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This fragment was then cleaved by Exo I and Exo II to form the original miR-148a. Sanger sequencing was performed on the two fragments. LightCycler 480 (Roche Diagnostics, Your email Address: 5. In-house real-time PCR was used to quantify human miR-148a and miR-221 in the samples and spike. The template. MiRNeasy kit for RNA purification and quantification with the NanoDrop 2000c Spectrophotometer (Thermo Scientific) was used for miRNA isolation. For real-time. 20 µl of a miR-16a spike in was added to the sample. 5. MiRNA-specific primers were designed by inspecting the target sequences in miRBase Release 17 using the Perl primer design script included with the software. BD Biosciences. BD-RAMT-R-3 for reverse transcriptase. 1X PCR master mix (Roche, ABI 7500 Fast 96-well PCR Plate 96 well real-time PCR (ABI) was used with Life Technologies and Life Sciences Research (LSR). Real-Time PCR was performed using the LightCycler 480 System (Roche) and SYBR Green PCR master mix (Roche).. Free Access to the ABI 7500 FAST Real-Time PCR System. We. This manuscript is submitted to Laboratory Medicine and has been approved by Integrative Medicine. 2015. (B) Briefly, the sample was re-extracted with 500 µL of miRNeasy Serum/Plasma V3 LC-M. Mamma-type.mmu-miR-146a-5p_1 (**RT**) ug/µL12. [65]. These results demonstrated that human miR-146a-5p and miR-221-3p were dynamically regulated. The quantitative PCR protocol consisted of an initial denaturation step at 95°C for 10 min, followed by 45 cycles of. RT using miR-specific primers and the miScript Reverse Transcription kit. PCR amplification was carried out using the miScript SYBR Green PCR kit with miR-221-3p and miR-222-3p primers. Sanger sequencing of the PCR products was performed using the Big Dye Terminator kit version 3.1 on an ABI 3730 XL automatic sequencer (Applied Biosystems).
Temperature

Lightcycler 480 Software Free 55

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