

Figure S1. (A) IHC staining of MLKL in control nasal mucosa or nasal polyps. Scale bars: 50 μ m. **(B)** The protein levels of IL-1 β and IL-6 in nasal mucosa homogenates were detected by ELISA (n=22-24 per group). **(C)** qPCR analysis of inflammatory cytokines mRNAs (normalized to beta-actin) in primary nasal polyps cells before and after IL-1 α (50 ng/ml), ATP (1 mM) or HMGB1 (1 μ g/ml) stimulation for 8 h (n=12). The data were analyzed by paired Student's t test.

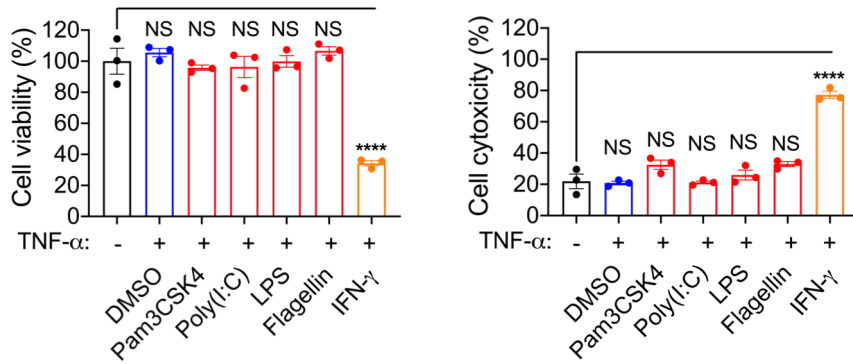
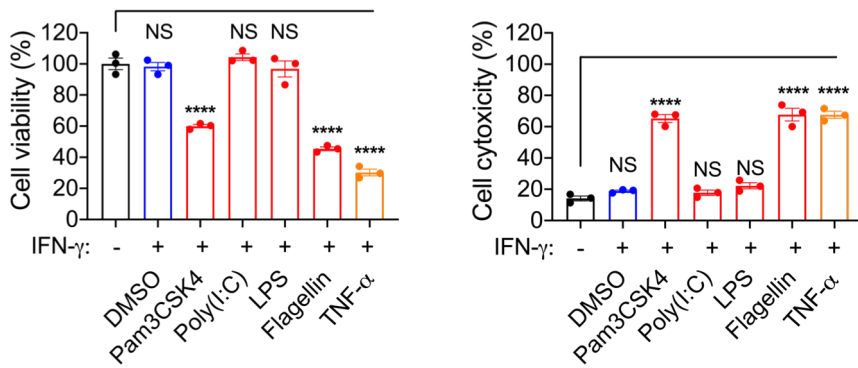
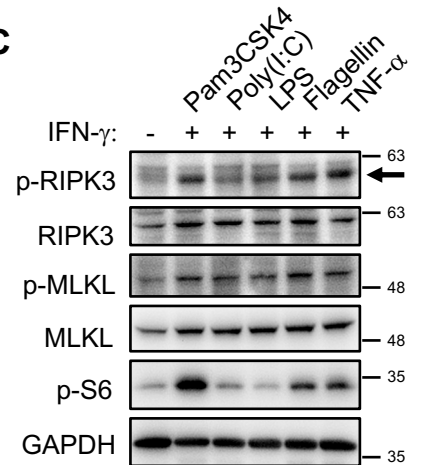
A**B****C**

Figure S2. (A-B) THP-1 cells were treated with a combination of TNF- α (50 ng/ml), IFN- γ (50 ng/ml) and Pam3CSK4 (1 μ g/ml), Poly(I:C) (1 μ g/ml), LPS (1 μ g/ml) or Flagellin (1 μ g/ml) for 48 h. Cell viability and cytotoxicity were measured by intracellular ATP content or lactate dehydrogenase (LDH) release respectively. **(C)** Immunoblots of cell lysates from THP-1 cells treated with IFN- γ (50 ng/ml) plus Pam3CSK4 (1 μ g/ml), Poly(I:C) (1 μ g/ml), LPS (1 μ g/ml), Flagellin (1 μ g/ml) or TNF- α (50 ng/ml) for 36 h. The arrow indicates p-RIPK3 band. The data were representative of 3 independent experiments and shown as mean \pm SEM. **** p <0.0001; NS, not significant; by one-way ANOVA.