SUPPLEMENTARY FIGURES



Supplementary Figure S1: The IKK inhibitor VII is the most potent IKK inhibitor. (A–D) BMMCs were pre-treated with different IKK inhibitors (as indicated) and stimulated with IL-33. Lysates were analyzed by westernblotting (A) or supernatants were collected and analyzed for IL-6 (B–D) (p < 0,001). (E) BMMCs were treated with cyclohexamide (340 µM) and stimulated with IL-3. Lysates were analyzed by westernblotting.

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Supplementary Figure S2: IKK2 mediates JNK-dependent mitogenic signaling. (A–C) $Jnk1^{-/-}$, $Jnk2^{-/-}$ and $Jnk3^{-/-}$ BMMCs were lysed and analyzed for the expression of JNK isoforms by westernblotting. (D) $Jnk1^{-/-}$ and $Jnk2^{-/-}$ BMMCs were analyzed for IL-3R α expression by flow cytometry.



Supplementary Figure S3: The cytokine production depends on *de novo* **protein-biosynthesis. (A)** BMMCs were pre-treated with SU6656 and stimulated with IL-33. Lysates were analyzed by westernblotting. (B) BMMCs were single stimulated with IL-33 or IL-33 in combination with IL-3. Total RNA was isolated, reverse transcribed and analyzed for the presence of IL-6 mRNA. (C) BMMCs were single stimulated with IL-33 or were sequentially (pre-stimulation with IL-3 for 30min: IL-3/IL-33 or pre-stimulation with IL-33: IL-33/IL-3) or simultaneously (IL-33=IL-3) stimulated. Supernatants were collected and analyzed for IL-6. (D, E) Cycloheximide-treated- (D), wt or *Myd88*⁴⁻ (E) BMMCs were single stimulated with IL-33 or IL-33 in combination with IL-3. Supernatants were collected and analyzed for IL-6.



Supplementary Figure S4: The IL-33- or IL-3/IL-33-induced cytokine production depends on Ca²⁺. (A, B) NF κ B-EGFP-MC/9 cells were pre-incubated with BAPTA-AM and stimulated with IL-33. Cells were analyzed for EGFP-production by flow cytometry (A) or supernatants were collected and analyzed for IL-6 (B). (C) BMMCs were treated with different ionomycin concentrations. Supernatants were collected and analyzed for IL-6 (D) NF κ B-EGFP-MC/9 cells were stimulated with IL-33, (D) IL-33 in combination with IL-3 (E) or were pre-treated with BAPTA-AM and stimulated with ionomycin (F). Cells were analyzed for EGFP production by flow cytometry (D–F). (G) Wt or *Orai1^{-/-}* BMMCs were stimulated with IL-33 in combination with IL-3. Supernatants were collected and analyzed for IL-6.