MagBeads FastDNA® Kit

Ready-to-Use for MPure-32™ aNAP System



Cat. No.: 117033600 (96 PREPS)

Quick-Start Manual

Revision 3.0 Dec 2023

Notes before starting:

- Add 2.5 mL Protease Dissolve Buffer to the Proteinase K, and store at -20~8°C after dissolve.
- Add 1.25 mL Protease Dissolve Buffer to the RNase A, and store at -20~8°C after dissolve.

Automation Extraction

A. Solid tissue (1-20mg)

- 1. Cut ~20 mg tissue into small pieces and transfer into a new 1.5 mL microcentrifuge tube. Add 200 μ L Buffer ATL and 20 μ L Proteinase K and incubate with shaker at 55 °C for 30 to 180 mins.
- 2. (Optional) Add 10 µL RNase A into the lysate and incubate at room temperature for 10 mins.
- 3. Add 200 µL Buffer AL into the samples, mix thoroughly by vortexing to yield homogenous solution.
- 4. Incubate at 70°C for 10 mins.

B. Anticoagulated blood or Plasma (200 µL)

- 1. Transfer 20 μL Proteinase K to a new 1.5 mL microcentrifuge tube.
- 2. Add 200 µL whole blood, plasma, or other body fluids into the tube. Shake to mix for 5 seconds.
- 3. Add 200 μ L Buffer AL into the samples. Invert the tube for 3 5 times, and vortex at maximum speed for 10s. Incubate at 70°C for 10 mins.

C. Saliva sample

- 1. Add 20 μL Proteinase K and 10 μL RNase A into a 1.5 mL microcentrifuge tube.
- 2. Transfer 450 µL saliva to the tube and shake to mix for 5 s.
- 3. Incubate at 55°C for 30 mins.

D. Culture cells

- 1. Collect cells ($<2 \times 10^6$) by centrifuging at 2,000 x g for 5 mins. Remove the supernatant.
- 2. Add 200 μ L Buffer PBS, 20 μ L Proteinase K, and 10 μ L RNase A into the sample. Resuspend the cells by vortexing.
- 3. Add 200 µL Buffer AL and vortex for 10 seconds. Incubate the mixture at 70°C for 10 mins.

E. Semen sample

- 1. Transfer 100 μL semen to a 1.5 mL microcentrifuge tube.
- 2. Add 100 μ L Buffer ATL, 10 μ L DTT Solution (1M), and 20 μ L Proteinase K into the samples. Shake at 55°C for 30 mins.
- 3. Add 200 µL Buffer AL into the sample, then vortex to mix and incubate at 70°C for 10 mins.

F. Swab DNA extraction

- 1. Transfer the swabs to a 2.0 mL microcentrifuge tube.
- 2. Add 500 μL ATL and 20 μL Proteinase K to the swab. Shake at 55°C for 15 to 30 mins.
- 3. Transfer the supernatant into a new 2.0 mL microcentrifuge tube.

G. Blood stains/Seminal stains

- 1. Transfer 3 slices (3mm) to the 2.0 mL microcentrifuge tube.
- 2. Add 250 μ L Buffer ATL and 20 μ L Proteinase K to the sample. Shake at high speed for 30 to 60 mins at 55 °C .
- 3. Add 250 µL Buffer AL into the samples, Shake at high speed for 10 mins at 70°C.
- 4. Centrifuge at 13,000 x g for 1 min.
- 5. Transfer 400 μL of the supernatant to a new 2.0 mL microcentrifuge tube.

H. FFPE Samples

- 1. Using a scalpel, trim excess paraffin off the sample block. Transfer 1 to 3 sections (5 $10 \mu m$) into a 1.5 mL microcentrifuge tube.
- 2. Remove Paraffin by xylene or Buffer DPS (not provided).
- 3. Add 200 μ L Buffer ATL and 20 μ L Proteinase K into the sample, mix well and incubate at 56°C for 60 mins, and 90°C for 60 mins.
- Cool the sample to room temperature and add 200 μL Buffer AL. Mix well before proceeding to extraction step.

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xtraction

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- Transfer 400 µL of supernatant carefully to well #1 or #7 of the pre-filled reagent.
- Place the reagent plate on MPure-32™ aNAP System and run the assay with the program named "FastDNA" which has the following setting:

		Time (a)				
Step Well	Process	Time (s)			Miving Speed	Temp (°C)
*****		Mix	Wait	Attract	Wilking Speed	icinp (c)
#1/#7	Bind	240	0	90	Medium	RT
#2/#8	Wash 1	120	0	60	Medium	RT
#3/#9	Wash 1	90	0	60	Medium	RT
#4/#10	Wash 2	90	0	60	Medium	RT
#5/#11	Wash 2	90	0	60	Medium	RT
#5/#11	Dry	0	300	0	-	RT
#6/#12	Elute	480	0	120	Medium	55
#1/#7	Magbeads Release	60	0	0	Medium	RT
	#2/#8 #3/#9 #4/#10 #5/#11 #5/#11	#1/#7 Bind #2/#8 Wash 1 #3/#9 Wash 1 #4/#10 Wash 2 #5/#11 Wash 2 #5/#11 Dry #6/#12 Elute	Well Process Mix #1/#7 Bind 240 #2/#8 Wash 1 120 #3/#9 Wash 1 90 #4/#10 Wash 2 90 #5/#11 Wash 2 90 #5/#11 Dry 0 #6/#12 Elute 480	#1/#7 Bind 240 0 #2/#8 Wash 1 120 0 #3/#9 Wash 1 90 0 #4/#10 Wash 2 90 0 #5/#11 Wash 2 90 0 #5/#11 Dry 0 300 #6/#12 Elute 480 0	Well Process Mix Wait Attract #1/#7 Bind 240 0 90 #2/#8 Wash 1 120 0 60 #3/#9 Wash 1 90 0 60 #4/#10 Wash 2 90 0 60 #5/#11 Wash 2 90 0 60 #5/#11 Dry 0 300 0 #6/#12 Elute 480 0 120	Well Process Mix Walt Attract Mixing Speed #1/#7 Bind 240 0 90 Medium #2/#8 Wash 1 120 0 60 Medium #3/#9 Wash 1 90 0 60 Medium #4/#10 Wash 2 90 0 60 Medium #5/#11 Wash 2 90 0 60 Medium #5/#11 Dry 0 300 0 - #6/#12 Elute 480 0 120 Medium

Transfer the eluted DNA into a clean 1.5 mL microcentrifuge tube. DNA is now ready for PCR and other downstream applications. Store the purified nucleic acid at -20°C for an extended

Note: If there are still Magnetic Beads remaining in eluted DNA, please centrifuge at 14,000 x g for 3-5 mins and transfer the supernatant into a clean 1.5 mL microcentrifuge tube.

Ordering Information



Automated extraction system from low to high throughput

Catalog No.	Product Name	Throughput
07EMC043	MPure-32™ aNAP System	Up to 32 samples
07EMC044	MPure-96 ™ aNAP System	Up to 96 samples





Instruments for lysing and homogenizing environmental and biological samples

Catalog No.	Product Name
116004500	FastPrép TM Classic
116005500	FastPrep-24 [™] 5G
116010500	FastPrep-96 TM
116012500	SuperFastPrep-2™
	116004500 116005500 116010500



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