















- Contamination any DNA (from the air, from handlers, reagents or previous PCR experiments in the same apparatus) can enter the system and can be amplified giving false results on DNA found at the scene.
- Error rate every 9000 nucleotides Taq polymerase will make a mistake. However after 30 cycles this increases to 1 in 300 as these are also copied mistakes that accumulate quickly.
- Compounds such as phenolics from plants, denim blue dye and haem products act as inhibitors in PCR.
- There are limits on amplification by PCR.

After 20 cycles it slows because:

- · Reagent concentrations limit the reaction
- Enzyme denatures after a few cycles through 95°C
- So much DNA is produced after a while when the DNA strands are separated they re-attach to each other rather than annealing.



- Each cycle only lasts 120 seconds so this is a rapid technique.
- Before PCR copies of genes were made by inserting them into replicating micro-organisms, a much slower process.
- PCR can amplify even the tiniest amount of DNA from a crime scene.









