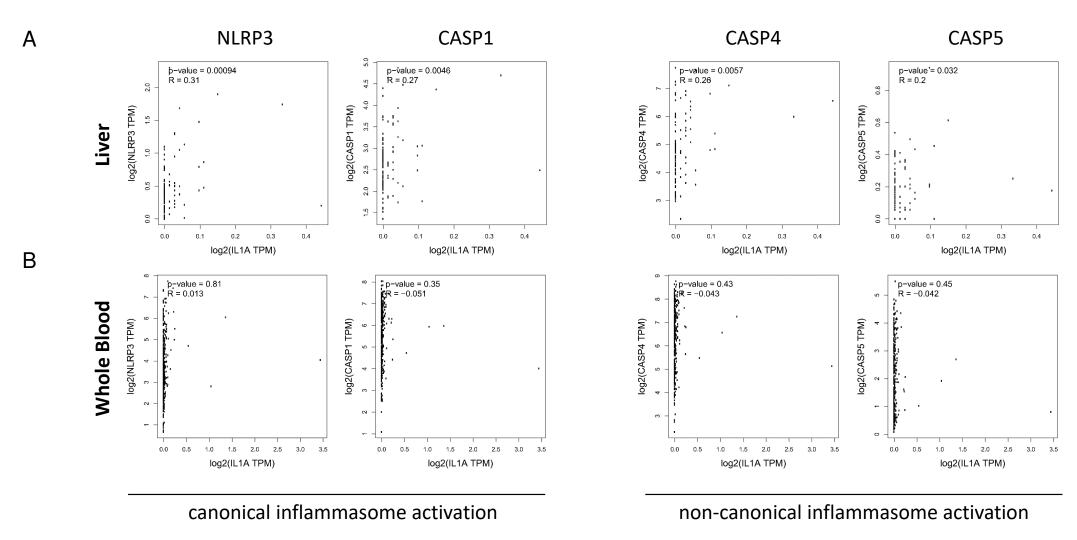
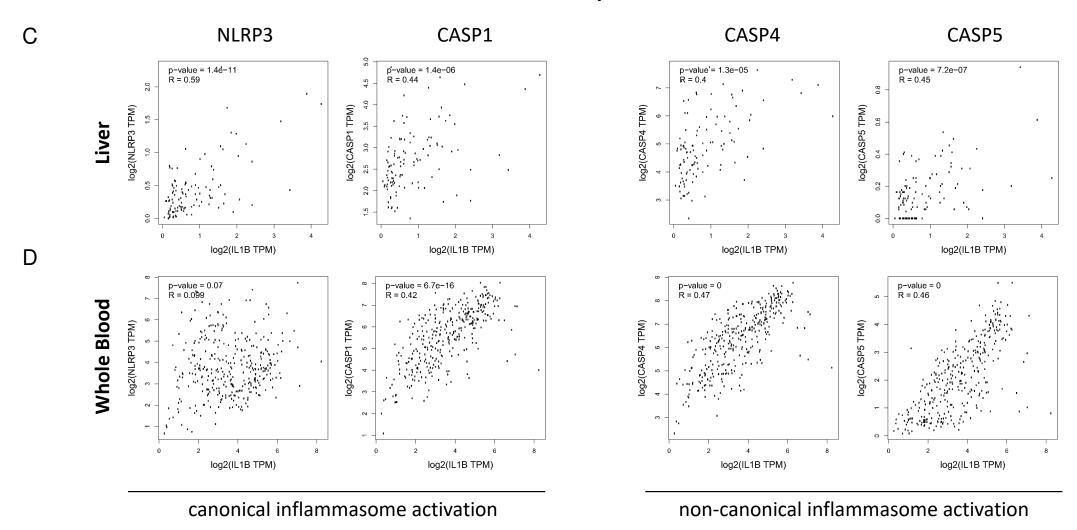
Supplementary Figure 1

IL-1α



Supplementary Figure 1

IL-1β



Supplementary Material 1.

- 2 Western blot of human liver. Total protein was harvested from human liver samples.
- 3 SDS-PAGE gels and nitrocellulose membranes were used as described previously
- 4 (Klein S. et al., 2012). Western blotting was performed as described previously in
- 5 (Uschner FE. et al., 2020). Membranes were incubated with the respective primary
- 6 antibodies: glyceraldehyde-3-phosphate dehydrogenase (GAPDH) (Santa Cruz
- 7 Biotechnology, CA, USA, sc-166545, Lot#B1716) diluted at 1:10.000 was used as
- 8 endogenous control to confirm equal protein loading, GSDMD (ab210070,
- 9 Lot:GR3258726-3) diluted at 1:1000 and cleaved N-terminal GSDMD (ab215203,
- Lot:GR3319820-1) diluted at 1:1000. The corresponding secondary peroxidase-
- couple antibody was added at 1:3000 dilution (Cell Signalling HRP-linked antibody).
- 12 After enhanced chemiluminescence (ECL Western, 32106, Thermo Scientific) digital
- detection was evaluated using ImageJ software (version 1.48v, NIH, USA) and results
- 14 were corrected for GAPDH levels.