Corrigendum

A simple method for site-directed mutagenesis using the polymerase chain reaction

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Incorrect details regarding PCR conditions were supplied for the Materials and Methods section of this paper. The correct details are published below.

**PCR conditions**

Template DNA (10 fmol) and primer sets (1 μM each) were incubated in a Perkin Elmer Cetus Thermal Cycler in 100 μl reaction volumes containing 10 mM Tris-HCl pH 8.3, 50 mM KCl, 2.5 mM MgCl₂, 10 μg gelatin, 0.2 mM of each of NTP and 2 units Taq polymerase (5).