

## Manual

# Anti-Inhibitor Kit

Kit for removing PCR inhibitors from DNA preparations.

| catalog # | size           |
|-----------|----------------|
| 1015-50   | 50 isolations  |
| 1015-250  | 250 isolations |

For research use only.

### Guarantee

A&A Biotechnology provides a guarantee on this product.

The company does not guarantee the correct performance of this kit in the event of:

- not adhering to the supplied protocol
- use of not recommended equipment or materials
- use of other reagents than recommended or which are not a component of the product
- use of expired or improperly stored product or its components

## Description

The principle of Anti-Inhibitor Kit is based on the ability to bound polyphenolic PCR inhibitors by absorption beads. After the addition of absorption beads to the column the DNA sample is loaded onto the minicolumn. In the next step the minicolumn is centrifuged. DNA which is contained in the filtrate is free of polyphenolic compounds known as potent inhibitors of many enzymatic reactions.

## Contents

| component        | 50 isolations | 250 isolations | storage  |
|------------------|---------------|----------------|----------|
| Minicolumns      | 50 pcs        | 250 pcs        | 15–25 °C |
| Absorption beads | 45 ml         | 210 ml         | 15–25 °C |

## Additional equipment and reagents

### Necessary

- 1.5 ml sterile Eppendorf tubes
- Microcentrifuge

## Preparation of minicolumns

1. Mix the bottles with the suspension of the absorption beads by inverting to get a homogeneous mixture.
2. Add **700 µl** of the absorption beads to the minicolumns.  
To transfer the dense suspension the absorption beads we suggest to cut off 2-3 mm of the end of a single pipette tip. The tip internal diameter should be larger than standard.
3. Close the minicolumns and centrifuge for **3 min at 8000 x g**.  
After centrifugation the surface is inclined.
4. Place the minicolumns into the 1.5 ml Eppendorf tubes (not included). The minicolumns are ready to use.

## Removal inhibitors from DNA samples

1. Apply **50 µl** of DNA samples containing inhibitors onto the surface of the absorbing particles placed in the minicolumns.
2. Centrifuge for **1 min at 8000 x g**.
3. Remove the minicolumns and close the tubes. The DNA obtained in the filtrate is ready for direct use for PCR or other enzymatic reactions. It may be stored until later use.



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version 2023-1

