AZUL Saliva DNA EXTRACTION KIT

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INTENDED USE

AZUL Saliva DNA Extraction Kit is an easy and efficient system for the isolation of total DNA from Saliva Samples.

SUMMARY AND EXPLANATION

This kit uses a silica-based spin column technology for isolating DNA from biological samples, thereby eliminating toxic phenol-chloroform extractions. The eluted DNA is suitable for all sensitive downstream applications such as qPCR and Next-Generation sequencing.

PRODUCT FEATURES

- Rapid purification of high-quality, ready-to-use DNA.
- No organic extraction or alcohol precipitation.
- Consistent and high yields.
- Complete removal of contaminants and inhibitors for reliable results.
- Kit formats for low- to high-throughput – options for automation of all kits.

PRECAUTIONS

- AZUL Saliva DNA Extraction kits are intended for use as supplied. Do not dilute or add other components to the AZUL Saliva DNA Extraction kit.
- Dispose of used reagents, debris, and consumables as hazardous waste according to established laboratory procedures.

DIRECTIONS FOR USE

1. In a microfuge tube, take around 300 µL - 1 mL of fresh Saliva samples or Saliva stored in mWRAPR Saliva Collection Tubes and add 300 µL - 700 µL of Lysis Buffer 1 (LB1).
2. To the tube, add 25 µL of Lysis Buffer 2 (LB2) and mix briefly by pipetting 2-3 times or vortex for 30 sec.
3. Add 50 µL of Proteinase K, mix well and incubate the tube at 56°C for 15 mins.
4. Centrifuge the tube at 13,000 rpm for 5 mins.
5. Carefully transfer the clear supernatant to a new 1.5 mL microfuge tube. Add 500 µL of Binding Buffer (BB) and mix the tube briefly by inverting it a few times.
6. Transfer 800 µL lysate to the spin column inserted in a collection tube and centrifuge at 12,000 rpm for 2 mins.
7. Discard the flow-through and place the purification column back into the collection tube. Repeat this step until the entire lysate has been transferred into the column and centrifuged.
8. Add 600 µL of Wash Buffer 1 (WB1) to the column and centrifuge at 12,000 rpm for 1 min.
9. Add 500 µL of Wash Buffer 2 (WB2) to the column and centrifuge at 12,000 rpm for 1 min to completely remove salts and impurities.
10. Transfer the purification column to a clean, sterile microfuge tube and add 30 µL -50 µL of Elution Buffer or DNase/RNase-free water to the center of the column.
11. Centrifuge the column at 12,000 rpm for 2 minutes.
12. Discard the purification column and store the eluted DNA at -20°C or -80°C until use.

KIT COMPONENTS

Components	For 50 preps	For 25 preps
Lysis Buffer 1 (LB1)	35 mL	20 mL
Lysis Buffer 2 (LB2)	2 mL	1 mL
Proteinase K	3 mL	2 mL
Binding Buffer (BB)	25 mL	15 mL
Wash Buffer 1 (WB1)	30 mL	15 mL
Wash Buffer 2 (WB2)	25 mL	15 mL
Elution Buffer (EB)	4 mL	2 mL
Spin Column	50 (Pouch pack)	25 (Pouch pack)

CAUTION

- Check the Binding Buffer and Lysis Buffer for any salt precipitation before every use.
- Re-dissolve any precipitate by warming the solution to 37°C, then cool it back to room temperature before use.
- During operation, always wear a lab coat, disposable gloves, protective goggles and mask.

KIT STORAGE AND STABILITY

- Store the kit at room temperature, Proteinase K and Lysis Buffer 2 at -20°C.
- Viable for 1 year if stored at appropriate conditions.

ORDERING INFORMATION

Please call us at +91 8088747968 or mail at hello@azooka.life for any queries or assistance.

Additional information can be found online at www.azooka.life

MANUFACTURED AT:
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