

Supplementary 1

The in vitro experiment results showed that iron overload significantly reduced cell viability, while EGCG could recover it. As important indicators of ferroptosis, the levels of Fe^{2+} , ROS, and C11-BODIPY were significantly increased under iron overload, while EGCG significantly decreased the C11-BODIPY level under FAC treatment. In addition, we used RSL3 to induce ferroptosis and found that both 50 and 100 μM of EGCG could significantly reduce the C11-BODIPY level. In vitro experiments also proved that EGCG could alleviate iron overload and RSL3-induced ferroptosis.

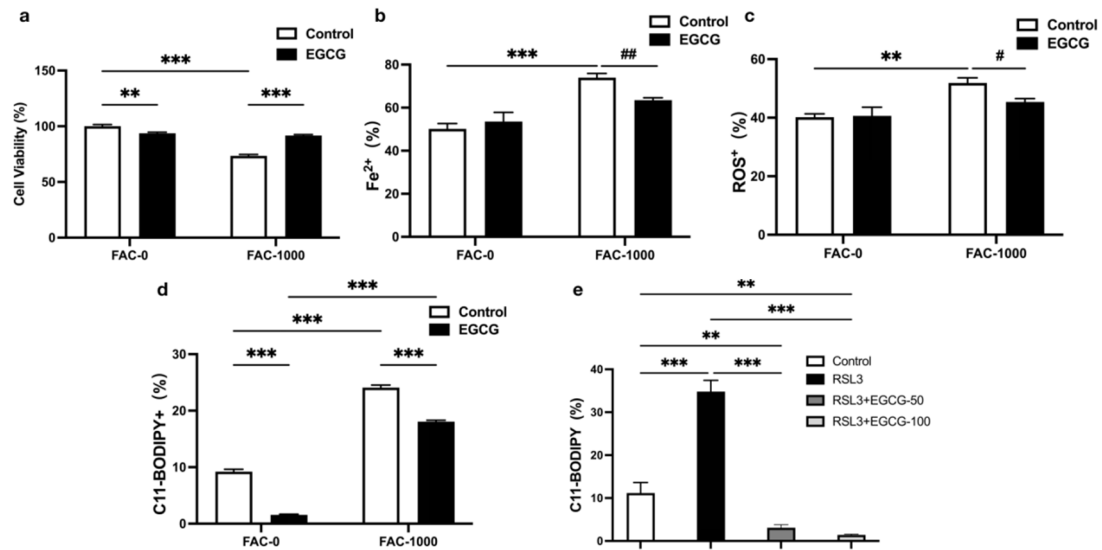


Figure S1. Iron overload induces ferroptosis, resulting in hepatocyte injury. (a–d) Cell viability, ferrous, ROS and C11-BODIPY level in the cells treated with FAC (n = 6) (Two-way ANOVA, $p < 0.05$ (*), $p < 0.01$ (**), $p < 0.001$ (***)). *t*-test: $p < 0.05$ (#), $p < 0.01$ (##)). (e) C11-BODIPY level in cells with RSL3 (n = 3) (One-way ANOVA, $p < 0.05$ (*), $p < 0.01$ (**), $p < 0.001$ (***)).