin re
GENETIC MARKERS	• Genetic HLA Cel



I. Influenzae, + S. pneumoniae gPCR ctiae + L. monocytogenes qPCR

2 + Varicella Zoster qPCR 7 + 8 gPCR + John Cunninghan (JC) Virus aPCR

H3N2 + H7N9) qPCR RSV aPCR CR

H. influenzae + M. catarrhalis gCPR sis + B. holmesii + B. parapertussis gPCR

EIEC) + Salmonella + Campylobacter + invasive gPCR olitica + Aeromonas gPCR ori qPCR

esistance + H. Pylori gPCR

iac detection gPCR





#### **Thermal Cycling Protocol for DNA:**



Step 1 Add 15 µl of rehydration buffer into each well

Add 5 µl of DNA/ RNA sample / positive control / negative control

Load the strips into the thermocycler and run the specified protocol

Interpretate results

#### **EASY PERFORMANCE:**

Just add: 15µl of buffer 5µl of sample

Step	Тетр	Time	Cycles
Initial denaturation	95ºC	2 min	1
Denaturation	95ºC	10 sec	
Annealing/Extension (Data collection*)	60ºC	50 sec	45



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