Control GFP-MC injected mice have a normal appearing epidermis with no intercellular edema and a thin stratum corneum (arrow). Scale bars for panels f, g, and h are 5µm.

Supplementary Figures

Supplementary Figure 1. IL-17A Gene transfer construct. a) Schematic representation of the minicircle DNA vector carrying the IL-17A construct. Left side illustrates the entire construct whereas on the right side the by-products of arabinose induction IL-17A MC episomal DNA (top) and bacterial backbone (bottom). **b)** Agarose gel electrophoresis showing the separation of individual fragments and corresponding sizes obtained from minicircle recombination.

Supplementary Figure 2. IL-17A induces osteoclast differentiation in a RANKL-dependent manner in vivo. Serum CTX-I concentration of GFP MC and IL-17A MC injected mice at a) 3 weeks and b) 26 weeks post gene transfer. c) Map of recombinant MC vector carrying the RANKL gene. d) Serum TRAP5b concentration after gene transfer of GFP MC and dose response of RANKL MC.

Supplementary Figure 3. IL-17A exacerbates synovial inflammation and bone destruction. Histological scoring comparison of C57BL/6 mice post-gene transfer of IL-17A or GFP, immunized and challenged with bovine collagen type II showing the degree of a) pannus formation b) reactive synovium c) leucocyte infiltration, d) cartilage proliferation, e) articular cartilage damage and f) bone erosion. The thick line represents the median value of histology scores. Both hind paws were analyzed for each mouse.

Supplementary 1







