# 2010 Geno/Grinder®

## APPLICATION NOTE

Quick DNA Extraction from Rice Seed (Wet)

With kind permission of RiceTec Inc, Alvin TX

### ABSTRACT

DNA extractions can be a very time consuming and tedious process. Finding a quick method in which DNA could be extracted and used for PCR is essential. Described below is a quick "dirty" method that produces a high enough concentration of DNA that can be used for PCR.

#### Sample Extraction

Samples are prepared using a 96 well 1ml assay block. Dispense one 5/32" stainless steel bead (4mm) into each well using the Grinding ball dispenser (SPEX SamplePrep cat. # 2100). Next, add one seed to each well. Dispense extraction buffer into each well and securely cap each well. After the samples have been capped, grind them in the Geno/Grinder2000 (SPEX Sample Prep 2000-115) at 500 strokes/minute for two minutes. Centrifuge for 1 min to bring all liquid to the bottom of the assay block. Incubate the samples in about 1" of water at 95°C for 20 minutes then place them on ice for approximately 10 minutes or until samples are cool to the touch. Centrifuge again for 1 minute. Add neutralizing extraction buffer and seal the assay block with sealing film. Centrifuge the samples for 10 minutes at 3000rpm. Transfer 300µL of the supernatant to a clean 96 well plate. DNA can be further purified with clean-up kits available on the market.

#### **Yields**

The total concentrations yielded from the samples range from 3-7ng/ $\mu$ L in a final volume of 200 $\mu$ L with purities of 1.5-1.8.

#### Conclusion

The method described above is sufficient for PCR and it takes less time than the standard chloroform extraction. Total time ranges from one to two hours.



:: APPLICATION NOTE SP025: Lysing / Homogenization

- : APPARATUS: Geno/Grinder<sup>®</sup>
- : APPLICATION: DNA Extraction from Rice Seeds



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