

Sample Preparation and Setup

Assumes 2x20s seed grind
in Waring blender

Note: Avoid lipid layer on
top and use caution
to avoid settled cottonseed

Note: Samples should be
transferred into blue dilution
tubes immediately and run
promptly



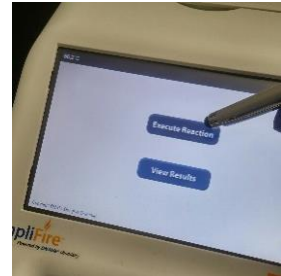
Add 600 µL of
MB4 buffer



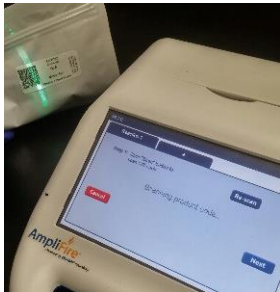
Add 2 scoops
of cottonseed



Vortex and heat
10 min at 95°C



Set up instrument:
Execute Reaction



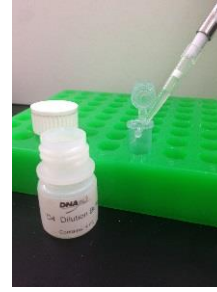
Scan barcode



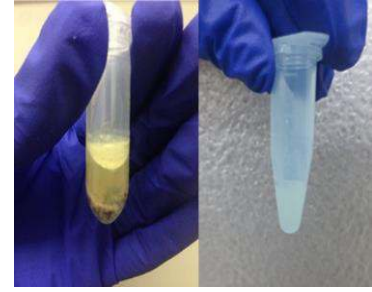
Input sample IDs



Vortex, then centri-
fuge 3 min at 10K x g



Add 150 µL D4
Dilution Buffer
to blue tube



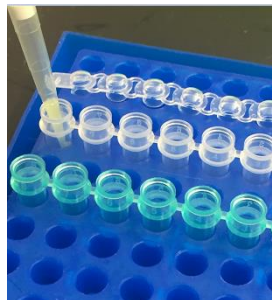
Add 50 µL supernatant to
blue tube, then vortex

Important: Use only the barcode associated with the kit

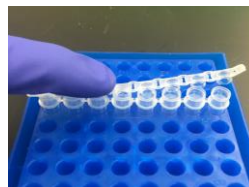
Assay Procedure



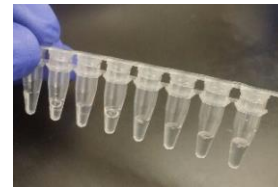
Add 25 µL sample
(blue tube) to
green 2x RB5
buffer strip



Add 50 µL of mixed
sample from green
strip to clear strip
(Master Mix)



Cap tubes tightly



Flick tube down to
remove any bubbles



Place strip
in AmpliFire
and start run



Place used assay
strip in sealable
pouch and close

Important: Change pipet tips and scoops between each sample

Important: Reaction tubes must be fully closed to prevent downstream contamination

Important: **Never** open reaction tubes post-assay