Use of Plasma Therapy for Severe Fever with Thrombocytopenia Syndrome Encephalopathy

Technical Appendix

Performance of Real-Time Reverse Transcription PCR to Detect Severe Fever with Thrombocytopenia Syndrome Virus RNA

We performed real-time reverse transcription PCR (RT-PCR), as described previously (1,2), to detect severe fever with thrombocytopenia syndrome virus RNA. RNA was extracted from serum with a viral RNA extraction kit (iNtRON Biotechnology, Gyeonggi, Republic of Korea). The real-time RT-PCR for severe fever with thrombocytopenia syndrome virus was performed in 25 μ L reaction mixtures containing 5 μ L of RNA template. We used a set of 2 primers (F-GGGTCCCTGAAGGAGTTGTAAA, R-

TGCCTTCACCAAGACTATCAATGT) and 1 probe (TexasRed-

TTCTGTCTTGCTGGCTCCGCGC-BHQ) for S segment, and a set of 2 primers (F-AAGAAGTGGCTGTTCATCATTATTG, R-GCCTTAAGGACATTGGTGAGTA) and 1 probe (FAM-TCATCCTCCTTGGATATGCAGGCCTCA-BHQ) for M segment. All assays were conducted at the following cycling conditions: 50°C for 30 min (1 cycle); 95°C for 10 min (1 cycle); 95°C for 15 s and 60°C for 45 sec (40 cycles).

References

- Kim WY, Choi W, Park SW, Wang EB, Lee WJ, Jee Y, et al. Nosocomial transmission of severe fever with thrombocytopenia syndrome in Korea. Clin Infect Dis. 2015;60:1681–3. <u>PubMed</u> <u>http://dx.doi.org/10.1093/cid/civ128</u>
- 2. Sun Y, Liang M, Qu J, Jin C, Zhang Q, Li J, et al. Early diagnosis of novel SFTS bunyavirus infection by quantitative real-time RT-PCR assay. J Clin Virol. 2012;53:48–53. PubMed http://dx.doi.org/10.1016/j.jcv.2011.09.031