GFast Gene RNA Purification kits - Sizes

BASIC



Fig. 1: FastGene® RNA Basic (6 Preps)



Fig. 2: FastGene® RNA Basic (50 Preps)



Fig. 3: FastGene® RNA Basic (250 Preps)

Premium



Fig. 1: FastGene® RNA Premium (6 Preps)



Fig. 2: FastGene® RNA Premium (50 Preps)



Fig. 3: FastGene® RNA Premium (250 Preps)

Ordering Information

Cat. No.	Product	Content
FG-80006	FastGene® RNA Basic Kit	6 Preps
FG-80050	FastGene® RNA Basic Kit	50 Preps
FG-80250	FastGene® RNA Basic Kit	250 Preps
FG-80RL025	FastGene® RNA Lysis Buffer	25 ml
FG-80RL125	FastGene® RNA Lysis Buffer	125 ml

FG-81006	FastGene® RNA Premium Kit	6 Preps
FG-81050	FastGene® RNA Premium Kit	50 Preps
FG-81250	FastGene® RNA Premium Kit	250 Preps

GFast Gene® RNA Isolation kits - Procedure

Stop	FastGene [®] RNA Basic		FastGene [®] RNA Premium		
Step	Standard protocol	Large input protocol	Standard protocol	Large input protocol	
Sample quantity	< 5 * 10 ⁶ cultured cells <10 mg animal tissue	< 10 ⁷ cultured cells <20 mg animal tissue	< 5 * 10° cultured cells <10 mg animal tissue	< 10 ⁷ cultured cells <20 mg animal tissue	
Resuspension lysis of the cells	350 μl buffer RL (with final concentration of 20 mM DTT or TCEP)	600 μl buffer RL (with final concentration of 20 mM DTT or TCEP)	350 μl buffer RL (with final concentration of 20 mM DTT or TCEP)	600 μl buffer RL (with final concentration of 20 mM DTT or TCEP)	
Filtration of cellular debris			Transfer lysate into a FastGene® RNA filter column Centrifuge at ≥ 10,000 x g for 1 min at room temp.		
Optimize RNA binding conditions	350 μl 70% ethanol Mix thoroughly	600 μl 70% ethanol Mix thoroughly	350 μl 70% ethanol Mix thoroughly	600 μl 70% ethanol Mix thoroughly	
RNA binding	Load mix onto FastGene® RNA binding column Centrifuge at ≥ 10,000 x g 1 min		Load mix onto FastGene [®] RNA binding column Centrifuge at ≥ 10,000 x g 1 min		
Protein elimination	Add 600 μ l of buffer RW1 Centrifuge at \geq 10,000 x g 30 s		Add 600 μ l of buffer RW1 Centrifuge at \geq 10,000 x g 30 s		
Desalination	\overrightarrow{A} dd 700 µl of buffer RW2 Centrifuge at ≥ 10,000 x g 30 s		Add 700 μ l of buffer RW2 Centrifuge at \geq 10,000 x g 30 s		
Removal of RW2	Centrifuge at full speed 1 min Transfer spin column to new 1.5 ml collection tube		Centrifuge at full speed 1 min Transfer spin column to new 1.5 ml collection tube		
Elution of RNA	Add 50 μl of buffer RE to membrane center Centrifuge at ≥ 10,000 x g		Add 50 μ l of buffer RE to membrane center Centrifuge at \geq 10,000 x g 1 min		
Optimize DNase I conditions			Add 5 μ l 10 x DNase I reaction buffer to the eluate		
DNA Digestion			Add 1 μl of DNase I to the mixture Incubate for 10 min		
RNA rebinding optimization			Add 250 μl of buffer RBD to the mixture Mix thoroughly by pipetting		
RNA binding			Transfer the mix into FastGene [®] RNA mini-elute column Centrifuge at \ge 10,000 x g 1 min		
Desalination Elimination of digested DNA			Add 700 μl buffer RW2 Centrifuge at ≥ 10,000 x g 30 s Transfer spin column in new 2	Add 700 μl buffer RW2 Centrifuge at ≥ 10,000 x g 30 s Transfer spin column in new 2 ml collection tube	
Removal of RW2			Centrifuge at full speed 1 min Transfer spin column in new 1.5 ml collection tube		
Elution of RNA			Add 10-50 μ l of buffer RE to the membrane center Centrifuge at \geq 10,000 x g 1 min		

RNA