



6T2S46

(For Professional Use Only) V2

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 total DNA from tissue. The purified DNA can be used with any downstream application which is qualitative or semi-quantitative assay. 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	50~100 mg tissue
Elution Volume	90~130 μL
Typical DNA yield	≥2 µg
Typical A260 / A280	≥1.7

4. Component Supplied with the Kit

Auto Tube	8 trays	Auto Tube with reagent buffers
Proteinase K	1.0 mL x 1	Proteinase K
Incubation Buffer	25 mL x 1	Tris buffer, surfactants, pH 8.0
Elution Buffer	20 mL x 1	Nuclease-Free Water
Base	2	A rack for 8 Auto Tubes
Protocol	1	Instruction guide for user
Strip	24	8-channel strip

5. Auto Tube Content

Well	Buffer	Volume (µL)	
1/7	Lysis Buffer	700	
2/8	Washing Buffer 1	800	
3/9	Magnetic Beads	800	
4 / 10	Washing Buffer 2	800	
5 / 11	Washing Buffer 2	800	
6 / 12	Elution Buffer	130	

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 be used for in vitro diagnostic use.
- 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.
- 7) 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 Pre-treatment, Storage and Transportation

■ Sample pre-treatment

- Place 50~100 mg minced tissue into 1.5 mL tube then add 200 μL incubation buffer and 10 μL Proteinase K then mix well.
- After incubated at 56°C for 1~2 hours on heater, centrifuged with 10000 RPM for 5 minutes.

Note: Vortex the sample for 10 seconds every 15 minutes to improve extraction efficiency.

Sample storage

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- 1) Animal tissue can be stored at
 - a. RT for 24 hours.
 - b. 2~8°C up to 7 days.
 - c. -20°C for long-term preservation.

■ Specimen transportation

Transportation of animal tissue specimen should follow 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 45°C.

- Preparing the Assembled Auto Tubes by inserting Auto Tubes into the Base completely.
- 2) Carefully remove the aluminum foil on the Auto Tubes.
- 3) Use micropipette to load 200 µL supernatant into well #1 / #7.
- 4) 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.
- 5) Push strips completely to the bottom of strip rack frame.
- 6) Close the door panel.
- Select the program "L-BNA-PK-AUTO". The parameters are given in following section.
- Once the program has ended, buzzer shall alarm. Take out Auto Tube carefully.
- 9) Carefully remove the Auto Tubes when the program is finished.
- 10) Use micropipette to transfer the purified nucleic acids from well #6 / #12 to a clean tube.
- 11) Discard used Auto Tubes and strips into the waste recycling bin.

11. Program

■ SLA-16 / 32 series

Progra	Program Name: L-B NA-PK-AUTO				Model: SLA-16/ 32 series			
Step	Well	Mixing (M)	Collect (S)	Rod	Mixing speed	Volume (μL)	Pause	Vapor (M)
1	3	1	90	ON	Medium	800	OFF	0
2	2	1	0	OFF	Medium	800	OFF	0
3	1	10	0	OFF	Low	900	OFF	0
4	2	0	90	ON	Medium	800	OFF	0
5	1	10	90	ON	Medium	900	OFF	0
6	2	5	90	ON	Medium	800	OFF	0
7	3	5	90	ON	Medium	800	OFF	0
8	4	5	90	ON	Medium	800	OFF	0
9	5	5	90	ON	Medium	800	OFF	10
10	6	10	120	ON	Medium	200	OFF	0
11	5	1	0	OFF	Medium	800	OFF	0
12	0	0	0	OFF	Medium	0	OFF	0

■ SLA-E13200 series

Program Name: L-BNA-PK-AUTO					Model: SLA-E13200 series				
Step	Well	Temp (°C)	Mixing (M)	Collect (S)	Rod	Mixing speed	Volume (μL)	Pause	Vapor (M)
1	3	45	1	90	ON	Medium	800	OFF	0
2	2	45	1	0	OFF	Medium	800	OFF	0
3	1	45	10	0	OFF	Low	900	OFF	0
4	2	45	0	90	ON	Medium	800	OFF	0
5	1	45	10	90	ON	Medium	900	OFF	0
6	2	45	5	90	ON	Medium	800	OFF	0
7	3	45	5	90	ON	Medium	800	OFF	0
8	4	45	5	90	ON	Medium	800	OFF	0
9	5	45	5	90	ON	Medium	800	OFF	10
10	6	45	10	120	ON	Medium	200	OFF	0
11	5	NA	1	0	OFF	Medium	800	OFF	0
12	0	NA	0	0	OFF	Medium	0	OFF	0

12. Reagent performance

Repeatability

Under repeatability conditions where nucleic acids are extracted with the same reagent kit on the same source samples by the same operator. The coefficient of variation of nucleic acid extraction concentration is less than 5%.

■ Reproducibility

A five-day reproducibility test was carried out with the same source samples for 5 consecutive days with the same reagent kit by different operators. The coefficient of variation of nucleic acid extraction concentration is less than 5%.

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	5 times

13. Explanation of Symbols

***	Manufacturer	[]i	Consult instructions for use
15°C- 35°C	Temperature limit	Σ	Contains sufficient for test
C€	CE mark	IVD	<i>In vitr</i> o diagnostic medical use
REF	Catalogue number	\triangle	Caution
LOT	Batch code	NON	Non-sterile
(2)	Do not re-use	类	Keep away from sunlight
~~	Date of manufacture	8	Use-by date

EC REP

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14. 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.