

Dnpep Genotyping Protocol

1. PCR amplification:
 - 60 ul reaction
 - 4 ul tail DNA
 - Annealing at 60°C
2. PCR Primers:
 - Sense: GACAAACTGGGTGGAGGTCCTC
 - Anti-sense: CTTGAGTTCACGGAAGCCAGCCTG
3. Run 10 ul on agarose gel
4. Clean remaining 50 ul with QIAquick PCR purification kit
5. Sequence purified PCR fragment with sense primer