

qPCR KITS





What is different in biotical qPCR KITS?

- All our qPCR kits are produced in the EU (SPAIN), are CE marked and have been validated under the new EU medical device EU regulatory mark for medical devices.
- The products are LYOPHILIZATED.
 - 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.



Customization options

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

INTERNAL CONTROL

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 **Cy5** 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.

biotical

Real Time PCR Product Catalogue



| MENINGITIS | • N. meningitis + H. Influenzae, + S. pneumoniae qPCR | | |
|---------------------------|--|--|--|
| | • E. coli + S. Agalactiae + L. monocytogenes qPCR | | |
| | | | |
| TRANSPLANTATION | • Herpes virus 1 + 2 + Varicella Zoster qPCR | | |
| | Herpes virus 6 + 7 + 8 qPCR | | |
| | • Poliomavirus BK + John Cunninghan (JC) Virus qPCR | | |
| | | | |
| RESPIRATORY VIRUS | • FLU (H1N1, H5N1, H3N2 + H7N9) qPCR | | |
| | • FLU A + FLU B + RSV qPCR | | |
| | • SARS-COV 2 qPCR | | |
| | | | |
| RESPIRATORY BACTERIA | • S. pneumoniae + H. influenzae + M. catarrhalis qCPR | | |
| | Bordetella pertussis + B. holmesii + B. parapertussis qPCR | | |
| | | | |
| GASTROINTESTINAL BACTERIA | Eschericia coli (EIEC) + Salmonella + Campylobacter + Shigella / Enteroinvasive qPCR | | |
| | Yersinia enterocolitica + Aeromonas qPCR | | |
| | Helicobacter pylori qPCR | | |
| | • E. Coli Typing qPCR | | |
| | | | |
| ANTIBIOTIC RESISTANCE | Clarithromycin resistance + H. Pylori qPCR | | |
| | Methicillin-resistant + Staphylococcus aureus qPCR | | |
| | | | |
| GENETIC MADKEDS | Le Constic HI A Coline detection appe | | |
| GENETIC MARKERS | Genetic HLA Celiac detection qPCR | | |



Thermal Cycling Protocol for DNA:



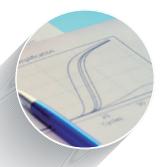
Step 1 Add 15 µl of rehydration buffer into each well



Step 2
Add 5 µl of DNA/
RNA sample /
positive control /
negative control



Step 3
Load the
strips into the
thermocycler and
run the specified
protocol



Step 4Interpretate results

EASY PERFORMANCE:

Just add: 15µl of buffer 5µl of sample

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



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