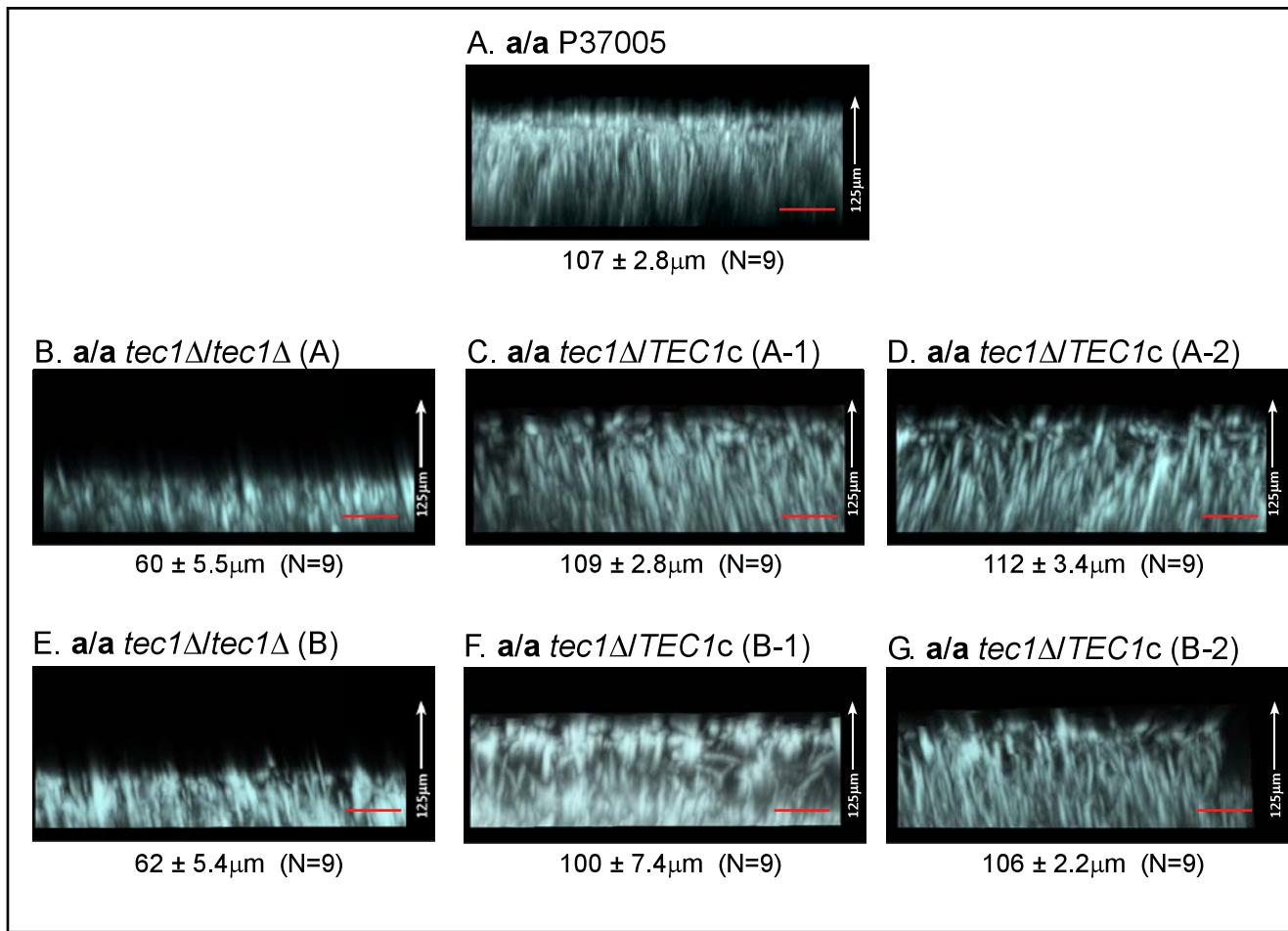


Supplemental Table S1. Primers used in this study.

Primer	Sequence (5' → 3')
P1	ATGTCAATTACTAAAACATA
P2	CTATGTTGTGACTGTTTACTTC
P3	ATGATGTCGCAAGCTACTCCT
p4	CTAAAACACTCACTAGTAAATCCTTC
TEC1-F1	CCCAATTCTC <u>AGGGCCCTAGTCAGGTAG</u>
TEC1-R1	GCAATTAAAGATT <u>GGATCCCCGCTAAACTAATG</u>
TEC1-F2	CACTTACTCA <u>TGCAGGATA</u> CATTAGTTAGC
TEC1-R2	CTCTAATACAAG <u>CCGCGGACAATTATGTAATCC</u>
TEC1-SR	GCAAAGAAGTGGAAACAAAAGAAACCC
CaSAT1-F	GGGCACTAAGCAGACAGCTCCTTGGC

Figure S1



Supplemental Figure Legend

Figure S1. Complementation of the *a/a* tec1 Δ /tec1 Δ mutant by introducing wild type TEC1 into its native location, rescues the aberrant mutant phenotype. Mutants *a/a* tec1 Δ /tec1 Δ (A) and *a/a* tec1 Δ /tec1 Δ (B) were complemented to generate *a/a* tec1 Δ /TEC1c (A-1) and (A-2), and mutants *a/a* tec1 Δ /TEC1c (B-1) and *a/a* tec1 Δ /TEC1c (B-2), respectively. Side views of a stack of 500 confocal microscopy scans are provided for each biofilm. A. *a/a* P37005. B. *a/a* tec1 Δ /tec1 Δ (A). C. *a/a* tec1 Δ /TEC1c (A-1). D. *a/a* tec1 Δ /TEC1c (A-2). E. *a/a* tec1 Δ /tec1 Δ (B). F. *a/a* tec1 Δ /TEC1c (B-1). G. *a/a* tec1 Δ /TEC1c (B-2). Thickness is presented as mean ± standard deviation (N=9). Scale bar = 20 μm.