Figure S1. CRLF1 knockdown efficiency was analyzed by RT-qPCR and western blotting. (A) RT-qPCR analysis of CRLF1 mRNA levels. Following transfection with si-CRLF1, the mRNA expression levels of CRLF1 were significantly reduced compared with si-NC groups. (B) Western blotting of CRLF1 protein levels. Representative blots and corresponding densitometric semi-quantification show a marked decrease in CRLF1 protein expression in the si-CRLF1 group. GAPDH was used as an internal loading control. Data are presented as the mean ± SD. ***P<0.001. CRLF1, cytokine receptor-like factor 1; NC, negative control; RT-qPCR, reverse transcription-quantitative PCR; si, small interfering.