

(For Professional Use Only) V5

# 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.

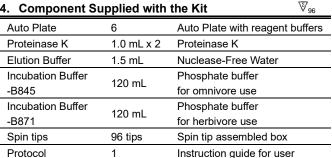
# 2. The basic principle

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



# 5. Auto Plate Content and Plate Position

Plate Position	Buffer	Volume (μL)
1	Lysis Buffer	600
2	Washing Buffer 1	800
3	Washing Buffer 2	800
4	Washing Buffer 2	800
5	Magnetic Beads	800
6	Elution Buffer	80
7	Spin tip	-

# 6. Kit Storage and Shelf Life

- Components under room temperature (15~35°C) can be stored until the expiration date labeled on the box.
- 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 reagent plate in an oven (preheated 42~60°C) from 5 to 10 minutes.
- 4) Avoid vigorous shaking, in order to avoid excessive formation of foam.
- Carefully remove aluminum foil to avoid splashing.
- 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 Plate / Auto Tube, and remember to mount the spin tips 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.
- 10) Reagent solution contains guanidine salt, avoid using bleach containing detergent.
- 11) 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

- 1) TANBead® Nucleic Acid Extraction System Model: Maelstrom 9600 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.
  - 2~8°C up to 7 days.
  - -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

- 1) Carefully remove the aluminum foil on the Auto Plates.
- 2) 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.
- 5) For Bacterial gDNA: Centrifuged at 10,000 x g for 1 minute. For Human gDNA: Skip this step and continue the next step.
- 6) Transfer the supernatant into wells of plate 1.

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

- 7) Follow the guide shown on the screen and place plates carefully. Make sure that the chamfer of the plate faces toward the same direction
- Carefully remove the Auto Plates when the program is finished.
- Use micropipette to transfer the purified nucleic acids from plate #6 to a clean tube.
- 10) Discard used Auto Plates and spin tips into the waste recycling

#### 11. Program

# ■ Maelstrom 9600

Program Name:6SC												
Plate		1		2		3	4	5	6	7		8
Volume(	μL)	90	0	800		800	800	800	150	-		-
Keep Temp.		45	5	45		40	-	-	50	-		
Action	١	Fo	r.	For		For.	For.	For.	For.	-		
Name		LE	3	WB	1	WB2	WB2	МВ	EB	-		TIP
Step	Pla	ate		emp. (°C)		lixing min)	Mixing (rpm)	Collec (sec)		Vapor (min)		ıse
1		5		-	0		3000	30	(	)	0	ff
2	:	1		60	10		3000	30	(	0		ff
3	:	2		50	1		3000	30	(	0		ff
4	;	3		40	1		3000	30	(	0		ff
5		4		-		1	3000	30	1	10		ff
6	(	6		65		5	3000	30	(	0		ff
7		5		-		0.1	3000	0		)	0	ff

# **△** Temperature set as "0" represents room temperature!

# ■ Maelstrom 9610

Program Name:6SC													
Plate	Plate 1		2	3	4	5		6		7	8		
Volume (μL)	9	900	800	800	800	800		150		-	-		
Keep Temp.		45	45	40	-	-		50		-	-		
Action		For.	For.	For.	For.	F	or.	For	For		-		
Name		LB	WB1	WB2	WB2	Ν	ИΒ	EB	EB -		TIP		
Step	Р	late	Temp. (°C)	Mixing (min)	Mixin (rpm		Col (se			apor min)	Pause		
1		5	-	0	0		3	0		0	Off		
2		1	60	10	3000	)	3	0		0	Off		
3		2	50	1	3000	)	3	0		0	Off		
4		3	40	1	3000	)	30 0		10		Off		
5		4	-	1	3000	3000		0	0 10		Off		
6		6	65	5	3000	3000		30 0		0	Off		
7		5	-	0.1	3000	3000		0		0 0		0	Off

### **⚠** Temperature set as "25" represents room temperature!

# 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 35°C	Temperature limit	Σ	Contains sufficient for test		
C€	CE mark	IVD	<i>In vitro</i> diagnostic medical use		
REF	Catalogue number	$\triangle$	Caution		
LOT	Batch code	NON	Non-sterile		
(3)	Do not re-use	类	Keep away from sunlight		
س	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.