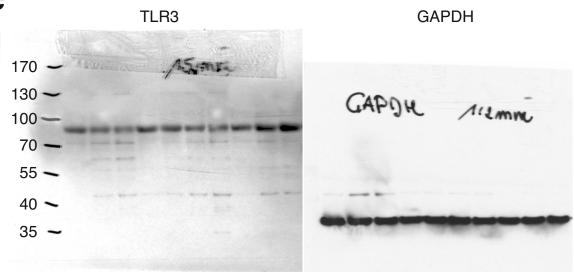
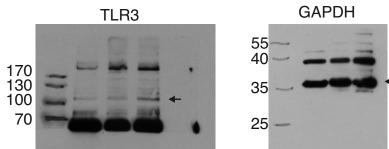
Fig.1c



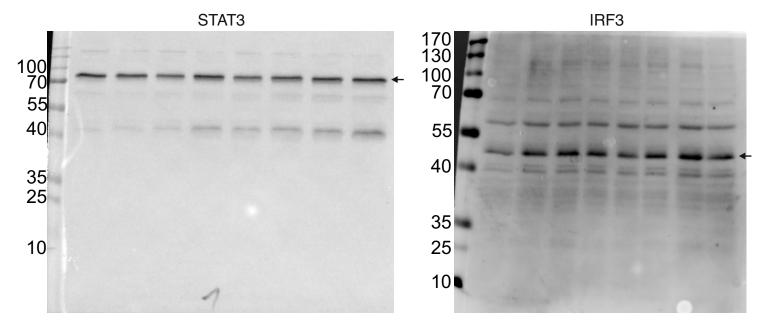
Membrane was first incubated with TLR3 antibody and later on with GAPDH antibody.



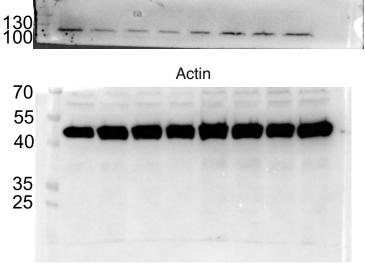


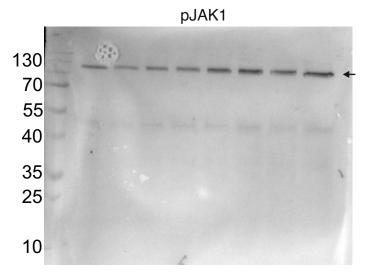
Membrane was cut between 55kDa and 70kDa. Upper part was incubated with TLR3 antibody, lower part with GAPDH antibody.

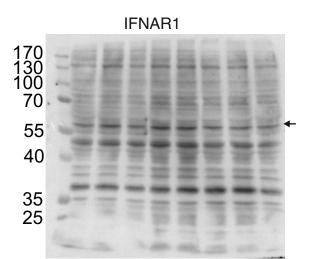
Fig.2e



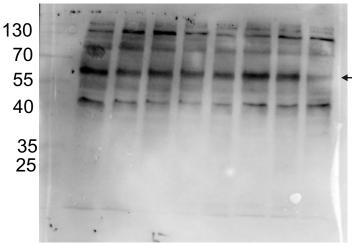




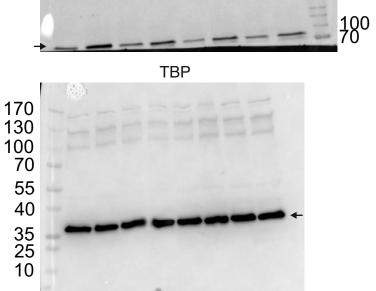




RUNX2



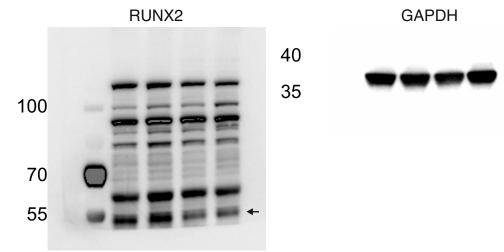




pIRF3
130
70
55
40
35
25
10

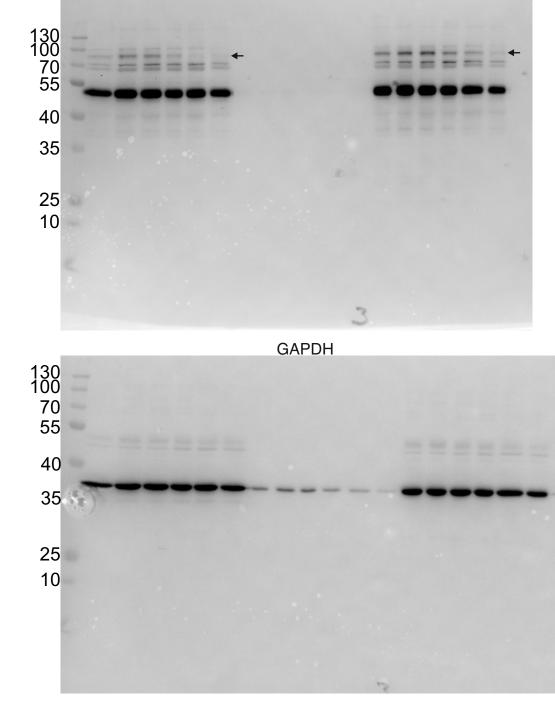
Subcellular fractionation was performed and phosphorylated proteins were detected in nuclear protein samples. TBP was used for nuclear fraction loading control.

Fig.2g



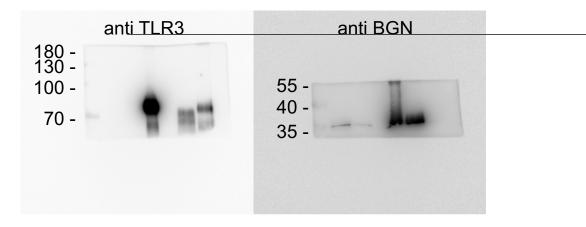




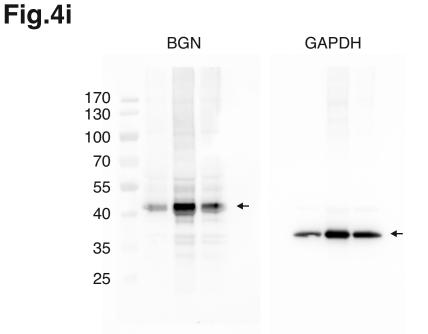


Membrane was first incubated with TLR3 antibody and later on with GAPDH antibody.

Fig.4g

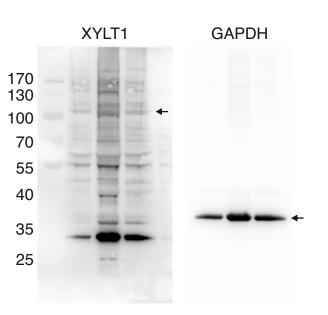


Membrane was cut between 55kDa and 70kDa. Upper part was used for detection of TLR3 ectodomain (~72kDa). Lower part was used for detection of purified SF21 BGN (~37kDa).

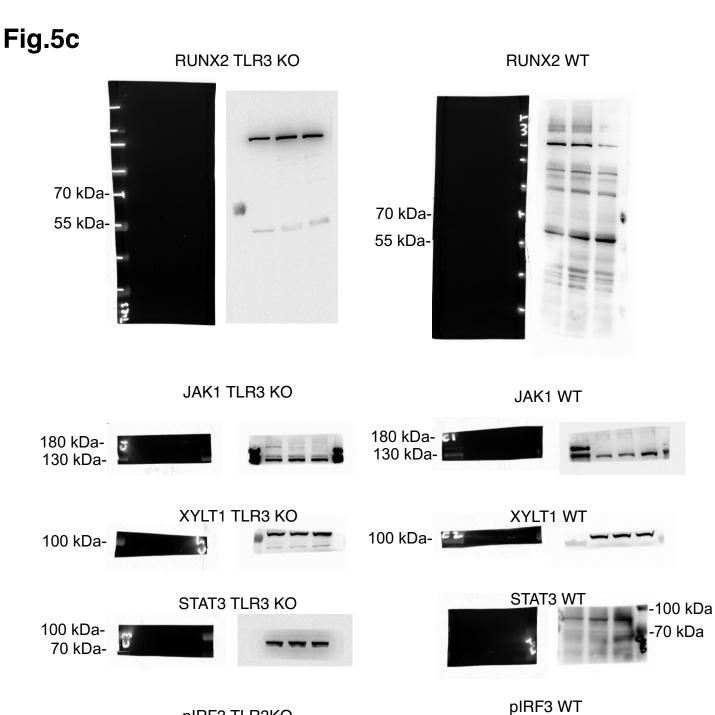


Membrane was first incubated with BGN antibody and later on with GAPDH antibody.

Fig.4j



Membrane was first incubated with XYLT1 antibody and later on with GAPDH antibody.





GAPDH TLR3 KO

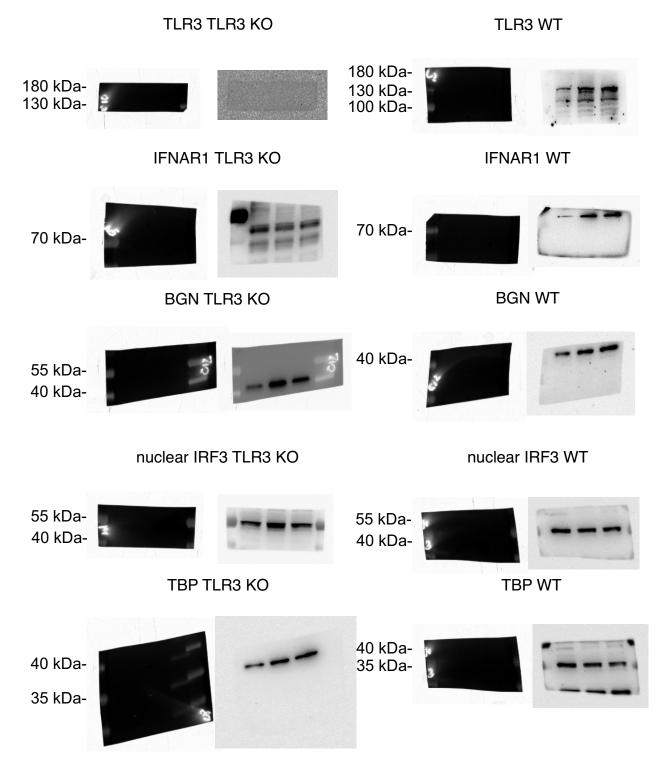
55 kDa-

40 kDa-









Subcellular fractionation was performed and IRF3 was detected as phosphorylated protein in nuclear protein samples. TBP was used for nuclear fraction loading control.

Fig.S1g

