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

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A pair of effectors encoded on a conditionally dispensable chromosome of *Fusarium oxysporum* suppress host-specific immunity

Yu Ayukawa^{1,2,7}, Shuta Asai^{1,3,7}, Pamela Gan¹, Ayako Tsushima^{1,5}, Yasunori Ichihashi^{1,6}, Arisa Shibata¹, Ken Komatsu², Petra M. Houterman⁴, Martijn Rep⁴, Ken Shirasu¹ & Tsutomu Arie²

Many plant pathogenic fungi contain conditionally dispensable (CD) chromosomes that are associated with virulence, but not growth in vitro. Virulence-associated CD chromosomes carry genes encoding effectors and/or host-specific toxin biosynthesis enzymes that may contribute to determining host specificity. *Fusarium oxysporum* causes devastating diseases of more than 100 plant species. Among a large number of host-specific forms, *F. oxysporum* f. sp. *conglutinans* (*Focn*) can infect Brassicaceae plants including Arabidopsis (*Arabidopsis thaliana*) and cabbage. Here we show that *Focn* has multiple CD chromosomes. We identified specific CD chromosomes that are required for virulence on Arabidopsis, cabbage, or both, and describe a pair of effectors encoded on one of the CD chromosomes that is required for suppression of Arabidopsis-specific phytoalexin-based immunity. The effector pair is highly conserved in *F. oxysporum* isolates capable of infecting Arabidopsis, but not of other plants. This study provides insight into how host specificity of *F. oxysporum* may be determined by a pair of effector genes on a transmissible CD chromosome.

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Pathogenic fungi often carry chromosomes that are not necessary for growth in the non-pathogenic state^{1,2}. Analogous to the well-characterized virulence plasmids in bacteria, the number of these dispensable chromosomes in individual isolates can vary. In plant pathogenic fungi, dispensable chromosomes that are associated with virulence are generally referred to as supernumerary, 'B', or conditionally dispensable (CD) chromosomes¹. When pathogenic fungi lack CD chromosomes, they can grow *in vitro*, but often exhibit attenuated or no virulence^{1,2}. The functions of CD chromosomes in some plant pathogenic fungi are associated with suppression or deactivation of host-specific factors. In *Fusarium solani*, for example, a CD chromosome carries phytoalexin detoxifying genes³. In contrast, the CD chromosomes of *Alternaria alternata* and *Cochliobolus carbonum* harbor host-specific toxin genes^{2,4}. Therefore, CD chromosomes can be crucial determinants of host specificity that are defined by phytotoxin activity or by defense against chemicals such as phytoalexins.

Fusarium oxysporum causes devastating diseases of more than 100 plant species, including economically important crops such as tomato, banana, and melon⁵. Individual isolates of *F. oxysporum* have different host ranges and are classified into *formae speciales* (ff. spp.) based on the susceptibility of plant species to infection. Although much is known about the genetics and pathology of *F. oxysporum*, the precise molecular mechanisms of host specificity remain unclear. So far, CD chromosomes have been identified in the tomato-infecting pathogen *F. oxysporum* f. sp. *lycopersici* (*Fol*) and in *F. oxysporum* f. sp. *radicis-cucumerinum* (*Forc*), a cucurbit-infecting pathogen^{6–8}. The *Fol* isolate 4287 and the *Forc* isolate 016 each contain a single virulence-associated CD chromosome that is transferable to other isolates^{7–9}. Horizontal transfer of the CD chromosomes from *Fol*4287 or *Forc*016 converts non-pathogenic *F. oxysporum* isolates into pathogens of their respective hosts^{6,7,9}. Part of this phytopathogenic conversion is often due to the expression of CD-encoded effectors that modulate host immunity against infection, such as Secreted In Xylem (SIX) effectors that are, as their name indicates, secreted into xylem elements during infection^{10,11}. A total of fourteen *SIX* genes (*SIX1* to *14*) have been identified from *Fol*¹⁰. The CD chromosome of *Fol*4287 contains all of the *SIX* genes except *SIX4*, which is not present in *Fol*4287^{6,12}, but is present in certain other *Fol* isolates. The CD chromosome of *Forc*016 contains *SIX6*, *SIX9*, *SIX11*, and *SIX13* homologs⁷. *SIX1*, *SIX3*, *SIX5*, and *SIX6* from

Fol are involved in overcoming resistance in tomato and the *SIX6* homolog from *Forc*016 is crucial for virulence in cucumber^{7,10}. However, their molecular mechanisms as virulence factors are as yet unknown.

Arabidopsis-infecting isolates of *F. oxysporum* are useful as a model pathosystem. There are at least three ff. spp. that cause disease on Arabidopsis: f. sp. *conglutinans*, f. sp. *matthiolarae*, and f. sp. *raphani*¹³. *F. oxysporum* f. sp. *conglutinans* (*Focn*) can also infect other Brassicaceae plants such as cabbage (*Brassica oleracea* var. *capitata*). The *SIX1* gene is required for full virulence on cabbage in *Focn*¹⁴, but the *Focn* factor(s) that are required for virulence on Arabidopsis have not been identified. We have previously shown that the *Focn* isolate Cong:1-1 (*Focn*Cong:1-1) harbors *SIX1*, *SIX4*, *SIX8*, and *SIX9* homologs on multiple chromosomes of different sizes¹⁵. Although these chromosomes are presumed to be conditionally dispensable in *Focn*, their status as CD chromosomes has not been established.

Here we report, through analyses of chromosome-deficient *Focn*Cong:1-1 strains and through horizontal chromosome transfer, that *Focn*Cong:1-1 has multiple CD chromosomes. Importantly, we identified individual CD chromosomes that are required for virulence on Arabidopsis, cabbage, or both. Furthermore, we identified a pair of effector genes on a CD chromosome that are required for suppression of Arabidopsis-specific phytoalexin-based immunity.

Results

Chromosome-level genome assembly of *Focn*Cong:1-1. We assembled the *Focn*Cong:1-1 genome sequence into 198 contigs with an N50 of 1.271 Mb. To improve contiguity, we further performed optical mapping using two restriction enzymes. The final assembly consisted of 22 scaffolds (SCs) with an N50 SC length of 4865 kb and a 99.1% complete BUSCO score (Table 1). For gene prediction, we generated transcriptome data from axenic culture and plant infections, resulting in a total of 21,781 genes, among which are eight presumptive effector genes (*SIX1*, *SIX4*, *SIX8*, *SIX9*, and *FOA1-FOA4*) that were previously known from Arabidopsis-infecting *F. oxysporum*¹⁶, as well as the homologous genes of *FOA1* and *FOA4*, which were named *FOA1b* and *FOA4b*, respectively. We did not detect homologs of any other *SIX* genes. To find unknown effectors, 1467 putative secreted proteins were screened for proteins with an effector-like structure using the EffectorP v1 and/or v2 algorithm^{17,18}. A total of 263 secreted proteins were predicted as effectors by both EffectorP v1 and v2. This prediction did not include *FOA1*, which is involved in the suppression of pattern-triggered immunity¹⁶, nor its homolog *FOA1b*. Therefore, a total of 265 proteins, including *FOA1* and *FOA1b*, were defined as high-confidence effector candidates (Table 1 and Supplementary Data 1).

The *F. oxysporum* genome is composed of core genomic regions that are conserved among *Fusarium* species, and additional accessory genomic regions that are conserved in certain isolates¹⁹. Comparative analysis with the *Fol*4287 genome as a reference indicated that (i) the *Focn*Cong:1-1 SCs have no homology with known accessory genomic regions in *Fol*4287 (chr01B; chr02B; chr03; chr06; chr14; chr15)⁶, (ii) similarly, there are genomic regions of *Focn*Cong:1-1 that have no homology with *Fol*4287, and (iii) the non-homologous genomic regions are enriched in transposable elements (TEs) (Fig. 1a). All known effector genes except *FOA4* are located in the TE-rich genomic region in *Focn*Cong:1-1 as follows: *SIX1* (in SC8), *SIX4* (SC9), *SIX8* (SC10), *SIX9* (SC3), *FOA1* (SC5), *FOA1b* (SC10), *FOA2* (SC9), *FOA3* (SC3), and *FOA4b* (SC10) (Supplementary Fig. 1). *FOA4* (SC12) may be a pseudogene since it is not expressed either *in vitro* or *in planta* (Supplementary Data 1 and 2). TEs are

Table 1 *Focn*Cong:1-1 genome statistics.

| Statistics | Value |
|---|--------------------------|
| Assembly stats | |
| Assembly size (Mb) | 68.8 (72.2) ^a |
| No. of scaffolds | 22 |
| No. of non-scaffolded contigs | 125 |
| Max scaffold length (kb) | 7,006 |
| N50 scaffold length (kb) | 4,865 |
| GC content (%) | 48.40 |
| BUSCO coverage (%) | 99.1 |
| Gene models | |
| Total no. of genes | 21,781 |
| Total no. of proteins | 22,094 |
| No. of genes encoding secreted proteins | 1,467 |
| No. of high-confidence effector candidates: | 265 ^{b,c} |
| Predicted by both EffectorP v1 and v2 | 263 |
| Predicted by either EffectorP v1 or v2 | 393 |

^aThe number in parentheses is total assembly size including non-scaffolded contigs.

^b*FOA1* and *FOA1b* are added to the list of effector candidates predicted by both EffectorP v1 and v2.

^cNote that some genes encode identical amino acid.

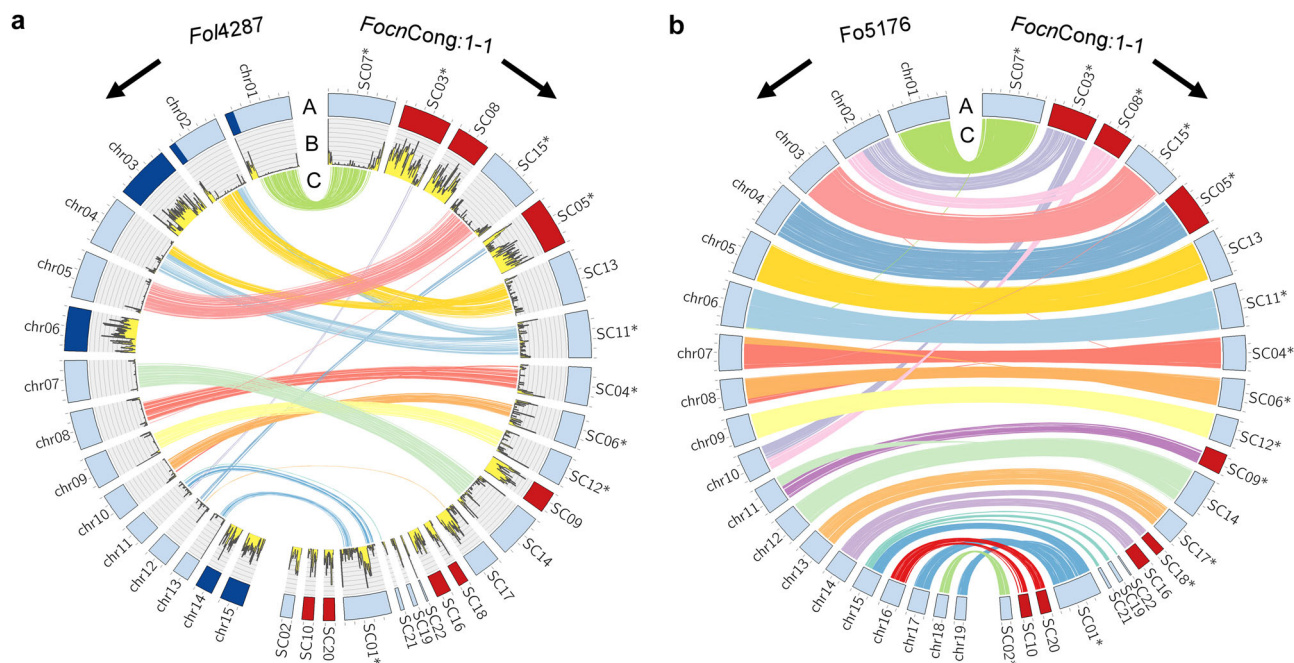


Fig. 1 Comparison of whole genome assemblies among *FocnCong:1-1*, *FoI4287*, and *Fo5176*. Whole genome assemblies were compared between *FoI4287* and *FocnCong:1-1* (**a**) and between *Fo5176* and *FocnCong:1-1* (**b**). Ring A: Circular representation of the pseudomolecules. Red and dark blue indicate dispensable genomic regions in *FocnCong:1-1* and known accessory regions in *FoI4287* (chr1B; chr2B; chr3; chr6; chr14; chr15)⁶, respectively. Light blue indicates the remaining regions. Ring B: Distribution of transposable elements (TEs) in 50 kb windows. Proportions of sequences of respective *FocnCong:1-1* SCs associated with TEs are shown in Supplementary Fig. 1. Ring C: Syntenic regions (>95% identity, 30 kb) between *FoI4287* and *FocnCong:1-1* assemblies (**a**) and between *Fo5176* and *FocnCong:1-1* assemblies (**b**). Asterisks indicate reverse-complemented scaffolds (SCs) for visual clarity. Ticks on bands represent 1 Mb.

suspected to be involved in the generation of genomic variations leading to environmental adaptation and, in the case of pathogens, they may have been involved in the acquisition of the ability to infect particular hosts²⁰. Therefore, chromosomes containing TE-enriched genomic regions have a high potential to be CD chromosomes.

Recently, a chromosome-level genome assembly of the Arabidopsis-infecting *F. oxysporum* isolate *Fo5176* was reported²¹. The genomes of *FocnCong:1-1* and *Fo5176* are very similar, sharing from 93.2% to 94.3% of their total scaffold/contig lengths (>95% identity, 10 kb). Synteny analysis revealed that (i) SC16 and SC18 of *FocnCong:1-1* correspond to chromosome 14 (chr14) of *Fo5176*, and (ii) SC10 and SC20 to chr16 (Fig. 1b), indicating that these SCs constitute, or contribute to the respective chromosomes. Due to the observations (i) and (ii) above, we refer to the chromosomes carrying these sequences as chr^{SC16/SC18} and chr^{SC10/SC20} in *FocnCong:1-1*, respectively.

***FocnCong:1-1* has multiple CD chromosomes.** To identify CD chromosomes of *FocnCong:1-1*, we generated chromosome-deficient strains by treatment with a mitosis inhibitor benomyl^{7,22}. For the parental strain, we utilized the previously generated strain *FocnCong:1-1* Δ *SIX4*, in which *SIX4*, located in SC9, had been replaced with a hygromycin B resistance gene (*hph*) cassette²³. After benomyl treatment, we obtained six hygromycin B-sensitive mutants (HS1 to HS6; Supplementary Figs. 2 and 3). To confirm the loss of dispensable genomic regions, we sequenced the genomes of *FocnCong:1-1* Δ *SIX4* and each of the HS mutants. As expected, *FocnCong:1-1* Δ *SIX4* maintained SC9 carrying *hph*, but all of the HS mutants had lost SC9 (Fig. 2a, b). We also found that (i) SC3 was absent in HS2, HS3, and HS4, (ii) SC5 and SC8 were lost only in HS6, (iii) chr^{SC10/SC20} was missing in HS3, HS4, HS5, and HS6, and (iv)

chr^{SC16/SC18} was lost only in HS4. In addition, duplication of SC2 and part of SC2 and SC17 occurred in HS1, HS5, and HS2, respectively (Fig. 2a).

Among the *FocnCong:1-1* HS mutants, there was no appreciable difference in colony size, but there was a significant difference in conidial formation (Fig. 2c, and Supplementary Figs. 2c and 4). *FocnCong:1-1* HS1 (Δ SC9) showed no difference in conidial formation or virulence on either Arabidopsis or cabbage compared to the parent strain Δ *SIX4*, indicating that SC9 is involved in neither conidial formation nor virulence (Fig. 2c, d and Supplementary Fig. 5). *FocnCong:1-1* mutants without SC3 (HS2, HS3, and HS4) had attenuated virulence on Arabidopsis and cabbage, but also had reduced ability to form conidia (Fig. 2c, d and Supplementary Fig. 5), suggesting that SC3 positively regulates conidial formation. To our surprise, loss of chr^{SC10/SC20} in *FocnCong:1-1* HS5 and HS6 increased conidial formation (Fig. 2c), but reduced virulence on Arabidopsis (Fig. 2d). Interestingly, *FocnCong:1-1* HS6 (Δ SC5/SC8/SC9/chr^{SC10/SC20}) lost virulence on both Arabidopsis and cabbage, whereas HS5 (Δ SC9/chr^{SC10/SC20}) retained virulence on cabbage, but not on Arabidopsis (Fig. 2d). These data indicate that chr^{SC10/SC20} is required for disease progression on Arabidopsis, but that SC5 and/or SC8 are involved only in causing disease in cabbage. Therefore, SC3, SC5, SC8, and chr^{SC10/SC20} are CD chromosomes affecting disease levels, with SC3 and chr^{SC10/SC20} being also associated with conidial formation.

***FocnCong:1-1* CD chromosomes are transferable.** We investigated whether *FocnCong:1-1* CD chromosomes are transferable under laboratory conditions, and what their effect might be on virulence. *FocnCong:1-1* HS6 lost multiple virulence-associated CD chromosomes (SC5, SC8, and chr^{SC10/SC20}) along with virulence on both Arabidopsis and cabbage (Fig. 2). Strain HS6

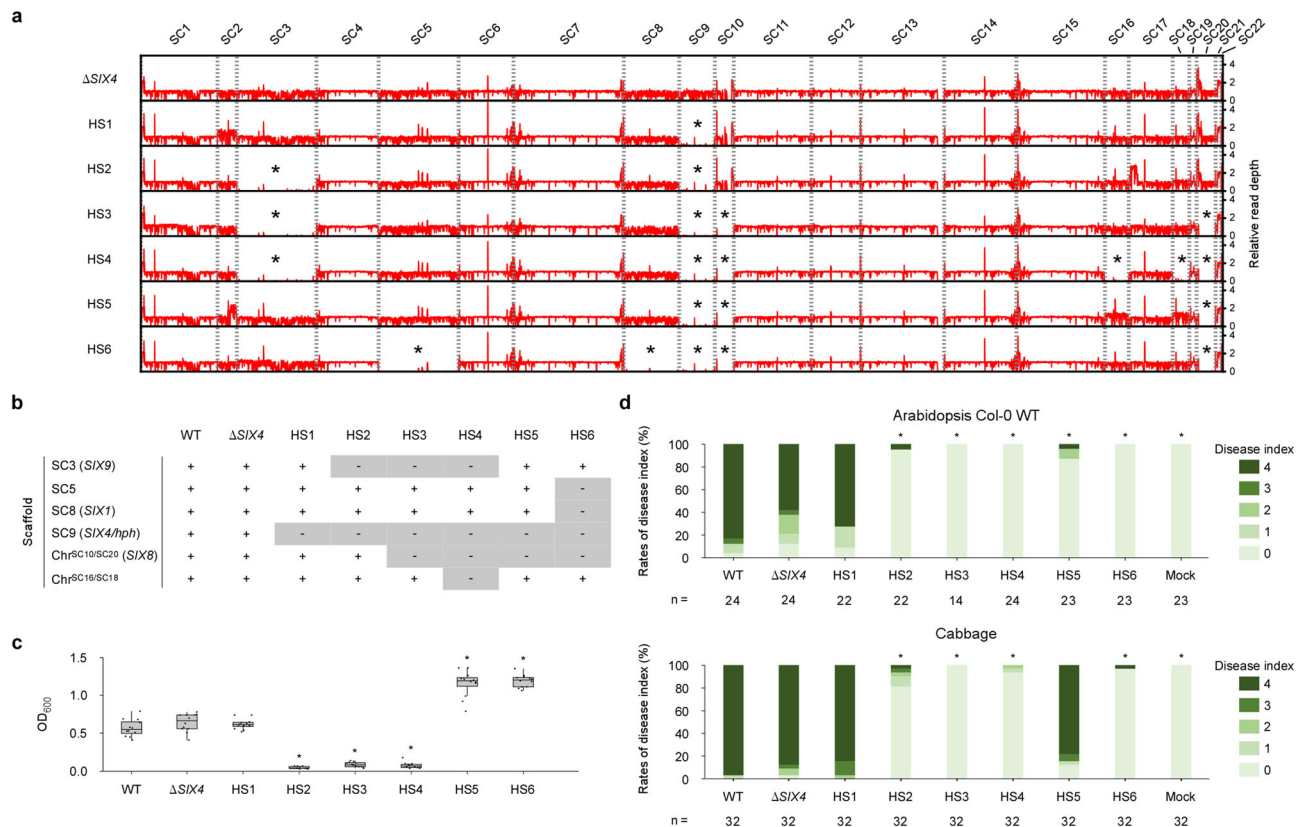


Fig. 2 Effects of loss of conditionally dispensable chromosomes on conidial formation and virulence in *FocnCong:1-1*. **a** Relative read mapping depths of *FocnCong:1-1* Δ SIX4 and HSs whole genomes. Asterisks indicate scaffold (SC)-level deletion. **b** Loss patterns of SC in *FocnCong:1-1* HSs. + and - represent maintained- and lost-SCs, respectively. *SIXs* located on particular SCs are shown in parentheses. **c** Conidial formation in *FocnCong:1-1* WT, Δ SIX4 and HSs. OD_{600} of conidial suspensions was measured from six colonies after 17 days of incubation on potato dextrose agar. Results of two independent experiments were combined and a total of twelve biological replicates per isolate are plotted. Boxplots indicate median value, estimated 25th and 75th percentiles, and whiskers represent 1.5 times the interquartile range. Asterisks represent significant differences from *FocnCong:1-1* Δ SIX4 ($*p < 0.0001$, Welch's t-test). **d** Virulence of *FocnCong:1-1* WT, Δ SIX4 and HSs to Arabidopsis and cabbage. Disease index was scored as described in Methods. Results of at least two experiments were combined. *n* denotes the number of plants investigated. Asterisks represent significant differences from *FocnCong:1-1* Δ SIX4 ($*p < 0.001$, Mann-Whitney U-test). Representative images of Arabidopsis and cabbage at 28 dpi are shown in Supplementary Fig. 5.

could therefore be used to determine the effects of chromosome transfer on virulence. A phleomycin-resistant *FocnCong:1-1* HS6 strain (HS6-BLE) was generated by introducing the phleomycin resistance gene (*ble*) (Supplementary Figs. 6a and 7). We co-incubated *FocnCong:1-1* HS6-BLE with hygromycin B-resistant *FocnCong:1-1* Δ SIX4 as a donor and selected four colonies (HCT1 to HCT4) that were resistant to both phleomycin and hygromycin B. There was no apparent difference in morphology or colony size (i.e., growth rate) in any of the presumptive *FocnCong:1-1* recipients (HCT1 to HCT4; Supplementary Fig. 6b). We confirmed chromosome transfer by contour-clamped homogeneous electric field (CHEF) electrophoretic karyotyping as well as PCR (Fig. 3a and Supplementary Figs. 6–8). *SIX1* (in SC8) and *hph* (in SC9) were detected in all *FocnCong:1-1* recipients, whereas *SIX8* (in SC10) and an SC20 marker (*FocnCong_v011766*) were detected only in HCT1 (Supplementary Fig. 6a). We did not detect the SC5 marker *FocnCong_v016149* in any recipient (Supplementary Fig. 6a). These results indicate that at least SC8, SC9, and chr^{SC10/SC20} are transferable. Conidial formation of the three *FocnCong:1-1* recipients HCT2, HCT3, and HCT4, which acquired SC8 and SC9, was comparable to that of HS6-BLE. In contrast, *FocnCong:1-1* HCT1, which received chr^{SC10/SC20}, SC8, and SC9, produced significantly fewer conidia than HS6-BLE, a phenotype similar to the donor Δ SIX4 (Fig. 3b). Because SC8 and SC9 are not involved in conidial formation (Fig. 2c), these results suggest a negative

involvement of chr^{SC10/SC20} in conidial formation. All *FocnCong:1-1* recipients (i.e., HCT1 to HCT4) that acquired SC8 and SC9 also regained virulence against cabbage (Fig. 3c and Supplementary Fig. 9). Because SC9 is not involved in virulence (Fig. 2d), we conclude that SC8 is necessary and sufficient for virulence to cabbage. In the case of Arabidopsis, *FocnCong:1-1* HCT1 showed higher virulence than the other recipients (HCT2, HCT3, and HCT4; Fig. 3c and Supplementary Fig. 9). Taken together with the pathology results of the chromosome-deficient *FocnCong:1-1* mutants (HS1 to HS6; Fig. 2), it is likely that chr^{SC10/SC20} is required for virulence on Arabidopsis.

Chr^{SC10/SC20} is involved in suppression of *CYP79B2/CYP79B3*-mediated immunity. A CD chromosome from the Arabidopsis-infecting anthracnose fungus *Colletotrichum higginsianum* has been reported to be involved in suppression of plant immunity that is dependent on tryptophan (Trp)-derived secondary metabolites²⁴. We investigated whether CD chromosomes of *FocnCong:1-1* encode products that also suppress specific immunity. For this experiment, we used the Arabidopsis double mutant *cyp79b2/cyp79b3* that lacks the ability to synthesize Trp-derived secondary metabolites²⁵. Among the chromosome-deficient *FocnCong:1-1* mutants (HS2 to HS6) with attenuated virulence to Arabidopsis Col-0 WT, only *FocnCong:1-1* HS5 (Δ SC9/chr^{SC10/SC20}) showed the same level of virulence on

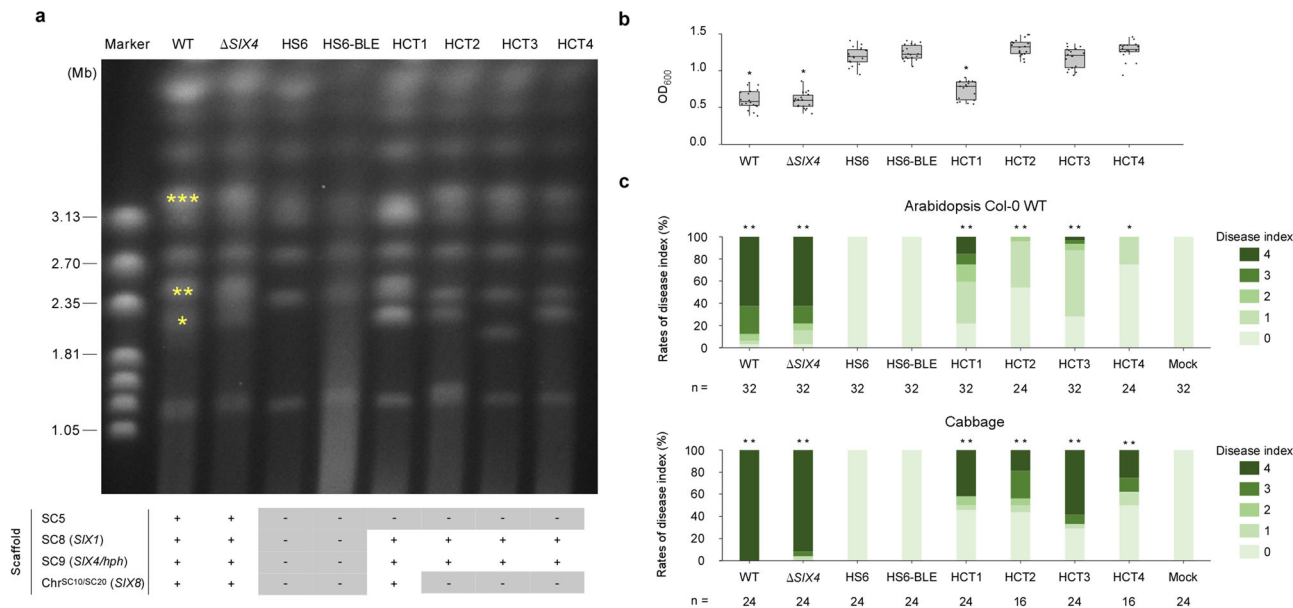


Fig. 3 Effects of chromosome transfer on conidial formation and virulence in *FocnCong:1-1* HS6. **a** Electrophoretic karyotype of *FocnCong:1-1* WT, Δ SIX4, HS6, HS6-BLE, and HCTs. Asterisks indicate chromosomes on which *SIX* genes are located¹⁵: **SIX4*, ***SIX8*, ****SIX1*. The table indicates the scaffold (SC) patterns confirmed by PCR as shown in Supplementary Fig. 6a. + and - represent maintained- and lost-SCs, respectively. *SIX*s located on SCs are shown in parentheses. **b** Conidial formation in *FocnCong:1-1* WT, Δ SIX4, HS6, HS6-BLE, and HCTs. OD₆₀₀ of conidial suspension was measured from six colonies after 17 days of incubation on potato dextrose agar. Results of three independent experiments were combined and a total of 18 biological replicates are plotted. Boxplots indicate median value, estimated 25th and 75th percentiles, and whiskers represent 1.5 times the interquartile range. Asterisks represent significant differences from *FocnCong:1-1* HS6-BLE (* $p < 0.0001$, Welch's t-test). **c** Virulence of *FocnCong:1-1* WT, Δ SIX4, HS6, HS6-BLE, and HCTs on Arabidopsis and cabbage. Disease index was scored as described in Methods. Results of at least two independent experiments were combined. *n* denotes the number of plants investigated. Asterisks represent significant difference from *FocnCong:1-1* HS6-BLE (** $p < 0.001$, * $p < 0.01$ Mann-Whitney U-test). Representative images of Arabidopsis and cabbage at 28 dpi are shown in Supplementary Fig. 9.

cyp79b2/cyp79b3 plants as was observed for its parent strain Δ SIX4 (Fig. 4a and Supplementary Fig. 10a, b). These results suggest that chr^{SC10/SC20} plays a key role in suppressing Trp-derived secondary metabolite-dependent immunity. *FocnCong:1-1* HS6 (Δ SC5/SC8/SC9/chr^{SC10/SC20}) was substantially less virulent on *cyp79b2/cyp79b3* plants. This is likely because SC5 or SC8 are involved in virulence other than through suppression of Trp-based immunity. In addition, we found that the *cyp79b2/cyp79b3* double mutant was resistant to all tested SC3-deficient *FocnCong:1-1* mutants (HS2 to HS4; Fig. 4a and Supplementary Fig. 10a, b), possibly due to some deficiency of conidial formation in these mutants (Fig. 2c).

To investigate which step or steps of infection the CD chromosomes contribute to, a histological analysis was performed using GFP-labeled *FocnCong:1-1* strains in Arabidopsis. *FocnCong:1-1* Δ SIX4-GFP always colonized xylem vessels of roots, often reaching stem elements in Arabidopsis WT, whereas *FocnCong:1-1* HS2-GFP lacking SC3 germinated on root surfaces but showed almost no colonization in xylem vessels of roots or stems (Fig. 4b), confirming its deficiency in growth in planta. *FocnCong:1-1* HS5-GFP (Δ SC9/chr^{SC10/SC20}) colonized root xylem vessels, but the frequency of stem colonization was low in Arabidopsis WT. In *cyp79b2/cyp79b3* double mutants, however, *FocnCong:1-1* HS5-GFP frequently colonized the stems as was observed for *FocnCong:1-1* Δ SIX4 in WT (Fig. 4b). These results suggest that chr^{SC10/SC20} is implicated in the ability to colonize beyond root xylem vessels into stems, and conversely that CYP79B2/CYP79B3 participate in inhibition of *FocnCong:1-1* colonization of stems.

To determine whether CYP79B2/CYP79B3-based antibiotics such as isothiocyanate, camalexin, or 4-hydroxyindole-3-carbonyl nitrile (4-OH-ICN)^{26,27} are associated with resistance, we

conducted bioassays with *FocnCong:1-1* HS5 (Δ SC9/chr^{SC10/SC20}) on three Arabidopsis mutants (*pen2*, *pad3*, and *cyp82c2*) that are unable to produce isothiocyanate, camalexin, and 4-OH-ICN, respectively^{26,27} (Fig. 4c). The *pad3* mutant, but not *pen2* or *cyp82c2*, was more susceptible to *FocnCong:1-1* HS5 than Col-0 WT (Fig. 4d and Supplementary Fig. 10c), suggesting that camalexin, but not isothiocyanate or 4-OH-ICN, is involved in resistance to *FocnCong:1-1*. Importantly, camalexin is produced in Arabidopsis, but not in cabbage²⁸. Because *FocnCong:1-1* HS5 (Δ SC9/chr^{SC10/SC20}) had attenuated virulence on Arabidopsis but full virulence on cabbage (Fig. 2d), chr^{SC10/SC20} is likely to contribute to suppression of Arabidopsis-specific immunity, specifically camalexin, to establish infection.

A pair of effectors are involved in virulence on Arabidopsis.

Because chr^{SC10/SC20} is likely to encode effectors that contribute to suppression of Arabidopsis-specific immunity, we searched for genes encoding potential effectors, and found a total of twelve effector candidate genes located on chr^{SC10/SC20} (Supplementary Data 1). Expression profiling revealed that *FocnCong_v001893* (*SIX8*) and *FocnCong_v001894* were highly expressed during infection (Fig. 5a and Supplementary Data 1). Interestingly, *SIX8* is adjacent to *FocnCong_v001894*, with an intergenic distance of 1057 bp on SC10 (Fig. 5b). The intergenic region contains a miniature impala inverted repeat (mimp-IR) sequence, which is related to TE sequences (Fig. 5b and Supplementary Fig. 11). A mimp-IR is also often located in the upstream regions of *SIX* and other effector candidate genes in *Fol*, *Forc*, and the melon-infecting pathogen *F. oxysporum* f. sp. *melonis*^{7,12,29}. To determine whether *SIX8* and *FocnCong_v001894* are involved in virulence on Arabidopsis, a genome fragment containing the *SIX8-FocnCong_v001894* locus was introduced into *FocnCong:1-1*

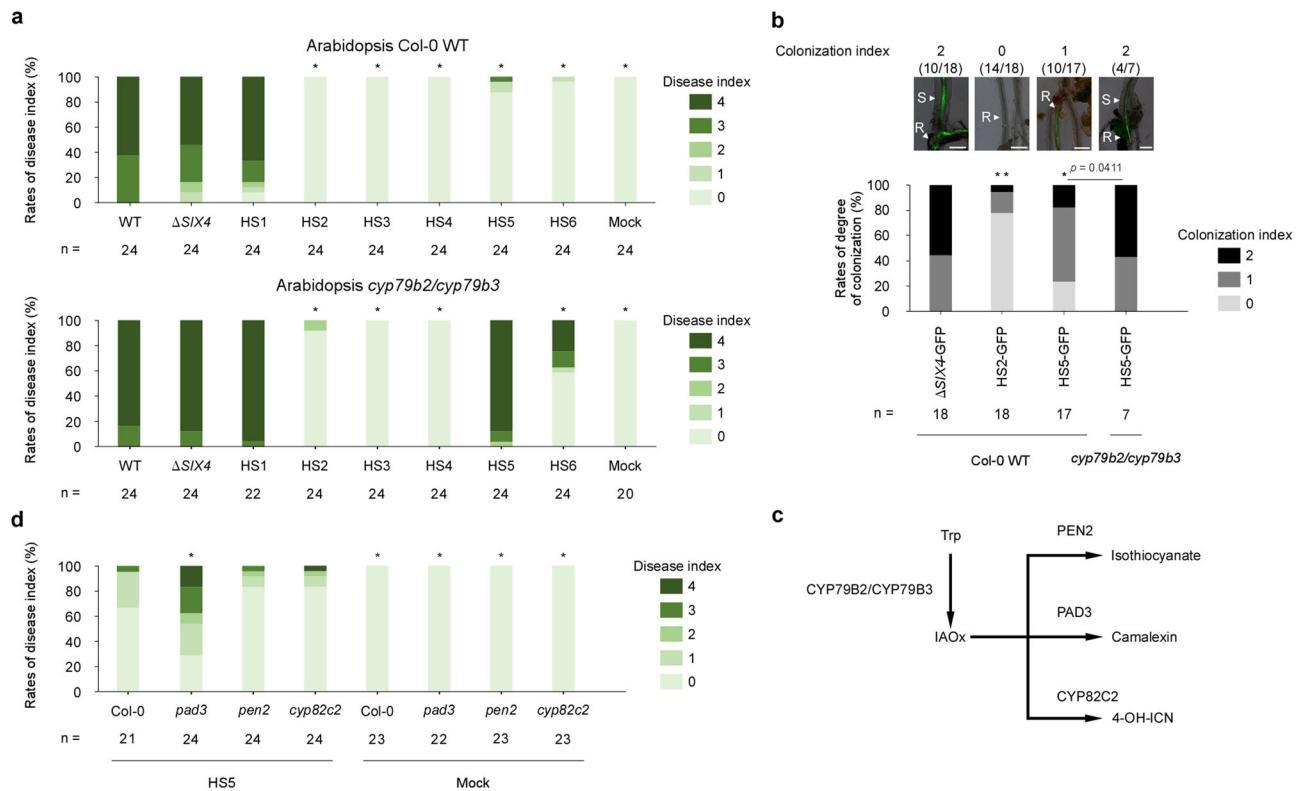


Fig. 4 Involvement of chr^{SC10/SC20} in suppression of CYP79B2/CYP79B3-mediated immunity. **a** Virulence of *FocnCong*:1-1 WT, Δ SIX4 and HSs on Arabidopsis Col-0 WT and the *cyp79b2/cyp79b3* double mutant. Disease index was scored as described in Methods. Results of three independent experiments were combined. *n* denotes the number of plants investigated. Asterisks represent significant difference from *FocnCong*:1-1 Δ SIX4 ($*p < 0.001$, Mann-Whitney U-test). Representative images of Arabidopsis at 28 dpi are shown in Supplementary Fig. 10a, b. **b** Infection phenotypes of *FocnCong*:1-1 Δ SIX4, HS2 and HS5 in Arabidopsis. Colonization index of Arabidopsis Col-0 WT and *cyp79b2/cyp79b3* double mutant inoculated with GFP-labeled *FocnCong*:1-1 Δ SIX4 (Δ SIX4-GFP), HS2 (HS2-GFP) or HS5 (HS5-GFP) at 12 dpi was scored from 0 to 2: 0, germination or colonization on root surface; 1, colonization in xylem vessels of roots; 2, transition from roots to stems. Results of at least two independent experiments were combined. *n* denotes the number of plants investigated. Asterisks represent significant differences from *FocnCong*:1-1 Δ SIX4-GFP ($**p < 0.001$, $*p < 0.01$, Mann-Whitney U-test). Representative images of root or stem of Arabidopsis at 12 dpi are shown above each bar. Scale bars indicate 200 μ m. S stems, R roots. **c** A simple scheme of biosynthesis of tryptophan (Trp)-derived defense compounds in Arabidopsis. IAox indole-3-acetaldoxime, 4-OH-ICN 4-hydroxy-indole carbonyl nitrile. **d** Virulence of *FocnCong*:1-1 HS5 on Arabidopsis Col-0 WT, *pad3*, *pen2*, and *cyp82c2* mutants. Disease index was scored as described in Methods. Results of three independent experiments were combined. *n* denotes the number of plants investigated. Asterisks represent significant difference from Arabidopsis Col-0 WT infected with *FocnCong*:1-1 HS5 ($*p < 0.01$, Mann-Whitney U-test). Representative images of Arabidopsis at 28 dpi are shown in Supplementary Fig. 10c.

HS5 (Δ SC9/chr^{SC10/SC20}) (Supplementary Figs. 12 and 13), which restored full virulence to *FocnCong*:1-1 HS5 (Fig. 5c and Supplementary Fig. 14a). In contrast, Arabidopsis WT was resistant to the other *FocnCong*:1-1 HS5 transformants that contained only *SIX8* or *FocnCong_v001894* (Fig. 5c and Supplementary Fig. 14a). It should be noted that virulence of knockout mutants that lack the *SIX8-FocnCong_v001894* locus in *FocnCong*:1-1 was significantly lower than for WT (Fig. 5d and Supplementary Figs. 14–16), suggesting that both *SIX8* and *FocnCong_v001894* are necessary for virulence on Arabidopsis. We therefore designated *FocnCong_v001894* as *Pair with SIX Eight1* (*PSE1*).

Genetic and functional conservation of the *SIX8* and *PSE1* loci.

Next, we investigated whether the *SIX8-PSE1* pair is conserved in Arabidopsis-infecting *F. oxysporum* isolates. Comparative analysis of highly contiguous and available genome assemblies of *F. oxysporum* isolates (Supplementary Table 1) showed that the *SIX8-PSE1* locus is completely conserved in Fo5176 and in the stock-infecting pathogen *F. oxysporum* f. sp. *matthiolae* (*Fomt*) PHW726, which can infect Arabidopsis^{13,30}, but not in isolates that cannot infect Arabidopsis (Fig. 6a, b). For example, the banana-infecting pathogen *F. oxysporum* f. sp. *cubense* (*Focb*)

tropical race 4 (TR4), which threatens banana production worldwide, has *SIX8* but not *PSE1*. In the other non-Arabidopsis-infecting isolates, except *Fol4287*, neither *SIX8* nor *PSE1* is present. *Fol4287* has multiple copies of *SIX8* and its homolog *SIX8b*^{12,31} but *PSE1* is not present in the published *Fol4287* gene annotation⁶. However, we found three loci similar to the *SIX8-PSE1* locus in chromosomes 2, 3, and 14 of *Fol4287*. At these loci, adjacent to *SIX8*, there is a *PSE1*-like gene (*PSL1*) differing in the C-terminal 10 amino acids (Fig. 6b and Supplementary Fig. 17). Furthermore, multiple *SIX8b* loci contain TEs inserted into adjacent *PSE1* sequences. For example, a transposase gene was found in the first intron of the *PSE1* homologs in two loci of chromosome 3 and another locus in chromosome 6 (Fig. 6b and Supplementary Fig. 18). Similarly, a presumptive transposase was found immediately upstream of the potential-but-unannotated *PSE1* homolog in another locus in chromosome 6 (Fig. 6b). Thus, TE insertion seems to have disrupted the *PSE1* adjacent to *SIX8b* in *Fol4287*.

To evaluate if the *SIX8-PSE1* locus in *Fomt*PHW726 and the *SIX8-PSL1* locus in *Fol4287* are able to function similarly in *FocnCong*:1-1, we cloned these loci and transformed them into the *FocnCong*:1-1 HS5 mutant (Δ SC9/chr^{SC10/SC20}; Supplementary

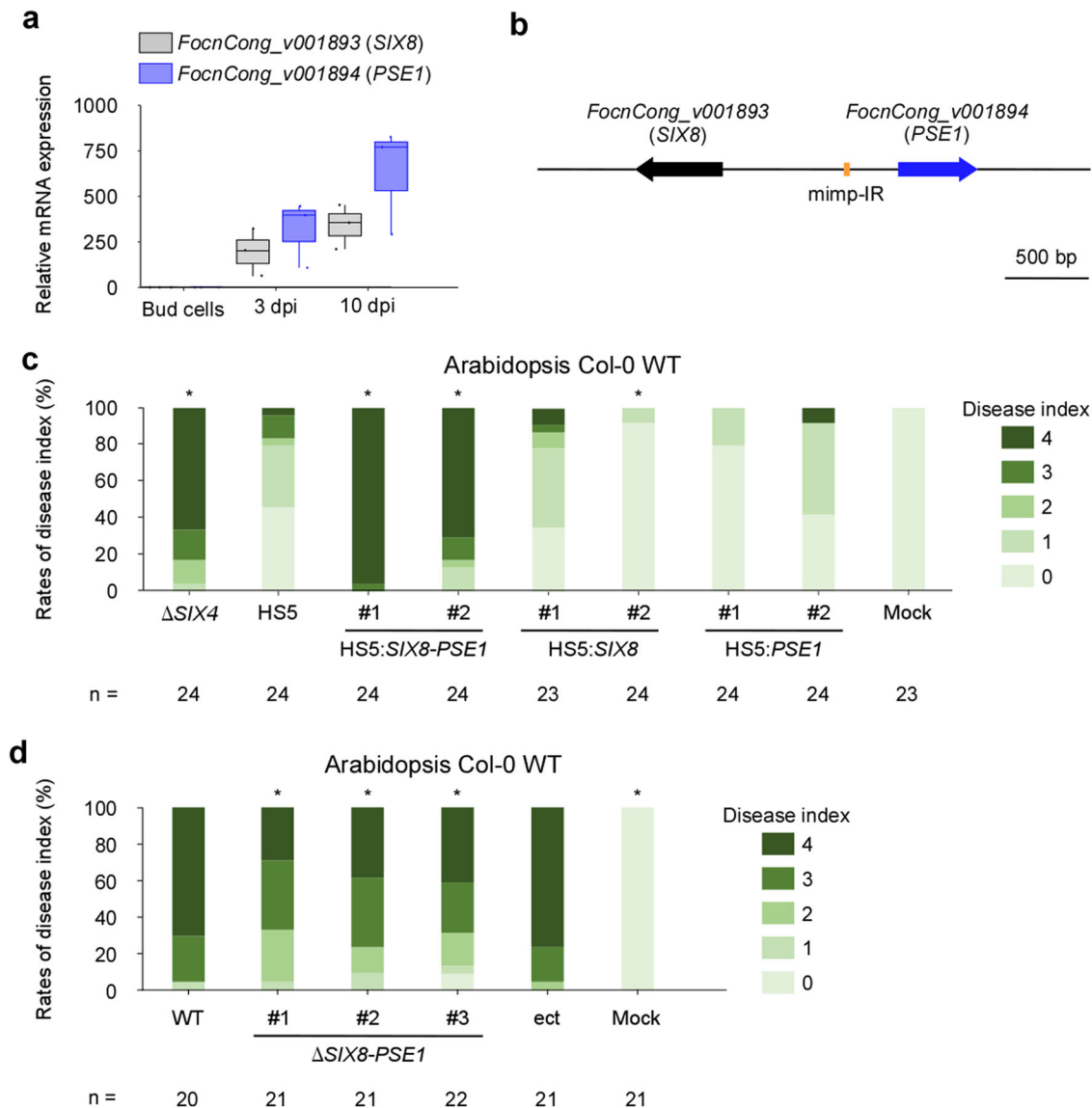


Fig. 5 The *SIX8-PSE1* locus is involved in virulence of *FocnCong:1-1* on Arabidopsis. **a** Relative transcript levels of *FocnCong_v001893* (*SIX8*) and *FocnCong_v001894* (*PSE1*) during infection of Arabidopsis Col-0 WT at 3 and 10 dpi. Data from three biologically independent samples are presented as fold changes compared with expression levels in bud cells. Expression levels were determined by qRT-PCR and normalized against *FocnCong:1-1* β -tubulin (*TUB2*). Boxplots indicate median value, estimated 25th and 75th percentiles, and whiskers represent 1.5 times the interquartile range. **b** Schematic representation of the *SIX8-PSE1* locus in *FocnCong:1-1*. *mimp-IR* miniature impala-like inverted repeat. **c** Virulence of *FocnCong:1-1* Δ *SIX4*, HS5, and HS5 transformants introduced with *SIX8* (HS5:*SIX8*), *PSE1* (HS5:*PSE1*) or both (HS5:*SIX8-PSE1*) on Arabidopsis Col-0 WT. Disease index was scored as described in Methods. *n* denotes the number of plants investigated. Asterisks represent significant difference from *FocnCong:1-1* HS5. (**p* < 0.001, Mann-Whitney U-test). Representative images of Arabidopsis at 28 dpi are shown in Supplementary Fig. 14a. **d** Disease index of Arabidopsis Col-0 WT challenged with *FocnCong:1-1* WT, Δ *SIX8-PSE1*, an ectopic transformant (ect) or water (mock) at 28 dpi was scored as described in Methods. Results of three independent experiments were combined. *n* denotes the number of plants investigated. Asterisks represent significant difference from WT (**p* < 0.05, Mann-Whitney U-test). Representative images of Arabidopsis at 28 dpi are shown in Supplementary Fig. 14b.

Figs. 19 and 20). Transformation with the *Fomt SIX8-PSE1* locus, but not the *Fol SIX8-PSL1* locus, restored full virulence to *FocnCong:1-1* HS5 in Arabidopsis Col-0 WT (Fig. 6c and Supplementary Fig. 21). These results suggest that the *SIX8-PSE1* locus is functionally distinct from *SIX8-PSL1* and is functionally conserved in Arabidopsis-infecting *F. oxysporum* isolates.

Discussion

Here we report the identification of a CD chromosome in *F. oxysporum* that is required for virulence on Arabidopsis. This CD chromosome encodes a pair of effectors (*SIX8* and *PSE1*) that are

involved in suppressing Arabidopsis-specific immunity, and are conserved in the other *F. oxysporum* isolates capable of infecting Arabidopsis. The mode of action potentially involves defense against, or suppression of, the phytoalexin camalexin. We also report that another CD chromosome is required for pathogenicity on cabbage. In addition, certain CD chromosomes are involved in conidial formation.

In plant pathogenic fungi, CD chromosomes associated with virulence are usually not involved in vegetative growth^{1,2}. In this sense, SC3 and chr^{SC10/SC20} in *FocnCong:1-1* are atypical CD chromosomes that affect conidial formation (Fig. 2c). Although the reduced virulence of SC3-deficient *FocnCong:1-1* mutants

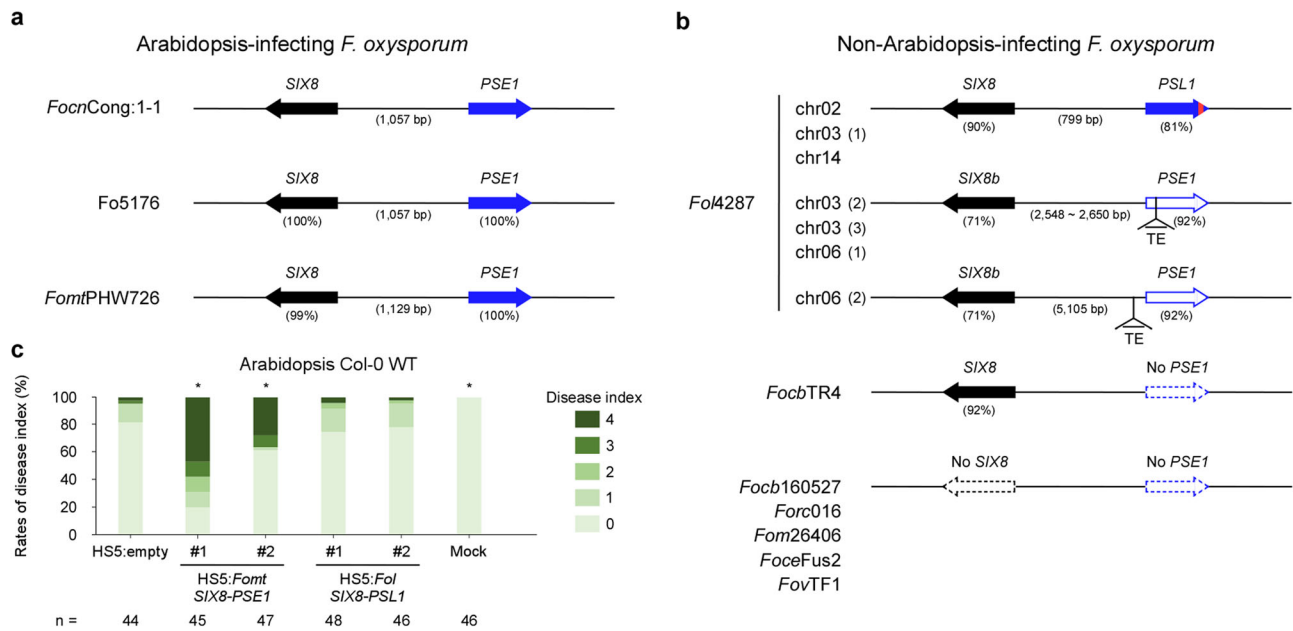


Fig. 6 Genetic and functional conservation of a pair of effectors, *SIX8* and *PSE1*, among *F. oxysporum* isolates. Comparison of the *SIX8-PSE1* loci in Arabidopsis-infecting (**a**) and non-Arabidopsis-infecting *F. oxysporum* isolates (**b**). A red region in the arrowhead of *PSL1* indicates a region of 10 amino acids that differ from *PSE1*. The amino acid level identity (%) to *FocnCong:1-1* *SIX8* or *PSE1* is shown in parentheses. *FocnCong:1-1*, *F. oxysporum* f. sp. *conglutinans* Cong:1-1; *FomtPHW726*, *F. oxysporum* f. sp. *matthiola* PHW726; *Fol4287*, *F. oxysporum* f. sp. *lycopersici* 4287; *FocbTR4*, *F. oxysporum* f. sp. *cubense* tropical race 4; *Focb160527*, *F. oxysporum* f. sp. *cubense* 160527; *Forc016*, *F. oxysporum* f. sp. *radicis-cucumerinum* 016; *Fom26406*, *F. oxysporum* f. sp. *melonis* 26406; *Focfus2*, *F. oxysporum* f. sp. *cepa* Fus2; *FovTF1*, *Fusarium oxysporum* f. sp. *vasinfectum* TF1. **c** Virulence of *FocnCong:1-1* HS5 transformants introduced with a hygromycin B resistance gene (*hph*) cassette (HS5:empty), the *Fomt* *SIX8-PSE1* locus (HS5:*Fomt* *SIX8-PSE1*) and the *Fol* *SIX8-PSL1* locus (HS5:*Fol* *SIX8-PSL1*) on Arabidopsis Col-0 WT. Disease index was scored as described in Methods. Results of six independent experiments were combined. *n* denotes the number of plants investigated. Asterisks represent significant difference from *FocnCong:1-1* HS5:empty. (**p* < 0.01, Mann-Whitney U-test). Representative images of Arabidopsis at 28 dpi are shown in Supplementary Fig. 21.

(HS2, HS3, and HS4) on Arabidopsis and cabbage (Fig. 2c, d) may be due to deficiency in the ability to form conidia, or to regulatory step(s) that has multiple unexplored phenotypic effects, we cannot exclude the possibility that yet-unknown effectors located on SC3 are implicated in virulence. Interestingly, SC3 contains a region partly syntenic to chromosome 11, which is a core chromosome of *Fol4287* (Fig. 1a). This syntenic region may contain dose-effective genes involved in conidial formation. In contrast to SC3, $\text{chr}^{\text{SC10/SC20}}$ negatively regulates conidial formation but positively contributes to virulence on Arabidopsis but not on cabbage (Fig. 2c, d), possibly representing a trade-off between vegetative growth and virulence to a particular host.

FocnCong:1-1 carries multiple CD chromosomes that have distinct virulence functions against specific hosts. For example, the CD chromosome $\text{chr}^{\text{SC10/SC20}}$ -deficient *FocnCong:1-1* HS5 is less virulent on Arabidopsis, but is able to develop severe disease on cabbage (Fig. 2d and Supplementary Fig. 5). This result may be explained by the fact that *FocnCong:1-1* HS5 maintains the CD chromosome SC8, which harbors a gene, *SIX1*, required for full virulence on cabbage¹⁴. Consistently, *FocnCong:1-1* HS6, which lacks both SC8 and $\text{chr}^{\text{SC10/SC20}}$, lost pathogenicity on both cabbage and Arabidopsis (Fig. 2d and Supplementary Fig. 5), and introduction of SC8 into HS6 restored virulence on cabbage (Fig. 3c and Supplementary Fig. 9). Thus, we conclude that $\text{chr}^{\text{SC10/SC20}}$ and SC8 are responsible for host-specific virulence on Arabidopsis and cabbage, respectively.

The target of the CD chromosome $\text{chr}^{\text{SC10/SC20}}$ effector is likely to be *CYP79B2/CYP79B3*-mediated immunity in Arabidopsis, because the loss of $\text{chr}^{\text{SC10/SC20}}$ attenuated virulence of *FocnCong:1-1* HS5 to WT, but not to the *cyp79b2/cyp79b3* double mutant (Fig. 4a and Supplementary Fig. 10). *CYP79B2/CYP79B3* had not previously been implicated in resistance to *F. oxysporum*.

For instance, Kidd et al.³² reported that susceptibility of *cyp79b2/cyp79b3* to *F. oxysporum* Fo5176 was not different from WT. Consistent with this report, our study shows that virulence of *FocnCong:1-1* on *cyp79b2/cyp79b3* is comparable to WT (Fig. 4a and Supplementary Fig. 10). Thus, only the use of CD chromosome-deficient mutants allowed us to uncover the involvement of *CYP79B2/CYP79B3* in resistance to *F. oxysporum*. Furthermore, histological analysis suggests that *CYP79B2/CYP79B3*-mediated immunity may be associated with inhibition of root-stem translocation of *FocnCong:1-1* (Fig. 4b). *CYP79B2/CYP79B3* is responsible for synthesis of Trp-derived secondary metabolites, including sulfur-containing compounds that are characteristic of the Brassicaceae²⁶. These sulfur-containing antimicrobial compounds differ among Brassicaceae species; for example, camalexin is produced in Arabidopsis, but not in cabbage²⁸. Our results suggest that *FocnCong:1-1* can overcome the Arabidopsis-specific immunity conferred by *PAD3*, a camalexin synthetic gene (Fig. 4d and Supplementary Fig. 10c), when the CD chromosome $\text{chr}^{\text{SC10/SC20}}$ that encodes the paired effectors *SIX8* and *PSE1* is present. This pair of effectors is highly conserved in Arabidopsis-infecting *F. oxysporum* isolates, but not in other isolates (Fig. 6), thus the presence of a particular CD chromosome that harbors these effector genes would contribute to the determination of host specificity.

In this study, *FocnCong:1-1* HSs were generated by treatment with the mitosis inhibitor benomyl. In the generation process, a genome rearrangement, but not just a chromosome loss, has occurred at least in HS1, HS2, and HS5 (Fig. 2a). We also investigated phenotypes in an additional HS mutant with the same karyotype as HS5 (HS5L: HS5-like mutant; Supplementary Figs. 22 and 23). Like HS5, HS5L showed virulence on *cyp79b2/cyp79b3* and *pad3* plants, but not on Col-0 WT plants. We cannot

rule out the possibility that these genome rearrangements affect phenotypes. In addition to the results of HS5L, however, the return of HS5 virulence on Arabidopsis in two independent HS5 transformants containing *FocnCong1-1 SIX8-PSE1* (Fig. 5c) supports the conclusion that the *SIX8-PSE1* pair is required for virulence on Arabidopsis.

We identified *SIX8* and *PSE1* as a gene pair adjacent but encoded on opposite DNA strands (head-to-head orientation) (Fig. 5b and Supplementary Fig. 11). Head-to-head orientation of effector genes has been documented for other *SIX* genes in *F. oxysporum*. For instance, in *Fol*, a pair of effector genes *SIX3* (also known as *AVR2*) and *SIX5* are also adjacently located in a head-to-head transcriptional orientation^{12,33,34}. Both *SIX3* and *SIX5* are required for not only full virulence in a susceptible host, but also disease resistance in tomato lines containing the resistance gene *I-2*^{33–35}, and the gene products are thus likely to function as a pair. The close head-to-head orientation may ensure coordinated transcription to produce both proteins at similar levels. Such system would be suitable for maintaining tight stoichiometry of two proteins in a complex. Indeed, *SIX5* interacts with *SIX3* at plasmodesmata in plant cells, facilitating cell-to-cell movement of *SIX3*^{33,34}. Unlike the *SIX3-SIX5* pair, however, we failed to detect direct interaction between *SIX8* and *PSE1* in a yeast two-hybrid assay (Supplementary Fig. 24). We cannot exclude the possibility that *SIX8* indirectly interacts with *PSE1*, e.g., via host target(s), or the yeast system may not be suitable for detecting interactions of these proteins. Alternatively, *SIX8* and *PSE1* may act independently. As bioinformatic analysis of *SIX8* and *PSE1* protein sequences gives no known domain annotations, identification of host targets of *SIX8* and *PSE1* will be required to clarify functions of the paired effectors. It is also notable that disruption or loss occurs in only *PSE1*, but not in *SIX8*, in certain non-Arabidopsis infecting *F. oxysporum* isolates. Perhaps *PSE1*, but not *SIX8*, is recognizable in plants that carry corresponding resistance proteins, leading to its disruption or loss to avoid detection.

In this work we demonstrate that the host range of *F. oxysporum* depends on CD chromosomes. In this respect, it is interesting that certain isolates, such as *Fol4287* and *Forc016*, have only a single virulence-associated CD chromosome, whereas *FocnCong1-1* has multiple CD chromosomes, each of which encodes host-specific effectors. Because the *FocnCong1-1* genome is very large (68.8 Mb) compared to most known *F. oxysporum* genomes, such as *Fol4287* (59.9 Mb)⁶ and *Forc016* (52.9 Mb)⁷, *FocnCong1-1* is likely to have expanded its host range by acquiring and maintaining additional CD chromosomes. Indeed, Masunaka et al.³⁶ have shown that a field isolate of *A. alternata* carrying two putative CD chromosomes has a wide host range. In that case, host-specific toxin genes on different chromosomes determine host range³⁶. In the case of *F. oxysporum*, host specificity can be determined, at least in part, by effectors, as seen in this study. Further functional analyses of the *SIX8-PSE1* paired effectors and their derivatives will be needed to dissect out the molecular mechanisms underlying effector-based host specificity in *F. oxysporum*.

Methods

Fungal strains and plants. Fungal strains used in this study are listed in Supplementary Table 2. For pre-incubation, all strains were incubated on potato dextrose agar (PDA; Nissui Pharmaceutical Co.) at 28 °C in the dark. For bud cell production, all strains were grown in NO₃ medium (0.17% yeast nitrogen base without amino acids, 3% sucrose and 1% KNO₃) at 120 strokes per minute (spm) for 4 days at 28 °C in the dark. For gene expression profiling (Supplementary Data 1 and 2), mycelia were harvested after 10 days of incubation on PDA at 28 °C. Bud cells were collected from NO₃ medium by filtration with a nylon mesh and centrifugation. Hyphae trapped with the nylon mesh were collected. Mycelia from PDA, bud cells, and hyphae were stored at -80 °C until RNA isolation.

Arabidopsis (Col-0 wild type, *pen2*, *pad3*, *cyp82c2*, and *cyp79b2/cyp79b3* mutants^{26,27}) and cabbage (cv. Shikidori and cv. Shosyu; Takii Seed) were cultured in pots containing autoclaved Super Mix A (Sakata Seed) and vermiculite (VS kakou). Arabidopsis was grown at 22 °C for 10 h under light and 14 h dark in a growth chamber. Cabbage was grown in a greenhouse.

Bioassays. For evaluation of disease severity, 14-day-old Arabidopsis and cabbage cv. Shikidori roots were injured with a forceps or a plastic peg and then irrigated with 1 ml of *FocnCong1-1* bud cell suspension (1 × 10⁷ cells/ml). Inoculated Arabidopsis plants were grown at 28 °C for 10 h under light and 14 h dark in a growth chamber. An Arabidopsis disease index was scored at 28 or 29 days post-inoculation (dpi) as: 0, no symptoms; 1, dwarf; 2, yellowing, vein clearing or wilting of one to a few leaves; 3, wilting of a whole plant; 4, dead. A cabbage disease index was also scored at 28 or 29 dpi as: 0, no symptoms; 1, yellowing lower leaves; 2, yellowing lower and upper leaves; 3, whole plant wilting; 4, dead.

For gene expression profiling (Supplementary Data 1 and 2), 20- or 21-day-old Arabidopsis and 17-day-old cabbage cv. Shosyu roots were irrigated with 1 ml of bud cell suspension (1 × 10⁷ cells/ml). At 3 dpi and 10 dpi, infected roots were washed with water to remove soil. The roots were stored at -80 °C until RNA isolation.

For observation of colonization of Arabidopsis by *FocnCong1-1*, roots of 14-day-old Arabidopsis were cut to approximately 1 cm lengths from the border between roots and stems and soaked in bud cell suspension (1 × 10⁷ cells/ml) for 1 min, then transferred to square plates containing soil. At 12 dpi, roots approximately 5 mm below soil surface were observed by an Olympus BX51 epifluorescence microscopy (Olympus) with excitation of 488 nm for GFP. Images were obtained with an Olympus DP74 digital camera (Olympus) and edited with cellSens (Olympus).

Fungal growth assays. *FocnCong1-1* strains were grown on PDA for 8 days at 28 °C in the dark from a freezer stock. For measurement of colony diameter, mycelium agar disks were collected from the growing edge of a colony using sterile plastic straws and placed in the center of fresh PDA plates. After 8 days, colony diameter was measured. For quantification of conidial formation, 17-day-old colonies were soaked in 10 ml of water and scraped with a colony spreader. Conidial suspensions were filtrated through a nylon mesh to remove mycelia and conidia were quantified at OD₆₀₀ with a WPA CO 8000 Cell Density Meter (WPA) or by counting using haemocytometer.

Plasmid construction. Primers used for plasmid construction are listed in Supplementary Table 3. To generate *SIX8-PSE1* locus complementation vectors, the *FocnCong1-1 SIX8-PSE1*, *Fomt SIX8-PSE1* and *Fol SIX8-PSL1* loci were amplified from genomic DNAs of *FocnCong1-1*, *FomtMAFF240332* and *Fol4287*, respectively, and cloned into pCR⁸/GW/TOPO^{*} vector using a pCR⁸/GW/TOPO^{*} TA Cloning^{*} Kit (Invitrogen) as described by the manufacturer. To introduce these loci into *FocnCong1-1* HS5, the complementation vector containing each locus was co-transformed with pCSN43 containing an *hph* cassette³⁷. For transformation vectors of *SIX8* or *PSE1*, an *hph* cassette was amplified from pCSN43³⁷ and assembled with *SIX8* or *PSE1*, which was amplified from the *FocnCong1-1 SIX8-PSE1* locus complementation vector as a template, using NEBuilder HiFi DNA Assembly Master Mix (New England Biolabs) as recommended by the manufacturer. The assembled fragments were cloned into pCR⁸/GW/TOPO^{*}.

To generate the *FocnCong1-1 SIX8-PSE1* locus disruption vector, the flanking regions of *SIX8* and *PSE1* were amplified from the *FocnCong1-1 SIX8-PSE1* locus complementation vector as a template and assembled with an *hph* cassette using NEBuilder HiFi DNA Assembly Master Mix (New England Biolabs). The assembled fragment was cloned into pCR⁸/GW/TOPO^{*}.

Constructs for yeast two-hybrid assays were generated from cDNAs of *SIX8* and *PSE1* without signal peptide sequences or a stop codon by amplification from cDNA generated from mRNA isolated from *FocnCong1-1*-infected Arabidopsis. Amplicons were inserted into pENTR⁺/D-TOPO^{*} (Invitrogen), and then into yeast expression vectors pDEST-DB and pDEST-AD³⁸ using Gateway[™] LR Clonase[™] II Enzyme Mix (Invitrogen) as described by the manufacturer.

Protoplast formation and transformation. For creation of GFP-labeled *FocnCong1-1* strains (Δ *SIX4*-GFP, HS2-GFP, and HS5-GFP), HS5 transformants (HS5: *SIX8-PSE1*, HS5: *SIX8*, HS5: *PSE1*, HS5: *Fomt SIX8-PSE1* and HS5: *Fol SIX8-PSL1*) and *SIX8-PSE1* knockout mutants (Δ *SIX8-PSE1*), which are listed in Supplementary Table 2, bud cells (4 × 10⁸) were incubated in 80 ml of potato dextrose broth (Difco) at 80 spm for 16 h at 28 °C. Mycelia germinated from the bud cells were collected by centrifugation (1800 g, 10 min) and washed with 1.2 M MgSO₄. Mycelial cell walls were digested with 25 ml 2% (w/v) Driselase (Sigma) and 2% (w/v) Lysing Enzymes (Sigma) in 1.2 M MgSO₄ and maintained at 80 spm for 3 h at 28 °C. Protoplasts were collected by filtration with a nylon mesh and centrifugation (1500 g, 10 min), and rinsed twice with 0.7 M NaCl. The protoplasts were resuspended in STC (1.2 M sorbitol, 50 mM CaCl₂, 10 mM Tris-HCl pH 7.5) and adjusted to 1 × 10⁸ cells/ml. For polyethylene glycol transformation, 30 µg plasmid DNA was added to 150 µl of the protoplast suspension as previously described³⁹. Transformants were selected and maintained on PDA containing

hygromycin B (100 µg/ml) or G418 (200 µg/ml) and verified by PCR using primers listed in Supplementary Table 3. Plasmid DNAs used for transformation are shown in Supplementary Table 4.

Genome sequencing and assembly. For PacBio sequencing, genomic DNA of *FocnCong:1-1* was isolated using CTAB and 100/G genomic tips (QIAGEN) as described in the 1000 Fungal genomes project (<http://1000.fungalgenomes.org>). The genome was sequenced on five PacBio RSII cells and assembled by the Hierarchical Genome Assembly Process (HGAP) v4 within SMRT Link (v5.1.0). Default values were kept and the expected genome size was set to 70 Mb.

For optical mapping, genomic DNA was isolated using a Blood and Cell Culture DNA Isolation Kit (Bionano Genomics) as described by the manufacturer. Genomic DNA was labeled with an NIRS Labeling Kit (Bionano Genomics) with *BspQI* and *BbvCI* as described by the manufacturer. The labeled DNA was scanned using a Bionano Irys platform. Bionano maps from two enzymes (*BspQI* and *BbvCI*) (Bionano Solve v3.2) were merged with PacBio sequence assemblies to produce long hybrid scaffolds. Completeness of gene space within the assembly was assessed through the presence of conserved single-copy genes using BUSCO version 3.0.2^{40,41}. Analysis with the Sordariomyceta data set (3725 genes) indicated the presence of 3690 genes (99.1%) in the assembly (Table 1). Whole-genome alignments were performed with nucmer (with $-maxmatch$) in MUMmer 3.23⁴².

For genome sequencing of *FocnCong:1-1* Δ *SIX4* and HSSs, genomic DNA was isolated using DNeasy Plant Mini Kits (QIAGEN). Illumina NovaSeq 6000 or HiSeq 2500 paired-end sequencing was used for *FocnCong:1-1* Δ *SIX4* and HSSs, except for HS3, using a library with a mean insert size of 550 bp. Illumina NextSeq 500 single-end sequencing was used for *FocnCong:1-1* HS3, from library preparation with a mean insert size of 350 bp. The Illumina sequence library was quality-filtered using the FASTX Toolkit 0.0.13.2 (Hannonlab) with parameters $-q20$ and $-p50$. Reads containing “N” were discarded. Quality-filtered libraries were aligned with the *FocnCong:1-1* genome using CLC Genomic Workbench 20 using default settings.

RNA extraction, cDNA synthesis, and qRT-PCR. Total RNAs were extracted using RNeasy Plant Mini Kit (QIAGEN). Total RNA (200–1000 ng) was used to generate cDNA in a 20 µl volume reaction according to the Invitrogen Superscript III Reverse Transcriptase protocol. cDNA was diluted 1:5, and 1 µl was used for a 10 µl qPCR reaction with 5 µl THUNDERBIRD SYBR Green mix (Toyobo) on an Mx3000P qPCR System (Agilent) using the following program: (1) 95 °C, 1 min, (2) [95 °C, 15 s, then 53 °C, 30 s, then 72 °C, 1 min] \times 40, (3) 95 °C, 1 min for *SIX8* and *PSE1*, or (1) 95 °C, 1 min; (2) [95 °C, 15 s, then 60 °C, 30 s, then 72 °C, 1 min] \times 40, (3) 95 °C, 1 min for *FocnCong:1-1* β -tubulin (*TUB2*), followed by a temperature gradient from 55 to 95 °C. Standard curves were generated using serial dilutions of cDNAs from *Arabidopsis* infected with *FocnCong:1-1* at 10 dpi for *SIX8* and *PSE1* and cDNAs from bud cells for *FocnCong:1-1* *TUB2*. *FocnCong:1-1* *TUB2* was used as a reference gene. Primers used for qPCR are listed in Supplementary Table 3.

RNA sequencing. Using the extracted RNA, strand-specific shotgun type of RNA library was prepared using the Breath Adapter Directional sequencing protocol⁴³. Briefly, mRNA was extracted and fragmented using magnesium ions at elevated temperature. The polyA tails of mRNA was primed by an adapter-containing oligonucleotide for cDNA synthesis. 5' adapter addition was performed by breath capture technology to generate strand-specific libraries. The final PCR enrichment was performed using oligonucleotides containing the full adapter sequence with different indexes. After cleanup and size selection, concentration of library was measured by microplate photometer Infinite²⁰⁰ PRO (TECAN) to pool libraries for Illumina sequencing systems. The libraries were sequenced on an Illumina NextSeq 500 platform. The Illumina sequence library was quality-filtered and aligned as above. Transcription levels for each transcript were calculated as TPM (transcripts per million).

Gene prediction and annotation. RNA sequencing data from *FocnCong:1-1* was aligned with the *FocnCong:1-1* genome using HISAT2 v.2.1.0⁴⁴ and used to guide gene model prediction using the BRAKER1 v1.9 pipeline⁴⁵. BRAKER1 was run with the repeat-softmasked genome created by RepeatMasker v4.0.7 (with $-engine$ ncbi $-species$ “ascomycota” $-xsmall$; <http://www.repeatmasker.org/>), using the fungal and softmasking options. Gene-coding sequences were annotated through BLASTp (E-value cutoff = 1E-6) searches against the July 2018 release of the SWISS-PROT database⁴⁶. Putative secreted proteins were identified through prediction of signal peptides using SignalP v.5.0⁴⁷ and removal of sequences with TMHMM v.2.0⁴⁸-predicted transmembrane domains. For effector prediction, putative secreted proteins were screened for proteins with an effector-like structure using EffectorP 1.0 and/or 2.0^{17,18}. In addition, BLASTp analyses (E-value cutoff = 1E-6) were performed for the fourteen *SIX* genes (*SIX1-SIX14*) and the four genes (*FOAI-FOA4*) known to be effectors in *Arabidopsis*-infecting *F. oxysporum*^{12,16}.

Analysis of repeat elements. Repeat element prediction was performed using the genome sequences of eight *F. oxysporum* strains in the NCBI database that had contig N50 values greater than 1 Mb (last accessed on November 24, 2019) as described in Gan et al.⁴⁹. Code used for this analysis is available at: https://github.com/pamgan/colletotrichum_genome.

The details of genome sequences used for this analysis are shown in Supplementary Table 5. Briefly, repeat sequences were predicted using RECON and RepeatScout via RepeatModeler open-1.0.11 (<http://www.repeatmasker.org>), TransposonPSI (<http://transposonpsi.sourceforge.net/>), LTR_retriever⁵⁰, and LTRPred⁵¹ (<https://github.com/HajkD/LTRpred>). Sequences that were longer than 400 bp from TransposonPSI, LTR_retriever, and LTRPred were combined and used as queries for BLASTx against RepBase⁵² peptide sequences bundled in RepeatMasker open-4.0.9-p2 (<http://www.repeatmasker.org>). Lastly, these sequences were used as queries for BLASTn against each fungal genome. Only sequences with more than five hits (BLASTn E-value cutoff = 1E-15) and/or with a hit to a RepBase peptide (BLASTx E-value cutoff = 1E-5) were retained for further analysis. Sequences from all sources were combined using VSEARCH v2.14.0⁵³, using 80% identity as the cutoff threshold. Consensus sequences were classified using RepeatClassifier (from RepeatModeler open-1.0.11). Known *Fusarium* repeat sequences registered in Dfam_Consensus-20181026 and RepBase-20181026 were extracted, except for those that were annotated as artefacts, simple repeats, or low complexity sequences. The custom repeat library was created by combining the consensus sequences and known *Fusarium* repeat sequences, and used as input for RepeatMasker open-4.0.9-p2. The “one code to find them all”⁵⁴ was used to reconstruct repeat elements.

Chromosome loss and transfer. A chromosome loss experiment was performed according to VanEtten et al.²². *FocnCong:1-1* Δ *SIX4* was incubated in M100 medium (1% glucose, 0.3% KNO₃, 6.25% salt solution) with benomyl (1.56, 3.13, or 6.25 µg/ml) at 120 spm for 4 days at 28 °C. The salt solution consists of 0.4% KH₂PO₄, 0.4% Na₂SO₄, 0.8% KCl, 0.2% MgSO₄·2H₂O, 0.1% CaCl₂, and 0.8% trace elements (0.006% H₃BO₃, 0.014% MnCl₂·4H₂O, 0.0844% ZnSO₄·7H₂O, 0.004% NaMoO₄·2H₂O, 0.006% FeCl₃, 0.04% CuSO₄·5H₂O). Hyphae were removed with a nylon mesh, and bud cells were collected by centrifugation at 1630 g for 10 min. Supernatant was discarded and the remnant with bud cells was spread on M100 plates containing 2% agar and 0.04% Triton X-100 (Wako), and the inoculated plate was overlaid with an autoclaved filter paper. Plates were incubated at 28 °C for 1 to 3 days, then the filter paper was transferred onto M100 medium containing hygromycin B (100 µg/ml) and incubated at 25 °C overnight. Hygromycin B-sensitive isolates were selected by comparing the plates, and then chromosome loss patterns were verified by PCR (Supplementary Fig. 2) using primers listed in Supplementary Table 3.

Chromosome transfer experiments were performed according to van der Does and Rep⁵⁵. A zeocin-resistant *FocnCong:1-1* HS6 (HS6-BLE) strain was generated by *Agrobacterium*-mediated transformation as previously reported⁵⁶ with *Agrobacterium tumefaciens* EHA105 harboring pRW1p⁵⁷. *FocnCong:1-1* Δ *SIX4* and HS6-BLE were co-incubated on PDA at 25 °C. Conidia were harvested from 7-day-old colonies, and conidial suspensions were spread on PDA containing hygromycin B (100 µg/ml) and phleomycin (100 µg/ml). Double drug-resistant colonies were selected, and then chromosome patterns were verified by PCR (Supplementary Fig. 6) using primers listed in Supplementary Table 3.

Contour-clamped homogeneous electric field (CHEF) gel electrophoresis.

CHEF gel plugs were made by resuspending protoplasts in STE (1 M sorbitol, 25 mM Tris-HCl pH 7.5, 50 mM EDTA). Protoplast concentration was adjusted to 4 \times 10⁸ cells/ml and added to the same amount of 1.2% low melting agarose gel (Bio-Rad) solution. Protoplast suspensions (2 \times 10⁸ cells/ml) containing 0.6% low melting agarose gel were added to 50-well disposable mold plates (Bio-Rad). Plugs were immersed in 10 ml of NDS (1% N-lauroyl sarcosinate sodium salt solution, 0.01 M Tris-HCl, 0.5 M EDTA) and incubated at 65 spm for 14 to 20 h at 37 °C. NDS was replaced with 0.05 M EDTA three times every 30 min. Plugs in 0.05 M EDTA were stored at 4 °C until use.

CHEF gel electrophoresis was done according to Inami et al.⁵⁸. Briefly, chromosomes were separated on 1% SeaKem[®] Gold Agarose (Lonza) in 0.5 \times TBE buffer at 4 to 7 °C for 260 h using a CHEF Mapper System (Bio-Rad). Switching time was 1200 to 4800 s at 1.5 V/cm with an included angle of 120°. The running buffer was exchanged every two or three days. Chromosomes of *Hansenula wingie* (Bio-Rad) were used as a DNA size marker. Gels were stained with 3 \times GelGreen (Biotium) to visualize chromosomes.

Yeast two-hybrid assays. For yeast two-hybrid assays, bait (pDEST-DB; DB) and prey vectors (pDEST-AD; AD) containing cDNA of *SIX8*, *PSE1* or empty vector controls were transformed into *S. cerevisiae* Y8930 and Y8800, respectively, with a slight modification of the method described by Lopez and Mukhtar et al.⁵⁹. Transformants carrying DB and AD were selected with synthetic defined (SD) media (0.67% yeast nitrogen base, 0.5% glucose, 0.01% adenine hemisulfate salt) supplemented with -Leu DO supplement (Clontech) (SD-Leu) and -Trp DO supplement (Clontech) (SD-Trp), respectively. Yeast transformants were mated in yeast extract peptone dextrose growth broth (1% yeast extract, 2% peptone, 2% glucose, 0.01% adenine hemisulfate salt) at 150 spm for 24 h at 28 °C. Diploid cells were selected with SD supplemented with -Leu/-Trp DO supplement (Clontech) (SD-Leu-Trp), and spotted on SD supplemented with -Leu/-Trp/-His DO supplement (Clontech) (SD-Leu-Trp-His) and SD-Leu-Trp with 1 mM 3-amino-1,2,4-triazole. Yeast colonies were observed after 72 h incubation.

Statistics and reproducibility. All statistical analyses were performed in EZR⁶⁰. Welch's *t*-test was used to analyze the statistical significance for continuous variables (e.g., OD₆₀₀ value of conidial suspensions), whereas Mann–Whitney *U*-test was used for evaluation of disease severity. The reproducibility was determined by using independent biological replicates as indicated in the figure legends. Individual values for data plots are included in Supplementary Data 3.

Reporting summary. Further information on research design is available in the Nature Research Reporting Summary linked to this article.

Data availability

The Whole Genome Shotgun project of *FocnCong*:1-1 has been deposited at DDBJ/ENA/GenBank under the accession RSAI00000000 (BioProject number PRJNA506492 and BioSample number SAMN10461798). The version described in this paper is version RSAI01000000. RNA sequencing data from culture medium and plant infections have been deposited in NCBI's Gene Expression Omnibus (GEO) and are accessible through GEO Series accession number GSE157823. The source data underlying Fig. 2c, d, 3b, c, 4a, b, d, 5a, c, d and 6c are provided as Supplementary Data 3. Other data are available by reasonable request.

Code availability

Code for repeat element prediction are available at: https://github.com/pamgan/colletotrichum_genome.

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References

- Covert, S. F. Supernumerary chromosomes in filamentous fungi. *Curr. Genet.* **33**, 311–319 (1998).
- Soyer, J. L., Balesdent, M. H., Rouxel, T. & Dean, R. A. To B or not to B: a tale of unorthodox chromosomes. *Curr. Opin. Microbiol.* **46**, 50–57 (2018).
- Miao, V. P., Covert, S. F. & VanEtten, H. D. A fungal gene for antibiotic resistance on a dispensable (“B”) chromosome. *Science* **254**, 1773–1776 (1991).
- Tsuge, T. et al. Host-selective toxins produced by the plant pathogenic fungus *Alternaria alternata*. *FEMS Microbiol. Rev.* **37**, 44–66 (2013).
- Edel-Hermann, V. & Lecomte, C. Current status of *Fusarium oxysporum* formae speciales and races. *Phytopathology* **109**, 512–530 (2019).
- Ma, L. J. et al. Comparative genomics reveals mobile pathogenicity chromosomes in *Fusarium*. *Nature* **464**, 367–373 (2010).
- van Dam, P. et al. A mobile pathogenicity chromosome in *Fusarium oxysporum* for infection of multiple cucurbit species. *Sci. Rep.* **7**, 9042 (2017).
- Vlaardingerbroek, I., Beerens, B., Schmidt, S. M., Cornelissen, B. J. & Rep, M. Dispensable chromosomes in *Fusarium oxysporum* f. sp. *lycopersici*. *Mol. Plant Pathol.* **17**, 1455–1466 (2016).
- Vlaardingerbroek, I. et al. Exchange of core chromosomes and horizontal transfer of lineage-specific chromosomes in *Fusarium oxysporum*. *Environ. Microbiol.* **18**, 3702–3713 (2016).
- de Sain, M. & Rep, M. The role of pathogen-secreted proteins in fungal vascular wilt diseases. *Int. J. Mol. Sci.* **16**, 23970–23993 (2015).
- Houterman, P. M. et al. The mixed xylem sap proteome of *Fusarium oxysporum*-infected tomato plants. *Mol. Plant Pathol.* **8**, 215–221 (2007).
- Schmidt, S. M. et al. MITEs in the promoters of effector genes allow prediction of novel virulence genes in *Fusarium oxysporum*. *BMC Genomics* **14**, 119 (2013).
- Diener, A. C. & Ausubel, F. M. *RESISTANCE TO FUSARIUM OXYSPORUM 1*, a dominant Arabidopsis disease-resistance gene, is not race specific. *Genetics* **171**, 305–321 (2005).
- Li, E. et al. A SIX1 homolog in *Fusarium oxysporum* f. sp. *conglutinans* is required for full virulence on cabbage. *PLoS One* **11**, e0152273 (2016).
- Kashiwa, T. et al. Sequencing of individual chromosomes of plant pathogenic *Fusarium oxysporum*. *Fungal Genet. Biol.* **98**, 46–51 (2017).
- Tintor, N., Paauw, M., Rep, M. & Takken, F. L. W. The root-invading pathogen *Fusarium oxysporum* targets pattern-triggered immunity using both cytoplasmic and apoplast effectors. *New Phytol.* **227**, 1479–1492 (2020).
- Sperschneider, J. et al. EffectorP: predicting fungal effector proteins from secretomes using machine learning. *New Phytol.* **210**, 743–761 (2016).
- Sperschneider, J., Dodds, P. N., Gardiner, D. M., Singh, K. B. & Taylor, J. M. Improved prediction of fungal effector proteins from secretomes with EffectorP 2.0. *Mol. Plant Pathol.* **19**, 2094–2110 (2018).
- Ma, L. J. Horizontal chromosome transfer and rational strategies to manage *Fusarium* vascular wilt diseases. *Mol. Plant Pathol.* **15**, 763–766 (2014).
- Dong, S., Raffaele, S. & Kamoun, S. The two-speed genomes of filamentous pathogens: waltz with plants. *Curr. Opin. Genet. Dev.* **35**, 57–65 (2015).
- Fokkens, L. et al. A chromosome-scale genome assembly for the *Fusarium oxysporum* strain Fo5176 to establish a model *Arabidopsis*-fungal pathosystem. *G3* **10**, 3549–3555 (2020).
- VanEtten, H., Jorgensen, S., Enkerli, J. & Covert, S. F. Inducing the loss of conditionally dispensable chromosomes in *Nectria haematococca* during vegetative growth. *Curr. Genet.* **33**, 299–303 (1998).
- Kashiwa, T. et al. An avirulence gene homologue in the tomato wilt fungus *Fusarium oxysporum* f. sp. *lycopersici* race 1 functions as a virulence gene in the cabbage yellows fungus *F. oxysporum* f. sp. *conglutinans*. *J. Gen. Plant Pathol.* **79**, 412–421 (2013).
- Plaumann, P. L., Schmidpeter, J., Dahl, M., Taher, L. & Koch, C. A dispensable chromosome is required for virulence in the hemibiotrophic plant pathogen *Colletotrichum higginsianum*. *Front. Microbiol.* **9**, 1005 (2018).
- Zhao, Y. et al. Trp-dependent auxin biosynthesis in *Arabidopsis*: involvement of cytochrome P450s CYP79B2 and CYP79B3. *Genes Dev.* **16**, 3100–3112 (2002).
- Bednarek, P. Sulfur-containing secondary metabolites from *Arabidopsis thaliana* and other Brassicaceae with function in plant immunity. *ChemBiochem.* **13**, 1846–1859 (2012).
- Rajniak, J., Barco, B., Clay, N. K. & Sattely, E. S. A new cyanogenic metabolite in *Arabidopsis* required for inducible pathogen defence. *Nature* **525**, 376–379 (2015).
- Sigareva, M. A. & Earle, E. D. Camalexin induction in intertribal somatic hybrids between *Camelina sativa* and rapid-cycling *Brassica oleracea*. *Theor. Appl. Genet.* **98**, 164–170 (1999).
- Schmidt, S. M. et al. Comparative genomics of *Fusarium oxysporum* f. sp. *melonis* reveals the secreted protein recognized by the *Fom-2* resistance gene in melon. *New Phytol.* **209**, 307–318 (2016).
- Thatcher, L. F., Manners, J. M. & Kazan, K. *Fusarium oxysporum* hijacks COI1-mediated jasmonate signaling to promote disease development in Arabidopsis. *Plant J.* **58**, 927–939 (2009).
- Fraser-Smith, S. et al. Sequence variation in the putative effector gene *SIX8* facilitates molecular differentiation of *Fusarium oxysporum* f. sp. *subense*. *Plant Pathol.* **63**, 1044–1052 (2014).
- Kidd, B. N. et al. Auxin signaling and transport promote susceptibility to the root-infecting fungal pathogen *Fusarium oxysporum* in *Arabidopsis*. *Mol. Plant. Microbe Interact.* **24**, 733–748 (2011).
- Ma, L. S. et al. The AVR2-SIX5 gene pair is required to activate I-2-mediated immunity in tomato. *New Phytol.* **208**, 507–518 (2015).
- Cao, L., Blekemolen, M. C., Tintor, N., Cornelissen, B. J. C. & Takken, F. L. W. The *Fusarium oxysporum* Avr2-Six5 effector pair alters plasmodesmatal exclusion selectivity to facilitate cell-to-cell movement of Avr2. *Mol. Plant* **11**, 691–705 (2018).
- Houterman, P. M. et al. The effector protein Avr2 of the xylem-colonizing fungus *Fusarium oxysporum* activates the tomato resistance protein I-2 intracellularly. *Plant J.* **58**, 970–978 (2009).
- Masanaka, A. et al. An isolate of *Alternaria alternata* that is pathogenic to both tangerines and rough lemon and produces two host-selective toxins, ACT- and ACR-toxins. *Phytopathology* **95**, 241–247 (2005).
- Staben, C. et al. Use of a bacterial hygromycin B resistance gene as a dominant selectable marker in *Neurospora crassa* transformation. *Fungal Genet. Rep.* **36**, Article 22 (1989).
- Ahmed, H. et al. Network biology discovers pathogen contact points in host protein-protein interactomes. *Nat. Commun.* **9**, 2312 (2018).
- Watanabe, S. et al. Mode of action of *Trichoderma asperellum* SKT-1, a biocontrol agent against *Gibberella fujikuroi*. *J. Pestic. Sci.* **32**, 222–228 (2007).
- Simao, F. A., Waterhouse, R. M., Ioannidis, P., Kriventseva, E. V. & Zdobnov, E. M. BUSCO: assessing genome assembly and annotation completeness with single-copy orthologs. *Bioinformatics* **31**, 3210–3212 (2015).
- Waterhouse, R. M. et al. BUSCO applications from quality assessments to gene prediction and phylogenomics. *Mol. Biol. Evol.* **35**, 543–548 (2018).
- Kurtz, S. et al. Versatile and open software for comparing large genomes. *Genome Biol.* **5**, R12 (2004).
- Ichihashi, Y., Fukushima, A., Shibata, A. & Shirasu, K. High impact gene discovery: simple strand-specific mRNA library construction and differential regulatory analysis based on gene co-expression network. *Methods Mol. Biol.* **1830**, 163–189 (2018).
- Kim, D., Langmead, B. & Salzberg, S. L. HISAT: a fast spliced aligner with low memory requirements. *Nat. Methods* **12**, 357–360 (2015).
- Hoff, K. J., Lange, S., Lomsadze, A., Borodovsky, M. & Stanke, M. BRAKER1: unsupervised RNA-seq-based genome annotation with GeneMark-ET and AUGUSTUS. *Bioinformatics* **32**, 767–769 (2016).
- Bairoch, A. & Apweiler, R. The SWISS-PROT protein sequence database and its supplement TrEMBL in 2000. *Nucleic Acids Res.* **28**, 45–48 (2000).
- Almagro Armenteros, J. J. et al. SignalP 5.0 improves signal peptide predictions using deep neural networks. *Nat. Biotechnol.* **37**, 420–423 (2019).
- Krogh, A., Larsson, B., von Heijne, G. & Sonnhammer, E. L. Predicting transmembrane protein topology with a hidden Markov model: application to complete genomes. *J. Mol. Biol.* **305**, 567–580 (2001).

49. Gan, P. et al. Telomeres and a repeat-rich chromosome encode effector gene clusters in plant pathogenic *Colletotrichum* fungi. *Environ. Microbiol.* <https://doi.org/10.1111/1462-2920.15490> (2021).
50. Ou, S. & Jiang, N. LTR_retriever: a highly accurate and sensitive program for identification of long terminal repeat retrotransposons. *Plant Physiol.* **176**, 1410–1422 (2018).
51. Benoit, M. et al. Environmental and epigenetic regulation of Rider retrotransposons in tomato. *PLoS Genet.* **15**, e1008370 (2019).
52. Bao, W., Kojima, K. K. & Kohany, O. Repbase update, a database of repetitive elements in eukaryotic genomes. *Mob. DNA* **6**, 11 (2015).
53. Rognes, T., Flouri, T., Nichols, B., Quince, C. & Mahe, F. VSEARCH: a versatile open source tool for metagenomics. *PeerJ* **4**, e2584 (2016).
54. Bailly-Bechet, M., Haudry, A. & Lerat, E. “One code to find them all”: a perl tool to conveniently parse RepeatMasker output files. *Mob. DNA* **5**, 13 (2014).
55. van der Does, H. C. & Rep, M. Horizontal transfer of supernumerary chromosomes in fungi. *Methods Mol. Biol.* **835**, 427–437 (2012).
56. Takken, F. L. W. et al. A one-step method to convert vectors into binary vectors suited for *Agrobacterium*-mediated transformation. *Curr. Genet.* **45**, 242–248 (2004).
57. Houterman, P. M., Cornelissen, B. J. & Rep, M. Suppression of plant resistance gene-based immunity by a fungal effector. *PLoS Pathog.* **4**, e1000061 (2008).
58. Inami, K. et al. A genetic mechanism for emergence of races in *Fusarium oxysporum* f. sp. *lycopersici*: inactivation of avirulence gene *AVR1* by transposon insertion. *PLoS One* **7**, e44101 (2012).
59. Lopez, J. & Mukhtar, M. S. Mapping protein-protein interaction using high-throughput yeast 2-hybrid. *Methods Mol. Biol.* **1610**, 217–230 (2017).
60. Kanda, Y. Investigation of the freely available easy-to-use software ‘EZR’ for medical statistics. *Bone Marrow Transpl.* **48**, 452–458 (2013).

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Author contributions

Y.A., S.A., P.G., A.T., I.Y., and A.S. conducted experiments. Y.A., S.A., K.K., P.M.H., M.R., K.S., and T.A. conceived and supervised the study. Y.A., S.A., K.S., and T.A. wrote the manuscript. All authors reviewed and approved the manuscript.

Competing interests

The authors declare no competing interests.

Additional information

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