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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 LddH 20. The Mixture Was Incubated At 22 $^{\circ}$ C For 1 H Andthenaddedinto100 μ L E. Coli DH5 α Chemically Com-petent Cells, Followed By Incubation On Ice For 30 Min And Heat 4th, 2024

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Supramolecular Chemistry, "chemistry Beyond The Molecule", Is Based On The Intermolecular Interactions, In Which The Building Blocks Are Linked Together By Non-covalent Bonds -6]. The Origin Of [1 Supramolecular Chemistry Is From The Living Biological Systems, Nucleic Acid Assembly, Phospholipid Membranes, Protein Folding, Ribosomes, And Microtubules Are The Representative Examples Of ... 2th, 2024

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Other SAMS Using Neighbour-Joining Method (Fig. 2). Based On 1000 Replicates, The Phylogenetic Tree Was Divided Into Fig 2: Unrooted Phylogenetic Tree Of Protein Sequences Of CcSAMS And Other SAMSs Constructed By The Neighbor-joining

Method. Bootstrap Majority Consensus Values O 2th, 2024

2. J. Sambrook And D. W. Russell, Molecular Cloning: A ...

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3, Respectively. The PCR Conditions, With TaKaRa LA Taq Polymerase And GC Buffer I, Were Essentially The Same As Those Described Previ-ously (10). The Nucleotide Sequence Of The Amplified DNA Fragment (507 Bp From The 5-end Of *Pld1*) Was 4th, 2024

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And Cloned Into The Same Sites Of PcDNA3.1/Hygro (Invitrogen, Carls-bad, CA), Resulting In PcDNA3.1-C4ST-1 (the Name Of C4ST-1 Was Given After The Determination Of Acceptor Specificity). The Second Gene Was Initially Identified In AA233362 And AA777237 Derived F 1th, 2024

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Gibson Cloning Procedure • Design Four PCR Primers With Overlap Between Insert/insert Or Insert/vector • Adjacent Segments Should Have Identical Sequences (A And B) • PCR Primers Should Contain 5' End That Is The Same As Adj 1th, 2024

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Proc. Natl. Acad. Sci. USA 92 (1995) With phenol, DNA was precipitated with ethanol and spooled onto a glass rod. Construction and screening of a cDNA 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 AZAP Express Arms (Stratagene); packaging was performed according to the ...
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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 3th, 2024

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Bled Using Vector NTI Express Software (Life Technologies, Carlsbad, CA) Based On The Obtained Partial Fragment And The 3'-/5'-full RACE Products. Amino Acid ... (Table 1) Were Designed Using Primer Express 3.0 To Develop A Real-time PCR Assay For Detecting Duck MAPK1 Expression. The Target 213-bp Amplicon Was 3th, 2024

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Gao Et Al. 3149 Table 1. Primers Used In This Study. Primers Name Sequence(5' To 3') SMART IVTM Oligonucleotide
AAGCAGTGGTATCAACGCAGAGTGGCCATTACGGCCGGG CDS III/3 ... 4th, 2024

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Yoshio Yazaki, And Hisamaru Hirai 1th, 2024

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ization Of Corkscrew (csw), Which Is Known To Be Maternally Required For Normal Determination Of Cell Fates At The Termini Of Drosophila Embryos [10], Showed It To Be A Member Of The Cytoplasmic Family Of PTPs [11]. Mutational Analysis Has Shown That Csw Works In Concert With D-raf To Positively Transduce The Torso Receptor Protein-tyrosine ... 2th, 2024

Modern Experimental Biochemistry, Molecular Cloning. A ...

4Boyer, R F (1986) Modern Experimental Biochemistry, Addison- Wesley Publishing Co, Reading, MA, USA -5Maniatis, T, Fritsch, E F, And Sambrook, J (1982) Molecular Cloning. A Laboratory Manual, Cold Spring Harbor Laboratory, New York, NY, USA 6Clark, John M, Switzer, R L (1977) Experimental Biochemistry, W HCited By: 20Publish Year: 1992Author: J R L Walker 4th, 2024

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