

FIG S1 Phagocytosis of strain LAC by PMNs in the absence (vehicle control) or presence of cytochalasin D as indicated. Human PMNs were incubated with 10 μ g/ml cytochalasin D for 15 min at 37°C prior to addition of bacteria. 30- μ l aliquots of each assay mixture was centrifuged onto slides with a CytospinTM instrument and cells were visualized by using a Wright-Giemsa stain. Images are representative of six experiments and were captured by using a 100× objective as described in Methods.

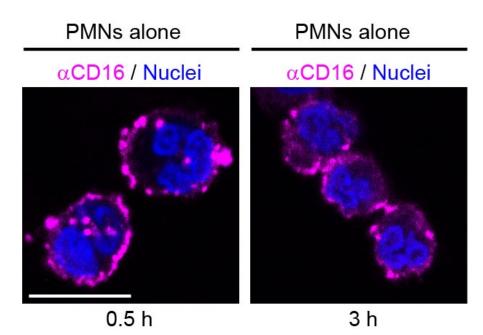


FIG S2 Plasma membrane labeling of control PMNs with anti-CD16 antibody (pink) following culture for (0.5 h) or 3 h, as indicated. Nuclei were stained with NucBlue as described in Methods. Images are representative of control PMNs in multiple experiments. Scale bar = $10 \mu m$.

Time (h)	CD16-	CD16+	CD16-	CD16+	CD16+
	FITC+	FITC+	FITC+	FITC+	FITC-
	αSa+	αSa+	αSa-	αSa-	(no <i>Sa</i>)
0.5	0	0	0	88.0 ± 2.8	12.0 ± 2.8
1	2.0 ± 1.6	3.5 ± 4.4	20.5 ± 12.5	61.0 ± 13.3	13.0 ± 11.4
2	2.5 ± 2.5	1.3 ± 2.5	23.0 ± 11.0	59.5 ± 12.5	13.8 ± 5.9
3	14.5 ± 12.8	5.5 ± 6.2	38.2 ± 8.3	27.8 ± 11.3	14.1 ± 9.4

TABLE S1 Quantification of microscopy data.

The percentage (%) of PMNs without *S. aureus* (CD16+/FITC–), or percentage of those that stain positive or negative (+ or –) with anti-CD16 antibody (CD16+ or CD16–), FITC-labeled *S. aureus* (FITC+), and/or *S. aureus* labeled with the combination of anti-*S. aureus* IgG + goat anti-rabbit IgG conjugated to AF594 (α Sa+ or α Sa–) was calculated for each timepoint. Fifty PMNs per sample were evaluated and scored. Results are presented as the mean ± SD of four experiments.