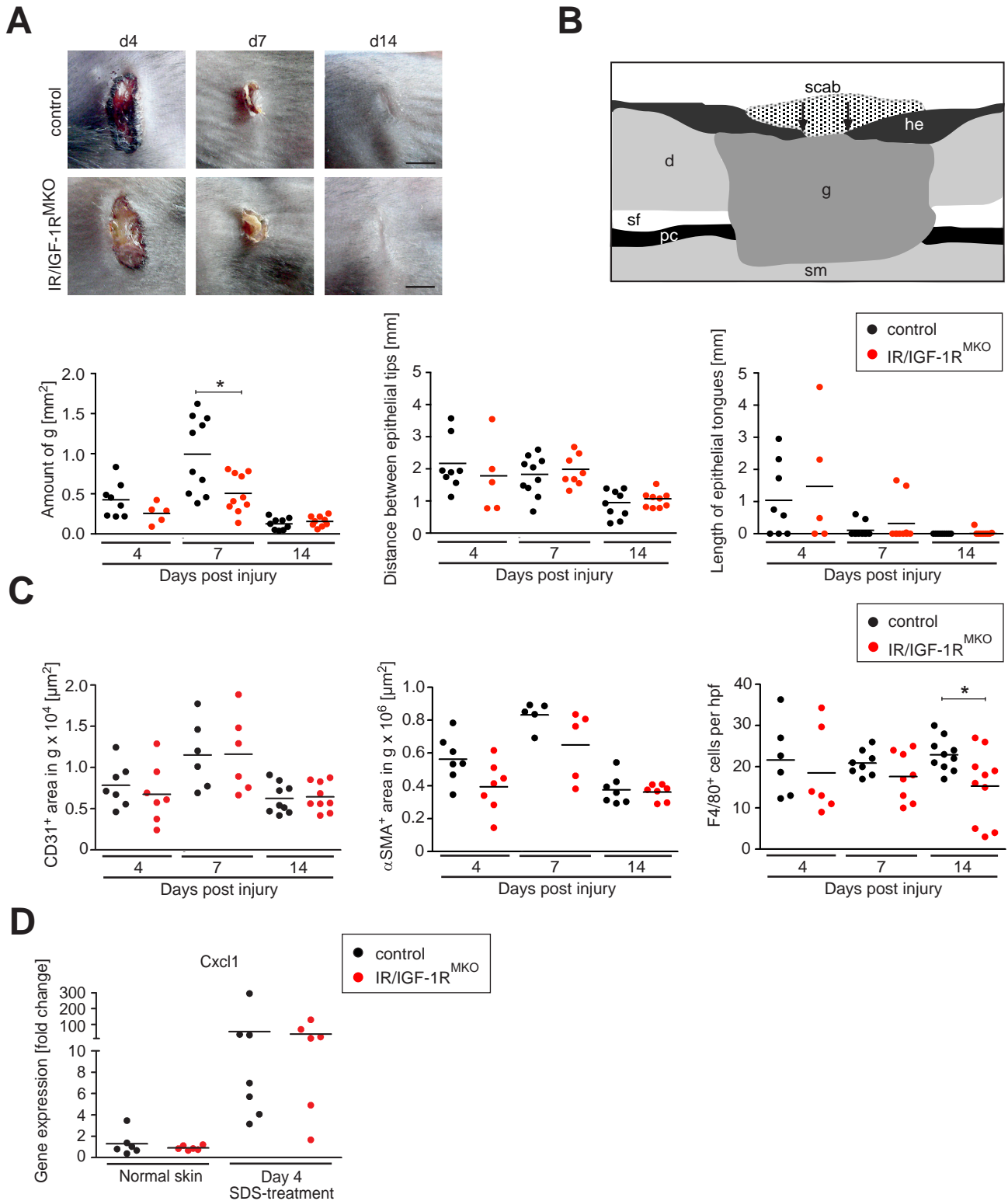
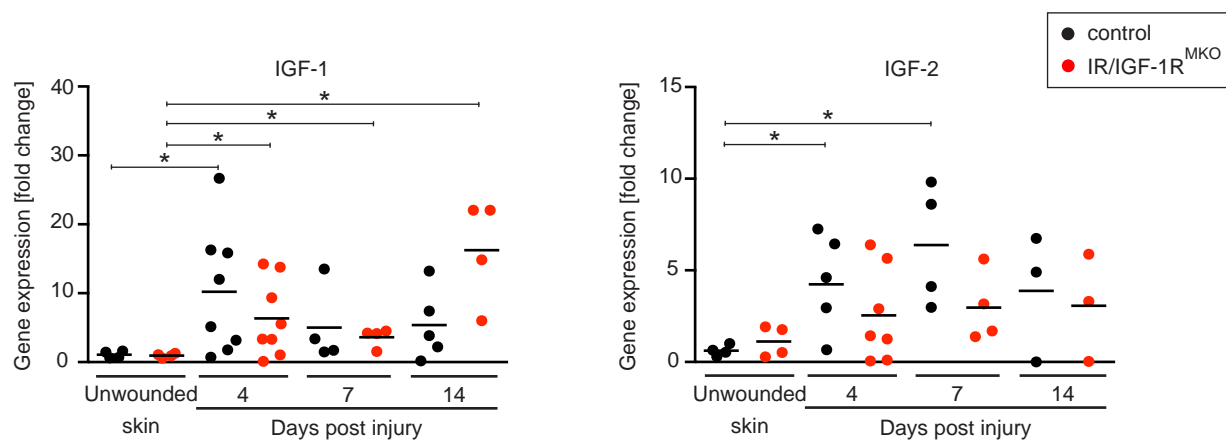


revised Supplemental Fig. 1

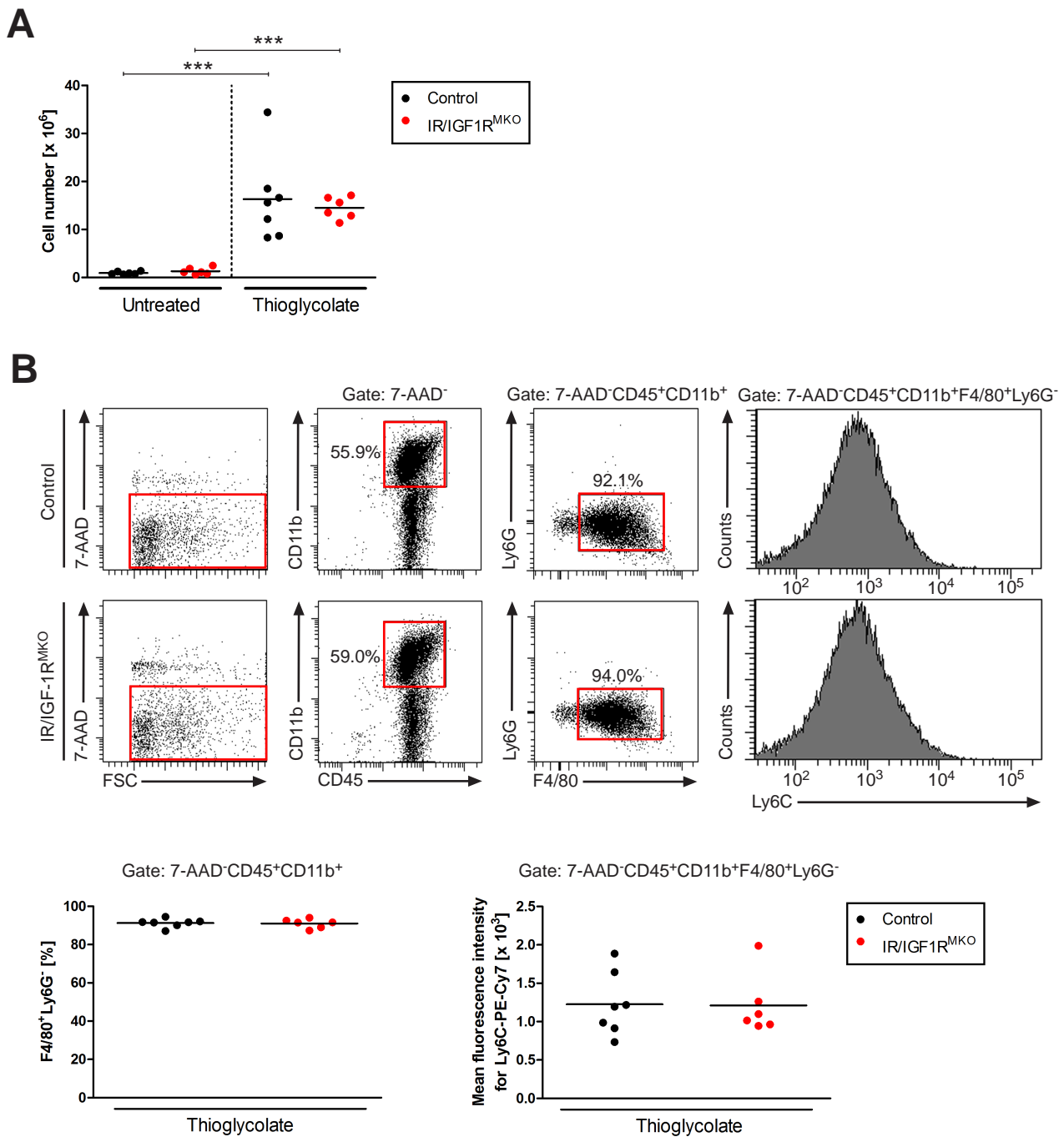


Supplemental Fig. 1: Myeloid cell-restricted IR/IGF-1R deletion has no major impact on overall wound closure kinetics. (A) Macroscopic appearance of wounds in IR/IGF-1R^{MKO} and control mice at different time points post injury; scale bar indicates 5 mm. (B) Upper right, schema illustrating skin wound histology around day 4 post injury. Morphometric analysis of H&E stained wound sections in control and IR/IGF-1R^{MKO} mice; e=epidermis, d=dermis, he=hyperproliferative epidermis, g=granulation tissue, pc=panniculus carnosus, sf=subcutaneous fat layer, sm=subcutaneous muscle layer. (C) Morphometric quantification of CD31⁺, αSMA⁺ area or F4/80⁺ cells (macrophages) within the granulation tissue at different time points post injury in IR/IGF-1R^{MKO} and control mice. (D) Relative gene expression of Cxcl1 in unwounded skin and wound tissues post injury as indicated. Each dot represents one wound; data are expressed as mean; * p-value < 0.05.

revised Supplemental Fig. 2



Supplemental Fig. 2: Expression of IGF-1 and IGF-2 is induced during skin wound healing. Relative gene expression of IGF-1 and IGF-2 in unwounded skin and wound tissues post injury as indicated; each dot represents one wound; * p-value <0.05.



Supplemental Fig. 3: IR/IGF-1R deficiency does not impact recruitment of macrophages into the peritoneal cavity after thioglycolate challenge. (A) Numbers of peritoneal cells in untreated IR/IGF-1R^{MKO} and control mice and 4 days after thioglycolate injection. (B) Upper panel, representative FACS analysis of peritoneal cells isolated from IR/IGF-1R^{MKO} and control mice 4 days after thioglycolate injection. 7-AAD⁻ cells were gated and analyzed for expression of CD45 and CD11b; 7-AAD⁻CD45⁺CD11b⁺ cells were gated and analyzed for expression of F4/80 and Ly6G; 7-AAD⁻CD45⁺CD11b⁺F4/80⁺Ly6G⁻ cells were gated and analyzed for expression of Ly6C. Lower panel, quantification of F4/80⁺Ly6G⁻ macrophages in the gate of CD45⁺CD11b⁺ cells and quantification of the mean fluorescence intensity of Ly6C-PE-Cy7 in the gate of 7-AAD⁻CD45⁺CD11b⁺F4/80⁺Ly6G⁻ cells. PE-Cy7, phycoerythrin-cyanine7; each dot represents one mouse; data are expressed as mean; *** p-value <0.001

Supplemental Table I: Primer pairs used in this study

| Target gene | Accession number | Primer sequence |
|----------------|------------------|---|
| IL-1 β | NM_008361.3 | forward: 5' - GGACCCCAAAAGATGAAGGGCTGC reverse: 5' - GCTCTTGTTGATGTGCTGCTGCG |
| IL-6 | NM_031168.1 | forward: 5' - ACACATGTTCTCTGGGAAATC reverse: 5' - AAGTGCATCATCGTTGTTTCATACA |
| iNOS | NM_010927.3 | forward: 5' - CCACCTTGGTGAAGGGACTGAGCT reverse: 5' - AGGGGCAAGCCATGTCTGAGACT |
| IL-10 | NM_010548.2 | forward: 5' - AGCCGGGAAGACAATAACTG reverse: 5' - CATTTCCGATAAGGCTTGG |
| GAPDH | NM_008084.2 | forward: 5' - CATGTTTGTGATGGGTGTGA reverse: 5' - AATGCCAAAGTTGTCATGGA |
| IGF-1 | NM_010512.4 | forward: 5' - GCGGTGCCCTTGAGACTCC reverse: 5' - CTGCGCATCCTCCCAAGTGC |
| IGF-2 | NM_010514.3 | forward: 5' - CACGGGGGAGCCTCTTCGGA reverse: 5' - TGGGGCAAGGGGAACAGCCT |
| TNF α | NM_013693 | forward: 5' - GACCCTCACACTCAGATCATCTTCT reverse: 5' - CCTCCACTTGGTGGTTTGCT |
| MIP1 α | NM_011337.2 | forward: 5' - CATATGGAGCTGACACCCCG reverse: 5' - CAGGAAAATGACACCTGGCTG |
| MCP1 | NM_011333.3 | forward: 5' - TCCACGTGTTGGCTCAGCCAG reverse: 5' - CCAGCCTACTCATTGGGATCATCTT |
| MCP3 | NM_013654.3 | forward: 5' - GCCCAATGCATCCACATGCTGCT reverse: 5' - TTCAGCGCAGACTTCCATGCCC |
| Relm- α | NM_020509 | forward: 5' - TATGAACAGATGGGCCTCCT reverse: 5' - GGCAGTTGCAAGTATCTCCAC |