

magneti**□** Quick **Plasmid Miniprep Kit**

Quick Start Guide

- Add 2.4ml of GPL-1 Buffer to hydrate the GPL-2 tube containing lyophilized pellets. Ensure the prepared GPL-2 Buffer is mixed well by vortexing.
 - Each GPL-2 tube provided contains 6 lyophilized pellets and is sufficient for 50 preps. The unused mixture should be kept at 4°C for storage and later use.

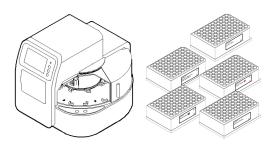


Pellet 1ml of bacterial cell sample and add to the empty Lysis plate provided. After centrifugation, resuspend bacterial pellet in 860µl of Lysis Buffer, then add 40µl of GPL-2 Buffer.

> Mix gently by pipetting up-down until solution is homogenous. Incubate for 5 mins at room temperature.

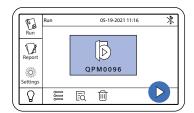


Place the Lysis, Wash #1, Wash #2, Binding Magnetic Nanoparticles, and Elution plates into purification system.



Place tip combs on Lysis plate or in dedicated tip plate, according to machine's specifications.

Run the OPM0096 Protocol.



When the protocol complete, remove the final Elution Buffer plate containing the purified DNA elution.

> Discard the tip comb and remaining plates from the purification system.

