

Figure S1. Light microscopy images of *E. coli* UTI89 grown under static or shaking conditions, UTI89 Δfim , UTI89 $\Delta fimH$ and UTI89 $\Delta fimH/pfimH$ incubated with canine blood with/without D-mannose (10× magnification).

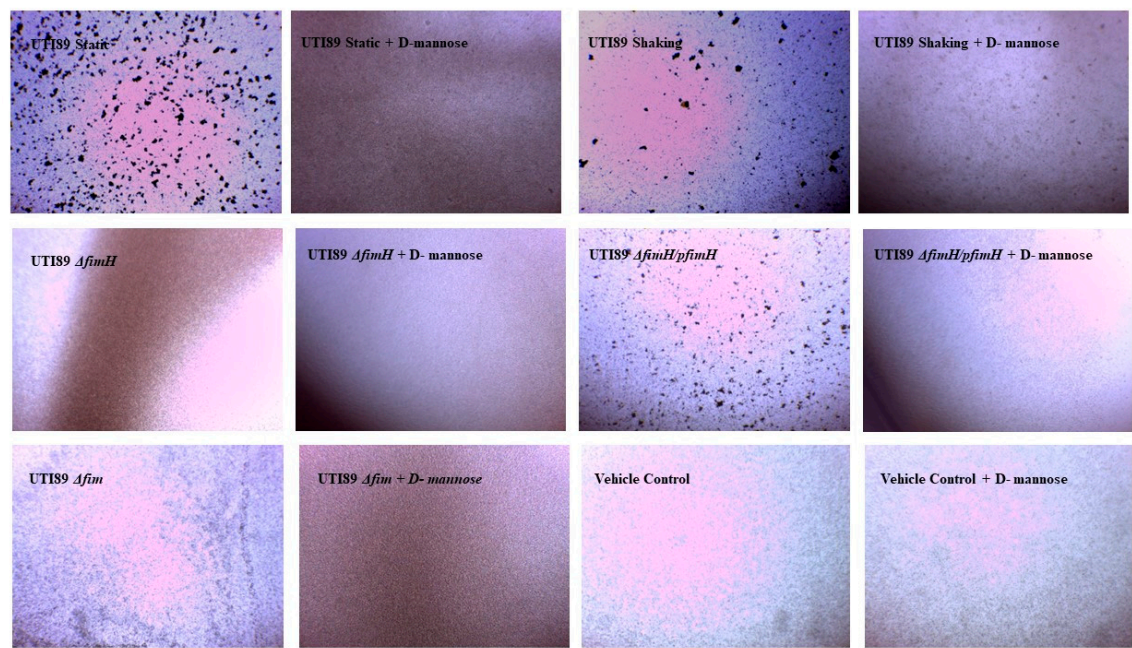


Table S1. Red Blood Cell Hemagglutination of various constructs of UTI89 with/without D-mannose.

	UTI89 Static	UTI89 Shaking	UTI89 $\Delta fimH$	UTI89 $\Delta fimH/pfimH$	UTI89 Δfim
Mannose (-)	+++	+	-	++	-
Mannose (+)	-	-	-	-	-
“-”: No agglutination “+”: Low agglutination “++”: Moderate agglutination “+++”: High agglutination.					

Figure S2. Inverted light microscopy images revealed that UTI89 grown statically has increased interaction with both MDA-MB-231 and MCF-7 breast cancer cells compared to the UTI89 grown under shaking conditions ($\times 20$ magnification; arrows point to the enlarged area).

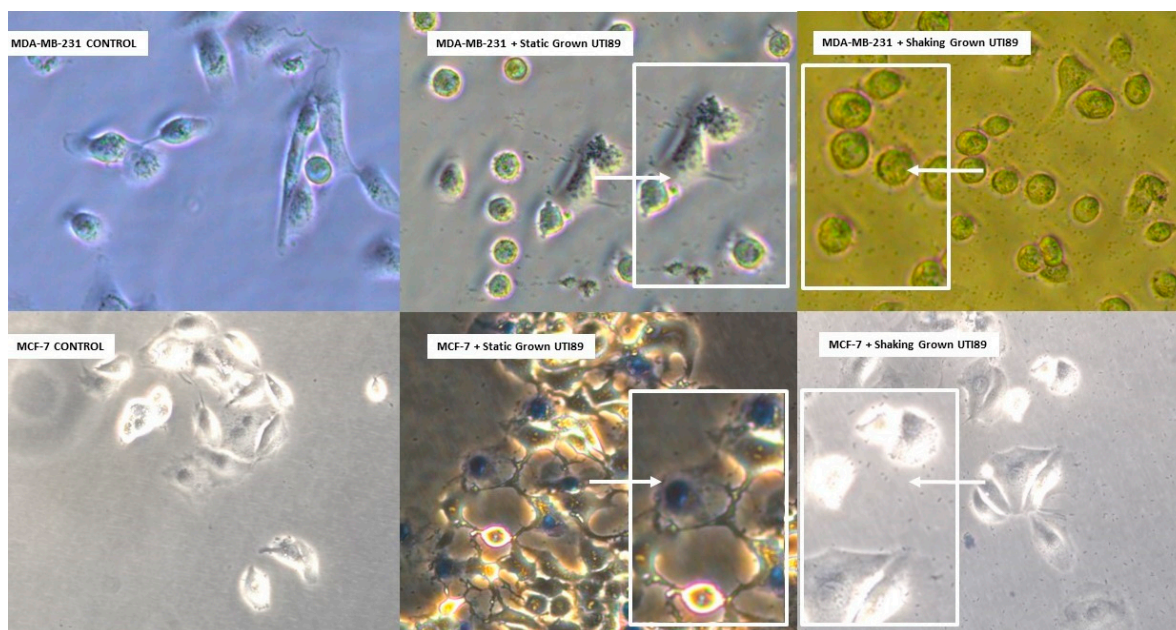


Figure S3. Inverted light microscopy images revealed significant cell surface area reduction in MDA-MB-231 cells upon incubation with both static and shaking UTI89 compared to MCF-7 breast cancer cells ($\times 20$ magnification).

