



# Corrigendum: Diversity of reductive dehalogenase genes from environmental samples and enrichment cultures identified with degenerate primer PCR screens

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## A corrigendum on

Diversity of reductive dehalogenase genes from environmental samples and enrichment cultures identified with degenerate primer PCR screens

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The PCR amplification conditions were inaccurate as originally published, omitting the number of cycles for the final amplification step—a total of 25 cycles should be conducted when annealing at 50°C.

The corrected methods are included below.

The following conditions were subsequently applied to all described samples: an initial denaturation at 95°C for 5 min, three cycles of denaturation at 95°C for 30 s, primer annealing at 38°C for 30 s, and elongation at 72°C for 90 s, three cycles of denaturation at 95°C for 30 s, primer annealing at 45°C for 30 s, and elongation at 72°C for 90 s, and 25 cycles of denaturation at 95°C for 30 s, primer annealing at 50°C for 30 s, and elongation at 72°C for 90 s, and a final extension at 72°C for 10 min.

**Conflict of Interest Statement:** The authors declare that the research was conducted in the absence of any

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