

(For Professional Use Only) V3

1. Intended Purpose

The TANBead® Nucleic Acid Extraction Kit is a nucleic acid purification kit based on magnetic bead technology by using with corresponding TANBead® Nucleic Acid Extractor, which can automatically isolate and purify high-quality microbial and host DNA from stool of human and other species. Moreover, additional step in pretreatment can be performed to enrich the desired extraction products of microbiome. The isolated DNA is ready for downstream applications such as PCR, Real-time PCR and microbiome profiling. The kit is intended for use by technicians, physicians, and biologists with well-trained in molecular biological techniques, the techniques of magnetic bead purification and in vitro diagnostic procedures. Any diagnostic results generated by using the sample preparation procedure in conjunction with any downstream diagnostic assay should be interpreted related to other clinical or laboratory findings. The kit is not limited to any specific disorder, condition, or other additional accompanying diagnostics. It is applicable for all population.

2. The basic principle

The silicon dioxide layer coated on the magnetic beads can adsorb the negatively charged molecules to purify nucleic acids from samples.

3. Specification

Starting Materials	50 mg stool
Elution Volume	50~80 μL
Typical DNA yield	≥2 µg

4. Component Supplied with the Kit

Auto Tube	8 trays	Auto Tube with reagent buffers
Proteinase K	1.0 mL x 2	Proteinase K
Elution Buffer	1.5 mL	Nuclease-Free Water
Incubation Buffer	120 mL	Phosphate buffer
-B845	120 IIIL	for omnivore use
Incubation Buffer	60 mL x 2	Phosphate buffer
-B871	00 IIIL X Z	for herbivore use
Strip	24	8-channel strip
Base	2	A rack for 8 Auto Tubes
Protocol	1	Instruction guide for user

5. Auto Tube Content

Well	Buffer	Volume (μL)
1 / 7	Lysis Buffer	600
2/8	Washing Buffer 1	800
3/9	Washing Buffer 2	800
4 / 10	Washing Buffer 2	800
5 / 11	Magnetic Beads	800
6 / 12	Elution Buffer	80

6. Kit Storage and Shelf Life

- Components under room temperature (15~35°C) can be stored until the expiration date labeled on the box.
- 2) The proteinase K is transported at room temperature. Upon received, please store proteinase K at 2~8°C.

7. Precautions

- 1) It can only be used for in vitro diagnostic.
- 2) Avoid using expired reagents.
- 3) When the temperature is below 20°C, place the Auto Plates / Auto Tubes in an oven (preheated 42~60°C) 5 to 10 minutes.
- Avoid vigorous shaking, in order to avoid excessive formation of foam.
- 5) Carefully remove aluminum foil to avoid splashing.
- 6) Do not expose the opened reagents or Auto Plates / Auto Tubes to air. The evaporation would lead to pH change, or effect on the extraction effectiveness.

- Please check the integrity of the Auto Plates / Auto Tubes and remember to insert the strips into the appropriate position of the suitable instrument before operating them.
- 8) Please wear a mask and disposable gloves when handling.
- 9) Use sterile consumables to avoid nuclease contamination.
- Reagent solution contains guanidine salt, avoid using bleach containing detergent.
- Avoid eyes, skin, and clothing contact with reagents. In case of any contact, flush with flowing water.
- 12) If any serious incident occurs, please report to the manufacturer and the competent authority of the member state in which the user and / or the patient is established.

8. Materials required, Not Supplied

- TANBead[®] Nucleic Acid Extraction System Model: SLA-16 / 32 / E13200 series (non-sterile)
- 2) Disposable gloves
- 3) Scissors, utility knives
- 4) Micropipette, disposable tips (10 μL / 200 μL / 1000 μL)
- 5) 1.5 mL microcentrifuge tube
- 6) 15 mL / 50 mL conical tube

9. Sample Collection, Transportation, and Storage

Sample collection and storage

- 1) Stool sample can be stored at
 - a. RT for 24 hours.

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- b. 2~8°C up to 7days.
- c. -20°C long-term preservation.

■ Specimen transportation

Transportation of stool specimen should be followed by specific infectious biological materials transportation-related law.

10. Nucleic Acids Extraction Protocol

Before operating, turn on the warm-up system of TANBead® Nucleic Acid Extractor, if it is equipped with temperature controller, please setting at 50°C

- 1) Carefully remove the aluminum foil on the Auto Tubes.
- Add 500 μL incubation buffer-B845 / B871 and 20 μL Proteinase K into 1.5 ml, tube

Note: Incubation buffer-B845: For Omnivore Use; Incubation buffer-B871: For Herbivore Use.

- 3) Add about 50 mg stool into 1.5 mL tube and mix well.
- 4) Incubate at 60°C for 10 minutes on heater.
- For Bacterial gDNA: Centrifuged at 10,000 x g for 1 minute.
 For Human gDNA: Skip this step and continue to the next step.
- 6) Transfer the supernatant into column #1 / #7 of Auto Tube.

Note: If samples are difficult to transferred, please use a cut off pipette tip and pipette gently.

- Push Assembled Auto Tubes completely to the bottom of the plate rack. Make sure that the chamfer of the plate is at the lower left.
- 8) Push strips completely to the bottom of strip rack frame.
- 9) Close the door panel.
- Select the program "6SC". The parameters are given in the following section.
- 11) When the program ends, buzzer shall alarm. Take out the auto tube carefully.
- 12) Use micropipette to transfer the purified nucleic acids from column #6 / #12 to a clean tube.
- Discard the used Auto Tubes and strips into the waste recycling bin

11. Program

■ SLA-16 / 32 series

Program Name: 6SC				Model: SLA-16 / 32 series				
Step	Well	Mixing (M)	Collect (S)	Rod	Mixing speed	Volume (μL)	Pause	Vapor (M)
1	5	0	60	On	Medium	800	Off	0
2	1	10	60	On	Low	800	Off	0
3	2	1	60	On	Medium	800	Off	0
4	3	1	60	On	Medium	800	Off	0
5	4	1	60	On	Medium	800	Off	10
6	6	5	60	On	Medium	80	Off	0
7	3	1	0	Off	Medium	800	Off	0
8	0	0	0	Off	Medium	0	Off	0

■ SLA-E13200 series

Program Name: 6SC					Model: SLA-E13200 series				
Step	Well	Temp (°C)	Mixing (M)	Collect (S)	Rod	Mixing speed	Volume (μL)	Pause	Vapor (M)
1	5	50	0	60	On	Medium	800	Off	0
2	1	50	10	60	On	Low	800	Off	0
3	2	50	1	60	On	Medium	800	Off	0
4	3	50	1	60	On	Medium	800	Off	0
5	4	50	1	60	On	Medium	800	Off	10
6	6	50	5	60	On	Medium	80	Off	0
7	3	N/A	1	0	Off	Medium	800	Off	0
8	0	N/A	0	0	Off	Medium	0	Off	0

12. Result

Nucleic acid product purified by TANBead® nucleic acid extraction kit can perform qualitative / quantitative analysis of specific genes by PCR, RT-PCR, Q-PCR or qRT-PCR. Please refer to the molecular diagnostic kit manual.

13. Reagent performance

Qualitative Analysis

A specific gene fragments can be amplified from nucleic acids products isolated from TANBead® nucleic acid extraction kit by PCR (Polymerase Chain Reaction) or RT-PCR (Reverse Transcription-PCR). This kit can work with different molecular biology reagents and apply for verity of molecular diagnosis.

■ The stability of extracted DNA

Storage Conditions	DNA stability
-80°C	Over 90 days
-20°C	28 days
4°C	14 days
25°C	2 days
Freeze-thaw	10 times

14. Explanation of Symbols

	Manufacturer	[]i	Consult instructions for use
15°C-	Temperature limit	Σ	Contains sufficient for test
CE	CE mark	IVD	In vitro diagnostic medical use
REF	Catalogue number	\triangle	Caution
LOT	Batch code	NON STERILE	Non-sterile
8	Do not re-use	紫	Keep away from sunlight
Μ	Date of manufacture	8	Use-by date

EC REP

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15. Post-market surveillance conclusion

After a risk assessment and clinical evaluation assessment, when weighing the benefits of medical device, patients, and the risks associated with the use of the device, the risk is acceptable. The postmarket surveillance report shows that no death or serious adverse events occurred.