

Supplementary Figure S1. *IFNG* rs1861494 SNP genotyping by ARMS-PCR method. (A) Primer sequences (two Reverse primers and one common Forward primer) designed to specifically amplify a 168 bp amplicon that discriminates both alleles of the *IFNG* rs1861494 SNP. (B) Image of an agarose gel displaying the PCR products obtained from three genotypically different individuals for the SNP under study is shown. PCR positive control: Human Growth Hormone (HGH) gene fragment (440 bp). *IFNG* rs1861494 genotypes were assessed from the presence/absence of PCR amplicon corresponding to the specific allele (A or G). (C) DNA sequencing of the amplicons obtained from three genotypically different individuals. Primers specificity of the rs1861494 SNP were confirmed by direct sequencing of the amplified *IFNG* gene fragment by Sanger method and a 100% concordance was obtained among the results obtained from ARMS-PCR and DNA sequencing.