Supplementary Fig. 1. Detection of the EBF2 gene variant (rs901176 SNP) was done by Sanger sequencing methods. We described genotyping using complementary sequences. The genotyping results by TaqMan methods are shown in (A), (C), and (E). Sanger sequencing chromatograms of rs901176 SNP in (B), (D), and (F). (A) and (B) represent the results of TT genotype; the red arrow mark above (B) indicates the TT homozygote. The results of the CT genotype of rs901176 are shown in (C) and (D); the blue arrow indicates a CT heterozygote. The results of the CC genotype are shown in (E) and (F); the green arrow represents a CC homozygote. EBF2, early B cell factor 2; SNP, single nucleotide polymorphism.