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Cloning In 10 μ L Volume Using 50 Ng Linearized PXST, 50 Ng Puri-fied Fragment, 5 U T4 DNA Ligase, 1 μ L 10 \times T4 Buffer And 6 μ L dH₂O. The Mixture Was Incubated At 22 $^{\circ}$ C For 1 H And then added into 100 μ L E. Coli DH5 α Chemically Com-petent Cells, Followed By Incubation On Ice For 30 Min And Heat Feb 9th, 2024

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Molecular Cloning, Expression, And Properties Of An ...3, Respectively. The PCR Conditions, With TaKaRa LA Taq Polymerase And GC Buffer I, Were Essentially The Same As Those Described Previously (10). The Nucleotide Sequence Of The Amplified DNA Fragment (507 Bp From The 5-end Of Pld1) Was Mar 5th, 2024
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DNA was precipitated with ethanol and spooled onto a glass rod. Construction and screening of a

expression library. *S. solfataricus* genomic DNA was partially digested with *Sau3A* and fragments from 0.5 to 2.5 kb were ligated to *Bam*HI-digested and dephosphorylated *λ*ZAP Express arms (Stratagene); packaging was performed according to the ...

May 19th, 2024 Isolation and molecular cloning of cellulase gene from ... and 1 kb plus DNA ladder, Invitrogen (catalogue no. 10787-018) was used as marker to determine the size of DNA fragment. DNA recovery The amplified DNA fragment verified with gel electrophoresis was extracted using TM PureLink Quick Gel Extraction Kit from Invitrogen (catalogue no. K2100-12). Four tubes of PCR reaction (25 Jan 17th, 2024).

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AAGCAGTGGTATCAACGCAGAGTGGCCATTACGGCCGGG CDS III/3 ... Apr 22th, 2024.

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