SUPPLEMENTARY FIGURES

Figure S1. (a, b) Relative mRNA expression for indicated genes in dorsal skin of WT, CIKS Δ KC (a) and IL-19 KO mice (b) after IMQ or control treatment as indicated (*p < 0.05; mean ± SEM; n = 6 mice per group).

Figure S2. (a) Immunofluorescence images of sections from IMQ- or control- treated dorsal skin from WT and IL-20RB KO mice stained with DAPI (blue), anti-K5 (red) and anti-Ki67 (green). Epidermal Ki67⁺ cells were quantitated by counting fluorescent dots in 200 × 200 µm areas from different sections per mouse (n=6-10 mice per group; Scale bar 100µm). (b, c) Relative mRNA expression for indicated genes in dorsal skin (b) and skin-draining LNs (c) of IMQ- or controltreated WT and IL-20RB KO mice (n = 8-12 mice per group). (d, e) Representative flow cytometric analyses of IMQ- or control-treated skin-draining LNs cells from WT and IL-20RB KO mice analyzed for expression of markers as shown. Numbers and percentages of IL-17A⁺ γ &T cells (d) and neutrophils (e) generated from flow cytometric analyses.(n = 6-12 mice per group) (*p < 0.05, **p < 0.01; mean ± SEM).

Figure S3. (a) CCR2 expression <u>on dendritic</u> epidermal T cells (DETC, TCR $\gamma\delta^{hi}$), dermal $\gamma\delta T$ cells (CD45⁺, TCR $\gamma\delta^{intermediate}$, TCR $\nu\gamma4^+$) from IMQ-treated WT and CCR2 KO mice. (b) Representative flow cytometric analyses of IMQ- or control-treated dorsal skin cells from WT and CCR2 KO mice analyzed for expression of markers as shown. Numbers and percentages of IL-17A⁺ $\gamma\delta T$ cells generated from flow cytometric analyses. (**p < 0.01; mean ± SEM; n = 3-6 mice per group).





