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## **Protocol For CDNA Synthesis And QRT-PCR CDNA Synthesis**

Mix Buffer/enzyme Components And Add To Primer/RNA Mix In A 0.2 MI Microfuge Tube. Place The Tube In The PCR Machine Programmed As Follows: 25°C, 10 Min, 50°C, 50 Min, 85°C, 5 Min. Add: 1 UI RNase H, Mix, Incubate At 37°C, 20 Min, Jan 2th, 2024

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Train, La Medicina Della Terra. Conoscere Se Stessi Con L'astrologia Page 5/9.

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1972 HOLMES ETAL. TABLE 1-Continued No. Ofstrains With Result Indicated

Obtained With Following Identification Parameters': Total Atmtdraig Taxon Or  
Parameter No. Of Automated Reading, 4 H Aut4matd R4ehding' Automated  
Reading, 24 H Manualreading, 24 H Identified Strains CI NI II CI NI II CI NI II CI NI II  
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2013 Method Material Required. Per Sample, For Normal BIOLOG: Sediment: -  
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