








































FastGene™ プラスミド ミニキット

ハイコピー／ローコピー プラスミドDNAの精製

FG-90402 100 preps , FG-90502 300 preps

	ハイコピープラスミド		ローコピープラスミド
	Fastプロトコル	スタンダードプロトコル	ローコピープロトコル
集菌	 培養液 1 ~ 3 mL >10,000 rpm 1 min  上清を除去	 培養液 1 ~ 5 mL >10,000 rpm 2 min  上清を除去	 培養液 5 ~ 10 mL >10,000 rpm 2 min  上清を除去
溶菌	 200 μ L mP1 ボルテックス 200 μ L mP2 転倒混合 室温 2 min 300 μ L mP3 転倒混合	 200 μ L mP1 ボルテックス 200 μ L mP2 転倒混合 室温 2 min 300 μ L mP3 転倒混合	 400 μ L mP1 ボルテックス 400 μ L mP2 転倒混合 室温 2 min 600 μ L mP3 転倒混合
ライセート 清澄化	 13,000 rpm 2 min 	 13,000 rpm 2 min 	 13,000 rpm 3 min 
カラム結合	 上清を分注 13,000 rpm 30 sec 	 上清を分注 13,000 rpm 30 sec 	 上清750 μ Lを分注 13,000 rpm 30 sec  } $\times 2$ 回
メンブレン 洗浄	 150 μ L mP4 + 300 μ L mP5 ^{*1} 13,000 rpm 3 min 	 (400 μ L mP4 13,000 rpm 30 sec) ^{*2}  600 μ L mP5 ^{*1} 13,000 rpm 30 sec 	 400 μ L mP4 13,000 rpm 30 sec  600 μ L mP5 ^{*1} 13,000 rpm 30 sec 
メンブレン 乾燥	 13,000 rpm 2 min 	 13,000 rpm 2 min 	
溶出	 50 μ L mP6 室温2 min 13,000 rpm 2 min 	 50 μ L mP6 室温2 min 13,000 rpm 2 min 	 50 μ L mP6* (*70°Cに加熱) 室温2 min 13,000 rpm 2 min 

*1: これらのバッファーは事前調製が必要です。 *2: スキップも可。詳細は取扱説明書をご確認ください。



日本ジェネティクス株式会社

<https://n-genetics.com> info@genetics-n.co.jp






















TEL 03 (3813) 0961

FAX 03 (3813) 0962

 **FastGene™ Plasmid mini kit**

Purification of high & low copy plasmid DNA

FG-90402 100 preps , FG-90502 300 preps

	High copy plasmid		Low copy plasmid
	Fast protocol	Standard protocol	Low copy protocol
Harvest of bacteria	 ON culture 1 - 3 mL >10,000 rpm ; 1 min Remove the supernatant	 ON culture 1 - 5 mL >10,000 rpm ; 2 min Remove the supernatant	 ON culture 5 - 10 mL >10,000 rpm ; 2 min Remove the supernatant
Lysis	 *1 200 µL of mP1 : Vortexing 200 µL of mP2 : Invert the tube 2 min at room temperature 300 µL of mP3 : Invert the tube	 *1 200 µL of mP1 : Vortexing 200 µL of mP2 : Invert the tube 2 min at room temperature 300 µL of mP3 : Invert the tube	 *1 400 µL of mP1 : Vortexing 400 µL of mP2 : Invert the tube 2 min at room temperature 600 µL of mP3 : Invert the tube
Lysate clarification	 13,000 rpm ; 2 min	 13,000 rpm ; 2 min	 13,000 rpm ; 3 min
Sample loading	 Load the supernatant 13,000 rpm ; 30 sec	 Load the supernatant 13,000 rpm ; 30 sec	 Load 750 µL of the supernatant 13,000 rpm ; 30 sec } ×2 times
Membrane washing	 150 µL mP4 + 300 µL mP5 *1 13,000 rpm ; 3 min	 (400 µL of mP4 13,000 rpm ; 30 sec) *2 600 µL of mP5 *1 13,000 rpm ; 30 sec	 400 µL of mP4 13,000 rpm ; 30 sec 600 µL of mP5 *1 13,000 rpm ; 30 sec
Membrane drying	 13,000 rpm ; 2 min	 13,000 rpm ; 2 min	 13,000 rpm ; 2 min
Elution	 50 µL of mP6 2 min at room temperature 13,000 rpm ; 2 min	 50 µL of mP6 2 min at room temperature 13,000 rpm ; 2 min	 50 µL of preheated (70°C) mP6 2 min at room temperature 13,000 rpm ; 2 min

*1 : need preparation before use. *2 : can be skipped. See detailed info in the handbook.



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