Hepatospecific ablation of p38α MAPK governs liver regeneration through modulation of inflammatory response to CCl₄-induced acute injury

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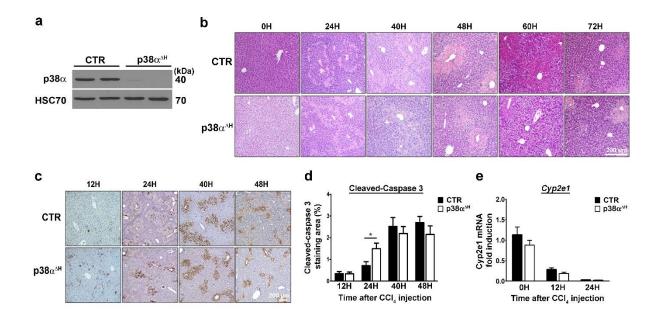
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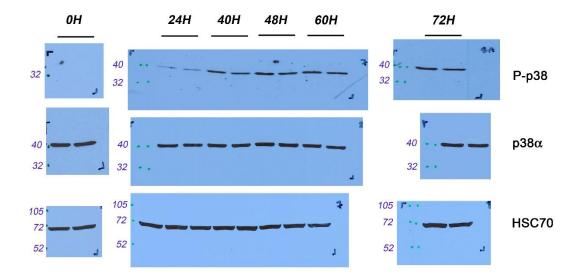
Supplementary figures



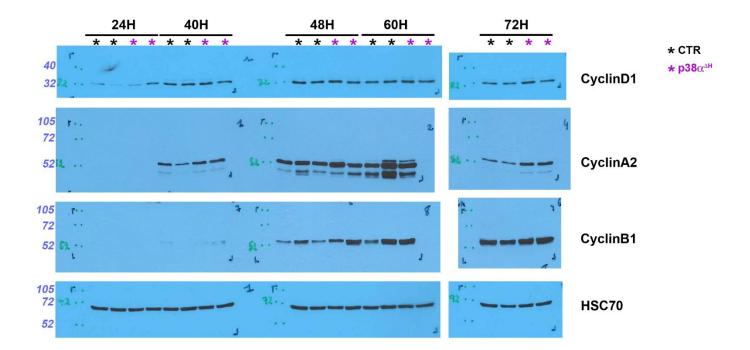
Supplementary Figure 1. Apoptosis and CCl₄ bioactivation are not impaired in p38 $\alpha^{\Delta H}$ mice during acute liver injury.

- a) Immunoblot analysis of p38 α isoform in CTR and p38 $\alpha^{\Delta H}$ liver samples in steady state conditions. HSC70 served as a loading control. Lanes show samples from independent biological replicates. The displayed figure was cropped and the original images are part of the supplementary data.
- **b)** Magnification of necrotic area related to Figure 2b, in CTR and $p38\alpha^{\Delta H}$ livers at different times after CCl₄ exposure.
- c) Representative Cleaved-Caspase 3 immunochemistry of CTR and p38 $\alpha^{\Delta H}$ liver sections at indicated time points after CCl₄ injection.
- d) Quantitative analysis of Cleaved-Caspase 3 labeled CTR and p38 $\alpha^{\Delta H}$ liver sections (percentage of brown area). Data represent the mean \pm SEM ($n \ge 4$ per group). *p<0.05(two-tailed t-test).

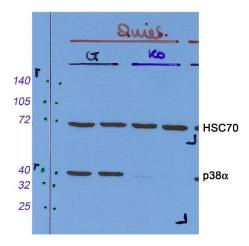
e) Relative mRNA level of Cyp2e1 measured by quantitative PCR in CTR and p38 $\alpha^{\Delta H}$ liver samples before (0H) and after CCl₄ injection (12H-24H). Gene expression levels were normalized to the abundance of 18s mRNA for each sample. Data represent the mean \pm SEM ($n \ge 6$ per group).



Supplementary Figure 2. Original pictures from Western blot membranes related to Figure 1b.



Supplementary Figure 3. Original pictures from Western blot membranes related to Figure 3e.



Supplementary Figure 4. Original pictures from Western blot membranes related to Figure Supplementary 1a.