- 2
- 3

Fig. S1



Fig. S1. Intracellular replication of *L. monocytogenes* in various cell types. The bars indicate mean values of \log_{10} CFU/ml and the error bars the SD. All experiments were performed at least two times with triplicate or duplicate data sets, of which one representative experiment is shown. Asterisks indicate significant differences between bacterial numbers in the respective cell type for the 4 or 8 h time point *vs.* the 0 h time point, as determined by a 2-sided *t* test with equal variance (** *P* ≤ 0.01; *** *P* ≤ 0.001).





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Fig. S2. Intracellular replication of indicated *F. tularensis* strains in J774 cells. The bars indicate mean values of \log_{10} CFU/ml and the error bars the SD. Experiments were performed two times with triplicate or duplicate data sets, of which one representative experiment is shown. Asterisks indicate a significant difference between the \log_{10} number of the CFU of the mutant in comparison to the parental strain LVS, as determined by a 2-sided *t* test with equal variance (* $P \le 0.05$; ** $P \le 0.01$).













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Fig. S4. Intracellular replication of indicated *F. tularensis* strains in ASC^{-/-} BMDM. The bars indicate mean values of \log_{10} CFU/ml and the error bars the SD. Experiments were performed three times with triplicate or duplicate data sets, of which one representative experiment is shown. Asterisks indicate a significant difference between the \log_{10} number of the CFU of the mutant in comparison to the parental strain LVS, as determined by a 2-sided *t* test with equal variance (* *P* ≤ 0.05; ** *P* ≤ 0.01; *** *P* ≤ 0.001).







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Fig. S5. Intracellular replication of indicated *F. tularensis* strains in MyD88^{-/-} BMDM. The bars indicate mean values of \log_{10} CFU/ml and the error bars the SD. Experiments were performed three times with triplicate or duplicate data sets, of which one representative experiment is shown. Asterisks indicate a significant difference between the \log_{10} number of the CFU of the mutant in comparison to the parental strain LVS, as determined by a 2-sided *t* test with equal variance (* *P* ≤ 0.05; ** *P* ≤ 0.01; *** *P* ≤ 0.001).

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Fig. S6. Intracellular replication of indicated *F. tularensis* strains in HeLa cells. The bars indicate mean55values of log_{10} CFU/ml and the error bars the SD. Experiments were performed two times with56triplicate or duplicate data sets, of which one representative experiment is shown. Asterisks indicate57a significant difference between the log_{10} number of the CFU of the mutant in comparison to the58parental strain LVS, as determined by a 2-sided t test with equal variance (* $P \le 0.05$; *** $P \le 0.001$).



- **Fig. S7.** Pictures representing cells microinjected with *F. tularensis* that fall into either of three
- 65 categories; (A) 0 20, (B) 20 100 and (C) 100 1000 bacteria/cell.



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Fig. S8. Microinjection of J774 cells (left panels) or BMDM (right panels) with GFP-expressing LVS and
non-labeled Δ*iglG* (top panels) or GFP-expressing LVS alone (bottom panels). Pictures were taken at

- 71 24 h after injection with a live-cell imaging microscope equipped with an EMCCD camera. Co-
- 72 localization of injected cells containing RD (red) and GFP expressing bacteria (green) resulted in
- 73 yellow signals. Representative pictures from one of two independent experiments are shown.

Fig. S9







infected by injection is represented by a dot that is assigned to the corresponding category 0 - 20, 20

79 – 100, and 100 – 1000 bacteria per cell.