

innuPREP Gel Extraction Kit from Analytik Jena

- Do not use more than 300 mg gel slice for one Spin Filter.
- Transfer the gel slice into a 1.5 mL or 2 mL microcentrifuge tube and add 650 μL Gel Solubilizer.
- Incubate for 10 minutes at 50 °C until the agarose gel slice is completely dissolved.
- Add 50 μL Binding Optimizer and mix the suspension by vortexing or pipetting up and down.
- Apply the sample onto the Spin Filter located in a Receiver Tube. Centrifuge at 11,000 x g for 1 minute.
- Discard the filtrate and re-use the **Receiver Tube**.
- Add 700 µL Washing Solution LS and centrifuge at 11,000 x g for 1 minute. Discard the filtrate and re-use the Receiver Tube.
- Repeat the former step.
- Centrifuge at maximum speed for 2 minutes to remove all traces of ethanol. Discard the Receiver Tube.
- Place the **Spin Filter** into an **Elution Tube**.
- Add 30 50 μL Elution Buffer (optionally prewarmed to 50 °C).
- Incubate at room temperature for 1 minute.
- Centrifuge at 11,000 x g for 1 minute.
- Note: A second elution step will increase the yield of extracted DNA:

From: Analytic Jena

