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Consensus Values O 18th, 2024

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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 27th, 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 19th, 2024

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From 0.5 To 2.5 Kb Were Ligated To BamHI- Digested And Dephosphorylated AZAPExpress Arms (Strat-Agene); 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 4th, 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 4th, 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 ... 26th, 2024

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