

M6TFS46

(For Research Use Only) V1

#### 1. Intended Use

This product is designed for isolating nucleic acid from various samples, which can be performed by using TANBead® Nucleic Acid Extractor and is intended for research use only.

# 2. Purpose

TANBead® Nucleic Acid Extraction Kit (M6TFS46) is dedicated to the isolation of DNA from small sample quantity or challenging sizes. Samples need to be treated by following the preparation steps, incubated in incubation buffer and treated with Proteinase K and DTT (Dithiothreitol). The lysates need to be added into well of tube and automatically processed by TANBead® Nucleic Acid Extractor. The highly extracted and purified DNA can be directly applied in such applications, including real-time PCR, short tandem repeat (STR) analysis, mitochondria DNA (mtDNA) analysis and other forensic tests.

#### 3. The basic principle

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

### 4. Specification

Starting Materials	Forensic samples (blood stain, dried blood spot, hair follicle, semen, cigarette, chewing gum, nail)
Elution Volume	90~130 μL

#### $\overline{\mathbb{V}}_{96}$ 5. Component Supplied with the Kit Auto Tube with reagent buffers Auto Tube 8 trays Proteinase K 1.0 mL x 2 Proteinase K Incubation 65 mL x 1 Tris buffer, surfactants, pH 8.0 Buffer Elution Buffer 1.5 mL x 1 Nuclease-Free Water A rack for 8 Auto Tubes Base 48 tips x 2 Spin tips Spin tip assembled box boxes Protocol Instruction guide for user 1

## 6. Auto Tube Content

Well	Buffer	Volume (μL)
1 / 7	Lysis Buffer	500
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

## 7. Kit Storage and Shelf Life

- 1) 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 receipt, please store proteinase K at 2~8°C.

## 8. Precautions

- 1) For research use only.
- 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) for 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 mount the spin tips into the appropriate position of the suitable instrument before operating them.

- 8) Before using, if the incubation buffer precipitates, please preheated over 40°C at least 5 minutes until the precipitates dissolve.
- 9) Please wear a mask and disposable gloves when handling.
- 10) Use sterile consumables to avoid nuclease contamination.
- Reagent solution contains guanidine salt, avoid using bleachcontaining detergent.
- 12) Avoid eyes, skin, and clothing contact with reagents. In case of any contact, flush with flowing water.
- 13) 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.

## 9. Materials required, Not Supplied

- TANBead® Nucleic Acid Extraction System Model: Maelstrom 8 series, Maelstrom 4800 series (non-sterile)
- 2) Disposable gloves
- 3) Scissors, utility knives
- 4) Micropipette, disposable tips (10  $\mu$ L / 200  $\mu$ L / 1000  $\mu$ L)
- 5) 1.5 mL microcentrifuge tube
- 6) 15 mL / 50 mL conical tube
- 7) 1M Dithiothreitol (DTT)

# 10. Sample Collection, Storage and Transportation

#### ■ Sample collection

Sample	Collection
Cigarette	Cut out 1/2 of the front filter and divide it into two pieces. Put both pieces (contain filter and outer paper) into a 1.5 mL tube.
Hair	Collect at least five hairs (0.5 - 1 cm with follicle) into 1.5 mL tube.
Blood stain	Collect stains with cotton swab and put the cotton part of swab into a 1.5 mL tube.
Dried blood spot	Collect 1 piece of dried blood spot (Φ = 6mm) into a 1.5 mL tube.
Semen stains	Collect stains with cotton swab and put the cotton part of swab into a 1.5 mL tube.
Chewing gum	Cut chewing gum into at least 10 mg and transfer them to a 1.5 mL tube.
Nail	Collect one nail and put it into a 1.5 mL tube.

# Specimen storage

- Forensic specimen should be analyzed as fresh as possible, if you need to storage specimen, follow the instruction:
  - a. RT for 24 hours.
  - b. 2~8°C up to 7 days
  - c. -20°C for long-term preservation

## Specimen transportation

Transportation of forensic specimen should follow specific forensic sample related regulation and keep specimen at RT during transportation.

## 11. Nucleic Acids Extraction Protocol

# 1) Sample pre-treatment

Sample	Treatment
Cigarette	Add 600 µL Incubation Buffer, 20 µL Proteinase K into 1.5 mL tube, then mix well.
Hair	Add 300 μL Incubation Buffer, 20 μL Proteinase K and 20 μL 1M DTT into 1.5 mL tube, then mix well.
Blood stain	Add 600 µL Incubation Buffer, 20 µL Proteinase K into 1.5 mL tube, then mix well.
Dried blood spot	Add 600 µL Incubation Buffer, 20 µL Proteinase K into 1.5 mL tube, then mix well.
Semen stains	Add 600 μL Incubation Buffer, 20 μL Proteinase K and 20 μL 1M DTT into 1.5 mL tube, then mix well.
Chewing gum	Add 300 µL Incubation Buffer, 20 µL Proteinase K into 1.5 mL tube, then mix well.
Nail	Add 300 μL Incubation Buffer, 20 μL Proteinase K and 20 μL 1M DTT into 1.5 mL tube, then mix well.

- 2) Incubate at 56°C, 900 rpm for at least 1 hour.
- 3) Preparing the Assembled Auto Tubes by inserting Auto Tubes into the Base completely.
- 4) Carefully remove the aluminum foil on the Auto Tubes.
- Add all lysate into well #1 / #7 of Auto Tube (Well filled with lysis buffer).
- 6) Set up spin tips.

**Maelstrom 8 series:** Handle to mount tips and make sure that there is no gap between the necks of spin tips and the spin shaft. **Maelstrom 4800 series:** Go to Tip page and press the mount tips region.

- 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) Select the program.

Maelstrom 8 series: Press "6TF-1" for input specimens at column #1 or "6TF-7" for input specimens at column #7.

Maelstrom 4800 series: Press "6TF".

The parameters are given in following section.

- 9) Carefully remove the Assembled Auto Tubes when the program is finished.
- 10) Use a micropipette to transfer the purified nucleic acids from well #6 / #12 to a clean tube.
- 11) Discard used Auto Tubes and spin tips into the waste recycling bin.

#### 12. Program

## **Maelstrom 8 series**

Program Name: 6TF-1 / 7							
Well	1/7 2/8 3/9 4/10 5/11 6/12						
Volume	<b>1100 (</b> μL)	800 (μL)	800 (μL)	800 (μL)	800 (μL)	<b>130 (</b> μL)	

Step	Well	Action	RPM	Time (Second)	CW/CCW (Second)	Temp.	Temp. Control
1	3/9	Mixing	3000	30	0	0	NO
2	3/9	Collection	0	30	0	0	NO
3	2/8	Mixing	3000	30	0	0	NO
4	1/7	Mixing	3000	420	0	70	YES
5	2/8	Collection	0	30	0	0	NO
6	1/7	Mixing	3000	480	0	0	NO
7	1/7	Collection	0	180	0	0	NO
8	2/8	Mixing	3000	120	0	0	NO
9	2/8	Collection	0	60	0	0	NO
10	3/9	Mixing	3000	60	0	0	NO
11	3/9	Collection	0	30	0	0	NO
12	4/10	Mixing	3000	60	0	0	NO
13	4/10	Collection	0	180	0	0	NO
14	5 / 11	Mixing	3000	60	0	0	NO
15	5 / 11	Collection	0	180	0	0	NO
16	5/11	Vapor	0	480	0	0	NO
17	6 / 12	Mixing	3000	300	0	0	NO
18	6 / 12	Collection	0	480	0	0	NO
19	3/9	Mixing	3000	10	0	0	NO

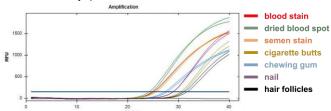
## Maelstrom 4800 series

Program N	Program Name: 6TF			Model: M	aelstrom 4	800 series	
Temp1	Temp2						
40	40						
Well	Name	Volume	Action	Mixing	Collect		
1/7	LB	1100	For.	Low	Low		
2/8	WB1	800	For.	Low	Low		
3/9	МВ	800	For.	Low	Low		
4 / 10	WB2	800	For.	Low	Low		
5 / 11	WB2	800	For.	Low	Low		
6 / 12	EB	130	For.	Low	Low		
Step	Well	Temp (°C)	Mixing (M)	Mixing Speed (RPM)	Collect (M)	Vapor (M)	Pause
1	3/9	-	0.5	3000	0.5	0	Off
2	2/8	-	0.5	3000	0	0	Off
3	1/7	70	7	3000	0	0	Off
4	2/8	-	0	500	0.5	0	Off

Step	Well	Temp (°C)	Mixing (M)	Mixing Speed (RPM)	Collect (M)	Vapor (M)	Pause
5	1/7	OFF	8	3000	3	0	Off
6	2/8	-	2	3000	1	0	Off
7	3/9	-	1	3000	0.5	0	Off
8	4/10	-	1	3000	3	0	Off
9	5/11	-	1	3000	3	8	Off

#### 13. Result

7 different samples (blood stain, dried blood spot, semen stain, chewing gum, nail, hair and cigarette butts) were purified by TANBead® nucleic acid extraction kit. Human GAPDH expression were detected by qPCR.



# 14. 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 acids 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 acids 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

# 15. Explanation of Symbols

***	Manufacturer	[]i	Consult instructions for use
15°C-	Temperature limit	Σ	Contains sufficient for test
REF	Catalogue number	$\triangle$	Caution
LOT	Batch code	NON	Non-sterile
$\otimes$	Do not re-use	*	Keep away from sunlight
سا	Date of manufacture	8	Use-by date
RUO	For research use only		