

Comparison of Taq DNA Polymerase from Vivantis and Supplier B

Amplification of MCS region with inserts 1.4kb from pTZ using M13 primer in a 50 μ l reaction.

5 μ l of PCR product is loaded per lane and electrophoresed in 1.0% TBE agarose gel.

2 different brands of Taq DNA Polymerase are tested in 2 units, 1 unit, 0.5 units and 0.25 units in a 50 μ l PCR reaction.

Experimental Date: 25th July 2016

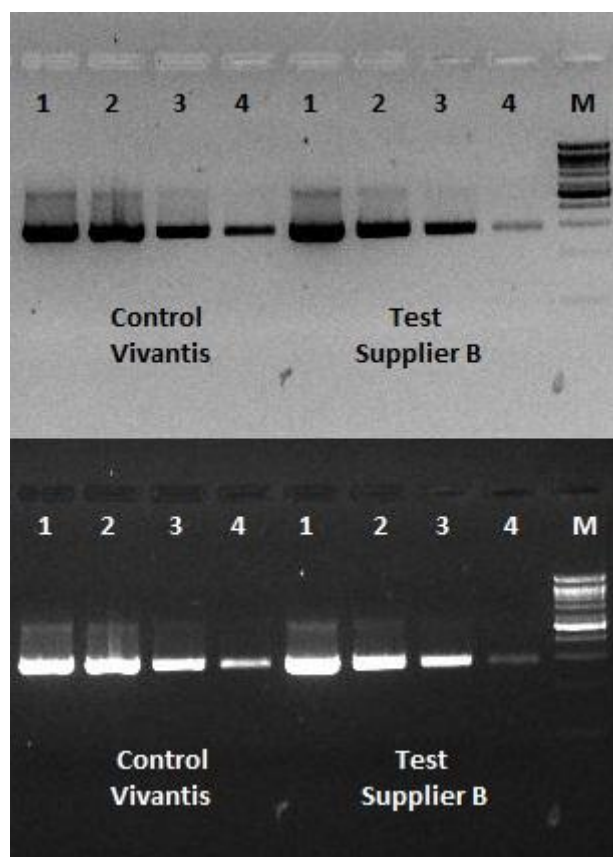


Figure 1: Amplification of pTZ template using M13 primer using different concentration of Taq polymerase. 5 μ l of PCR product is loaded per lane. Expected PCR product size amplified is 1.4kb.

Legend:

- 1: Amplification using 0.3ng/ μ l of pTZ DNA template, 0.2 μ M of primers and 2U of Taq
 - 2: Amplification using 0.3ng/ μ l of pTZ DNA template, 0.2 μ M of primers and 1U of Taq
 - 3: Amplification using 0.3ng/ μ l of pTZ DNA template, 0.2 μ M of primers and 0.5U of Taq
 - 4: Amplification using 0.3ng/ μ l of pTZ DNA template, 0.2 μ M of primers and 0.25U of Taq
- M: VC 1kb DNA ladder



Conclusion:

Both Taq DNA Polymerase are performed up to standard; the performance of Vivantis's Taq DNA Polymerase is equivalent to the performance of Supplier B's Taq DNA Polymerase.

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25th July 2016

Pairing Nature with Scientific Discoveries

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