

Order Information

Model No./ Description	Specification (Tests/ Kit)	Sample Type	Sample Volume	Certificate
Da059 I DNA/RNA Purification Kit	48 tests/kit	Serum, Plasma, Throat swab, nasopharyngeal secretion, Cervical Exfoliated Cell, etc.	200ul	CE, NMPA

Component

Description	Specification	Quantity
Spin column	48/bag	1
Collection Tube	96/bag	2
Lysis Solution	11 mL/bottle	1
Inhibitor Remover	16.6mL/bottle	1
Deionized Solution	11 mL/bottle	1
Eluent	6mL/bottle	1
Protease K	2.7mL/bottle	1
Carrier RNA	155ug/tube	2



https://en.daangene.com/

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DNA/RNA Purification Kit
(Spin Column)
DA059 I



Extensive applications, classic and trustworthy.

Features

★ Economical

No need extraction instrument.

★ Safe and non-toxic

No phenol or chloroform, environmental friendly and healthy.

★ Efficient removal of contaminants

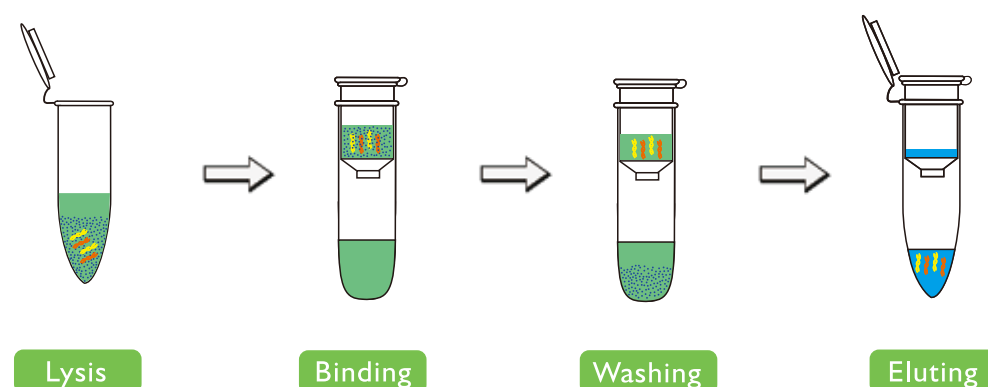
Purified DNA can be used in any downstream applications such as PCR, probe hybridization, electrophoresis, sequencing.

★ Suitable for different kinds of samples

Such as whole blood, serum, plasma, all kinds of swabs, secretion, sputum, tissue, bronchoalveolar lavage fluid.

Principle

Lysis buffer with strong protein denaturant, can quickly dissolve protein and release nucleic acid. Under the action of lysis buffer and ethanol, the released nucleic acid can be combined to the silicone membrane. Then by the action of inhibitor remover and deionized solution, the protein, inorganic salt ions and other organic impurities would be removed. Pure nucleic acid can be eluted readily with elution buffer.



Procedure



01 Lysis

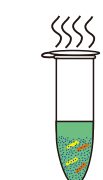
Add 50 μ L protease K
200 μ L sample
200 μ L lysis buffer(with
Carrier RNA)



Vortex, 15s



High-speed centrifugation
for 10s



Incubate 72°C, 10min

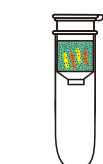


02 Binding

Add 250 μ L ethanol



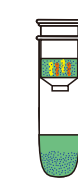
Vortex, 15s



Transfer the mixture into
the spin column



12000g, 1 min



03 Washing

Add 500 μ L inhibitor remover(with Et)
12000g, 1 min



Add 500 μ L deionized solution(with Et)
12000g, 1 min



Add 500 μ L deionized solution(with Et)
12000g, 1 min



14000g, 3 min



Open the lid
Incubate 72°C, 2 min



04 Eluting

Add 50 μ L elution buffer
Incubate RT, 1 min



14000g, 1 min



Purified Nucleic Acid