Suggestions

* Pipet. tech: Press down - go in sample - slow release up + down

* Always put master mix in b4 adding DNA → mix really well

* ~~Should have Blank Measurement~~

* Make a Negative Control - run it out on gel
  Primer/loading dye mix 22.5 µL + 2.5 µL water (water instead of DNA)
  Should have no bands on gel → if you do → have contam.

* digest PCR products w/ HaeIII - incubate for an hour