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Ase), 2.5 µl Of Each Of The Two Primer Stock Solu-tions (10 µmol/liter), 5 P.l C, T, A, G Nucleotide Mix (Boehringer Mannheim, Indianapolis, IN, 2 µmol/ Liter For Each Nucleotide), And 29 µl Sterile Distilled Water. Reactions Were Amplified Through 35 Cycles At The Following Parameters: One Minute At 95°C,File Size: 251KB Mar 3th, 2024

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C. For N-terminal Fusion Only Primer, Including The ATG Of The Open Reading Frame Is Optional. The Reading Frame Of The Gene Must Match The Reading Frame Through AttB1. L. The Reading Frame Of Th Feb 4th, 2024

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Primer Pairs For QPCR Should Amplify Unique Target Sequences Between 70-150 Bp Long, That Are Areas Of Low Secondary Structure (GC Content

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