





















# FastGene™ Plasmid mini kit

## 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 100preps

FG-90502 300preps

	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 1～3ml >10,000rpm 1min 上清を除去	 培養液 1～5ml >10,000rpm 2min 上清を除去	 培養液 5～10ml >10,000rpm 2min 上清を除去
溶菌	 200μl mP1 <sup>※1</sup> ボルテックス 200μl mP2 転倒混合 室温 2min 300μl mP3 転倒混合	 200μl mP1 <sup>※1</sup> ボルテックス 200μl mP2 転倒混合 室温 2min 300μl mP3 転倒混合	 400μl mP1 <sup>※1</sup> ボルテックス 400μl mP2 転倒混合 室温 2min 600μl mP3 転倒混合
ライセート 清澄化	 13,000rpm 2min	 13,000rpm 2min	 13,000rpm 3min
カラム結合	 上清を分注 13,000rpm 30sec	 上清を分注 13,000rpm 30sec	 上清750μlを分注 13,000rpm 30sec           } ×2回
メンブレン 洗浄	 150μl mP4 + 300μl mP5 <sup>※1</sup> 13,000rpm 3min	 (400μl mP4 13,000rpm 30sec) <sup>※2</sup> 600μl mP5 <sup>※1</sup> 13,000rpm 30sec	 400μl mP4 13,000rpm 30sec 600μl mP5 <sup>※1</sup> 13,000rpm 30sec
メンブレン 乾燥		 13,000rpm 2min	 13,000rpm 2min
溶出	 50μl mP6 室温2min 13,000rpm 2min	 50μl mP6 室温2min 13,000rpm 2min	 50μl mP6* (*70℃に加熱) 室温2min 13,000rpm 2min





















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

# FastGene™ Plasmid mini kit

## Purification of high & low copy plasmid DNA

FG-90402 100preps

FG-90502 300preps

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

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