Supplement Data.



Figure S1 DT-13 exhibited no obvious effect on unactivated ECs in vitro and in vivo

Figure S1 DT-13 exhibited no obvious effect on unactivated ECs *in vitro* and *in vivo*. (A) The cytotoxicity of DT-13 detected by MTT assay. After 48 h of incubation with various concentrations of DT-13, the viability of the treated cells was determined by MTT assays. No significant cytotoxic effects of DT-13 on HUVECs were observed with concentrations up to 1 μ M. (B) HUVECs were pretreated with 1 μ M DT-13 without TNF- α stimulation. No significant difference between the control and the DT-13-treated group on HUVECs adhesion was noted. Bar=20 μ m. (C) No differences were noted between the control group and DT-13-treated group regarding the number of mouse monocyte WEHI3 cells bound to the endothelium of mouse aortas. Bar=100 μ m (D) DT-13 exhibited no significant effect on the inhibition of leukocyte migration in normal mice. The data represent the mean \pm SD of three experiments.

Figure S2 DT-13 inhibited TNF-α-induced the activation and degradation of IκB-α in HUVECs.



Figure S2 DT-13 inhibited TNF- α -induced the activation (A) and degradation (B) of I κ B- α in HUVECs. HUVECs were pretreated with DT-13 (0.01, 0.1 or 1 μ M) for 1 h followed by TNF- α (10 ng/ml) exposure. The expression of phospho-I κ B- α (B) and I κ B- α (C) were detected by western blotting. The data represent the mean \pm SD of three experiments. [#]*P*<0.05, ^{##}*P*<0.01 vs. the control group; **P*<0.05, ***P*<0.01vs. the TNF- α group.

Figure S3 The over-expression of NF-KB-p65.



Figure S3 The over-expression of NF-\kappaB-p65. HUVECs were transfected with 1 µg of pNF- κ B–M98 or pcDNA for 43 h, cells were pretreated with or without DT-13 at 1 µM for 1 h before TNF- α (10 ng/ml) induction for 4 h.The expression of p65 was detected by western blotting. The data represent the mean ± SD of three experiments. ^{##}*P*<0.01 vs. corresponding groups.