

## Supplementary Material

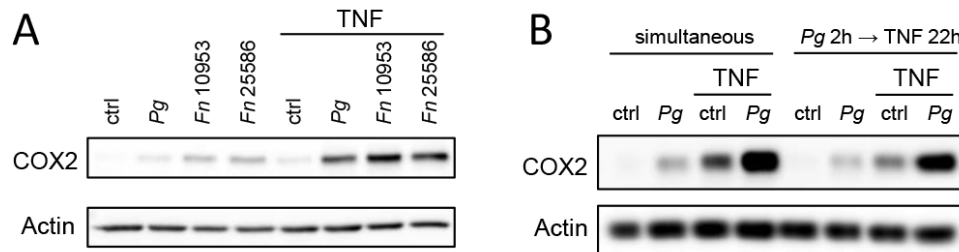
**Supplementary table S1.** Composition of *F. alocis* culture medium

| Component                   | Amount per L |
|-----------------------------|--------------|
| Beef extract                | 5 g          |
| Yeast extract               | 5 g          |
| Bio-tryptone                | 15 g         |
| Glucose                     | 2 g          |
| L-cysteine                  | 0.6 g        |
| Arginine                    | 17.42 g      |
| Sodium chloride             | 5 g          |
| Magnesium sulfate           | 0.1 g        |
| Potassium phosphate dibasic | 6 g          |
| Sodium carbonate            | 0.6 g        |
| L-cysteine                  | 0.5 g        |
| hemin                       | 10 mg        |
| Vitamin K                   | 0.5 mg       |

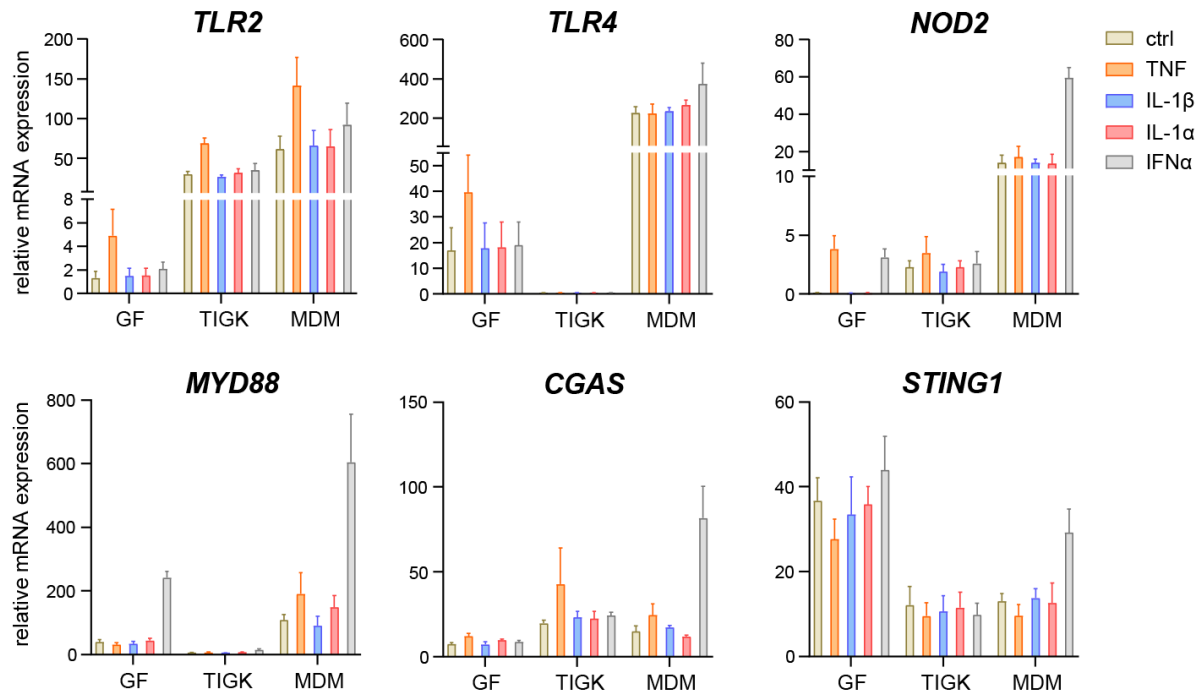
**Supplementary table S2.** Sequences of primers used for qPCR analyses.

| Gene         | Forward primer          | Reverse primer          |
|--------------|-------------------------|-------------------------|
| <i>COX2</i>  | AGCCCTTCCTCCTGTGCCT     | AATCAGGAAGCTGCTTTTTACCT |
| <i>IL8</i>   | GCTCTGTGTGAAGGTGCAGT    | CCAGACAGAGCTCTCTTCCA    |
| <i>TLR2</i>  | CGGAATGTCACAGGACAGCA    | TACCACAGGCCATGGAAACG    |
| <i>TLR4</i>  | ACCATCATTGGTGTGTCGGTC   | AGCCAGCAAGAAGCATCAGG    |
| <i>MYD88</i> | CTAAGAAGGACCAGCAGAG     | GAAGCATCAGTAGGCATCA     |
| <i>NOD2</i>  | GATTGGCTGCCTTCCTTCTA    | GAGCGTCTCTGCTCCATCAT    |
| <i>CGAS</i>  | GCCAGTAGTGCTTGGTTTCC    | GTTCCCCGAAAGAAGAATCC    |
| <i>STING</i> | TCAAGGATCGGGTTTACAGC    | TGGCAAACAAAGTCTGCAAG    |
| <i>RPLP0</i> | GCGTCCTCGTGGAAGTGACATCG | TCAGGGATTGCCACGCAGGG    |

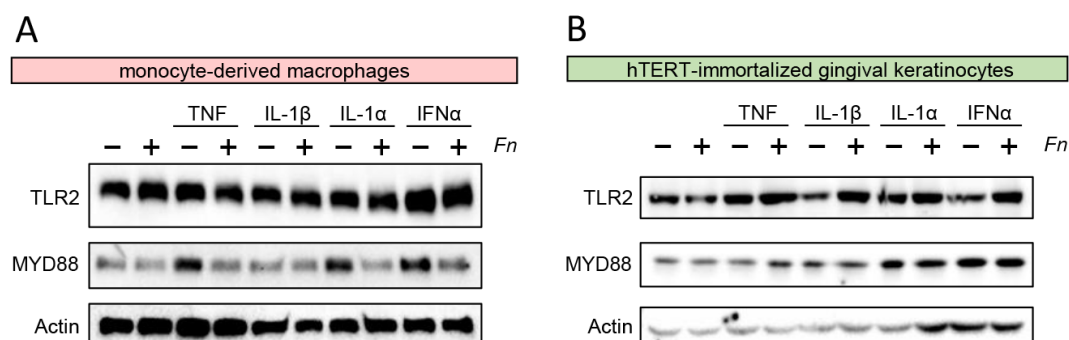
## Supplementary Figures



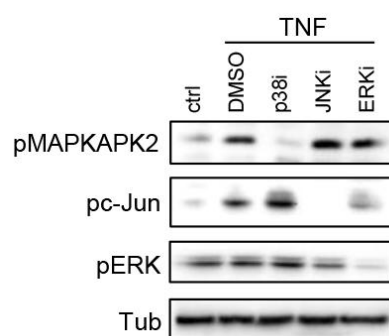
Supplementary figure S1. (A) Western blot analysis of COX-2 in GFs infected with *P. gingivalis* (*Pg*) or two different strains of *F. nucleatum* (*Fn*) (ATCC 10953 or ATCC 25586) in the absence (ctrl) or presence of 10 ng/ml TNF for 24 h. (B) Western blot analysis of GFs that were subjected to two different protocols of *P. gingivalis* (*Pg*) infection and TNF stimulation: cells were either infected and stimulated simultaneously or were infected with *P. gingivalis* for 2 h, followed by washing and stimulation with 10 ng/ml TNF in DMEM containing 2% FBS and antibiotics (2.5 mg/ml gentamicin and 2 mg/ml metronidazole) for 24 h. Actin was used as loading control and results shown are representative of three independent experiments.



Supplementary figure S2. qPCR analysis of *TLR2*, *TLR4*, *NOD2*, *MYD88*, *CGAS*, and *STING1* expression in GFs, MDMs, and TIGKs (n=3) stimulated with TNF (10 ng/ml), IL-1 $\beta$  (10 pg/ml), IL-1 $\alpha$  (10 pg/ml) or IFN $\alpha$  (100 U/ml) for 4 h. Data are shown as mean relative mRNA expression calculated using the  $\Delta C_t$  method to allow for comparison between different cell types.



Supplementary figure S3. Western blot analysis of TLR2 and MYD88 in (A) MDMs and (B) TIGKs infected with *F. nucleatum* (*Fn*) in the absence or presence of TNF (10 ng/ml), IL-1 $\beta$  (10 pg/ml), IL-1 $\alpha$  (10 pg/ml) or IFN $\alpha$  (100 U/ml) for 24 h. Actin was used as loading control and results representative of two independent experiments are shown.



Supplementary figure S4. Western blot analysis of phospho (p)MAPKAPK2, pc-Jun, and pERK in GFs pretreated with MAP kinase inhibitors (p38i, JNKi, ERKi) for 60 min prior to stimulation with 10 ng/ml TNF for 30 min. Tubulin was used as loading and results shown are representative of two independent experiments.