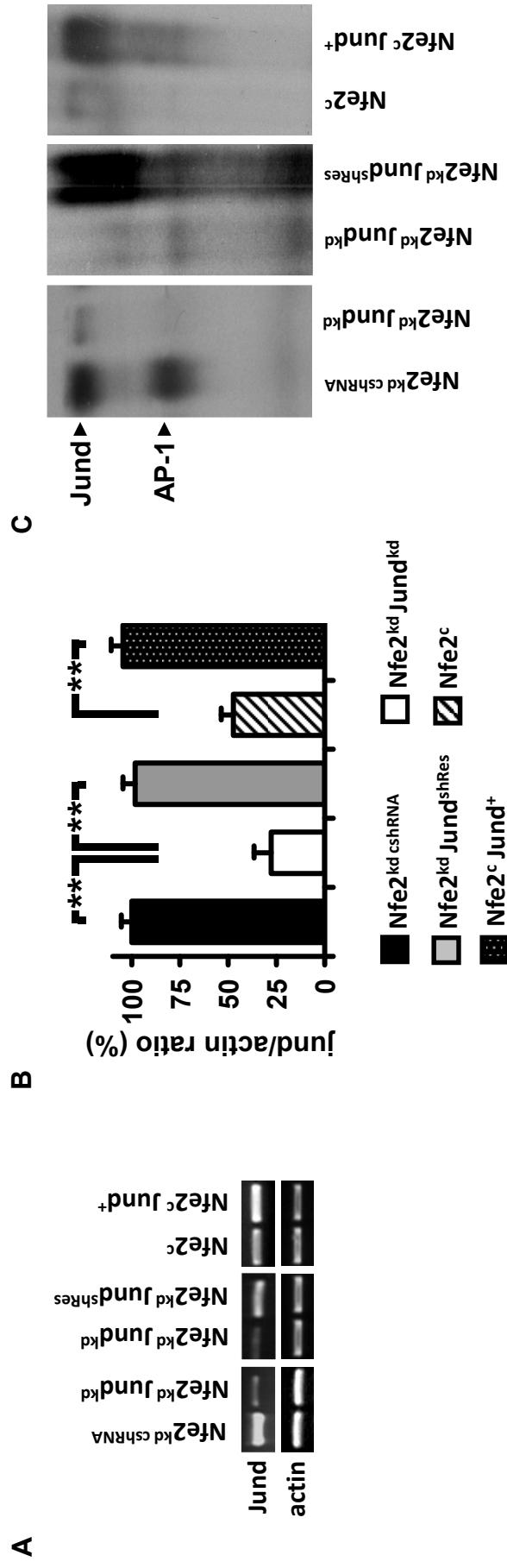


Supplementary Figure S1, related to Figure 2

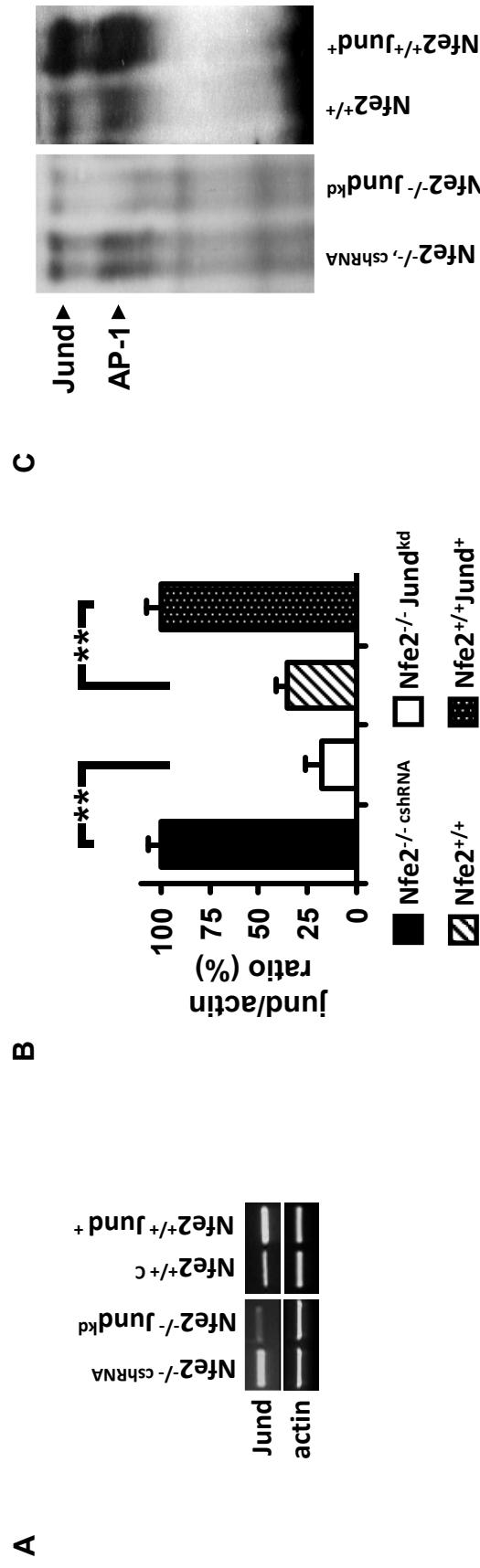


Supplementary Figure S1: Efficiency of Jund knock down and knock in vitro

Infection of Nfe2^{kd} trophoblast cells with JunD expressing lentiviral particles (Nfe2^{kd} Jund^{kd}) reduces JunD mRNA levels (a, b) and binding activity of JunD (c) when compared to Nfe2^{kd} cells transfected with non-specific shRNA (Nfe2^{kd} cshRNA). Reduction of JunD expression or JunD binding activity is not observed when Nfe2^{kd} Jund^{kd} trophoblast cells are transiently infected with a shRNA resistant JunD mutant (Nfe2^{kd} Jund^{shRes}, a-c). Infection of control trophoblast cells (Nfe2^c) with JunD overexpressing lentiviral particles (Nfe2^c Jund⁺) increases JunD expression and JunD binding activity (a-c).

Representative images of semi-quantitative RT-PCR (a), bar graph summarizing results of RT-PCR analyses (b), and representative images of supershift EMSA for JunD (c). mean value \pm s.e.m.; **P < 0.01 (T-test).

Supplementary Figure S2, related to Figure 3



Supplementary Figure S2: Efficiency of Jund knock down and Jund overexpression *in vivo*

Following transduction of Nfe2^{-/-} blastocysts with JunD specific shRNA expressing lentiviral particles (Nfe2^{-/-}, JunD kd) JunD mRNA levels (a, b) and JunD binding activity (c) are reduced in placental extracts obtained at day 14.5 p.c. Conversely, overexpression of JunD using the lentiviral approach (Nfe2^{+/+}, JunD⁺) results in enhanced JunD mRNA levels and JunD binding activity in placental extracts obtained at day 14.5 p.c. Representative images of semiquantitative RT-PCR (a), bar graph summarizing results of RT-PCR analyses (b), and representative images of supershift EMSA for JunD (c). mean value ± s.e.m.; **P < 0.01 (T-test).