Additional file 3: Supplementary figures S1, S2, S3 and S4

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Fig. S1. Pulse and persistent stimulation by a high $\text{TNF}\alpha$ dose (10ng/ml). Amplitude of nuclear to cytoplasmic NF- κ B oscillations in single cells, and nuclear NF- κ B averaged over 100 cells compared with the experiment on mouse embryonic fibroblast (Hoffmann et al. *Science* 2002, **298**:1241-1245). Panel A, 5, 15, 30 and 60 min. long pulse stimulation. Panel B, persistent stimulation. Nuclear NF- κ B averaged over 100 cells compared with data from I κ B ε and I κ B β deficient and wild-type fibroblast (Hoffmann et al. *Science* 2002, **298**:1241-1245).



Fig. S2. The role of A20 negative feedback control in persistent $\text{TNF}\alpha$ stimulation. Model predictions versus experiment for wild type and A20-/- mouse embryonic fibroblast (Lee et al. *Science* 2000, **289**:2350-2354).



Fig. S3. The role of A20 negative feedback control in pulse $\text{TNF}\alpha$ stimulation. Experiment on 3T3 (Werner et al. *Science* 2005, **309**:1857-1861) stimulated by a 45 min. pulse of 1= ng/ml $\text{TNF}\alpha$, versus model predictions for $\text{TNF}\alpha=1$ ng/ml and $\text{TNF}\alpha=10$ ng/ml (a better agreement with the experiment is achieved when in the numerical simulation $\text{TNF}\alpha$ concentration is set at 10ng/ml). Panel A, IKK activity of wild type cells in response to 45-min stimulation. Panel B, IKK activity of A20-/- cells. Panel C, nuclear NF- κ B of wild-type cells. Panel D, nuclear NF- κ B of A20-/- cells.



Fig. S4. Nuclear to cytoplasmic NF- κ B oscillations during persistent treatment by 10ng/ml TNF for 5 levels of total NF- κ B, 10000, 20000, 50000, 100000, 200000 molecules. The figure shows weak dependence of oscillation period to the total amount of NF- κ B, in agreement with the single cell experiment (Nelson et al. *Science* 2005, **308**:52b).