

Supplementary Figure 1.

a, Density plots of Cell Ranger outputs for 3 dpi MaFIA wound cells treated with vehicle or AP20187.

b, Expression of key marker genes used to annotate all cell types in the MaFIA wound.

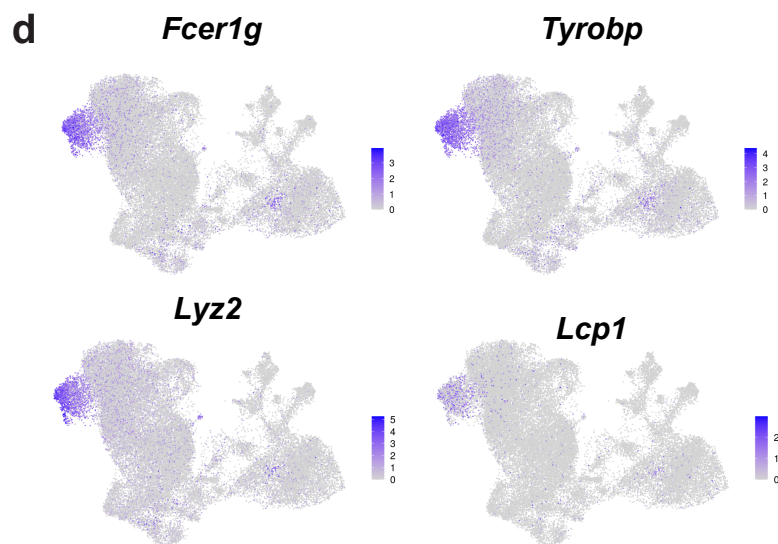
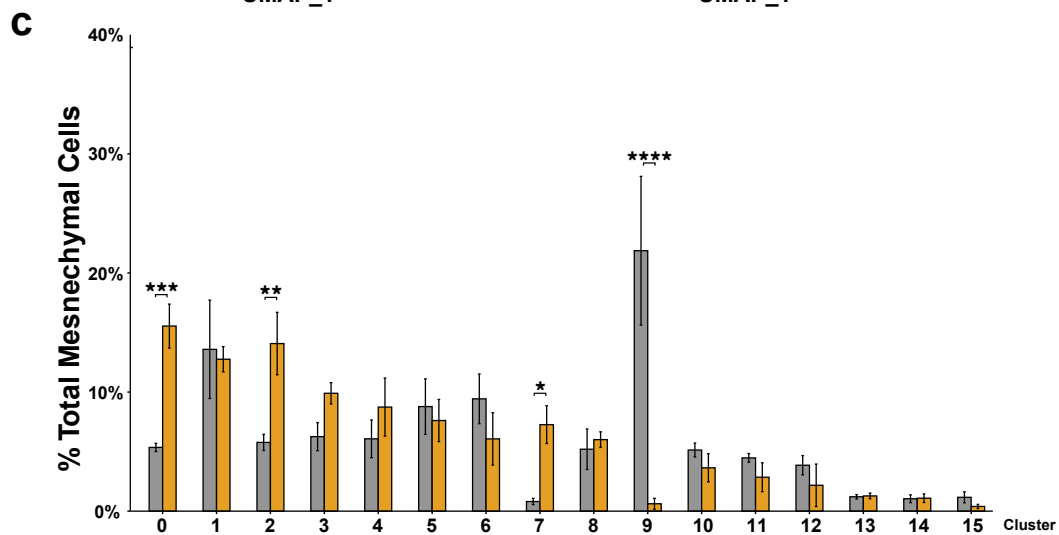
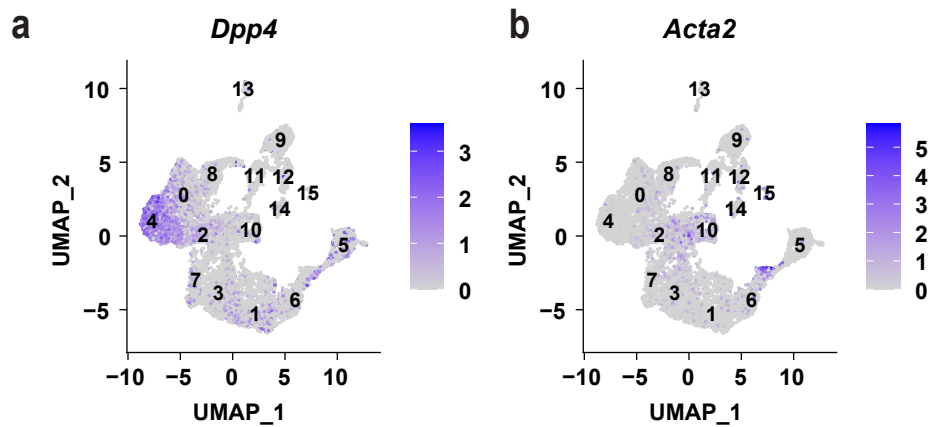
c, Expression of the key marker genes recovered by differential expression analysis for all cell types in the MaFIA wound.

d, Split UMAP of wounds treated with vehicle and AP20187 (n = 3 per condition); macrophage population highlighted in yellow.

e, Quantitative analysis of macrophage percentage over total cells (2-way repeated measures ANOVA).

f, Quantitative analysis of the percentage of each cell type over total cells (excluding macrophages) in the wounds treated with vehicle and AP20187 (2-way repeated measures ANOVA).

*P < 0.05, **P < 0.01, ***P < 0.001, ****P < 0.0001. Data are presented as mean ± s.d.



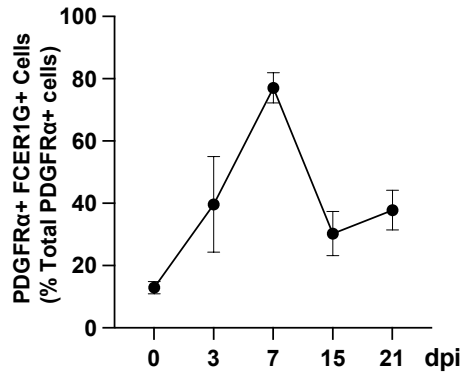
Supplementary Figure 2.

a, Expression of *Dpp4* in mesenchymal subclusters.

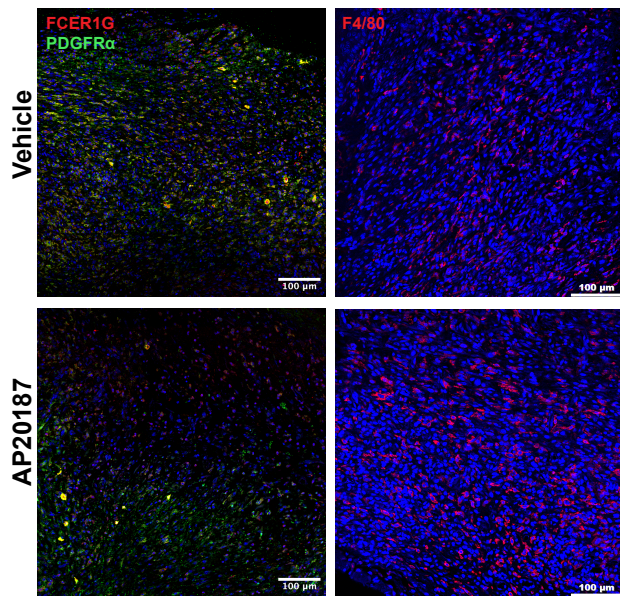
b, Expression of *Acta2* in mesenchymal subclusters.

c, Quantitative analysis of the percentage of each mesenchymal cell subtype over total mesenchymal cells (2-way repeated measures ANOVA).

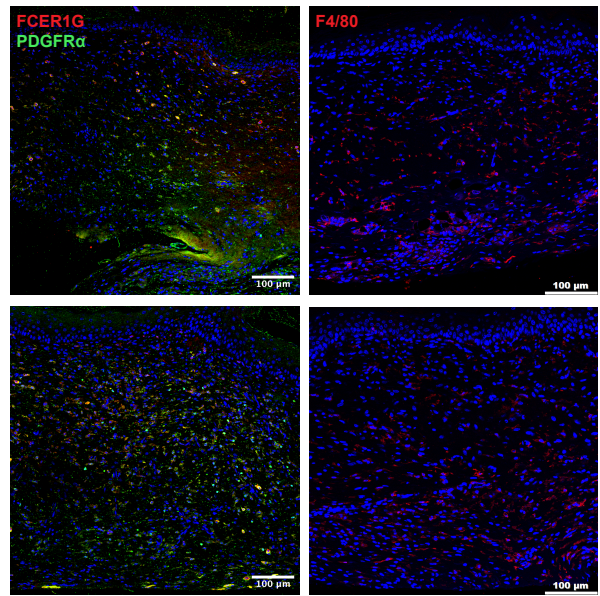
d, Expression of *Fcer1g*, *Tyrobp*, *Lyz2* and *Lcp1* in a published time-course scRNA-seq data on wound healing.

a**b**

7 dpi

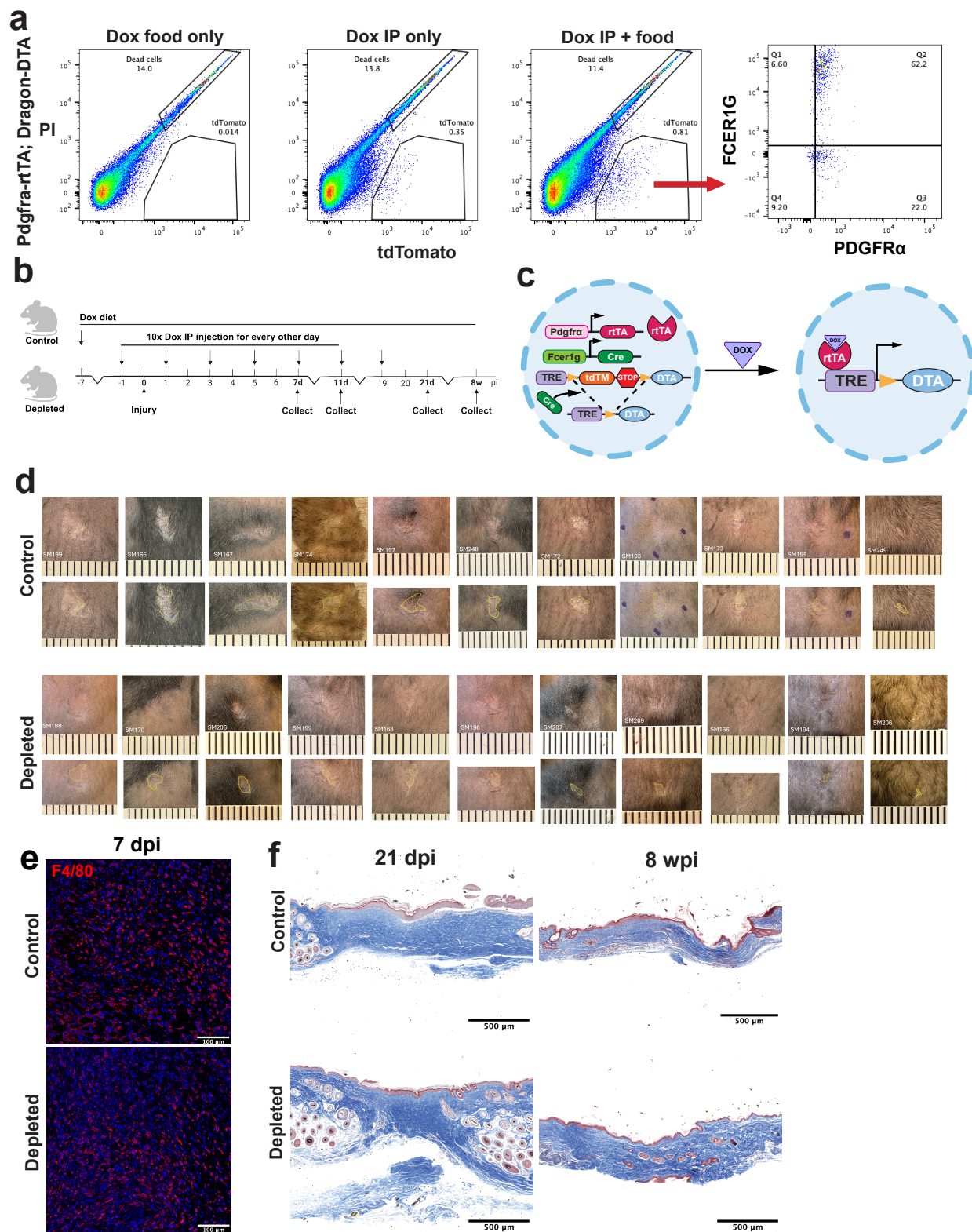
**c**

14 dpi

**Supplementary Figure 3.**

a, Percentage of FCER1G⁺ PDGFR α ⁺ cells over PDGFR α ⁺ cells over time during wound healing (n = 2 for unwounded, 15 dpi and 21 dpi; n = 3 for 3 dpi and 7 dpi).

b-c, Representative immunofluorescence images of FCER1G, PDGFR α , F4/80 staining of 7 dpi (**b**) and 14 dpi (**c**) wounds from MaFIA treated with vehicle and AP20187 (n=3 per condition for 7 dpi wounds; n=3 for 14 dpi vehicle-treated wounds; n=2 for 14 dpi AP20187-treated wounds).



Supplementary Figure 4.

a, Left: FACS analysis of PI and tdTomato expression in skin cells collected from 7dpi *Pdgfra*-rtTA; Dragon-DTA mice treated with Dox food only, or Dox IP only, or Dox food and IP; Right: FACS analysis of FCER1G and PDGFR α expression of tdTomato⁺ PI⁻ skin cells from *Pdgfra*-rtTA; Dragon-DTA mice treated with Dox food and IP.

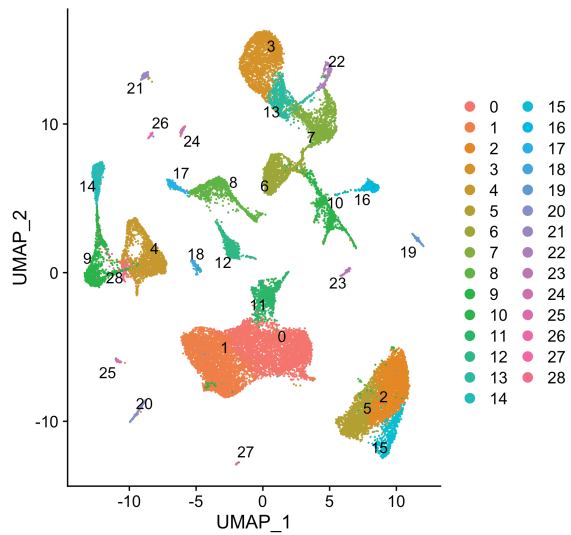
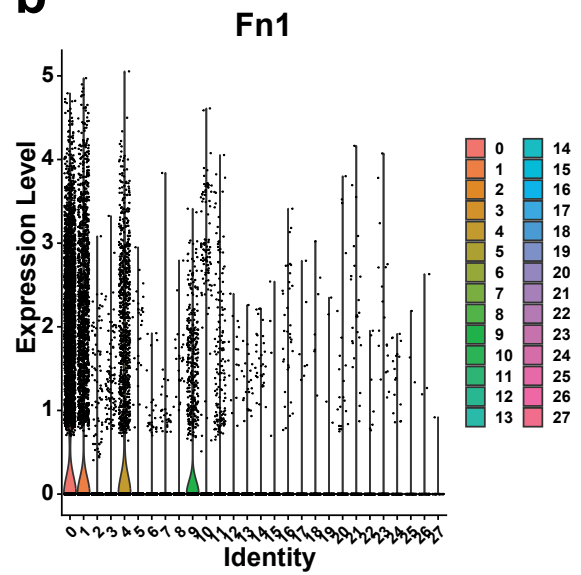
b, Schematic of experiment design.

c, Schematic of the *Pdgfra*-rtTA; *Fcer1g*Cre; Dragon-DTA mouse, which allows DTA-mediated ablation of *Fcer1g*-and *Pdgfra*-expressing cells upon Dox administration.

d, Macroscopic appearance of scars in control and depleted groups at 8wpi. Top row: raw photo; Bottom row: scars contoured for measurement.

e, Representative immunofluorescent images of 7 dpi wounds stained with F4/80 in control and depleted groups (control: n = 3, depleted: n = 4).

f, Masson's trichrome-stained sections from control and depleted groups at 21 dpi and 8 wpi (n=11 per time point).

a**b****Supplementary Figure 5.**

a, UMAP of cell clusters in MaFIA wounds at 3 dpi with *Fcer1g*-expressing population mapped back as cluster 28.

b, Violin plot showing *Fn1* expression across all clusters.

Cell type	Cluster	Markers
Basophil	21	Prss34, Mcpt8, Gata2
Dendritic/Langerhans cell	11,20	Cd74, Cd80, Cd86,
Endothelial cell 1	16	Sox17, Emcn, Pecam1
Endothelial cell 2	26	Prox1, Ccl21a, Pecam1
Epithelial cell	3,6,7,13,22	Krt15, Krt17, Sox9
Macrophage	0,1	Pf4, Adgre1, Itgam
Mast cell	18	Kit, Fcer1a, Cpa3
Melanocyte	23	Pax3, Dct, Tyrp1
Mesenchymal cell	4,9,14,24,25	Col1a1, Col1a2, Pdgfra, Col5a1, Loxl1, Lum, Fbln1, Cd34
Mitotic cell	10	Ube2c, Pclaf, Birc5
Muscle	19	Acta1, Trdn, Tnni2
Natural killer cell	17	Klrb1c, Gimap4, Ncr1
Neutrophil	2,5,15	Mmp8, Retnlg, Ly6g
T cell	8,12	Cd3g, Cd3e, Trdc

Supplementary Table 1. Markers used to annotate cell types in 3 dpi MaFIA wound single-cell data.