

innuPREP Plasmid Mini Kit

Protocol 2: Isolation of plasmid DNA from 5-15 ml bacterial culture

Recommended steps
before starting

 Prepare Washing Solution B according to the instruction

1.	Starting material	<i>E. coli</i> overnight culture	■ > 5-15 ml
2.	Pellet cells	\$	Centrifuge: max. speed, 8 minDiscard supernatant completely
3.	Resuspend cells		 Add 550 µl Resuspension Buffer Vortex Transfer solution to 2.0 ml tube
4.	Lysis Don't vortex!	*	 Add 550 µl Lysis Buffer Mix: invert tube 6–8 times Lysis time: ≤ 5 min
5.	Neutralization	*	 Add 750 µl Neutralization Buffer Mix: invert tube 6–8 times Centrifuge: max speed, 8 min
6.	Binding of DNA Re-use Receiver Tube	***	 Add Spin Filter to Receiver Tube Add 850 µl of clarified sample to Spin Filter 11,000 x g (~11,000 rpm): 1 min Discard filtrate and load residual sample to Spin Filter 11,000 x g (~11,000 rpm): 1 min



7. Washing

Re-use Receiver Tube





- Add 650 µl Washing Solution A
- 11,000 x g (~11,000 rpm): 1 min
- Add 750 µl Washing Solution B
- 11,000 x g (~11,000 rpm): 1 min

8. Remove Ethanol

Re-use Receiver Tube





- Discard filtrate
- Add Spin Filter to Receiver Tube
- Centrifuge: max speed, 3 min

9. Elution





- Add Spin Filter to a 1.5 ml reaction tube
- Add 100 μl Elution Buffer P
- Incubation: 3 min @ RT
- 11,000 x g (~11,000 rpm): 1 min

Order No.: Spin Filter without cap

845-KS-5040010 10 reactions 845-KS-5040050 50 reactions 845-KS-5040250 250 reactions 845-KS-5040500 500 reactions

Spin Filter with cap

845-KS-5041010 10 reactions 845-KS-5041050 50 reactions 845-KS-5041250 250 reactions 845-KS-5041500 500 reactions

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