

SUPPLEMENTAL INFORMATION

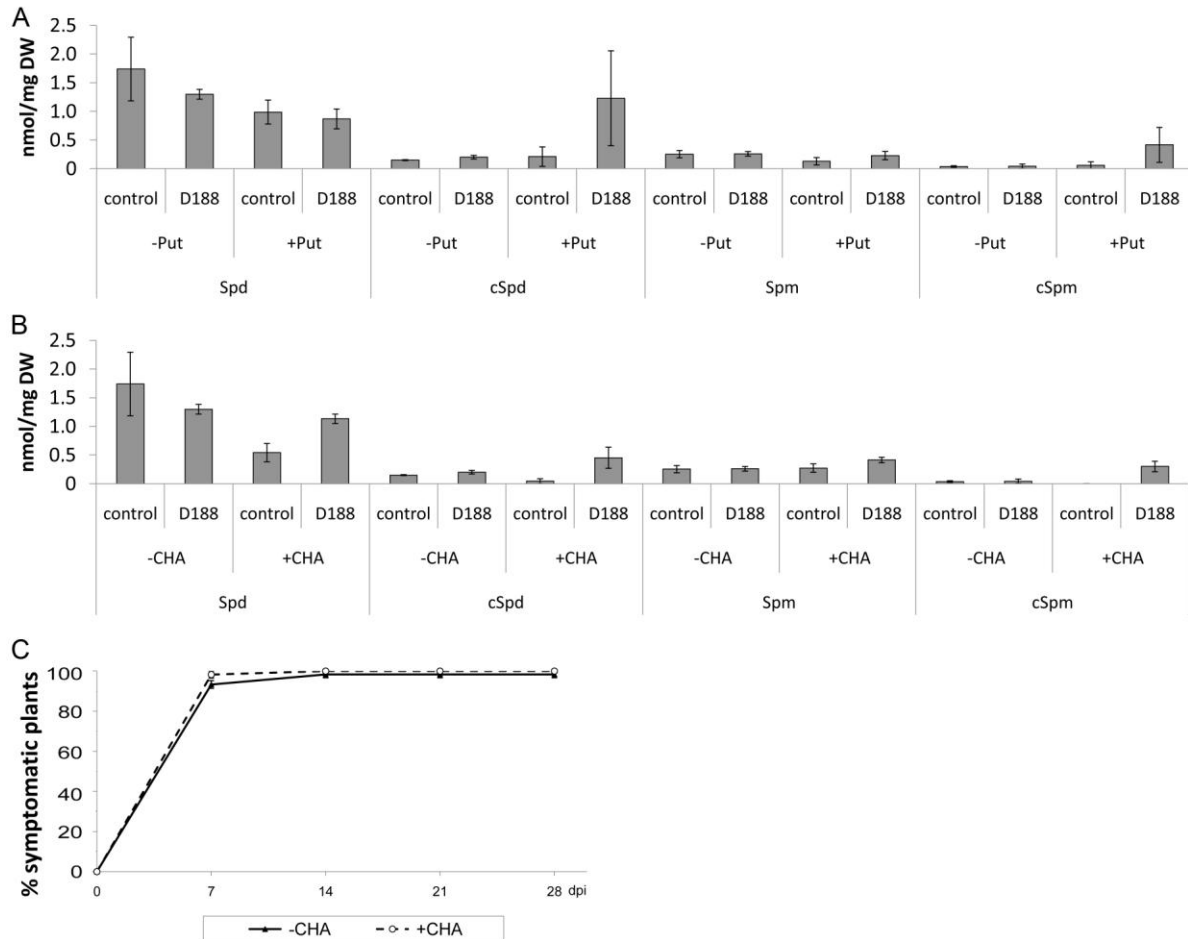


Figure S1. Effect of modulating the Put content on polyamine levels and symptom development in *R. fascians*-infected *Arabidopsis* Col-0.

Error bars represent standard errors (n=3). Student's *t*-tests did not reveal any statistical differences. A, Free and conjugated Spd and Spm levels measured at 14 dpi. Plants were placed on 1 mM Put 10 days before infection. B, Free and conjugated Spd and Spm levels measured at 14 dpi. Plants were placed on the SPDS inhibitor CHA (1 mM) just before infection. C, Percentage of symptomatic plants at different time points after infection.

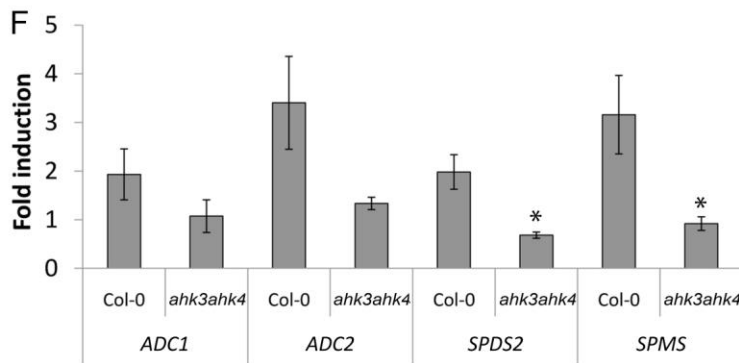
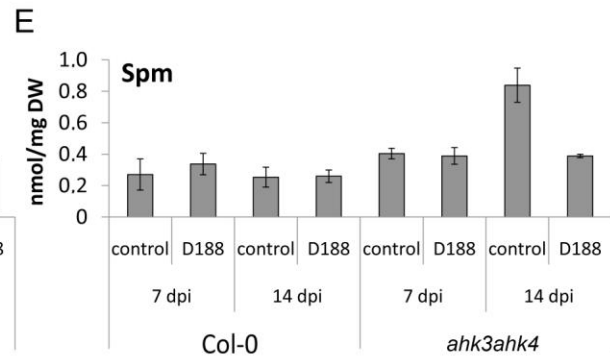
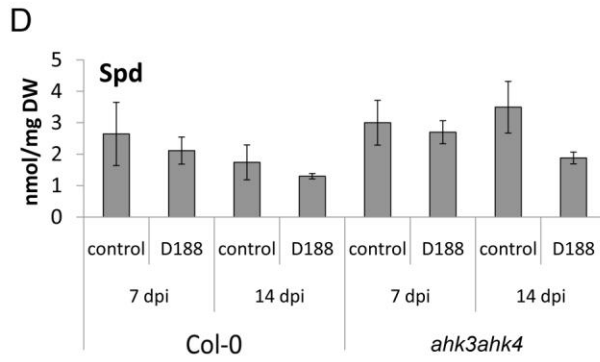
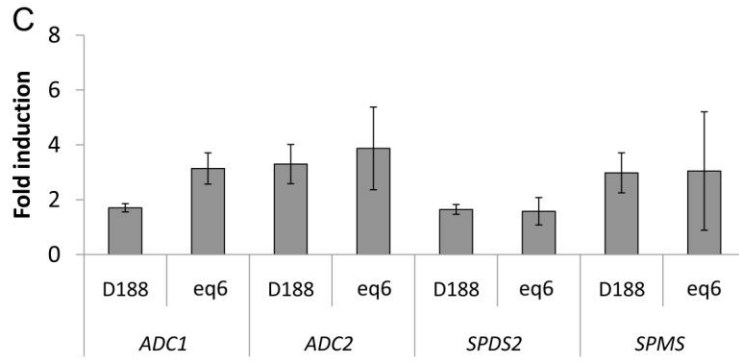
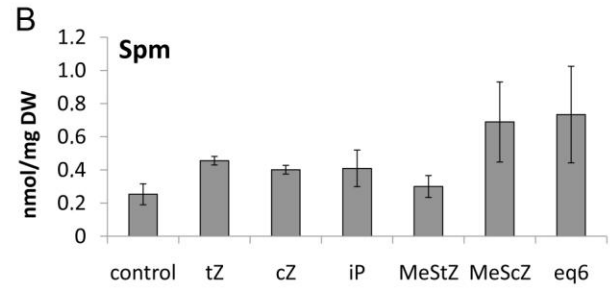
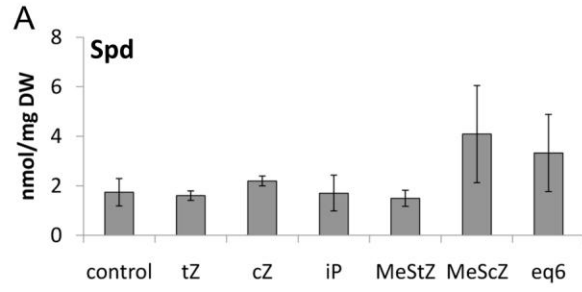


Figure S2. Importance of cytokinins on Spd and Spm production.

Error bars represent standard errors (n=3). Student's *t*-tests did not reveal any statistical differences (A-E). A-B, Free Spd (A) and (B) Spm levels in Col-0 plants treated with 10 μ M of different cytokinins or with an equimolar mix of 1 μ M each of iP, tZ, cZ, and their 2MeS derivatives (eq6) for 14 days. C, Fold-change in transcript levels of polyamine biosynthesis genes in D188-infected (D188 versus mock-infected) and eq6-treated (treated versus non-treated) Col-0 plants at 14 dpt. D-E, Free Spd (D) and Spm (E) levels in D188- and mock-infected (control) Col-0 and *ahk3ahk4* plants at 14 dpi. F, Fold-change in transcript levels of polyamine biosynthesis genes in D188 versus mock-infected Col-0 and *ahk3ahk4* plants at 14 dpi. Asterisks mark statistical differences ($p < 0.05$) between wild-type and mutant plants according to a Student's *t*-test.

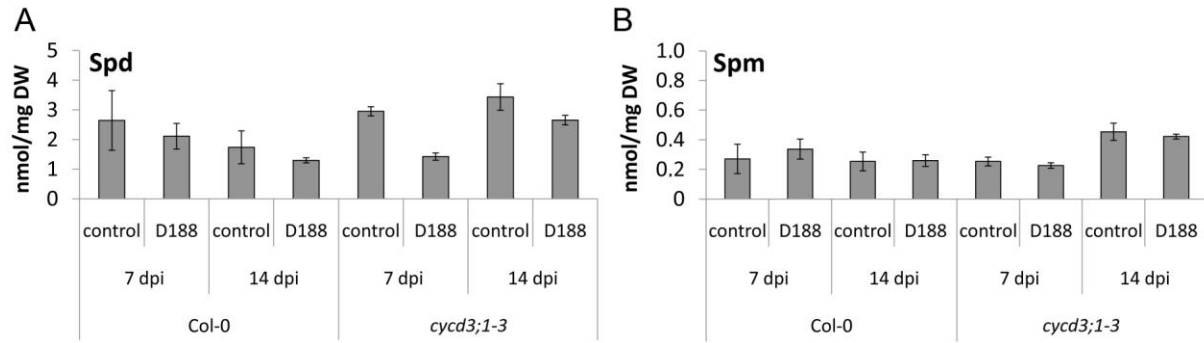


Figure S3. Free Spd (A) and Spm (B) levels in D188- and mock-infected (control) Col-0 and *cycd3;1-3* plants at 14 dpi.

Error bars represent standard errors (n=3). Student's *t*-tests did not reveal any statistical differences.

Table S1. *In vitro* and *in planta* bacterial growth and *fasA* gene expression in the presence and absence of Put.

To assess the effect of exogenous Put on bacterial growth, the turbidity of cultures (OD₆₀₀) and the colony forming units (CFU/mg fresh weight (FW)) present in infected *Arabidopsis* plants were determined, both upon growth in the presence or absence of 1 mM Put. We also monitored the expression of the *fasA* gene involved in cytokinin production in cultures of the GUS reporter strain D188-pSPIP*fasAgus* (Pertry et al., 2010) incubated under optimal conditions for *fas* gene expression (see “Material and Methods”) and in bacterial cells present in plant tissues, again in the presence or absence of 1 mM Put. No significant differences were measured for either of these parameters using Student’s *t*-tests.

Treatment	OD₆₀₀	CFU/mg FW	GUS activity
<i>In vitro</i>			
Pyr	0.837 ± 0.018		612.34 ± 124.36
His + Pyr	0.834 ± 0.011		7740.21 ± 517.57
Put + Pyr	0.893 ± 0.017		506.01 ± 111.50
Put + His + Pyr	0.997 ± 0.061		9404.64 ± 824.89
<i>In planta</i>			
Control		3615.22 ± 1401.66	40.06 ± 9.50
+Put		3477.66 ± 1177.51	58.08 ± 8.22

Table S2. Primers used for qRT-PCR amplifications.

Gene	AGI	Primer sequence	Reference
<i>ACT2</i>	At3g18780	Forward: GGCTCCTCTTAACCCAAAGGC	Simón-Mateo et al. (2006)
		Reverse: CACACCATCACCAGAATCCAGC	
<i>ADC1</i>	At2g16500	Forward: CCAAGGTGTGTATCCTGTGAAAT	Jubault et al. (2008)
		Reverse: AGCTTCTAAACCGAATCGAAAAC	
<i>ADC2</i>	At4g34710	Forward: GCGATGGACCACACAGCTTT	Jubault et al. (2008)
		Reverse: AGAACATCCGCTGAGGACTGA	
<i>AIH</i>	At5g08170	Forward: TCGAGAATGCAAGAGAGATCGTT	Jubault et al. (2008)
		Reverse: CATTTTCGGCGACGGAAGTA	
<i>CPA</i>	At2g27450	Forward: GATCAAGTCGAAAAGGCAAAGCT	Jubault et al. (2008)
		Reverse: CCATCCATAGTAAGAAGCACCTTGT	
<i>SPDS1</i>	At1g23820	Forward: AATCACCACCTCTCACAAACCC	Jumtee et al. (2008)
		Reverse: TCGGTGGCAGAGGTTTCTTTA	
<i>SPDS2</i>	At1g70310	Forward: TTGCCCGTGAAGAGACCTAGA	Jubault et al. (2008)
		Reverse: TCCACCGTTCTCTGTTTCCAT	
<i>SPMS</i>	At5g53120	Forward: TGGCTCCATACTCATCTTATTGAA	Jubault et al. (2008)
		Reverse: CGCATAGTGAACACTTTTGAATG	
<i>ACL5</i>	At5g19530	Forward: CCATCATTTGCGGACACATG	Jubault et al. (2008)
		Reverse: GAGACGAAAGAAGGAGCGTTTAGA	
<i>SAMDC1</i>	At3g02470	Forward: TCTTTGAGCCAAGCATCTTTCA	Jubault et al. (2008)
		Reverse: GCAGCAGGTGTAAGAATTTTCATCA	
<i>SAMDC2</i>	At5g15950	Forward: TCTCCGAGATCTACCTTGAAATG	Jubault et al. (2008)
		Reverse: GATTCCCTATTCTTCTCGTCCT	
<i>SAMDC3</i>	At3g25570	Forward: GGTGGATAGGGTTCGGTTTG	Cui et al. (2010)
		Reverse: GGTGAGCAACATTCAACAGTC	
<i>SAMDC4</i>	At5g18930	Forward: GGTGACCGTTACTCAACTATCCA	This work
		Reverse: CGAAGCTCGCGTAGCTAAA	
<i>ARR5</i>	At3g48100	Forward: TTTAAAAGCTCAAAGATTCACACACA	Jasinski et al. (2005)
		Reverse: ATCAGCAAAGAAGCCGTAATGT	
<i>CYCD3;1</i>	At4g34160	Forward: CGTTCGTAGACCACATTATCAGGAG	Depuydt et al. (2009)
		Reverse: CGGAGATTACAGAGAGGAGGAGAC	
<i>CDKB1;1</i>	At3g54180	Forward: GGTGGTGACATGTGGTCTGTTGG	Boudolf et al. (2004)
		Reverse: CGCAGTGTGAAACACCCGG	

References

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