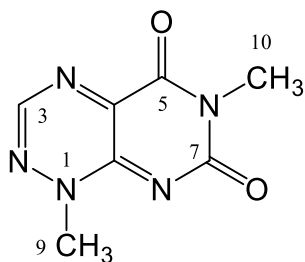


Supplementary Note 1. Physicochemical data and antimicrobial activity of the symbiont-produced compounds

Toxoflavin (1)



ESI(+) m/z 194 (M+H)⁺, HRESI(+)-MS m/z 194.0674 (calcd. for C₇H₈N₅O₂ 194.0673)

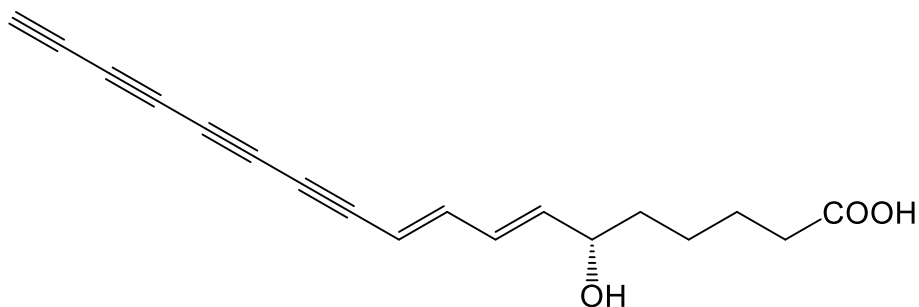
UV (PDA): λ_{\max} = 258, 397 nm

NMR: d₆-DMSO, ¹H NMR 600 MHz, ¹³C NMR 150 MHz

Carbon	¹³ C	¹ H (mult., <i>J</i> in Hz)
3	144.7	8.96 (s)
4a	146.5	-
5	154.1	-
7	159.0	-
8a	150.9	-
9	42.4	3.94 (s)
10	28.2	3.24 (s)

Antimicrobial activity (1 mg mL⁻¹): An inhibition zone of 22 mm against *Brevibacillus laterosporus* was measured (procedure described in Methods section, *Antimicrobial bioassays*).

Caryoynencin (2)

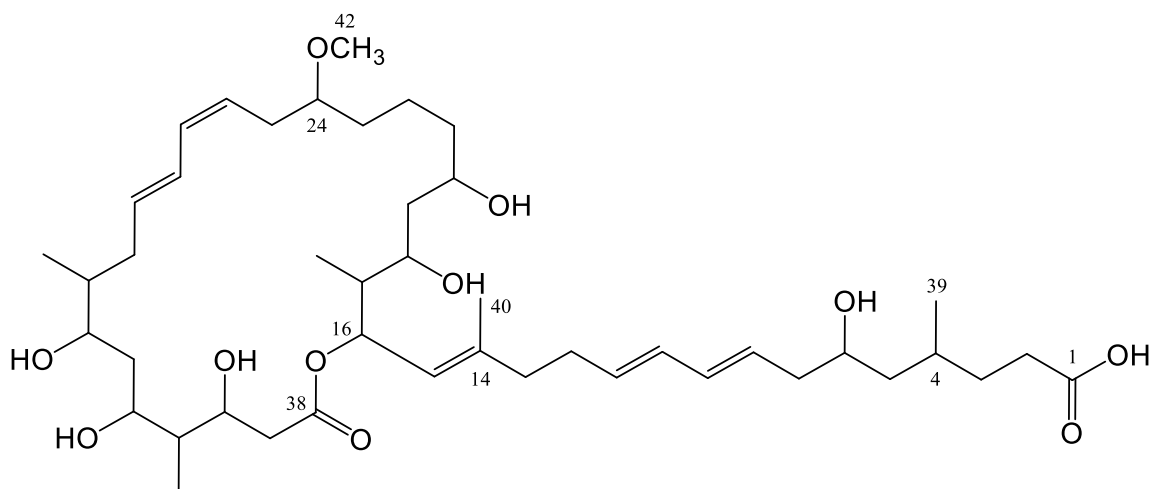


ESI(-) m/z 279 (M-H)⁻, HRESI(-)-MS m/z 279.1031 (calcd. for C₁₈H₁₇O₃ 279.1027)

UV (PDA): λ_{\max} = 294, 239, 280, 268, 358 nm.

Antimicrobial activity: active against *P. lilacinum* (see Methods, *Antimicrobial bioassays*) (Supplementary Fig. 21b).

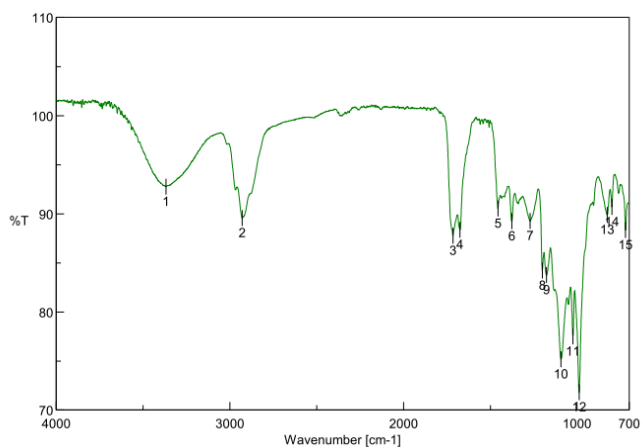
Lagriene (3)



ESI(-) m/z 777 (M-H)⁻, HRESI(-)-MS m/z 777.5172 (calcd. for C₄₄H₇₃O₁₁ 777.5158)

UV (PDA): λ_{\max} = 231 nm

IR-spectrum:



NMR: d₆-DMSO, ¹H NMR 600 MHz, ¹³C NMR 150 MHz

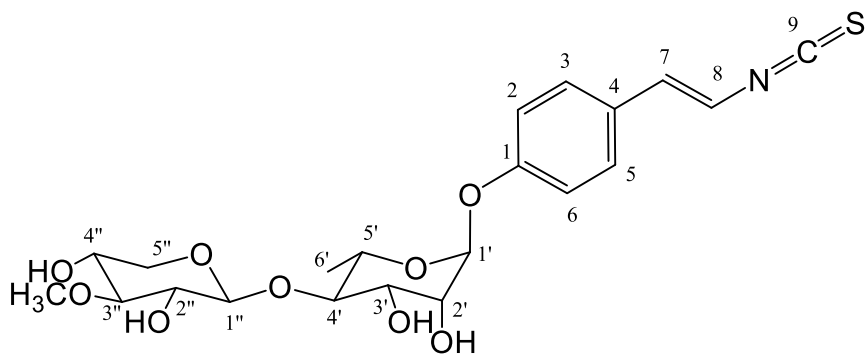
Carbon	¹³ C	¹ H (mult., <i>J</i> in Hz)
1	174.1	-
2	31.4	2.16 (m)
3	32.4	1.47 (m) 1.32 (m)
4	28.2	1.60 (m)
5	43.8	1.26 (m) 1.03 (m)
6	67.5	3.49 (m)
7	41.5	2.08 (m)
8	129.3	5.55 (m)
9	131.6	5.95 (m)
10	130.7	5.95 (m)
11	131.2	5.48 (m)
12	30.2	2.27 (m) 2.11 (m)
13	38.9	2.06 (m) 2.02 (t, 7.2)
14	139.1	-
15	122.9	5.00 (d, 9.5)
16	72.1	5.26 (m)
17	41.8	1.78 (m)
18	66.5	3.77 (s)
19	39.3	1.45 (m) 1.32 (m)
20	66.8	3.62 (m)
21	38.3	1.48 (m)
22	20.6	1.35 (m) 1.21 (m)
23	31.9	1.36 (m)
24	79.6	3.21 (m)
25	30.1	2.38 (m) 2.27 (m)
26	125.1	5.26 (m)
27	130.1	6.00 (m)
28	126.6	6.29 (m)

29	133.6	5.72 (m)
30	36.5	2.18 (m)
31	37.7	1.48 (m)
32	71.7	3.53 (m)
33	39.3	1.45 (m) 1.32 (m)
34	69.2	3.84 (m)
35	43.6	1.33 (m)
36	68.6	3.81 (m)
37	39.6*	2.45 (m) 2.15 (m)
38	170.4	-
39	18.8	0.80 (m)
40	16.6	1.68 (s)
41	10.5	0.71 (d, 7.0)
42	55.6	3.22 (s)
43	13.1	0.79 (m)
44	9.7	0.76 (d, 7.0)

* signal overlapping with solvent signal

Antimicrobial activity (1 mg mL⁻¹): Inhibition zones of 23 mm against *Bacillus thuringiensis*, 30 mm against *Mycobacterium vaccae*, 19 mm against vancomycin-resistant *Enterococcus faecalis*, and 18 mm against methicillin-resistant *Staphylococcus aureus* were measured (procedure described in Methods section, *Antimicrobial bioassays*).

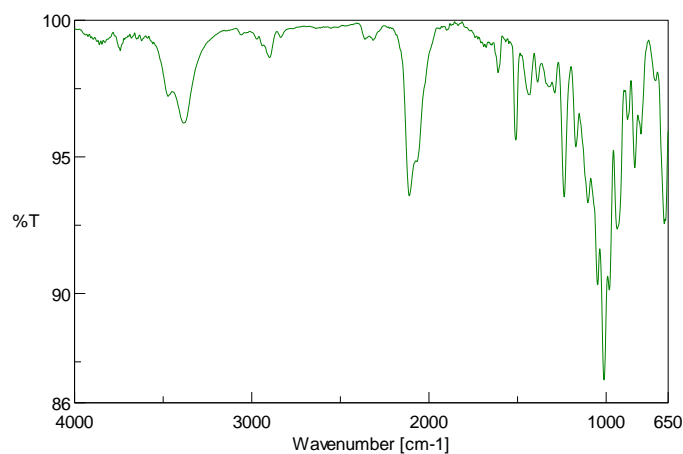
Sinapigladioside (4)



ESI(-) m/z 468 (M-H)⁻, HRESI(-)-MS m/z 468.1334 (calcd. for C₂₁H₂₆NO₉S 468.1334)

UV (PDA): λ_{\max} = 312 nm.

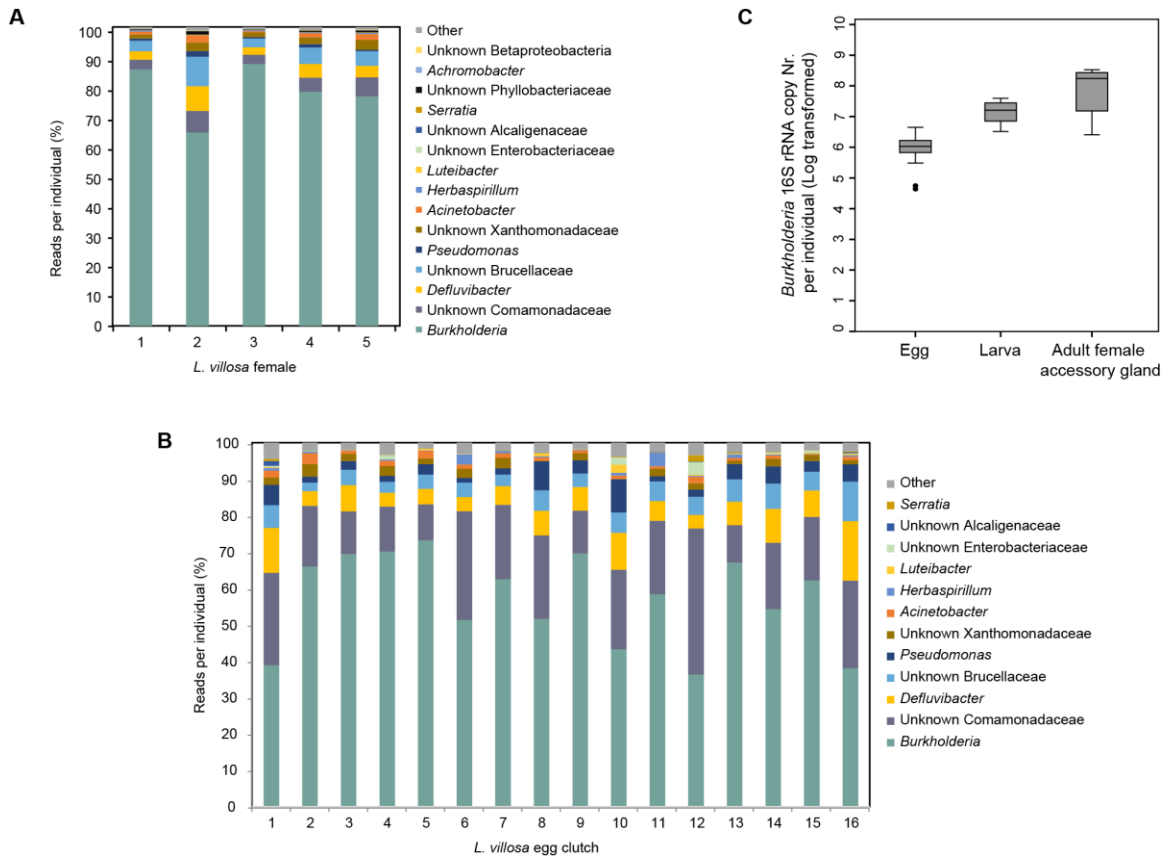
IR-spectrum:



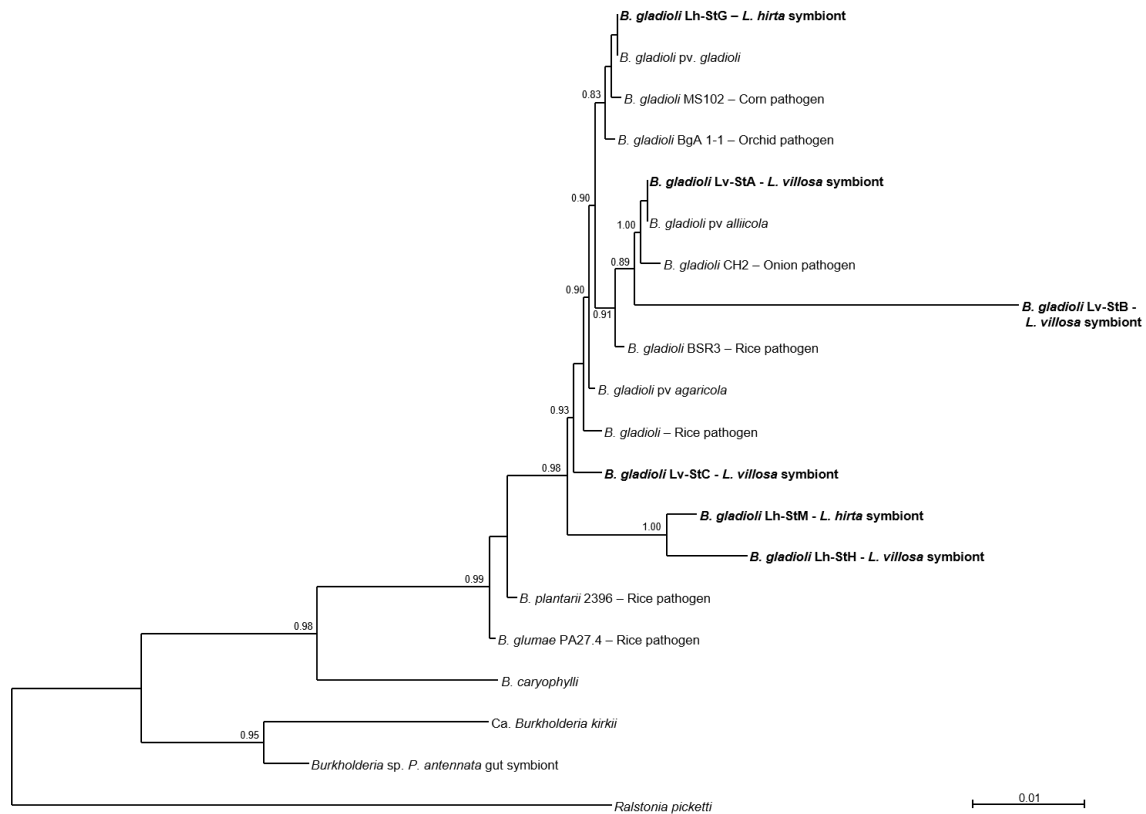
NMR: d_6 -DMSO, ^1H NMR 600 MHz, ^{13}C NMR 150 MHz

Carbon	^{13}C	^1H (mult., J in Hz)
1	156.4	-
2	116.7	7.04 (d, 8.8)
3	128.0	7.43 (d, 8.8)
4	127.3	-
5	128.0	7.43 (d, 8.8)
6	116.7	7.04 (d, 8.8)
7	131.7	6.86 (d, 13.9)
8	114.0	7.00 (d, 13.9)
9	131.5	-
1'	97.5	5.46 (d, 1.6)
2'	69.3	3.92 (m)
3'	68.9	3.75 (m)
4'	81.8	3.48 (m)
5'	67.9	3.59 (m)
6'	17.3	1.20 (d, 6.2)
1''	104.0	4.25 (d, 7.8)
2''	72.7	3.10 (m)
3''	86.2	2.93 (t, 9.0)
4''	68.8	3.42 (m)
5''	65.7	3.75 (m) 3.17 (t, 10.9)
3''-OCH₃	60.0	3.50 (s)
2'-OH	-	5.25 (d, 4.3)
3'-OH	-	4.32 (d, 2.5)
2''-OH	-	5.35 (d, 5.4)
4''-OH	-	5.15 (d, 5.5)

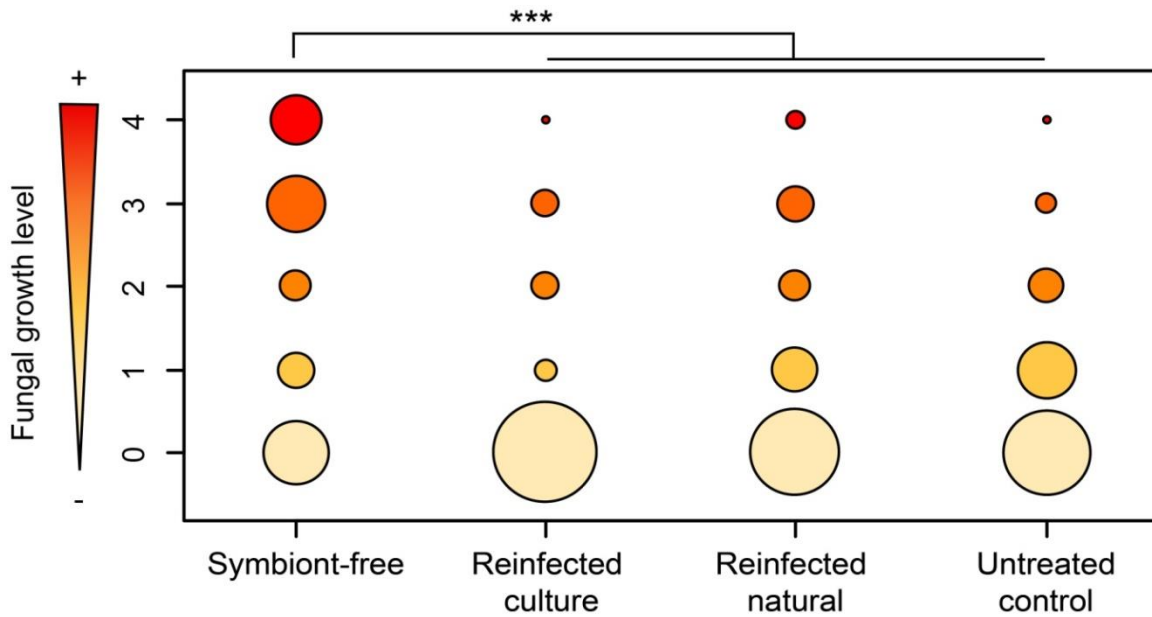
Antimicrobial activity (1 mg mL^{-1}): Inhibition zones of 30 mm against *P. lilacinum*, 32 mm against *Aspergillus fumigatus* and 25 mm against *Penicillium notatum* were measured (procedure described in Methods section, *Antimicrobial bioassays*) (Supplementary Fig. 21a).



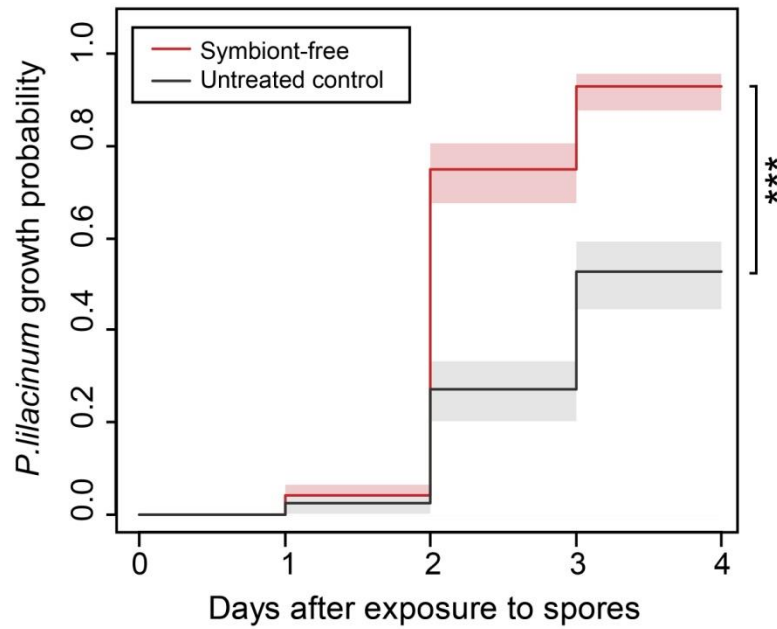
Supplementary Fig. 1. *Burkholderia* is consistently the most abundant taxon in *Lagria villosa* eggs, larvae and the accessory glands associated with the reproductive tract of adult females. Microbial composition as revealed by 454 pyrosequencing of partial 16S rRNA gene sequences carried out on (A) the accessory glands of six field-collected *L. villosa* females and (B) 16 egg clutches laid by field-collected *L. villosa* females, based on 97% similarity OTU clustering as described in the methods section (Microbial community analysis); (C) Quantification of *Burkholderia* symbionts by qPCR using a 172 bp region of the 16S rRNA gene as described in the methods section (Horizontal Transmission experiment) in 15 egg clutches (abundance per individual egg is represented), six larvae between 32 and 43 days old, and eight accessory glands from adult females (abundance for a single gland per individual is represented). The center value of the boxplots represents the median, and the whiskers denote minimum and maximum values.



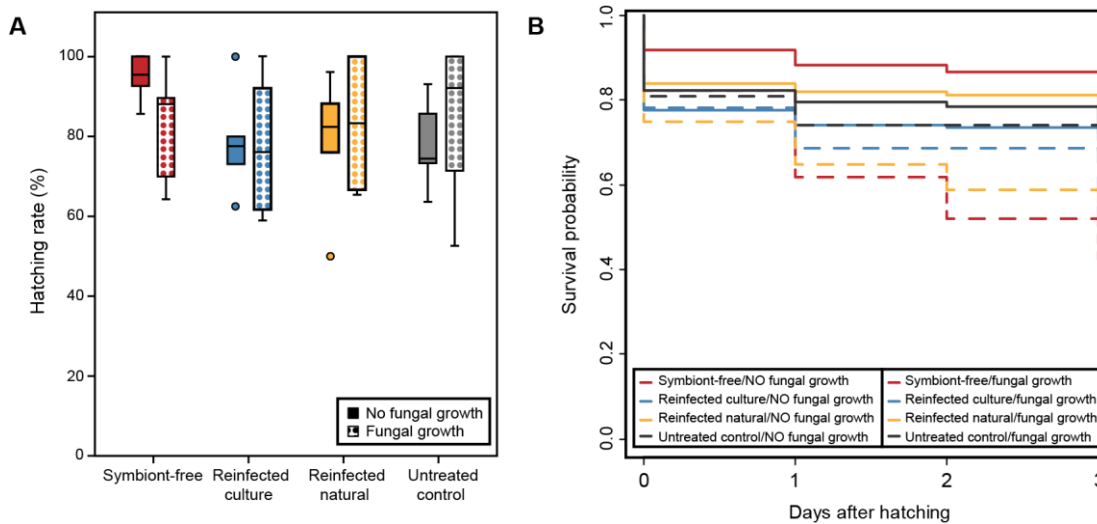
Supplementary Fig. 2. *L. villosa* beetles, as well as the related species *L. hirta*, carry at least three symbiotic *B. gladioli* strains. Phylogenetic reconstruction based on an approximately-maximum-likelihood algorithm of selected *Burkholderia* using partial 16S rRNA gene sequences (1,169 bp), showing the placement of the *L. villosa* - associated strains relative to other *Burkholderia*. Local support values above 0.7 are reported at the nodes. References to sequences extracted from public databases are listed in Supplementary Table 1.



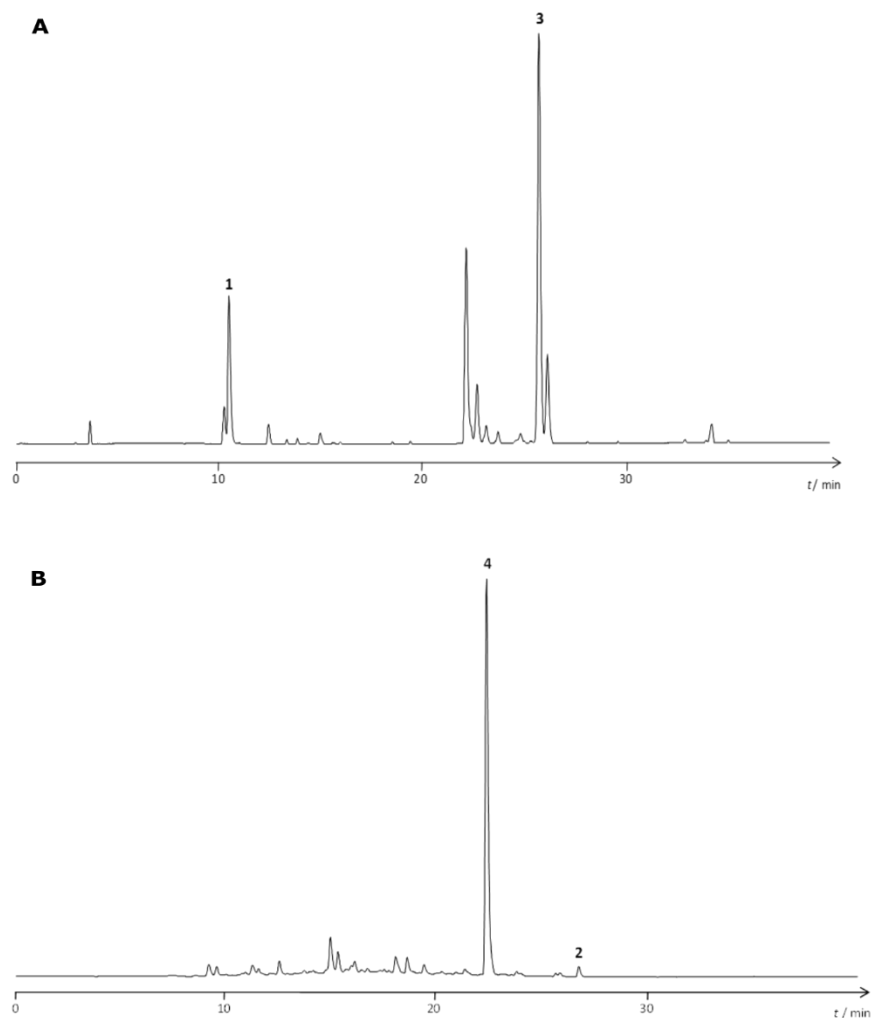
Supplementary Fig. 3. When present, *P. lilacinum* reaches higher biomass on *L. villosa* eggs in the absence of the *Burkholderia* symbionts. Fungal growth was estimated qualitatively during blind monitoring of 720 eggs (180 eggs per treatment from 6 independent clutches) as described in the methods section (*Fungal inhibition on eggs and survival assays*), assigning the level of growth to one of the following categories (0 = no visible growth, 1 = minor growth directly on surface and barely noticeable, 2 = multiple mycelia in contact with surface, 3 = considerable growth on surface, 4 = surface completely covered by mycelia). For statistical analysis, a generalized linear mixed model with a Poisson distribution and clutch as random factor was used (*** $p < 0.001$). The radius of each circle corresponds to the number of eggs in the respective category.



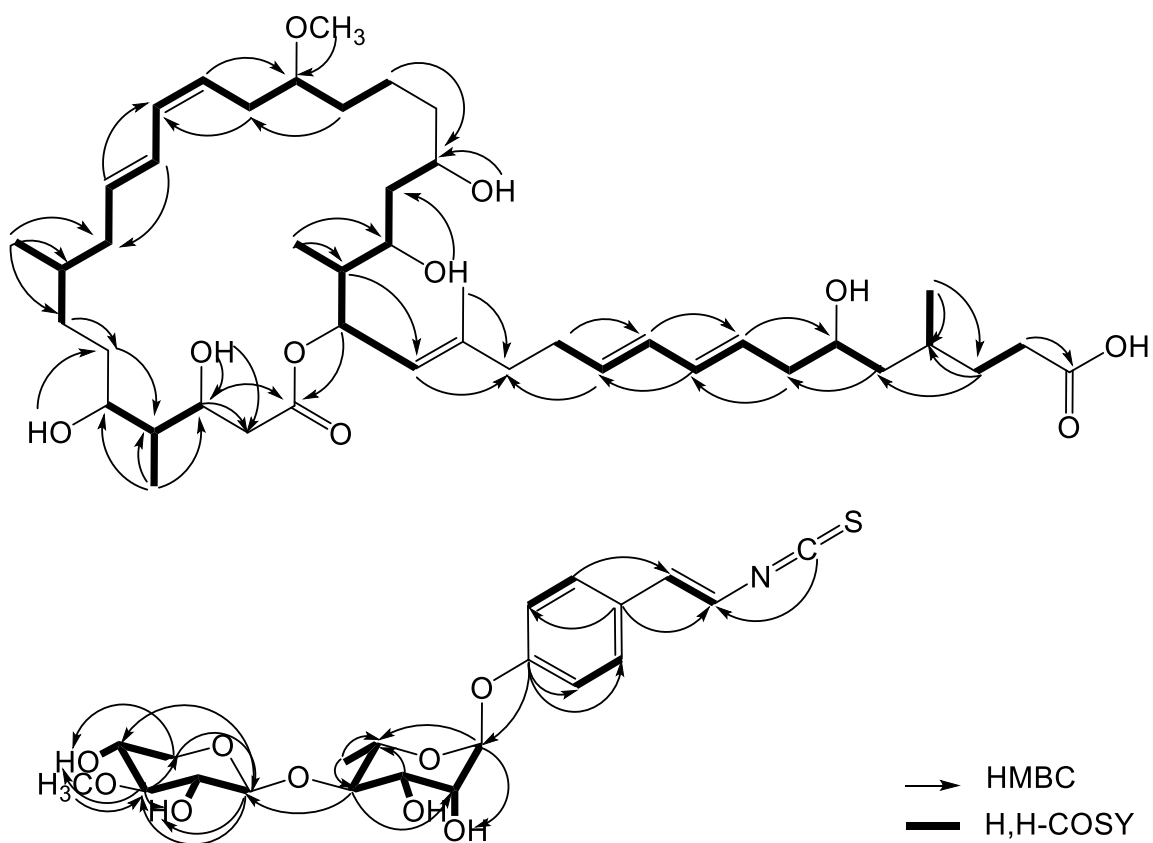
Supplementary Fig. 4. Eggs laid by symbiont-free *L. villosa* females are less protected against *P. lilacinum* fungal growth in comparison to their symbiotic counterparts. Symbiont-free females were obtained by rearing from surface-sterilized eggs, while symbiotic females were taken from the normal beetle culture. For each treatment, six clutches and 30 eggs per clutch were tested as described in the methods section excluding reinfection procedures (*Fungal inhibition on eggs and survival assays*). For statistical analysis, a Cox mixed effects model including clutch as a random factor was used (*** $p < 0.001$). Estimated survival curves (Kaplan-Meier) and the corresponding standard error are plotted.



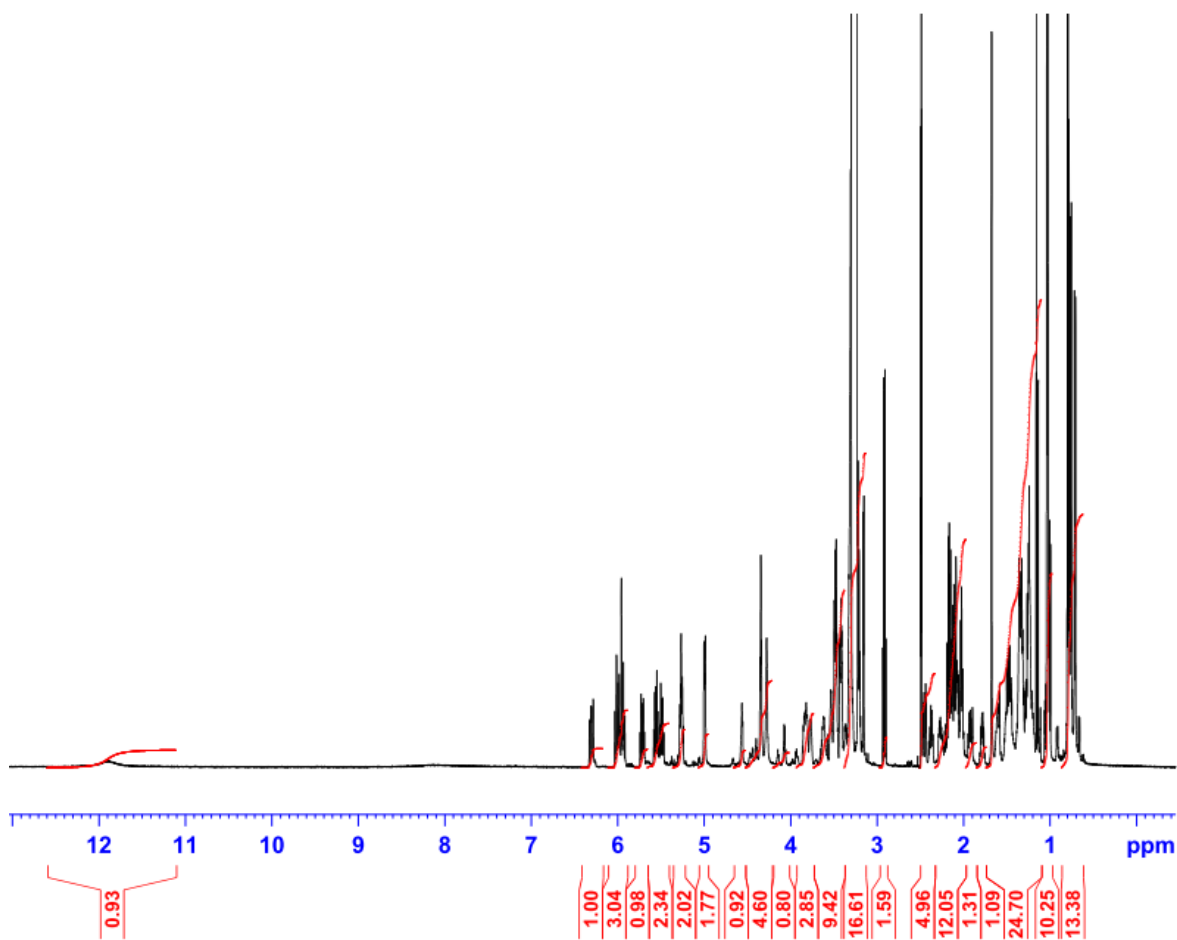
Supplementary Fig. 5. The detrimental effect of *P. lilacinum* growth on *L. villosa* eggs does not significantly reflect on hatching rate, but it causes increased larval mortality in the first days after hatching. (A) There was no statistically significant effect of either treatment or fungal growth level on hatching rate of the six egg clutches (Generalized linear mixed model with a Poisson distribution, $p > 0.05$). The center value of the boxplots represents the median, the whiskers denote minimum and maximum values, and circles represent outliers (more than 1.5 interquartile ranges from nearest box edge). (B) Fungal growth on the eggs has a negative effect on the survival of the larvae during the first days after hatching and affects individuals from the treatments differently, with aposymbionts showing the most pronounced effect (Cox Mixed-Effects Model; Fungus, $p < 0.001$; Treatment:Fungus, $p < 0.05$). Survival curves based on Kaplan-Meier estimates are plotted. For both (A) and (B), sample sizes were as follows: Symbiont-free: no fungal growth 60, fungal growth 118; Reinfected culture: no fungal growth 147, fungal growth 32; Reinfected natural: no fungal growth 111, fungal growth 68; Untreated control: no fungal growth 107, fungal growth 73.



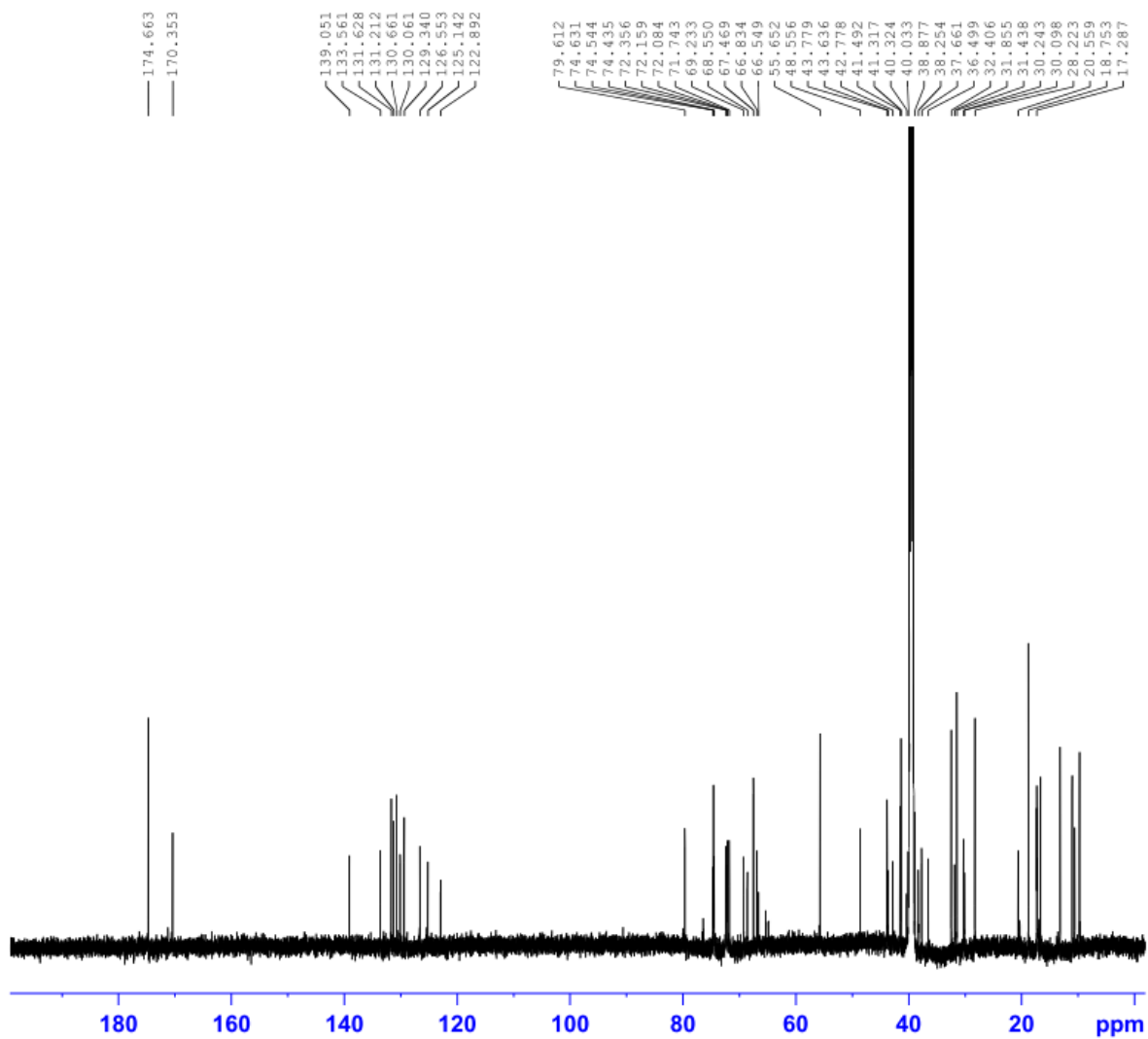
Supplementary Fig. 6. HPLC profiles of crude extracts of *B. gladioli* Lv-StA liquid cultures indicating the production of compounds (1-4) *in vitro*. (A). Chromatogram (PDA total scan) of the crude extract of *B. gladioli* Lv-StA cultured on MGY medium. Numbers indicate production of toxoflavin (1) and lagriene (3). (B) HPLC profile (294 nm) of the crude extract of *B. gladioli* Lv-StA cultured on PDB medium. Numbers indicate production of caryoynencin (2) and sinapigliadioside (4).



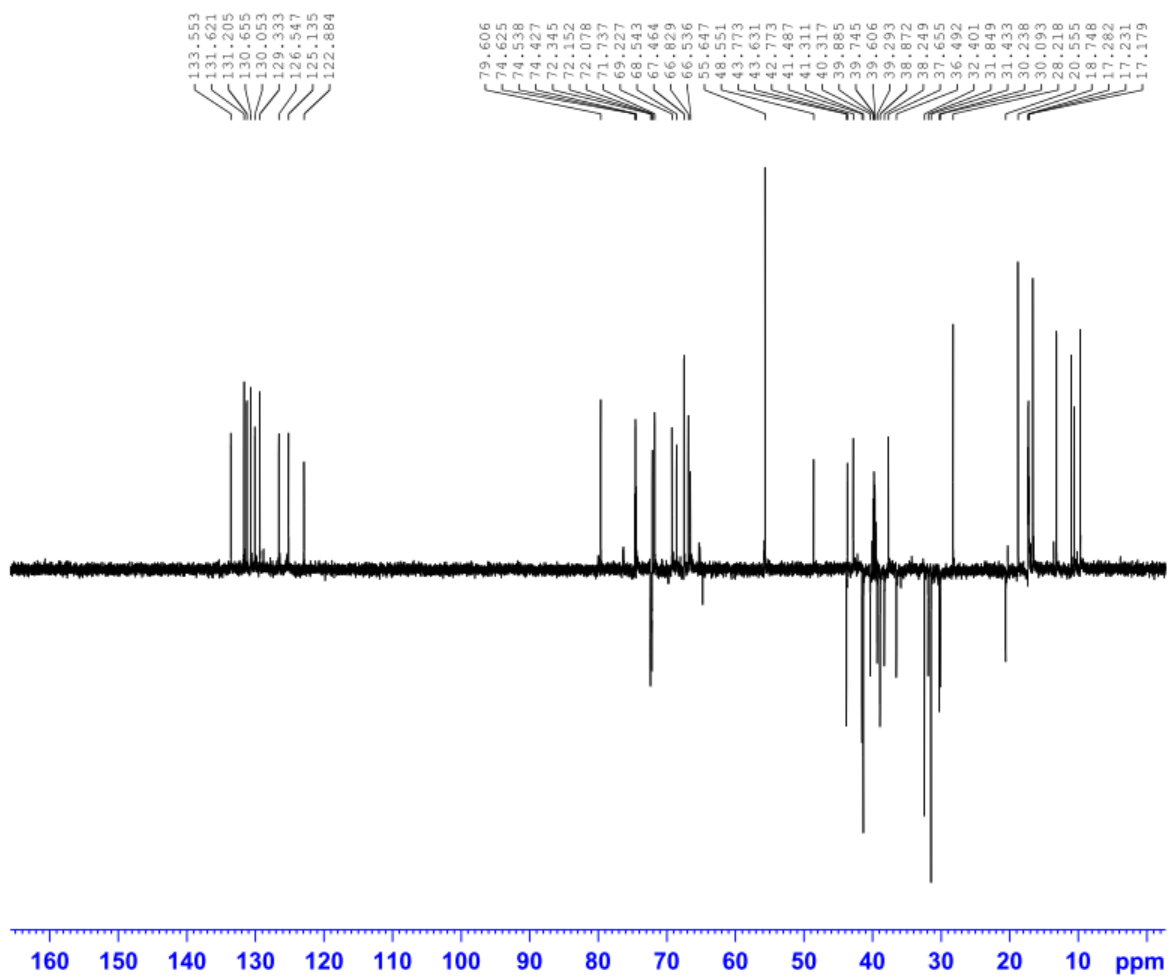
Supplementary Fig. 7. Key 2D NMR couplings of lagriene (3) and sinapigladioside (4).



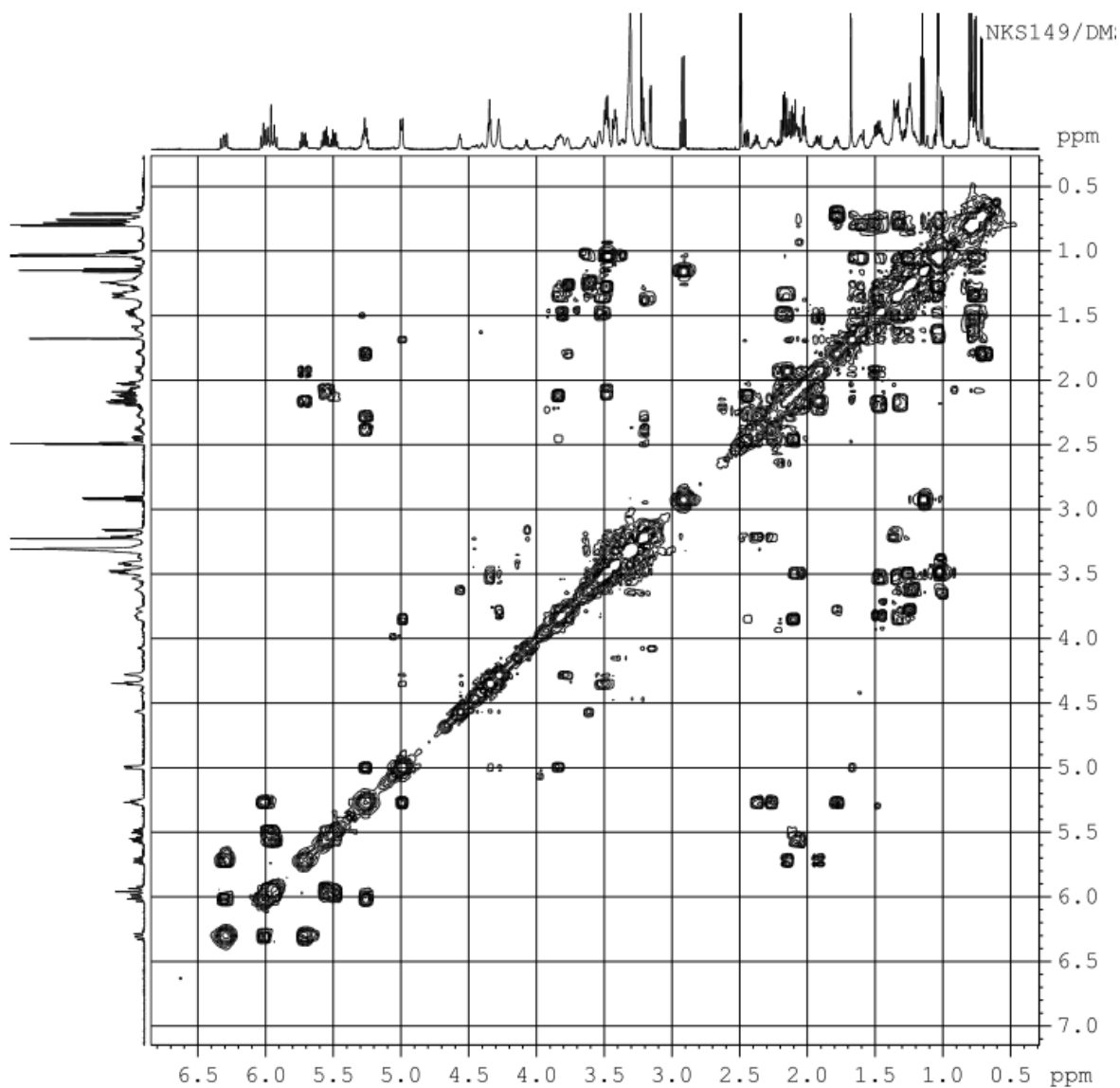
Supplementary Fig. 8. ^1H NMR spectrum of lagriene (3).



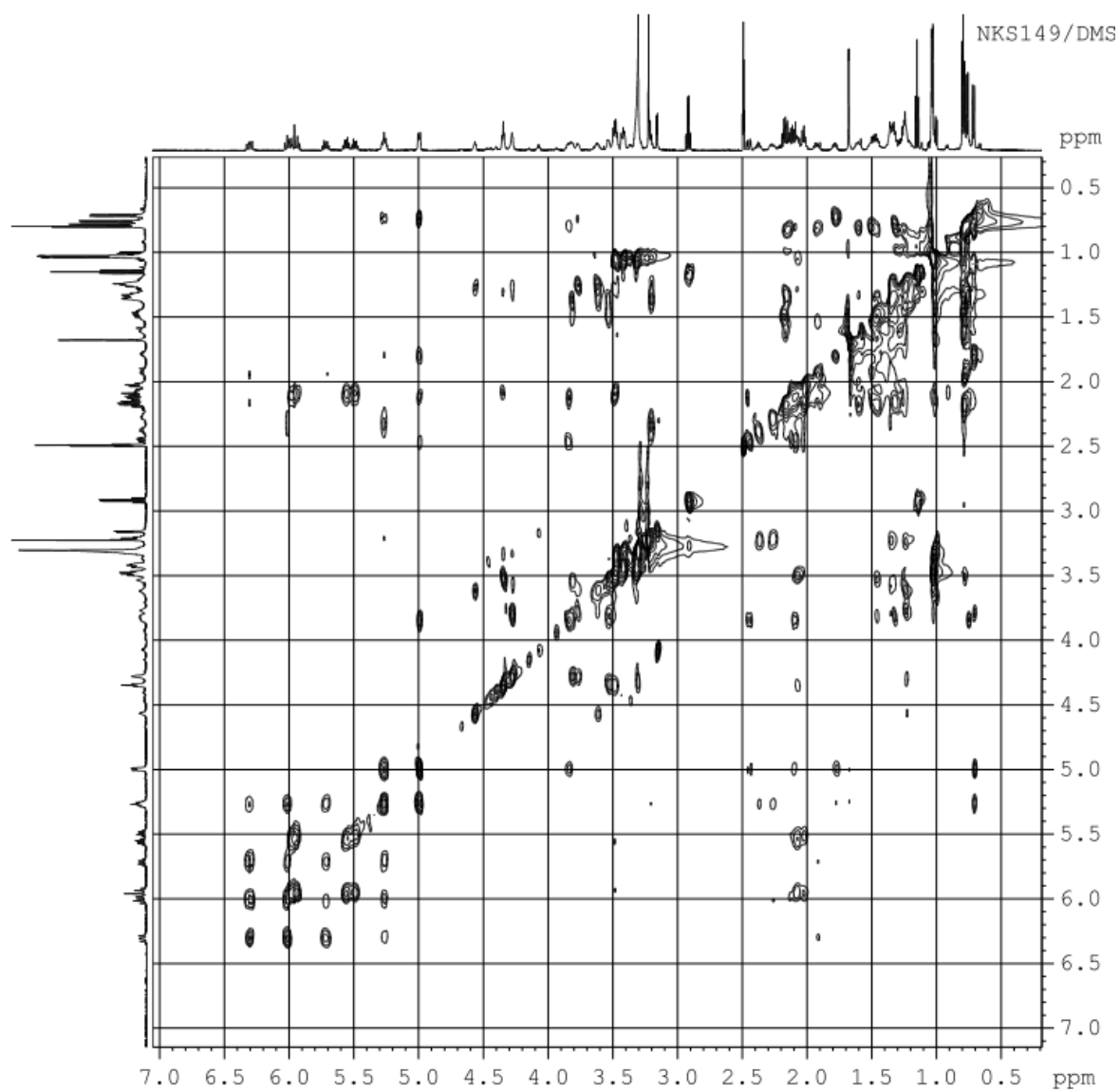
Supplementary Fig. 9. ^{13}C NMR spectrum of lagriene (3).



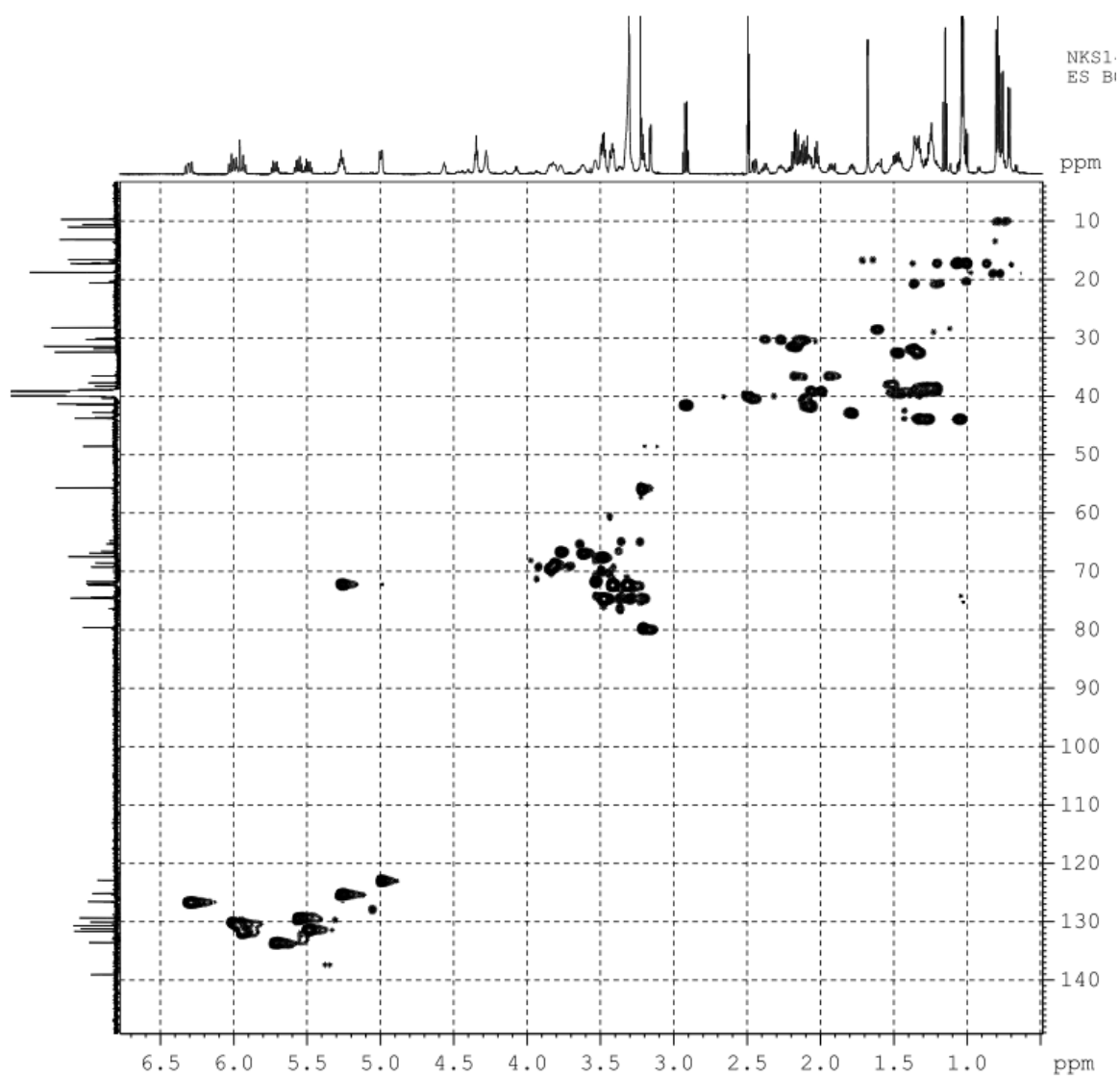
Supplementary Fig. 10. DEPT135 NMR spectrum of lagriene (3).



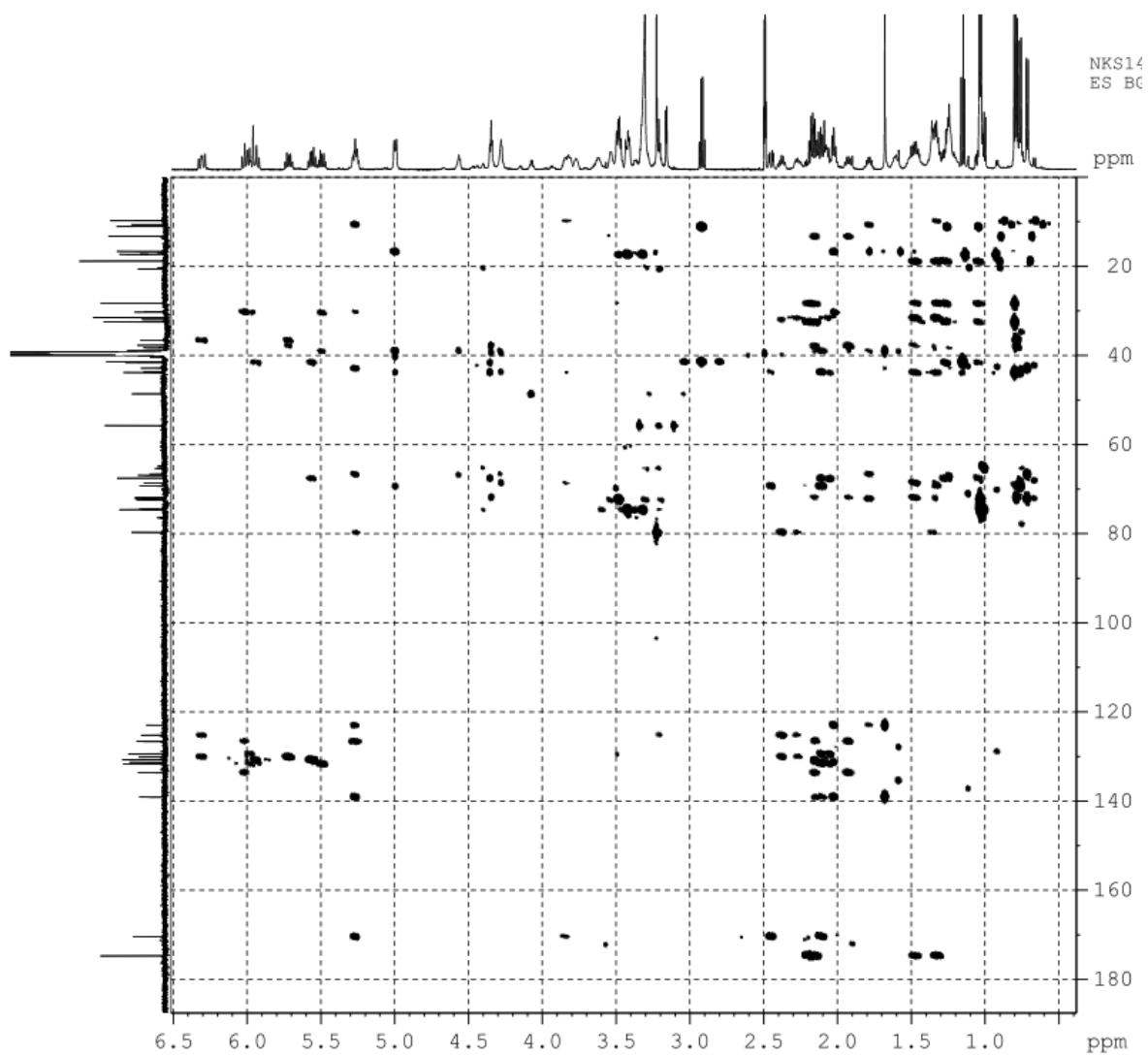
Supplementary Fig. 11. H,H-COSY spectrum of lagriene (3).



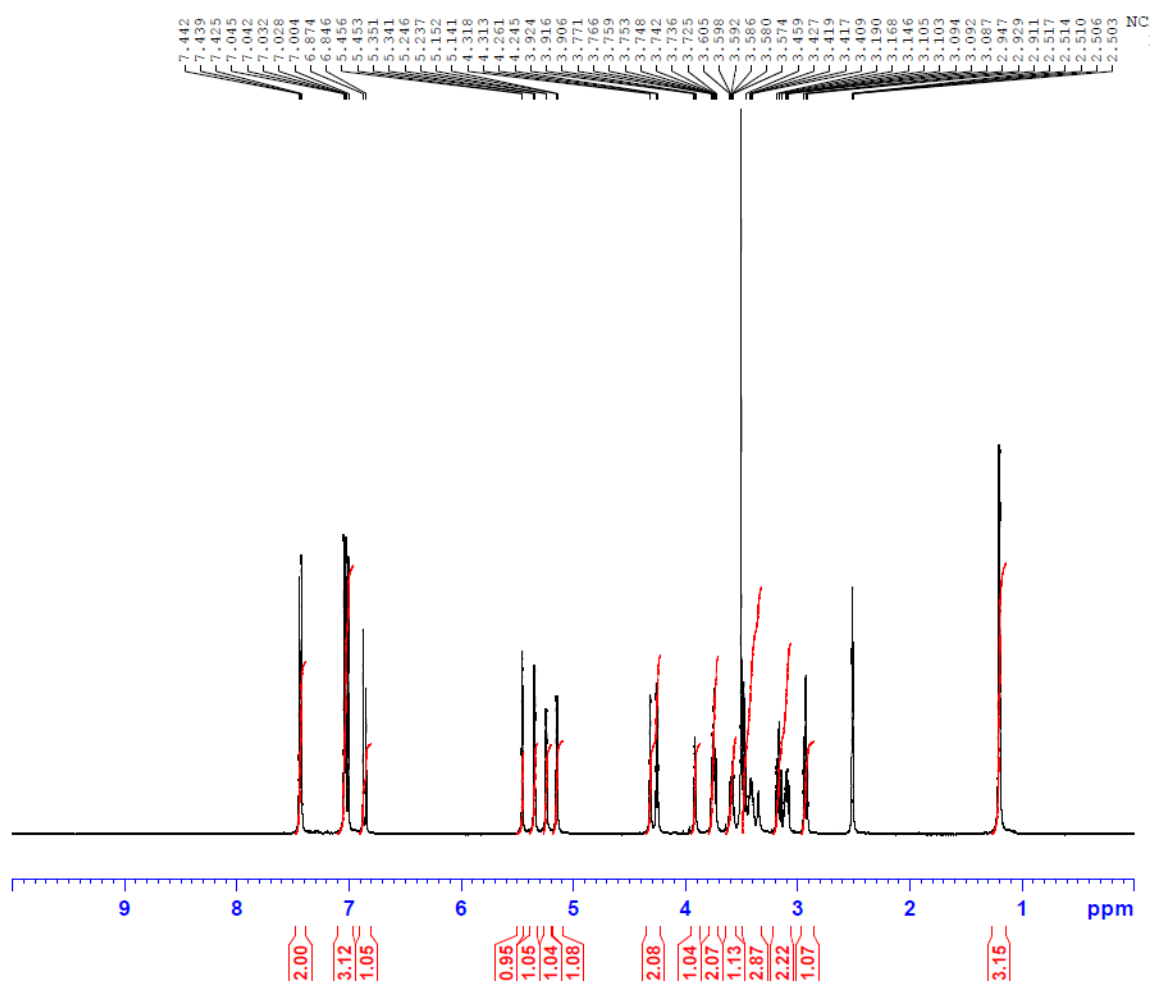
Supplementary Fig. 12. TOCSY spectrum of lagriene (3).



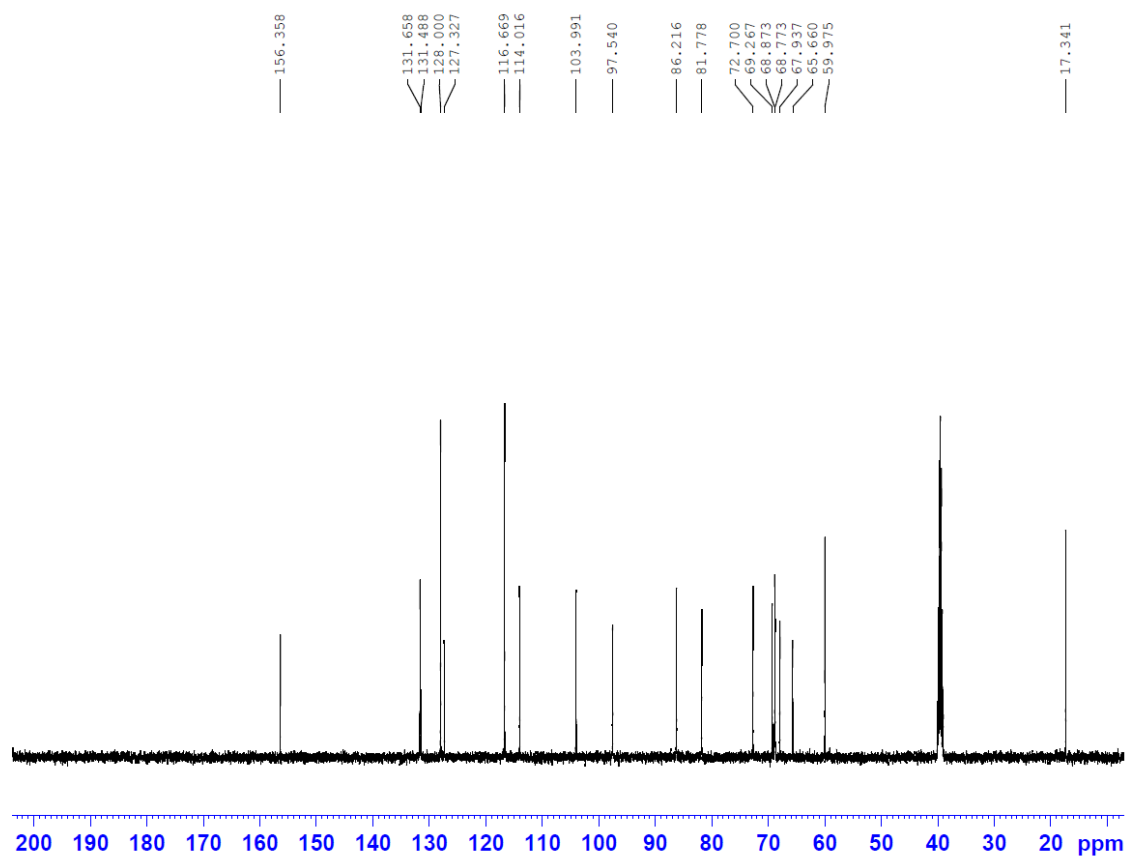
Supplementary Fig. 13. HSQC spectrum of lagriene (**3**).



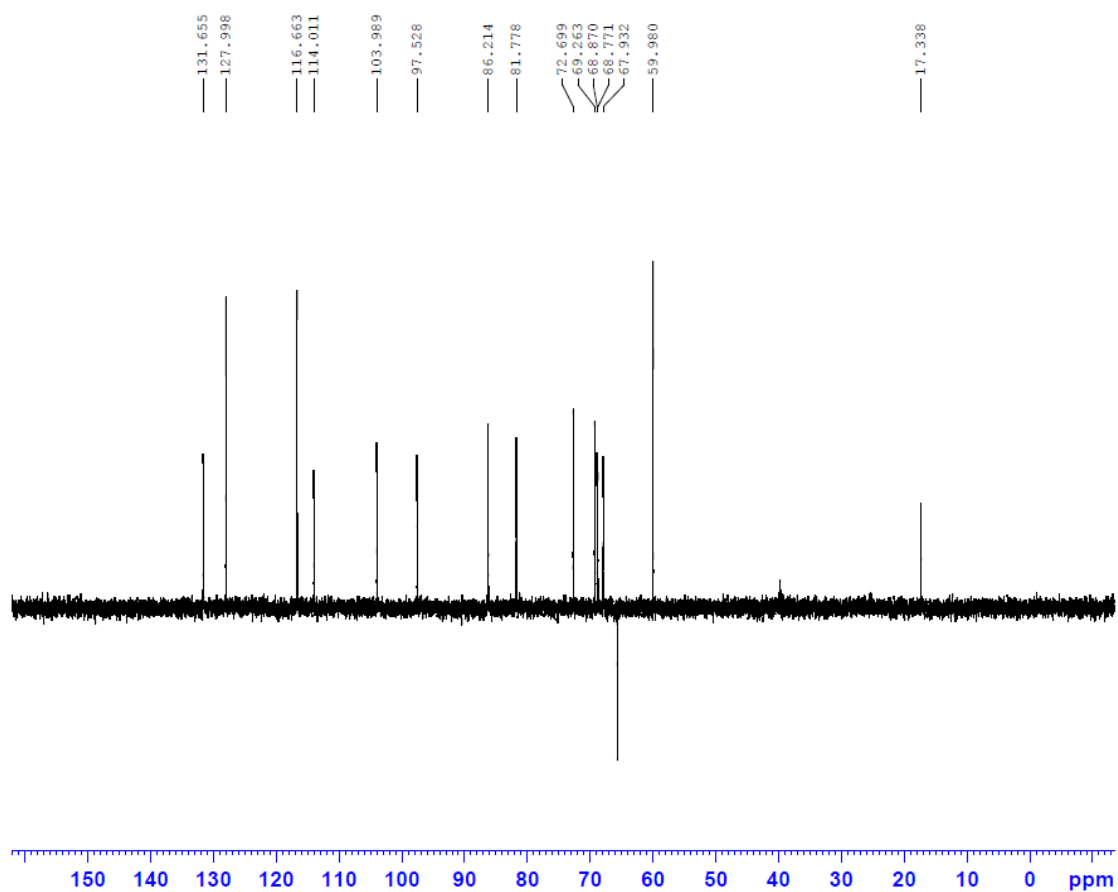
Supplementary Fig. 14. HMBC spectrum of lagriene (3).



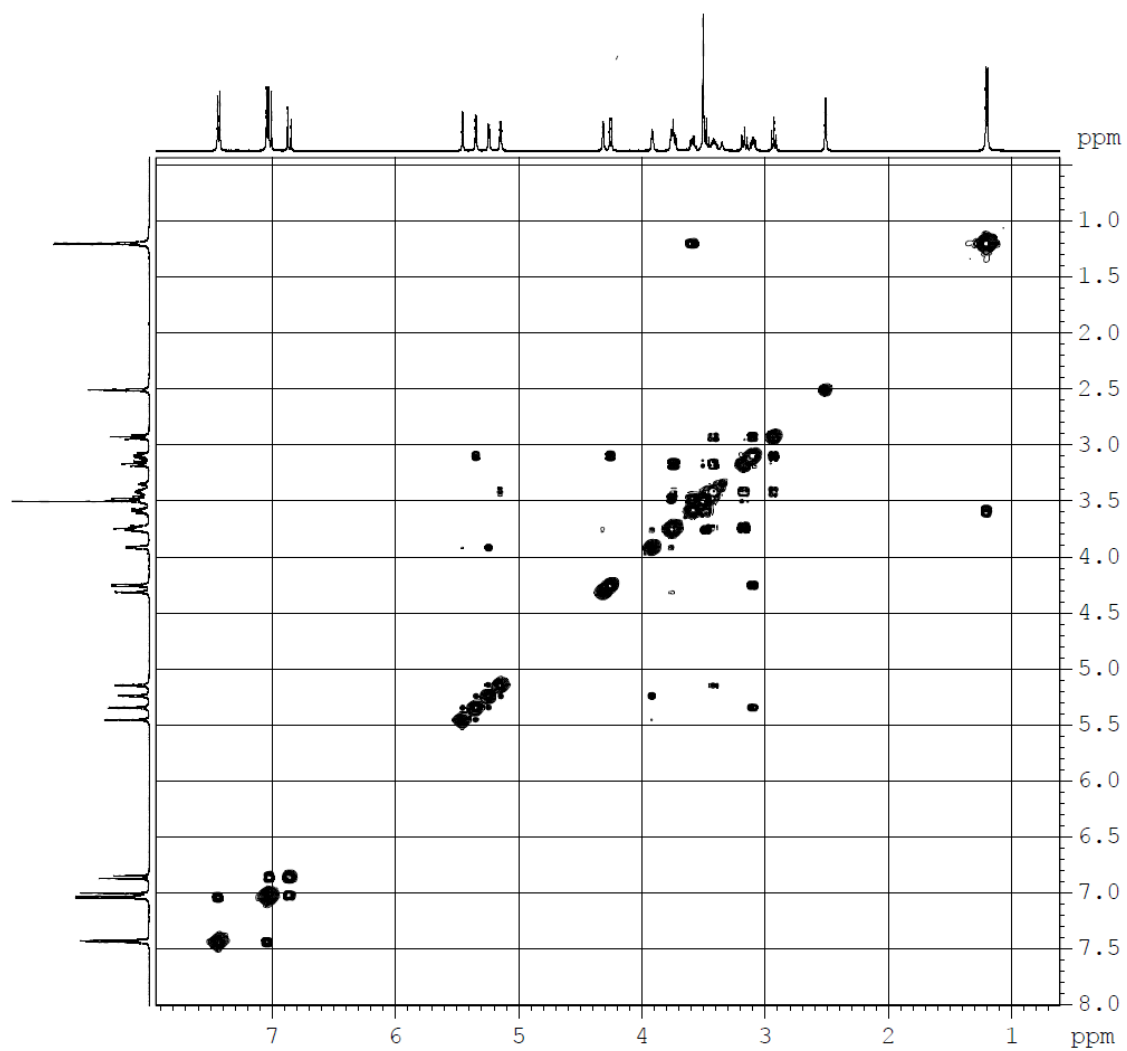
Supplementary Fig. 15. ^1H NMR spectrum of sinapigladioside (4).



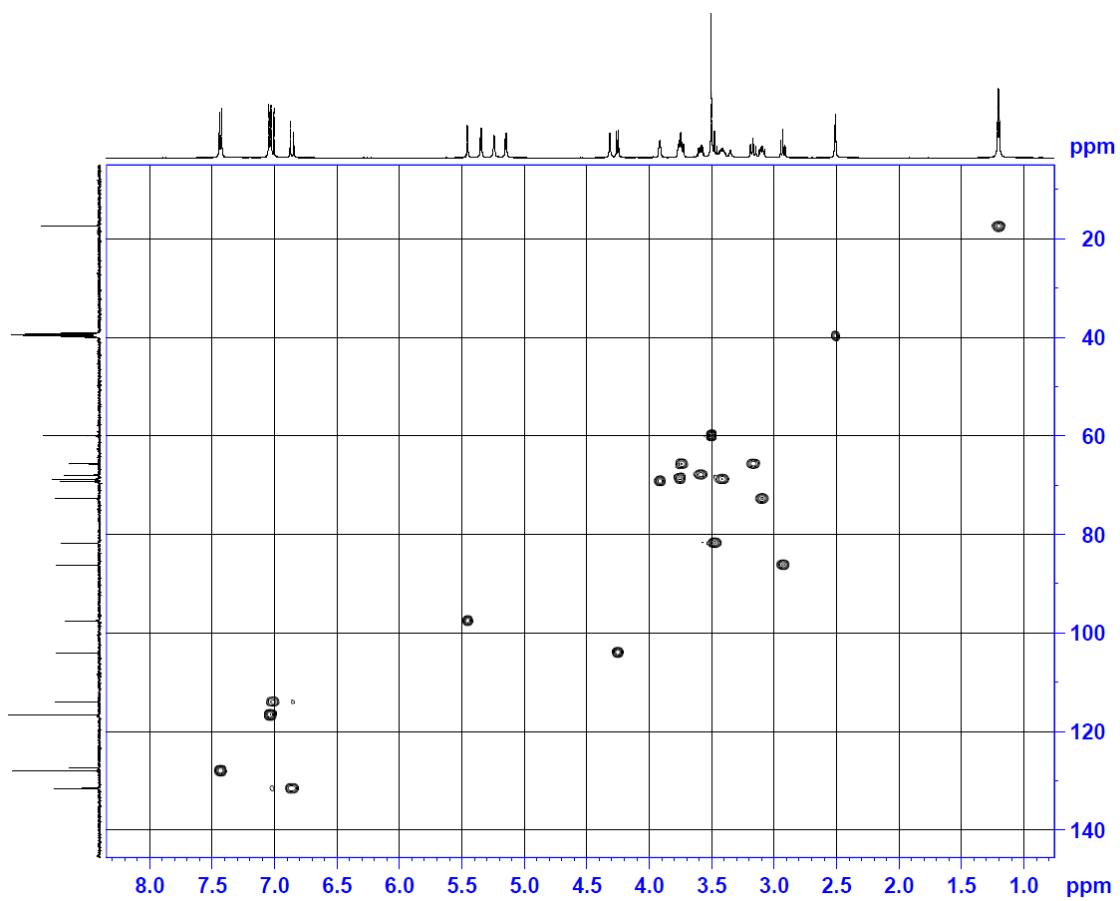
Supplementary Fig. 16. ^{13}C NMR spectrum of sinapigladioside (4).



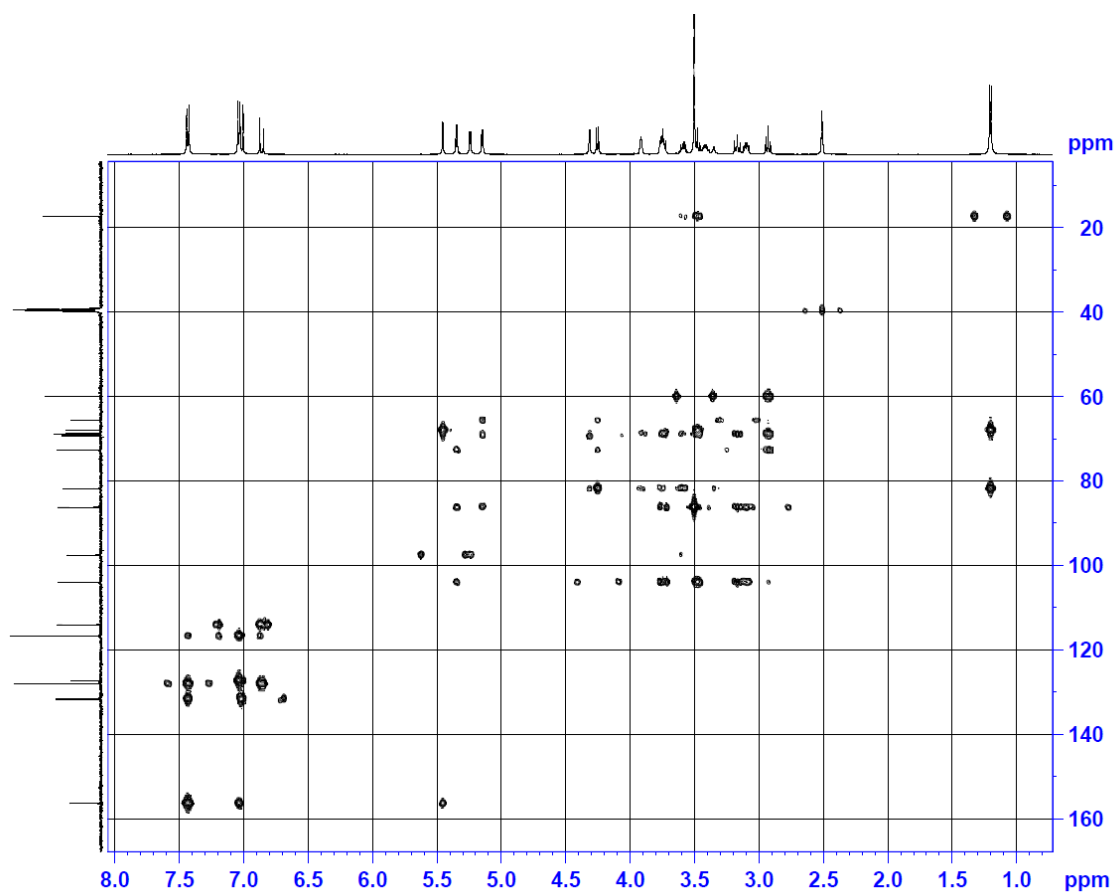
Supplementary Fig. 17. DEPT135 NMR spectrum of sinapigladioside (4).



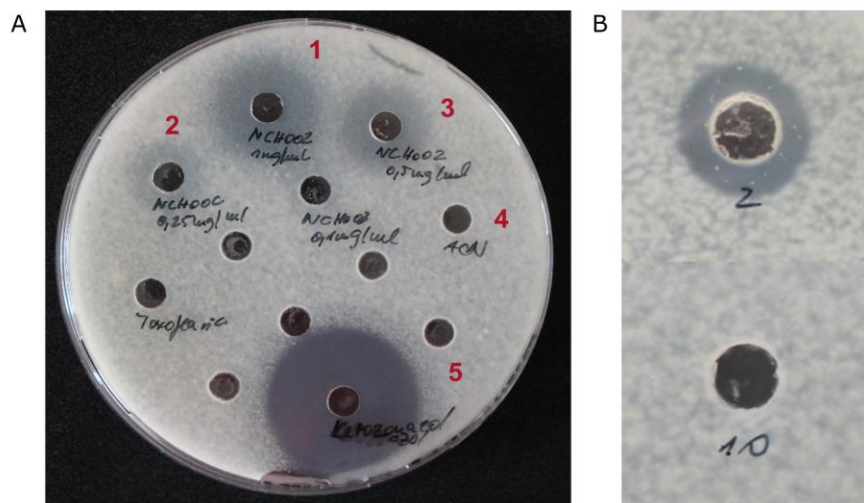
Supplementary Fig. 18. H,H-COSY spectrum of sinapigladioside (4).



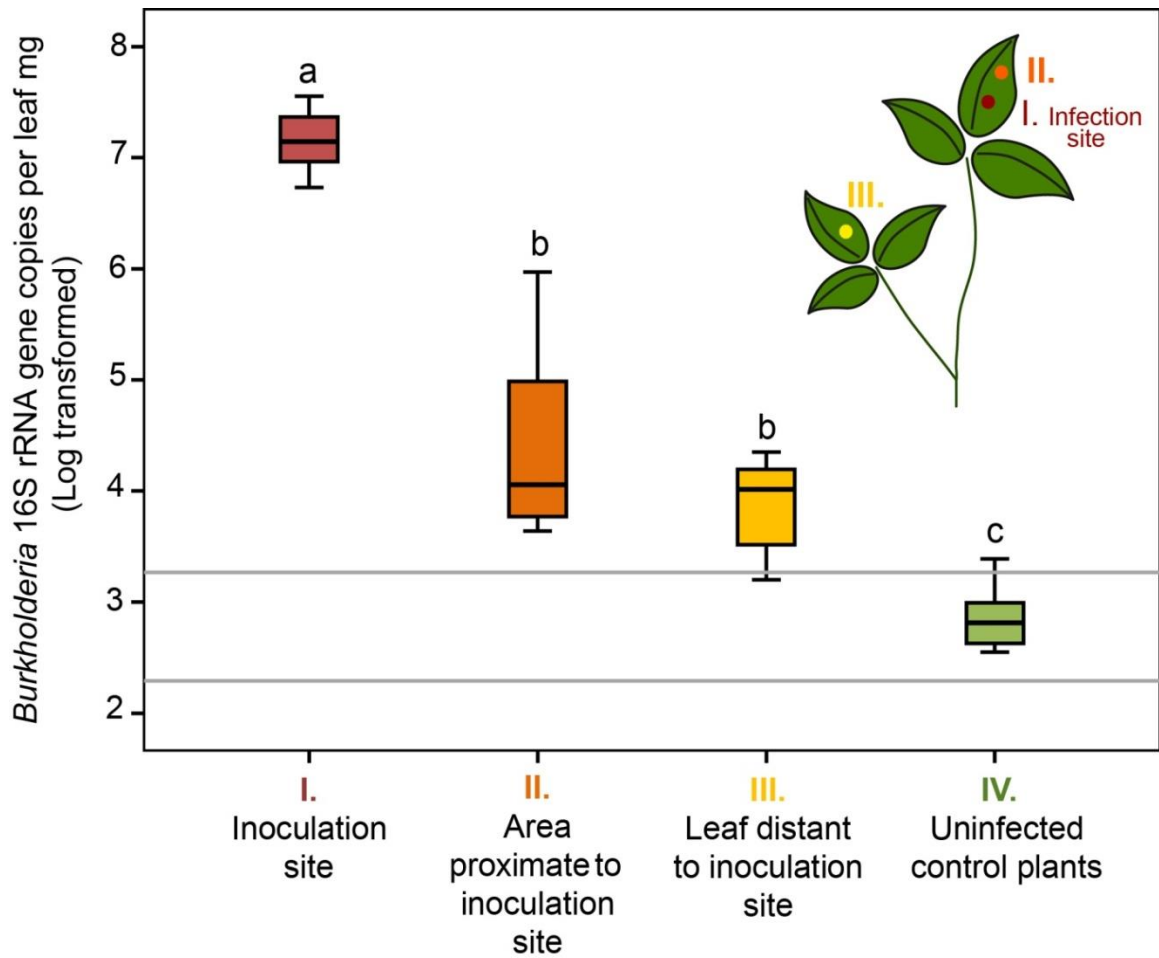
Supplementary Fig. 19. HSQC spectrum of sinapigladioside (4).



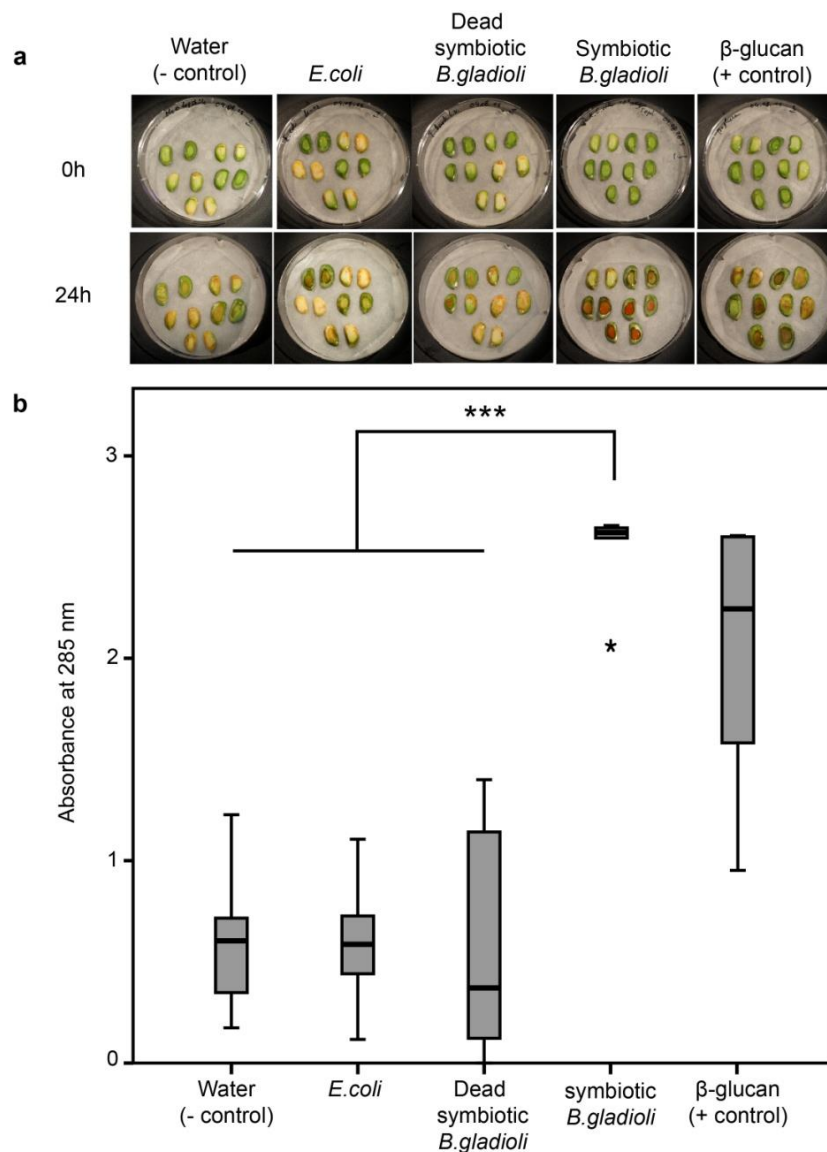
Supplementary Fig. 20. HMBC spectrum of sinapigladioside (4).



Supplementary Fig. 21. The *B. gladioli* symbionts produce metabolites that are active against *Purpureocillium lilacinum*, a natural fungal antagonist of *L. villosa*. (A) Sinapigladioside produced by *L. villosa* StA exhibits antifungal activity against *P. lilacinum*. Wells 1-3 correspond to sinapigladioside at 1, 0.5 and 0.25 mg mL⁻¹ respectively, well 4 to the negative control (solvent = acetonitrile) and well 5 to the positive control (Ketoconazol). (B) A caryoynencin-containing fraction inhibits the growth of *P. lilacinum* *in vitro* (upper picture). Negative control (solvent = methanol) (lower picture).



Supplementary Fig. 22. *Burkholderia* symbionts of *L. villosa* can establish a systemic infection in soybean plants. 38 days after infection, *B. gladioli* Lv-StA were found in higher concentration in plant tissues adjacent to the infection site. Significant titers of bacteria were also detected in areas proximate to the inoculation site, as well as in distant leaves (ANOVA, $n = 18$ per group, $p < 0.001$; Tukey test), demonstrating dispersal within the plant, probably via the vascular tissues. Bacteria were quantified using qPCR of a 167 bp region of the 16S rRNA gene as described in the methods section (*Plant fitness effect upon *B. gladioli* infection*). Gray lines correspond to the maximum and minimum values obtained for negative controls in the same qPCR run. Different letters above boxes represent significant differences according to an ANOVA with Tukey post-hoc tests. The center value of the boxplots represents the median, and the whiskers denote minimum and maximum values.



Supplementary Fig. 23. *Burkholderia* symbionts of *L. villosa* elicit a defense response in soybean cotyledons. Cotyledon assay determining glyceollin production by soybean upon inoculation with *B. gladioli* symbionts isolated from *L. villosa*, in comparison with negative (water, *E. coli* and dead *B. gladioli*) and positive controls (β -glucan). (A) Red coloration after 24h is indicative of glyceollin production as observed in *B. gladioli* and β -glucan treatments; (B) UV-spectrophotometric measurements at 285 nm support significant differences in glyceollin amounts in the different treatments (Kruskal Wallis test with Dunn post-hoc test, $n = 8$ cotyledons per group, *** $p < 0.001$). The center value of the boxplots represents the median, the whiskers denote minimum and maximum values, and the star represents an extreme outlier (more than 3 interquartile ranges from nearest box edge).

Supplementary Table 1. 16S rRNA gene sequences from *Burkholderia* symbionts of Lagriinae, and selected references used for phylogenetic reconstruction (Fig. 2b and Supplementary Fig. 2).

<i>Sequence Id.</i>	<i>Category^a</i>	<i>Strain</i>	<i>Accession</i>
<i>Lagria nigricollis</i> Symbiont	Lagriinae symbiont	Ln	KT888028
<i>Lagria okinawana</i> Symbiont	Lagriinae symbiont	Lo	KT888029
<i>Lagria rufipennis</i> Symbiont	Lagriinae symbiont	Lr	KT888030
<i>Lagria villosa</i> Symbiont	Lagriinae symbiont	Lv-StA	KT888027
<i>Lagria villosa</i> Symbiont	Lagriinae symbiont	Lv-StB	KU358661
<i>Lagria villosa</i> Symbiont	Lagriinae symbiont	Lv-StC	KU358661
<i>Lagria hirta</i> Symbiont	Lagriinae symbiont	Lh-StG	KT888026
<i>Lagria hirta</i> Symbiont	Lagriinae symbiont	Lh-StH	KU574041
<i>Lagria hirta</i> Symbiont	Lagriinae symbiont	Lh-StM	KU574046
<i>Ecnolagria</i> sp. Symbiont	Lagriinae symbiont	Ec	KT888031
<i>Burkholderia gladioli</i> BSR3 - Rice pathogen	Plant pathogen - <i>B.gladioli</i> clade	BSR3	NR_102847
<i>Burkholderia gladioli</i> MS102 - Corn pathogen	Plant pathogen - <i>B.gladioli</i> clade	MS 102	EU053154
<i>Burkholderia gladioli</i> CH-2 - Onion pathogen	Plant pathogen - <i>B.gladioli</i> clade	CH-2	AY500138
<i>Burkholderia gladioli</i> st3 - Orchid pathogen	Plant pathogen - <i>B.gladioli</i> clade	strain 3	DQ090078
<i>Burkholderia gladioli</i> - Rice pathogen	Plant pathogen - <i>B.gladioli</i> clade	321gr-6	DQ355169
<i>Burkholderia gladioli</i> pv. <i>agaricicola</i>	Fungal pathogen - <i>B.gladioli</i> clade	CFBP 3580	GU936678
<i>Burkholderia gladioli</i> pv. <i>alliicola</i>	Plant pathogen - <i>B.gladioli</i> clade	CFBP 2422	GU936679
<i>Burkholderia gladioli</i> pv. <i>gladioli</i>	Plant pathogen - <i>B.gladioli</i> clade	CFBP 2427	GU936677
<i>Burkholderia plantarii</i>	Plant pathogen	-	U96933
<i>Burkholderia plantarii</i> - 2396	Plant pathogen	2396	AB183679
<i>Burkholderia plantarii</i> - NBRC104888	Plant pathogen	NBRC104888	AB682222.1
<i>Burkholderia glumae</i> 336gr-1 - Rice pathogen	Plant pathogen	336gr-1	DQ355164.1
<i>Burkholderia glumae</i> - 99gr-4b - Rice pathogen	Plant pathogen	99gr-4b	DQ355167
<i>Burkholderia glumae</i> - PA27.4 - Rice pathogen	Plant pathogen	PA27.4	EF193641.1
<i>Burkholderia caryophylli</i>	Plant pathogen	ATCC 25418	AB021423
<i>Burkholderia endofungorum</i>	Fungal endosymbiont- plant pathogen	HKI 456T	AM420302
<i>Burkholderia rhizoxinica</i>	Fungal endosymbiont- plant pathogen	HKI 454	AJ938142
<i>Burkholderia cepacia</i>	Plant and animal pathogen - BCC	ATCC 25416	U96927
<i>Burkholderia vietnamiensis</i>	Opportunistic animal pathogen - BCC	LMG 10929	AF097534
<i>Burkholderia oklahomensis</i>	Animal pathogen - Pseudomallei group	C6786	DQ108388
<i>Burkholderia fungorum</i>	PBE	LM16225	AF215705
<i>Burkholderia kururiensis</i>	PBE	-	AB024310
<i>Burkholderia nodosa</i>	PBE	Br3437	AY773189
<i>Burkholderia terrae</i>	PBE	KMY02	AB201285
'Ca. <i>Burkholderia kirkii</i> '	PBE	835462	AF475068
<i>Burkholderia</i> sp. [<i>P.antennata</i> symbiont]	Insect-associated	PAN136	AB558189
<i>Burkholderia</i> sp. [<i>P.bicoloripes</i> symbiont]	Insect-associated	PBI_clone1	AB558203.1
<i>Burkholderia</i> sp. [<i>R. pedestris</i> symbiont]	Insect-associated	RPE64	AB558208
<i>Burkholderia</i> sp. [<i>Tetraponera binghami</i> gut isolate]	Insect-associated	-	AF459796
<i>Ralstonia pickettii</i>	Outgroup	12J	NC010678

^aExcept for Lagriinae symbionts and insect-associated strains, the categorization is based on previous studies^{1,2}

Supplementary Table 2. Toxoflavin (1) (*tox*) biosynthetic genes and predicted functions of proteins.

Gene	Length (bp)	Putative protein	Homologous protein	Accession number	Identity/ Similarity
<i>toxA</i>	612	S-Adenosylmethionine- dependent methyltransferase	ToxA (<i>B. glumae</i>)	BAA92862.2	98%/98%
<i>toxB</i>	543	GTP cyclohydrolase II	ToxB (<i>B. glumae</i>)	BAB88913.1	98%/98%
<i>toxC</i>	1692	WD-repeat protein	ToxC (<i>B. glumae</i>)	BAB88914.2	97%/98%
<i>toxD</i>	933	TRP-2	ToxD (<i>B. glumae</i>)	BAB88915.1	89%/91%
<i>toxE</i>	1170	Deaminase	ToxE (<i>B. glumae</i>)	BAB88916.2	92%/94%
<i>toxR</i>	600	LysR family transcriptional regulator	ToxR (<i>B. glumae</i>)	BAC77727.1	88%/90%

Supplementary Table 3. Caryoynencin (2) (*cay*) biosynthetic genes and predicted functions of proteins.

Gene	Length (bp)	Putative protein	Homologous protein	Accession number	Identity/ Similarity
<i>cayA</i>	1557	Fatty acyl-AMP ligase	CayA (<i>B. caryophylli</i>)	AIG53814.1	76%/85%
<i>cayB</i>	930	Fatty acid desaturase	CayB (<i>B. caryophylli</i>)	AIG53817.1	85%/89%
<i>cayC</i>	975	Fatty acid desaturase	CayC (<i>B. caryophylli</i>)	AIG53820.1	85%/91%
<i>cayD</i>	324	Phosphopantetheine attachment site	CayD (<i>B. caryophylli</i>)	AIG53823.1	81%/93%
<i>cayE</i>	1101	Fatty acid desaturase	CayE (<i>B. caryophylli</i>)	AIG53826.1	88%/94%
<i>cayF</i>	975	Alpha/Beta hydrolase	CayF (<i>B. caryophylli</i>)	AIG53829.1	74%/83%
<i>cayG</i>	1197	Cytochrome P450 monooxygenase	CayG (<i>B. caryophylli</i>)	AIG53832.1	79%/90%

Supplementary Table 4. Lagriene (3) (*lag*) biosynthetic genes and predicted functions of proteins.

Gene	Length (bp)	Description	Domains
<i>lagD</i>	16284	Polyketide synthase	KS-ACP-ACP ER KS-KR-ACP KS-DH-KR-ACP KS-DH-KR
<i>lagE</i>	11178	Polyketide synthase	ACP KS-KR-ACP KS-ACP-ACP KS-KR
<i>lagF</i>	9927	Polyketide synthase	MT-ACP KS-KR-ACP KS-KR-ACP KS
<i>lagG</i>	14991	Polyketide synthase	DH-KR-ACP KS-KR-ACP KS-MT-ACP KS-KR-ACP KS
<i>lagH</i>	10800	Polyketide synthase	DH-ACP KS-KR-ACP KS-DH-KR-MT-ACP
<i>lagI</i>	17523	Polyketide synthase	ER KS-KR-ACP KS-KR-MT-ACP KS-KR-ACP KS-KR TE

Supplementary Table 5. Proteins encoded upstream and downstream of the *lagD-lagI* gene cluster.

Gene	Length (bp)	Putative protein	Homologous protein	Accession number	Identity/ Similarity
<i>orf-16</i>	1353	MATE efflux protein	sce3194 (<i>Sorangium cellulosum</i> Soce56)	CAN93353.1	68%/82%
<i>orf-15</i>	975	Malonyl CoA-acyl carrier protein transacylase	sce3195 (<i>Sorangium cellulosum</i> So ce56)	CAN93354.1	47%/63%
<i>orf-14</i>	855	4'-Phosphopantetheinyl transferase	sce5058 (<i>Sorangium cellulosum</i> So ce56)	CAN95221.1	41%/56%
<i>orf-13</i>	1173	Malonyl CoA-acyl carrier protein transacylase	Malonyl CoA-acyl carrier protein transacylase (<i>Burkholderia pseudomallei</i>)	3G87_A	51%/62%
<i>orf-12</i>	1401	Amidase	sce3176 (<i>Sorangium cellulosum</i> So ce56)	CAN93335.1	56%/69%
<i>orf-11</i>	360	Alpha/beta-hydrolase	sce3177 (<i>Sorangium cellulosum</i> So ce56)	CAN93336.1	58%/76%
<i>orf-10</i>	3234	Hypothetical protein	SorM (<i>Sorangium cellulosum</i> So ce12)	ADN68497.1	38%/51%
<i>orf-9</i>	747	Enoyl-CoA hydratase	sce3179	CAN93338.1	75%/83%
<i>orf-8</i>	789	Enoyl-CoA hydratase	(<i>Sorangium cellulosum</i> So ce56) sce3180	CAN93339.1	63%/78%
<i>orf-7</i>	1260	3-Hydroxy-3-methylglutaryl CoA synthase	(<i>Sorangium cellulosum</i> So ce56) sce3181/EtnO	CAN93340.1	76%/86%
<i>orf-6</i>	1227	Beta-ketoacyl synthase	(<i>Sorangium cellulosum</i> So ce56) sce3182/EtnP	CAN93341.1	64%/76%
<i>orf-5</i>	246	Acyl carrier protein	(<i>Sorangium cellulosum</i> So ce56) sce3183	CAN93342.1	68%/81%
<i>orf-4</i>	1395	Malonyl CoA-acyl carrier protein transacylase	(<i>Sorangium cellulosum</i> So ce56) sce3184	CAN93343.1	68%/82%
<i>orf-3</i>	264	Acyl carrier protein	(<i>Sorangium cellulosum</i> So ce56) sce3185	CAN93344.1	60%/81%
<i>orf-2</i>	609	Malonyl CoA-acyl carrier protein transacylase	(<i>Sorangium cellulosum</i> So ce56) sce3186/EtnB	CAN93345.1	55%/70%
<i>orf-1</i>	1968	Asparagine synthase	(<i>Sorangium cellulosum</i> So ce56) sce3187/EtnC (<i>Sorangium cellulosum</i> So ce56)	CAN93346.1	66%/79%
<i>orf+1</i>	486	Hypothetical protein	-	-	-
<i>orf+2</i>	924	Hypothetical protein	-	-	-
<i>orf+3</i>	609	Glutathione-S-transferase	Glutathione-S-transferase (<i>Yersinia pestis</i>)	4G9H_A	47%/65%
<i>orf+4</i>	954	LysR family transcriptional regulator	LysR family transcriptional regulator (<i>Neisseria meningitidis</i>)	3HHG_A	34%/54%
<i>orf+5</i>	2124	TonB-dependent siderophore receptor	TonB-dependent siderophore receptor (<i>Pseudomonas fluorescens</i>)	3QLB_A	24%/37%
<i>orf+6</i>	858	Molybdate ABC transporter	Molybdate-binding protein (<i>Xanthomonas axonopodis</i> pv. <i>citri</i>)	3GZG_A	26%/39%
2400/ <i>orf+7</i>	1092	Oxidoreductase	Luciferase-like monooxygenase (<i>Bacillus cereus</i>)	3RAO_A	36%/54%

Supplementary Table 6. Microbial strains used for *in vivo* and *in vitro* activity bioassays.

Species	Strain	Source	Location	Accession Number/ Database Identifier	Database
<i>Purpureocillium lilacinum</i>	LV1	<i>Lagria villosa</i> egg	MPI-CE, Jena, Germany	KY630747, KY630748, KY630749	GenBank
<i>Trichoderma harzianum</i>	LESF555	<i>Achromyrmex heyeri</i> fungal garden	Sentinela do Sul, Rio Grande do Sul, Brazil	KT279018, KT278950, KT278904	GenBank
<i>Beauveria bassiana</i>	LESF477	<i>Atta</i> sp. female alate	Botucatu, Sao Paulo, Brazil	CRM 1216	CRM - UNESP
<i>Candida albicans</i>	BMSY 212	Jena Microbial Resource Collection	Jena, Germany	STI50163	JMRC - Friedrich Schiller University
<i>Penicillium notatum</i>	JP 36	Jena Microbial Resource Collection	Jena, Germany	SF011781	JMRC - Friedrich Schiller University
<i>Aspergillus fumigatus</i>	ATCC46645	Jena Microbial Resource Collection	Jena, Germany	STH00450	JMRC - Friedrich Schiller University
<i>Sporobolomyces salmonicolor</i>	SBUG 549	Jena Microbial Resource Collection	Jena, Germany	ST035974	JMRC - Friedrich Schiller University
<i>Bacillus thuringiensis</i>	DSM2048	Jena Microbial Resource Collection	Jena, Germany	STI11409	JMRC - Friedrich Schiller University
<i>Bacillus subtilis</i>	ATCC 6633	Jena Microbial Resource Collection	Jena, Germany	STI10880	JMRC - Friedrich Schiller University
<i>Brevibacillus laterosporus</i>	ATCC 31932	Jena Microbial Resource Collection	Jena, Germany	STI11381	JMRC - Friedrich Schiller University
<i>Staphylococcus aureus</i>	SG 511	Jena Microbial Resource Collection	Jena, Germany	STI10760	JMRC - Friedrich Schiller University
<i>Escherichia coli</i>	SG 458	Jena Microbial Resource Collection	Jena, Germany	ST033699	JMRC - Friedrich Schiller University
<i>Pseudomonas aeruginosa</i>	SG 137	Jena Microbial Resource Collection	Jena, Germany	ST033772	JMRC - Friedrich Schiller University
<i>Pseudomonas aeruginosa</i>	K 799/61	Jena Microbial Resource Collection	Jena, Germany	ST033771	JMRC - Friedrich Schiller University
<i>Staphylococcus aureus</i>	134/93 (MRSA)	Jena Microbial Resource Collection	Jena, Germany	STH00435	JMRC - Friedrich Schiller University
<i>Enterococcus faecalis</i>	1528 (VRE)	Jena Microbial Resource Collection	Jena, Germany	ST033700	JMRC - Friedrich Schiller University
<i>Mycobacterium vaccae</i>	IMET 10670	Jena Microbial Resource Collection	Jena, Germany	STI10670	JMRC - Friedrich Schiller University

Supplementary References

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