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# Genome-wide identification of fitness determinants in the *Xanthomonas campestris* bacterial pathogen during early stages of plant infection

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## Summary

- Plant diseases are an important threat to food production. While major pathogenicity determinants required for disease have been extensively studied, less is known on how pathogens thrive during host colonization, especially at early infection stages.
- Here, we used randomly barcoded-transposon insertion site sequencing (RB-TnSeq) to perform a genome-wide screen and identify key bacterial fitness determinants of the vascular pathogen *Xanthomonas campestris* pv *campestris* (*Xcc*) during infection of the cauliflower host plant (*Brassica oleracea*). This high-throughput analysis was conducted in hydathodes, the natural entry site of *Xcc*, in xylem sap and in synthetic media.
- *Xcc* did not face a strong bottleneck during hydathode infection. In total, 181 genes important for fitness were identified in plant-associated environments with functional enrichment in genes involved in metabolism but only few genes previously known to be involved in virulence. The biological relevance of 12 genes was independently confirmed by phenotyping single mutants. Notably, we show that XC\_3388, a protein with no known function (DUF1631), plays a key role in the adaptation and virulence of *Xcc* possibly through c-di-GMP-mediated regulation.
- This study revealed yet unsuspected social behaviors adopted by *Xcc* individuals when confined inside hydathodes at early infection stages.

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## Introduction

Pests and pathogens of crops cause significant losses in yield and quality. The search for new control strategies relies on our understanding of host–pathogen interactions. *Xanthomonas* bacteria cause disease on more than 400 plant species and are responsible for important losses on various economically important crops worldwide, including rice, citrus, cassava, banana and tomato (Büttner & Bonas, 2010). Among them, *Xanthomonas campestris* pv *campestris* (*Xcc*), the causal agent of black rot disease, is the major bacterial pathogen of *Brassica* crops such as cauliflower, cabbage, mustard and radish (Vicente & Holub, 2013). Transmissible by seeds, *Xcc* survives as an epiphyte and enters the plant through leaf organs called hydathodes. Hydathodes are plant organs located at the leaf margins where they mediate guttation of xylem sap-derived fluid when evapotranspiration is limited (Cerutti *et al.*, 2017, 2019). Hydathodes are composed of three

main tissues: stomata-like water pores within the epidermal layer, an inner epithem composed of small thin-walled parenchyma and a hypertrophied xylem (Cerutti *et al.*, 2017, 2019). Three days after inoculation, *Xcc* switches to necrotrophic behavior and destroys the inner epithem tissue of hydathodes (Cerutti *et al.*, 2017; Luneau *et al.*, 2022a). Simultaneously, *Xcc* accesses xylem vessels and progressively spreads systemically in the plant.

*Xcc* deploys a large arsenal of virulence factors to successfully infect the host plant and complete its life cycle. Among them, two secretion systems are essential for pathogenicity: the type II secretion system (T2SS) which exports enzymes such as plant cell wall-degrading enzymes to the extracellular space and the type III secretion system (T3SS), which translocates type III effector (T3E) proteins into the host cells to suppress immune responses and hijack host metabolism (Büttner & Bonas, 2010; Tang *et al.*, 2021). In addition, *Xcc* produces lipopolysaccharides and exopolysaccharides (EPS) named xanthan that protect bacterial cells against

environmental stresses and support biofilm formation (An *et al.*, 2020). Quorum sensing coordinates bacterial behavior, including biofilm dispersal, and is required for disease (An *et al.*, 2020). *Xcc* also possesses a wide range of two-component systems that ensure perception of environmental signals and initiation of adaptive response (Qian *et al.*, 2008). Together, these traits contribute to the fitness of the bacterium during the infection. However, while previous studies have investigated in depth the virulence factors deployed by *Xcc* to implement its pathogenic lifestyle at late stages of infection, very few have looked at the genetic determinants of fitness at early stages of plant infection (An *et al.*, 2020; Timilsina *et al.*, 2020; Luneau *et al.*, 2022a). *Xcc* colonization of hydathodes is associated with sedentary behavior, activation of pathogenicity determinants (T3SS) and expression of high-affinity transporters for nutrient uptake (e.g. phosphate, sulfate, nitrate; Luneau *et al.*, 2022a). Nevertheless, transcriptome profiling only provides gene expression levels without evaluating their *in planta* contribution to bacterial fitness.

Transposon insertion mutagenesis coupled with next-generation sequencing technologies now allow the study of bacterial fitness at a genomic scale (Cain *et al.*, 2020; van Opijnen & Levin, 2020). Randomly barcoded-transposon insertion site sequencing (RB-TnSeq) allows the high-throughput evaluation of gene contributions to fitness using a saturated library of bacteria mutagenized by barcoded transposons (Wetmore *et al.*, 2015). Such TnSeq approaches have been widely used to study animal pathogens and have led to the discovery of infectious processes including virulence factors, genes required for transmission between hosts or antibiotic resistance (reviewed in Cain *et al.*, 2020; van Opijnen & Levin, 2020) but have only recently been applied to plant pathogens. To date, TnSeq screens of genes contributing to *in planta* fitness have been performed in *Pantoea stewartii* (on corn; Duong *et al.*, 2018), *Dickeya dadantii* (on chicory; Royet *et al.*, 2019), *Agrobacterium tumefaciens* (on tomato; Gonzalez-Mula *et al.*, 2019; Torres *et al.*, 2022), *Pseudomonas syringae* (on bean and pepper; Helmann *et al.*, 2019, 2020), *Ralstonia solanacearum* (on tomato; Su *et al.*, 2021) and *Xanthomonas hortorum* (on lettuce; Morinière *et al.*, 2022). While these studies provided diverse insights specific to each pathosystem, they all highlighted the importance of metabolic capacities and secretion systems for optimal growth *in planta*.

Here, RB-TnSeq was used to identify the genetic basis of *Xcc* adaptation to the environments it colonizes within the host, including hydathodes, xylem sap and synthetic media. We identified essential genes for *Xcc*, as well as genes that contribute specifically to fitness *in planta*. We show that *XC\_3388* encoding a hypothetical protein is important for adaptation and pathogenicity of *Xcc* inside cauliflower hydathodes, probably through c-di-GMP regulation, which highlights the importance of social behaviors during plant infection.

## Materials and Methods

### Bacterial strains and culture conditions

The *Xcc* 8004::GUS-GFP strain (Cerutti *et al.*, 2017) was used as a recipient for the RB-TnSeq library and all deletion mutants

(Supporting Information Table S1). For competition assays, we used the *Xcc* 8004::GUS\*-GFP\* variant, which contains point mutations inactivating the catalytic sites of both reporter proteins as described (Luneau *et al.*, 2022b). *Xcc* was cultivated in MOKA rich medium (Blanvillain *et al.*, 2007), MME glucose-poor medium (Arlat *et al.*, 1991) or MME glucose complemented with dropout – leucine/tryptophan/histidine (LTH) supplements instead of casamino acids (10.5 g l<sup>-1</sup> K<sub>2</sub>HPO<sub>4</sub>, 4.5 g l<sup>-1</sup> KH<sub>2</sub>PO<sub>4</sub>, 1 g l<sup>-1</sup> (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub>, 1 mM MgSO<sub>4</sub>, 0.15% (w/v) DropOut-LTH, 20 mM glucose) at 28°C under agitation at 200 rpm or on MOKA-agar plates.

Deletion mutants were constructed with the *SacB* double recombination method using pK18 derivatives (Table S4; Schäfer *et al.*, 1994). Complemented strains were obtained by genomic integration *in trans* of selected genes under the constitutive *ptac* promoter using pK18\_CompR3 plasmid derivatives (Table S4; Luneau *et al.*, 2022a). Primers designed for all constructions are listed in Table S4. *Escherichia coli* strain TG1 was used as the plasmid donor and strain TG1 pRK2073 as helper for triparental plasmid conjugation into *Xcc* (Figurski & Helinski, 1979; Ditta *et al.*, 1980). *Escherichia coli* TG1 strains were cultivated at 37°C on LB-agar or in LB under shaking at 200 rpm. When appropriate, we used the antibiotics rifampicin (50 µg ml<sup>-1</sup>), kanamycin (50 µg ml<sup>-1</sup>) and spectinomycin (40 µg ml<sup>-1</sup>) and the fungicide pimaricin (30 µg ml<sup>-1</sup>).

### Plant material and xylem sap harvest

Four-week-old *Brassica oleracea* var *botrytis* cv Clovis (L.) F<sub>1</sub> cauliflower plants grown under glasshouse conditions were inoculated in the second true leaf and placed in growth chambers (8 h light; 22°C; 70% relative humidity). Xylem sap was collected from decapitated cauliflower stems as described (Dugé de Bernonville *et al.*, 2014).

### Construction of the RB-TnSeq library and transposon insertion site sequencing

The RB-TnSeq barcoded transposon insertion library was constructed by conjugating the *E. coli* APA752 donor library containing the barcoded *mariner* plasmid pKMW3 into *Xcc* 8004::GUS-GFP. Mating was performed overnight at a 1 : 1 ratio on nutrient agar (2 g l<sup>-1</sup> yeast extract, 5 g l<sup>-1</sup> peptone, 5 g l<sup>-1</sup> NaCl containing 300 µM diaminopimelic acid). The mixture was pooled into a single sample, resuspended in MOKA, plated on MOKA + 50 µg ml<sup>-1</sup> kanamycin plates and incubated for 2 d at 28°C to select for insertion mutants. Approximately 600 000 mutant clones were obtained, each one representing one transposition event. These clones were pooled together in 120 ml MOKA kanamycin and 40 ml of 80% glycerol, aliquoted in 1 ml samples and frozen at -80°C. Genomic DNA from *Xcc* mutant library samples was sequenced and barcodes were mapped as previously described (Wetmore *et al.*, 2015).

### Identification of essential genes

The genome of *Xcc* strain 8004 (NCBI accession no. CP000050, Qian *et al.*, 2005) was annotated using transcriptomic datasets

(Luneau *et al.*, 2022a) and used for all RB-TnSeq analyses (<https://doi.org/10.25794/reference/id52ofys>).

Essential genes were predicted as described (Price *et al.*, 2018). Genes that possess insertion sites (TA site) but that lack insertions in our library are likely to be essential in the condition used to build the library (MOKA-rich medium). Briefly, for each protein-coding gene, we computed the total read density in TnSeq (reads/nucleotides across the entire gene) and the density of insertion sites within the central 10–90% of each gene (sites/nucleotides). We did not consider the DNA barcodes in this analysis of essential genes. Genes that could have no insertion by chance because of their short length (375 nucleotides or shorter) were excluded from the study.

### Fitness screening in hydathodes

For each biological replicate, a 2 ml aliquot of the library was thawed and grown overnight at 28°C with shaking at 200 rpm in MOKA + 50 µg ml<sup>-1</sup> kanamycin. The culture was washed twice by serial centrifugations (6000 g, 10 min) and resuspended in 1 mM MgCl<sub>2</sub>. After measuring the OD<sub>600</sub>, we pelleted 10<sup>9</sup> *Xcc* cells by centrifugation at 13 000 g for 5 min and stored them at -80°C to be used as T<sub>0</sub> samples. Hydathode inoculation was achieved using a protocol adapted from Cerutti *et al.* (2017). Briefly, cauliflower leaves were dipped for 15 s inside the *Xcc* suspension adjusted to an OD<sub>600</sub> = 0.1 (10<sup>8</sup> cfu ml<sup>-1</sup>) in 1 mM MgCl<sub>2</sub> + 0.5% (v/v) Tween 80. Plants were then placed in growth chamber conditions and covered for 24 h with a mini-glasshouse lid to reach 100% relative humidity. The lid was removed and the infection was allowed to proceed for 3 or 6 d. *Xcc* populations were retrieved by macrodissection of hydathodes (16 per leaf) using a 1.5 mm diameter Biopsy Punch (Miltex®; Electron Microscopy Sciences, Hatfield, PA, USA) and bead-ground in 1 mM MgCl<sub>2</sub> for 2 min at 30 Hz. In total, we collected and pooled *c.* 1650 hydathodes for each of the four biological replicates. With an average of 35 000 barcodes per hydathode, this theoretically represents an average of 57 × 10<sup>6</sup> mutants per replicate. Each experiment covers the library *c.* 100 times. DNA was extracted using the Promega Wizard® Genomic DNA Purification Kit in which we added a protein degradation step by incubating the samples with 400 µg ml of proteinase K at 37°C for 1 h before protein precipitation. Samples were stored at -80°C until preparation for barcode sequencing.

### Bottleneck assays in hydathodes

Hydathode inoculation and sampling were performed as described above. After 3 d in the plant, eight hydathodes per leaf were harvested and individually bead-ground in 200 µl of 1 mM MgCl<sub>2</sub> for 2 min at 30 Hz. From this suspension, 10 µl was used to perform serial dilution, plated on MOKA-agar + kanamycin 50 µg ml<sup>-1</sup> and counted to check that each hydathode was well infected and contained the same amount of bacterial cells (*c.* 10<sup>5</sup> CFU per hydathode). We performed four biological replicates of three plants each for a total of 96 hydathodes. Individual hydathode homogenates were spotted on sterile filters, placed on

MOKA-agar + kanamycin 50 µg ml<sup>-1</sup> and incubated for 3 d at 28°C. Grown populations were individually resuspended by mixing filters with 1.5 ml of MOKA and pelleted (6000 g, 10 min). DNA was extracted from each pellet and used for barcode sequencing as described above.

### Barcode sequencing and calculation of gene fitness

Barcode sequencing, mapping and analysis to calculate the relative abundance of barcodes were done using the RB-TnSeq methodology and computational pipeline developed by Wetmore *et al.* (2015); the code is available at <https://bitbucket.org/berkeleylab/feba/>. For each experiment, fitness values (*|f|*) for each gene were calculated as a log<sub>2</sub> ratio of relative barcode abundance after library growth in a given condition divided by relative abundance in the time<sub>0</sub> sample. Fitness values were normalized across the genome so the typical gene has a fitness value of 0. Insertions outside coding sequences were not considered.

We considered genes to be important for *Xcc* fitness when the mutants were strongly affected (i.e. gene fitness *|f|* ≥ 1), and when this phenotype was deemed robust (i.e. *|t*-score| ≥ 3) (Wetmore *et al.*, 2015; Helmann *et al.*, 2019). We retained genes that match those criteria in at least three of the four independent biological replicates. We verified the correlation between biological replicates of RB-TnSeq assays to show experimental reproducibility (Fig. S1). Enrichment analysis considering gene ontology was conducted with R (v.4.0.4) using the TOPGO package v.2.40.0 using Fisher's exact test and *P*-value < 0.01 (Alexa *et al.*, 2006).

### Competition assays

We mixed the *Xcc* strain 8004::GUS\*-GFP\* (hereafter WT\*) and deletion mutants in the 8004::GUS-GFP background to assess the fitness of the latter as previously described (Luneau *et al.*, 2022b). Briefly, we inoculated cauliflower hydathodes with the same protocol as for the RB-TnSeq assay except that we adjusted the inoculum suspensions to OD<sub>600</sub> = 0.05 for each strain (final OD<sub>600</sub> = 0.1) in 1 mM MgCl<sub>2</sub> + 0.5% (v/v) Tween 80. To maximize the throughput of the validation assays, we chose to inoculate only one plant for each of the mixtures. We plated inoculation suspensions to use as T<sub>0</sub> reference samples as well as 6 d postinoculation (dpi) samples consisting of 16 pooled and ground hydathodes. The competitive index (CI) of each strain was calculated as the ratio of the relative frequencies of mutant vs reference strains in the 6 dpi infected sample compared to the T<sub>0</sub> inoculum (Taylor *et al.*, 1987; Macho *et al.*, 2007). Hence, we consider a CI > 1 as a gain of fitness of the mutant compared to the WT\* reference strain and a CI < 1 as a loss of fitness. We performed at least four independent biological replicates of the competition assays for all strains tested.

### Pathogenicity assays

We evaluated bacterial aggressiveness of the mutant strains by piercing-inoculation in the midrib of cauliflower leaves using a needle dipped in a *Xcc* inoculum adjusted at OD<sub>600</sub> = 0.1. We



assessed disease progression at 10 dpi according to the annotation scale presented by Luneau *et al.* (2022a) in three independent biological replicates of five individual plants each.

### Internal growth curves in hydathodes

Bacterial populations in hydathodes were determined after dip-inoculation as described above. After 6 d, hydathodes were sampled using a cork borer (1.5 mm in diameter), individually ground using a Tissue Lyser MM 400 grinder (Retsch, Haan, Germany), twice for 30 s at a frequency of 30 Hz, in 1.2 ml deep-well plates containing five or six glass beads (2 mm in diameter) per well and 200 µl of sterile water. The homogenates were serially diluted in sterile water and 5 µl drops were spotted three times on MOKA plates supplemented with rifampicin and pimarcin. Plates were incubated at 28°C for 48 h and colonies were enumerated in spots containing one to 30 colonies. Bacterial densities in leaves were calculated as log CFU cm<sup>-2</sup>.

### Phenotyping assays

Motility, exopolysaccharide production and protease activity were assessed as previously described (Luneau *et al.*, 2022a). For amylase activity assays, 5 µl of an overnight culture adjusted to  $4 \times 10^8$  CFU ml<sup>-1</sup> was spotted on plates containing 15 ml of MOKA agar with 0.125% potato starch (Prolabo; VWR, Radnor, PA, USA) supplemented with 30 µg ml<sup>-1</sup> pimarcin. Plates were incubated at 28°C and imaged 24 h after inoculation. Diameters of colonies and halos of degradation were measured by washing plates using distilled water and staining using lugol solution until a strong blue coloration and starch degradation halo appeared. Calculation of enzymatic activity was performed as follows:  $\text{PrA} = [\pi(r_{\text{halo}})^2 - \pi(r_{\text{colo}})^2] / [\pi(r_{\text{colo}})^2]$  where  $r$  is the radius. Each experiment was biologically replicated at least three times.

For biofilm visualization, strains were grown overnight in MOKA-rich medium, washed twice with sterile water before being resuspended in MME minimal medium supplemented with 20 mM glucose at a final concentration of  $10^8$  cells ml<sup>-1</sup>. Five milliliters of each strain suspension was distributed in six-well plates (VWR tissue culture plate) containing a sterile borosilicate coverslip in the bottom. One plate per time point is required. Plates were incubated without agitation at room temperature. Before imaging, 3 µl of propidium iodide was added to each well and incubated for 10 min. Biofilms were visualized using a spinning disk microscope with a 60× immersion lens.

## Results

### A saturated library of *Xcc* transposon-insertion mutants identifies 365 essential genes

To screen for genes involved in the adaptation of *Xcc* strain 8004 to environments encountered during its life cycle, a library of randomly barcoded insertion mutants was constructed using a *mariner* transposon (Wetmore *et al.*, 2015; Fig. 1a). High-throughput transposon insertion site sequencing allowed the mapping of each

insertion site in the *Xcc* genome and their association with specific barcode sequences. After filtering out chimeric reads and nonunique transposon barcodes and computationally excluding insertions located outside the main body of the gene (the central 10–90% of the coding sequence length), we identified 192 986 mutants for BarSeq-based fitness analysis (see [Materials and Methods](#) section). The mutant library encompassed 51 853 of the 77 290 possible *mariner* insertion sites (TA dinucleotides) in the *Xcc* 8004 genome (NCBI accession no. CP000050, Qian *et al.*, 2005). Because TnSeq analyses depend on the quality of the annotations, we used an improved annotation of the genome (Luneau *et al.*, 2022a).

With a four-fold coverage and one insertion every 100 bp on average, the mutagenesis was saturated and allowed the study of 3665 of 4617 annotated coding sequences (CDS) (Luneau *et al.*, 2022a).

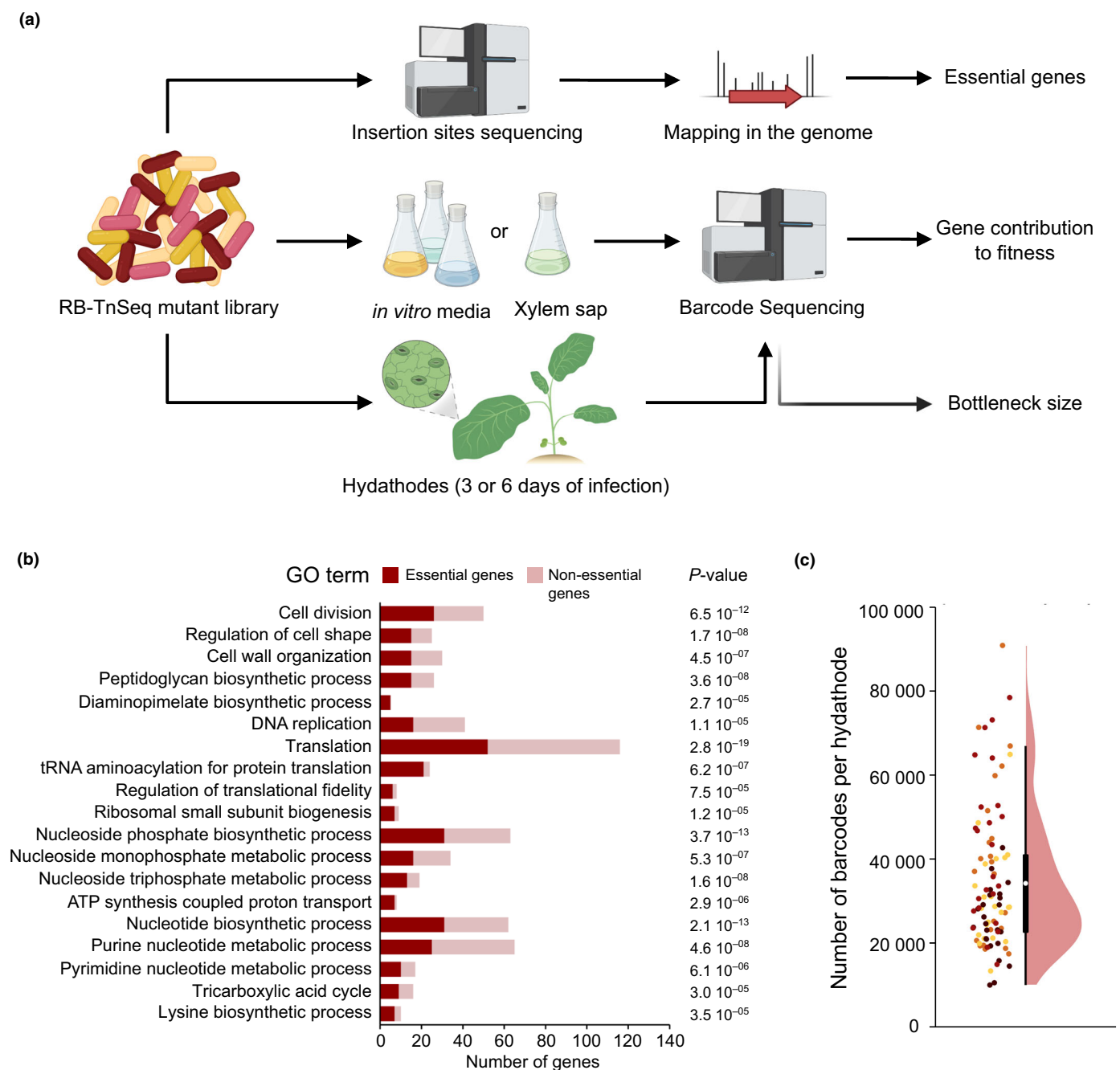
Among those 4617 *Xcc* genes, 80 do not have a TA site in their sequence and cannot be disrupted by a *mariner* transposon. In total, 365 genes were identified as essential for viability in *Xcc* strain 8004 based on both the presence of TA sites localized in the gene body and the absence or very low abundance of transposon insertions (see [Methods S1](#); [Table S2](#)). A Gene Ontology (GO) enrichment analysis highlighted that core cellular functions related to DNA replication, translation, cell envelope, cell division and nucleotide metabolism were most significantly enriched among those essential genes (Fig. 1b). The lysine biosynthesis pathway was the only essential amino acid biosynthetic pathway for *Xcc* growth in the rich medium used to construct the library. The use of such TnSeq library data for essentiality of genes in defined environmental conditions significantly improved functional annotation of the genome of the *Xcc* strain 8004 and identified potential targets for disease control.

### Hydathode pores do not impose a biologically significant infection bottleneck for *Xcc*

Before any TnSeq analysis, it was important to ensure that a significant part of the mutant library can enter host tissues to minimize sampling artifacts. We thus determined the inoculation bottleneck in *Brassica oleracea* var. *botrytis* (cauliflower), the host of isolation of strain 8004 (Turner *et al.*, 1984). Robust hydathode infection was achieved by dip inoculation in the suspension of the mutant library, and the number of distinct barcodes present in each hydathodes was determined by BarSeq (see [Materials and Methods](#) section). A mean of 34 582 distinct barcodes was found per hydathode (Fig. 1c), giving a low-end estimate of the infection bottleneck. This result indicates that, in our experimental setup, the entry into hydathodes is not limiting infection as only a few *Xcc* cells per hydathode are sufficient to initiate an infection that leads to disease (Robeson *et al.*, 1989).

### In total, 181 genes impact *Xcc* multiplication in xylem sap and hydathodes

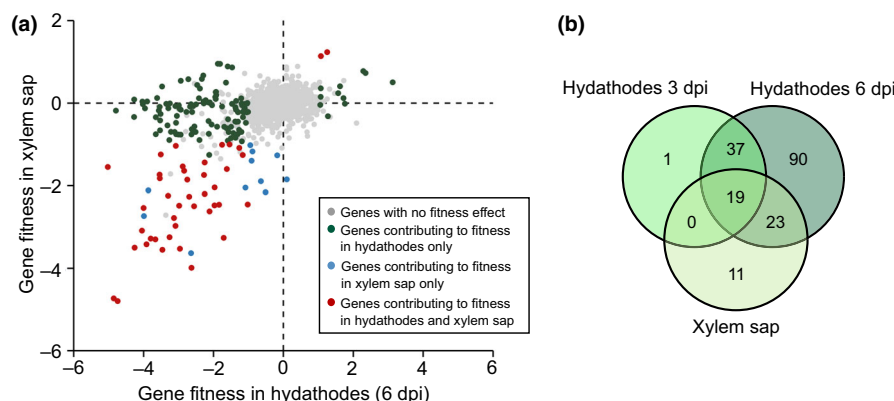
Because *Xcc* is a vascular pathogen colonizing xylem vessels, we first screened for fitness determinants supporting *Xcc* growth in



**Fig. 1** RB-TnSeq approach to identify *Xanthomonas campestris* pathovar *campestris* (Xcc) fitness determinants. (a) Experimental workflow. Barcoded-transposon insertion sites in the Xcc 8004::GUS-GFP genome were determined by transposon insertion site sequencing of the RB-TnSeq mutant library and used to identify essential genes. Barcode sequencing of the bacterial population was performed to quantify the relative abundance of all mutants before and after growth of the library in synthetic media, xylem sap and hydathodes of *Brassica oleracea* var *botrytis* cv Clovis F<sub>1</sub> cauliflower plants. Individual gene fitness values were obtained in each condition. Counting the number of distinct barcodes present within individual hydathodes indicates the bottleneck size for hydathode infection. Created with BioRender.com. (b) Summary of Gene Ontology (GO) functional categories enriched among Xcc essential genes (red) compared to nonessential genes (pink). Enrichment significance (*P*-value) was calculated using Fisher's exact test. (c) Infection bottleneck in hydathodes determined by counting the number of distinct barcodes found within single cauliflower hydathodes (*n* = 96) 3 d after dip inoculation with the RB-TnSeq library. Each dot represents a single hydathode. The color represents the four biological repetitions. The boxplot (black) and densityplot (pink) show the distribution of barcode counts across all samples. The white dot indicates the mean value. The central box of the boxplot shows the central 50% of values (from the first quartile to the third quartile) while the whiskers indicate the values located within 1.5 times the interquartile range.

xylem sap (see Methods S1). Xylem sap contains low concentrations of mineral ions, amino acids, organic acids and simple sugars (Dugé de Bernonville *et al.*, 2014) and was thus used as a

proxy for the environment encountered in the xylem. This *ex planta* RB-TnSeq fitness analysis in cauliflower xylem sap identified 53 Xcc genes (Fig. 2; Table S3a). Fifty-one genes had negative



**Fig. 2** RB-TnSeq screening of *Xanthomonas campestris* pathovar *campestris* (*Xcc*) fitness determinants during growth in plant-associated environments. (a) Comparison of gene contribution to fitness during growth within hydathodes and in xylem sap of *Brassica oleracea* var *botrytis* cv Clovis F<sub>1</sub> cauliflower plants. Genes with no fitness effect or that do not fit our robustness requirement are in gray ( $|l| \geq 1$ ,  $|t\text{-score}| \geq 3$ ). Genes found as differentially represented in hydathodes only are in green, in xylem sap only are in blue and in both conditions are in red. Values used here are the mean of four biological replicates. (b) Venn diagram showing the number of genes contributing to *Xcc* fitness during growth in hydathodes at 3 or 6 d postinoculation and in xylem sap.

fitness scores, indicating that the corresponding genes are needed for growth in xylem sap, while only two had positive fitness scores. These 51 genes are mainly involved in amino acid and sugar metabolism consistent with the limited nutritional value of xylem sap. Comparing ‘plant-associated environments’ together, only 11 of the 53 genes important for fitness in xylem sap were specific to this condition (Fig. 2b; Table S3). Most of these 11 genes are involved in central metabolic pathways and are important for growth in synthetic media (Table S3). (p)ppGpp metabolism was specifically important in xylem sap, where *Xcc* probably faces nutrient-scarce conditions requiring a stringent response (Irving *et al.*, 2021; Bai *et al.*, 2022).

Before reaching the xylem, *Xcc* infects hydathodes where it resides as a biotroph for 3 d and becomes a necrotroph at 6 dpi (Cerutti *et al.*, 2017). An *in planta* RB-TnSeq fitness analysis was conducted at 3 and 6 dpi to dissect early infection processes and adaptive responses inside hydathodes (Fig. 1a). Together, 170 genes significantly contributed to *Xcc* fitness (Fig. 2; Table S3b, c). Despite the low generation number at 3 dpi (three generations), all but one of those early fitness determinants were again identified at 6 dpi, indicating that similar selective forces apply during those two infection stages (0–3 dpi and 4–6 dpi; Fig. 2b). A GO term enrichment analysis revealed that functions related to the biosynthesis of cofactors, purine (inosine monophosphate), lipopolysaccharides (LPS) and multiple amino acids as well as phosphate uptake and gluconeogenesis were significantly associated with *Xcc* fitness in hydathodes.

To distinguish genes specifically important for growth *in planta* from those important for growth *in vitro*, a fitness analysis was performed in MME minimal medium supplemented with glucose as the carbon source as well as in MOKA-rich medium, originally used to build the library (Fig. S2a; Table S3e). Among the 170 genes identified to be important for fitness in hydathodes, 51 are also important for *in vitro* growth, including the 19 genes identified to be important for *Xcc* fitness in both hydathodes and xylem sap (Fig. 2b; Table S3d). These genes are

therefore not specific to *in planta* environments (Table S3e). Those 19 genes were associated with biosynthesis of purine (inosine monophosphate) and vitamins (NADH and folate), and TCA cycle-related pathways. Among amino acid biosynthetic pathways, all except cysteine, methionine, proline and valine were required for optimal fitness in hydathodes (Table S3b). In xylem sap, only proline biosynthesis was important.

### Expression of virulence determinants represents a measurable fitness cost during hydathode infection

Very few genes important for *Xcc* fitness in hydathodes are related to virulence functions (Y-Q. He *et al.*, 2007; Fig. S3). Those genes are involved in EPS biosynthesis and the T2SS, but genes for the T3SS, T3Es or quorum sensing signal biosynthesis were not identified. A gain of fitness was observed in mutants for major virulence regulators, including the HrpG and HrpX transcriptional activators of the T3SS, the RpfC and RpfG two-component system involved in quorum sensing signal perception, and many transcriptional regulators involved in adaptive responses to the environment such as XibR (iron metabolism; Pandey *et al.*, 2016), RavR (oxygen tension; He *et al.*, 2009), HpaR (extracellular proteases production; Wei *et al.*, 2007), VemR (motility and EPS production; Tao & He, 2010) and Clp (EPS and extracellular enzyme production; Y-W. He *et al.*, 2007) (Table 1). Such cross-complementation at the population level highlights the fitness cost of producing virulence factors for the individual and reveals social behaviors of *Xcc* within hydathodes.

### RB-TnSeq enables the identification of genes with subtle and significant contributions to *in planta* fitness or pathogenicity

We assessed individually the fitness of seven mutants corresponding to eight genes with reduced fitness in the RB-TnSeq experiment (Fig. 3; Table 2). The fitness deficit measured by RB-

**Table 1** Genes whose mutation results in a gain of fitness inside hydathodes.

Gene ID <sup>a</sup>	Locus tag <sup>b</sup>	Fitness browser ID <sup>c</sup>	Gene name	Hyd 3 dpi	Hyd 6 dpi	Xylem sap	MME Glucose DROPOUT –LTH	MME glucose	MOKA
XC_0486	XCC8004_a06231	Xcc-8004.624.1	<i>clp</i>	1.13	2.29	0.78	0.24	0.2	0.29
XC_0522	XCC8004_a06761	Xcc-8004.673.1	–	0.2	1.07	0.15	0.04	–0.05	0.04
–	XCC8004_a27231	Xcc-8004.2699.1	–	1.23	1.25	1.23	0.03	0.02	0.28
XC_2228	XCC8004_a27921	Xcc-8004.2768.1	<i>ravR</i>	0.03	1.06	–0.04	0.13	0.06	0.02
XC_2252	XCC8004_a28231	Xcc-8004.2799.1	<i>vemR</i>	0.35	1.07	0.36	–0.42	–0.28	–0.1
XC_2272	XCC8004_a28471	Xcc-8004.2823.1	<i>fliQ</i>	1.39	1.57	0.24	0.07	0.02	0.02
XC_2333	XCC8004_a29211	Xcc-8004.2898.1	<i>rpfC</i>	1.25	2.35	0.73	–0.28	–0.07	0.26
XC_2335	XCC8004_a29231	Xcc-8004.2900.1	<i>rpfG</i>	1.65	3.13	0.5	–0.12	0.06	0.21
XC_2827	XCC8004_a35451	Xcc-8004.3512.1	<i>hpaR</i>	0.73	1.62	0.41	0.04	–0.01	0.27
XC_3076	XCC8004_a38371	Xcc-8004.3806.1	<i>hrpX</i>	0.31	1.72	0.12	–0.11	–0.06	0
XC_3077	XCC8004_a38401	Xcc-8004.3809.1	<i>hrpG</i>	0.77	1.76	–0.02	–0.03	–0.18	0.01
XC_3388	XCC8004_a42701	Xcc-8004.4210.1	–	0.3	1.27	–0.32	–1.24	–0.9	–0.61
XC_3760	XCC8004_a47281	Xcc-8004.4669.1	<i>xibR (xerR)</i>	0.75	1.08	1.14	0.18	0.02	0.31

Fitness values in red for mutants with a gain of fitness and in blue for a loss of fitness; complete fitness data can be found in Supporting Information Table S2. Hyd, hydathodes.

<sup>a</sup>Gene ID corresponds to *Xanthomonas campestris* pathovar *campestris* (Xcc) 8004's CDS annotation given by Qian *et al.* (2005).

<sup>b</sup>Locus\_tag corresponds to the identification tag attributed when the new annotation was performed using RNAseq data published by Luneau *et al.* (2022a).

<sup>c</sup>Fitness browser ID corresponds to the gene name used for fitness analysis.

TnSeq was independently confirmed and complemented upon coinoculation in cauliflower hydathodes with the WT strain (1 : 1 ratio) for six mutants except for *XC\_1113* (Fig. 3a). This experiment supports the RB-TnSeq results (Spearman correlation coefficient  $r=0.92$ ,  $P=0.0011$  for genes with reduced fitness). Alone, five mutants also reached reduced population levels in hydathodes at 6 dpi, while the  $\Delta XC_{1431}$  and  $\Delta XC_{1113}$  mutants behaved similarly to the WT (Fig. 3b). Upon wound-inoculation into cauliflower leaves, five mutants but not  $\Delta glgB1$  also showed reduced pathogenicity at 6 dpi which could be complemented (Fig. 4a).

We also focused on three genes of unknown function (*XC\_0522*, *XC\_3388* and *XCC8004\_a27231-41*) whose mutation conferred a fitness gain in the RB-TnSeq screen at 6 dpi. Competition assays in a 1 : 1 ratio against the WT strain failed to reproduce the gain of fitness observed in RB-TnSeq (Fig. 3c). However, a significant gain of fitness for the  $\Delta hrpX$  and  $\Delta 27231-41$  (knockout of the overlapping *XCC8004\_a27231* and *XCC8004\_a27241* genes) mutants could be measured at a 1 : 100 ratio (mutant : WT, Fig. 3d) indicating that these mutants can indeed behave like cheaters. The failure to reproduce the gain of fitness for  $\Delta XC_{0522}$  and  $\Delta XC_{3388}$  mutants may thus be due to the 1 : 100 ratio because it remains much higher than the ratio of these mutants inside our RB-TnSeq mutant library. Among those three genes,  $\Delta 27231-41$  and  $\Delta XC_{3388}$  deletion mutants had a reduced pathogenicity, though to a lesser extent than the *hrpX* mutant which was avirulent (Fig. 4b).

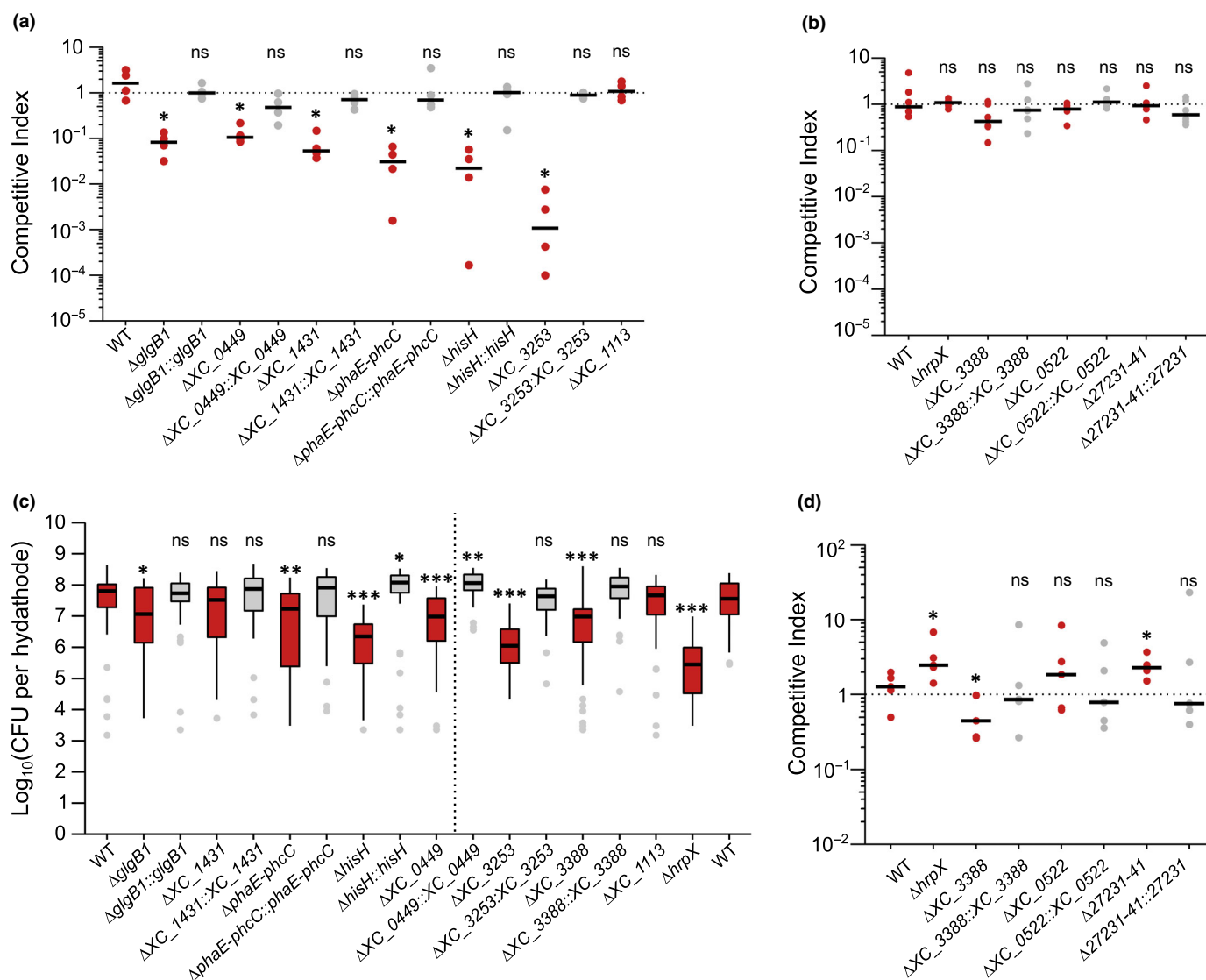
Overall, these results indicate that a gene's contribution to fitness does not strictly correlate with a measurable contribution to virulence nor *in planta* growth (Figs 3, S4b) and that the RB-TnSeq screen was able to robustly identify genes with subtle contributions to fitness and pathogenicity.

## Functional analysis of genes encoding proteins of unknown function involved in pathogenicity

Two putative transcriptional MarR and TetR family regulators (*XC\_0449* and *XC\_1431* respectively) and two proteins of unknown function (*XC\_3253* and *XC\_3388*) which contribute to fitness and pathogenicity during plant infection were selected for their potential roles in the regulation of gene expression. Transcriptome analyses of the mutants and complemented strains grown in MOKA-rich medium were conducted. The *XC\_0449* regulon includes only 11 genes with function involved mainly in amino acid biosynthesis whereas *XC\_1431* seems to regulate only five genes. Most of these genes belong to the same operon (*XC\_1432* to *XC\_1436*) coding for a multidrug efflux system, suggesting that *XC\_1431* is involved in stress responses (Table S4). RNAseq profiling of  $\Delta XC_{3253}$  showed a deregulation of 35 genes, in part associated with stress responses and motility, which was confirmed by plate motility assays (Fig. S5a). Also, 10 genes code for hypothetical proteins and seven genes are involved in sulfate metabolism and biosynthesis of sulfur amino acids such as methionine and cysteine (Table S4). The  $\Delta XC_{3253}$  strain also shows a decrease of extracellular protease activity (Fig. S5b). All of these results partially explain the strong phenotype of this mutant in pathogenicity and fitness. However, the exact function of this protein remains unclear. Because the *XC\_3253* gene is located downstream of an LPS biosynthesis operon, we hypothesize that this gene could be involved in cell envelope formation.

The  $\Delta XC_{3388}$  mutant displayed pleiotropic physiological alterations including hypermotility on swimming plates, reduced extracellular amylase activity, low EPS production and disturbed biofilm formation (Fig. 5). These observations are consistent with the 'rough' colony phenotype visible on agar plates (Fig. S6a). In





**Fig. 3** Contribution of *Xanthomonas campestris* pathovar *campestris* (*Xcc*) genes identified by RB-TnSeq to fitness during infection of cauliflower hydathodes. (a, b) Competition assays at a 1 : 1 ratio of mutants (red boxes) associated with a loss (a) or gain (b) of fitness in RB-TnSeq or complemented mutants (gray boxes) against the *Xcc* strain 8004::GUS\*-GFP\* (hereafter WT\*). Dots represent independent biological replicates ( $N \geq 4$ ) consisting of 16 pooled hydathodes of *Brassica oleracea* var *botrytis* cv Clovis F<sub>1</sub> cauliflower plants each. (c) Internal growth curves in individual hydathodes 6 d after dip-inoculation (three independent biological replicates,  $n \geq 37$  hydathodes in total per strain). Strains separated by the dotted line were inoculated in different experiments. The central box of the boxplot shows the central 50% of values (from the first quartile to the third quartile) while the whiskers indicate the values located within 1.5 times the interquartile range. The median is represented as a black horizontal. Outliers are shown as grey dots. (d) Competition assays at a 1 : 100 ratio (Mutant/WT\*) of *Xcc* mutants associated with a gain of fitness in RB-TnSeq. Dots represent independent biological replicates ( $N \geq 4$ ) consisting of 16 pooled hydathodes each. Statistical significance of growth differences between each strain and the WT was assessed with the Wilcoxon test (ns: not significant; \*,  $P < 0.05$ ; \*\*,  $P < 0.01$ ; \*\*\*,  $P < 0.001$ ).

addition, metabolic fingerprinting on Biolog plates of the  $\Delta$ XC\_3388 mutant identified altered metabolic signatures on several carbon sources including glucose, glutamate, maltose and arbutin (Fig. S6b). These observations were independently confirmed by growth experiments in minimal medium supplemented with these carbon sources (Fig. S6c). These results indicate a broad physiological role for XC\_3388 in *Xcc* adaptation to environmental conditions.

To better evaluate the extent and basis of  $\Delta$ XC\_3388 mutant phenotypes, we compared the transcriptomes of the  $\Delta$ XC\_3388

mutant with the *Xcc* 8004 WT strain and the complemented strain  $\Delta$ XC\_3388::3388 (Fig. 5e; Table S4). In support of the observed phenotypes (Fig. 5a–d), the XC\_3388 mutation strongly induces the expression of chemotaxis and flagellar motility genes while repressing EPS biosynthesis genes. We also noted a decreased expression of genes encoding ribosomal proteins, possibly as a stress response. Importantly, several signaling pathways were significantly deregulated in  $\Delta$ XC\_3388, including the higher expression of many genes encoding GGDEF domain-containing proteins and diguanylate cyclases, which mediate

**Table 2** Candidate genes selected for investigation based on the RB-TnSeq data.

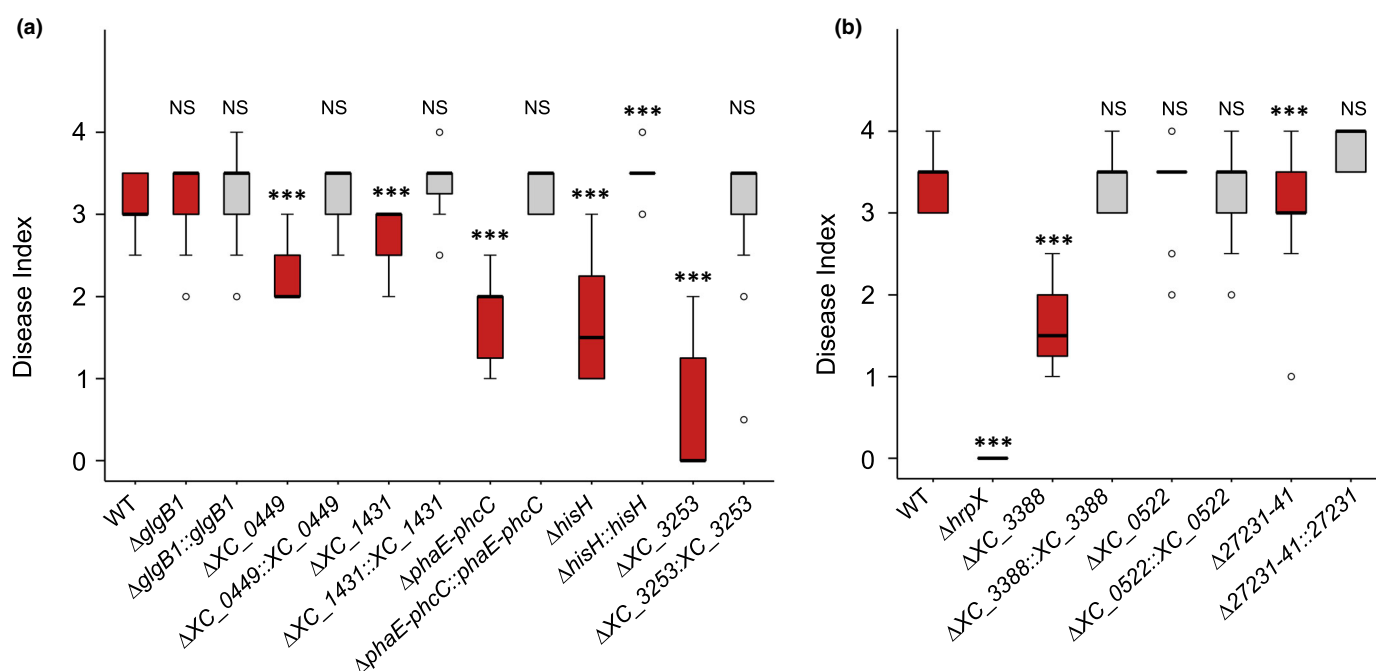
Gene ID <sup>a</sup>	Locus tag <sup>b</sup>	Fitness browser ID <sup>c</sup>	Predicted function	Gene name	Hyd 3 dpi	Hyd 6 dpi	Xylem sap	MME glucose dropout –LTH	MME glucose	MOKA
XC_0143	XCC8004_a01841	Xcc-8004.186.1	Glycogen branching protein	<i>glgB1</i>	–1.97	–2.67	–0.09	–0.08	–0.15	–0.08
XC_0449	XCC8004_a05761	Xcc-8004.577.1	MarR family transcriptional regulator	–	–1.51	–2.14	–0.44	–0.24	–0.22	0.11
XC_0522	XCC8004_a06761	Xcc-8004.673.1	PbsX family transcriptional regulator	–	0.20	1.07	0.15	0.04	–0.05	0.04
XC_1113	XCC8004_a14101	Xcc-8004.1413.1	Ferric enterobactin receptor	<i>bfeA</i>	–0.47	–1.94	0.10	–0.01	–0.05	0.09
XC_1431	XCC8004_a17891	Xcc-8004.1792.1	Transcriptional regulator	–	–0.98	–2.04	–0.05	–0.48	–0.18	0.10
XC_1958	XCC8004_a24301	Xcc-8004.2420.1	PHA synthase subunit	<i>phaE</i>	–1.99	–2.94	–0.77	0.44	0.31	0.10
XC_1959	XCC8004_a24311	Xcc-8004.2421.1	Poly (3-hydroxybutyric acid) synthase	<i>phcC</i>	–1.79	–3.26	–0.90	0.44	0.22	0.03
–	XCC8004_a27231	Xcc-8004.2699.1	Hypothetical protein	–	1.23	1.25	1.23	0.03	0.02	0.28
XC_2377	XCC8004_a29681	Xcc-8004.2945.1	Imidazole glycerol phosphate synthase subunitHisH	<i>hishH</i>	–2.07	–4.03	0.04	–4.30	0.18	0.20
XC_3076	XCC8004_a38371	Xcc-8004.3806.1	HrpX protein	<i>hrpX</i>	0.31	1.72	0.12	–0.11	–0.06	0.00
XC_3253	XCC8004_a40621	Xcc-8004.4026.1	Hypothetical protein	–	–1.61	–4.04	–3.09	–0.63	–0.16	0.05
XC_3388	XCC8004_a42701	Xcc-8004.4210.1	Hypothetical protein	–	0.30	1.27	–0.32	–1.24	–0.90	–0.61

Fitness values in red for mutants with a gain of fitness and in blue for a loss of fitness; complete fitness data can be found in Supporting Information Table S2. Hyd, hydathodes.

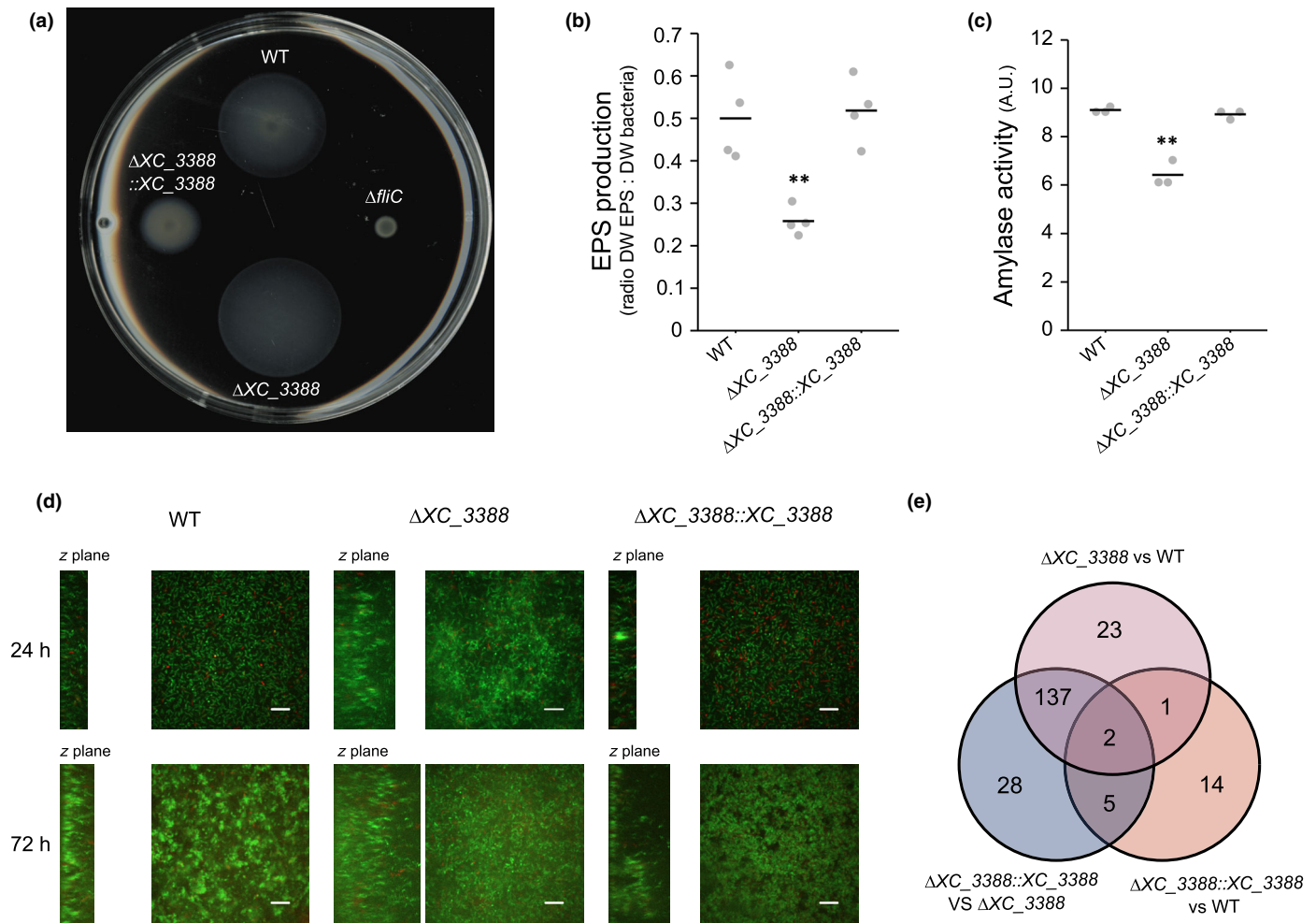
<sup>a</sup>Gene ID corresponds to *Xanthomonas campestris* pathovar *campestris* (Xcc) 8004's CDS annotation given by Qian *et al.* (2005).

<sup>b</sup>Locus\_tag corresponds to the identification tag attributed when the new annotation was performed using RNAseq data published by Luneau *et al.* (2022a).

<sup>c</sup>Fitness browser ID corresponds to the gene name used for fitness analysis.



**Fig. 4** Pathogenicity of *Xanthomonas campestris* pathovar *campestris* (Xcc) mutants in genes identified by RB-TnSeq during infection of cauliflower hydathodes. Severity of disease symptoms caused by knockout (red boxes) and complemented (gray boxes) strains of candidate genes 10 d after piercing inoculation into *Brassica oleracea* var *botrytis* cv Clovis F<sub>1</sub> cauliflower leaf midvein. Strains are grouped based on the phenotype of associated mutants that displayed either a loss (a) or a gain (b) of fitness by RB-TnSeq analysis at 6 dpi in hydathodes. Statistical significance of differences in symptoms severity between each strain and the 8004 WT was determined with the Wilcoxon test (\*\*\*,  $P < 0.001$ ; ns, not significant). The central box of the boxplot shows the central 50% of values (from the first quartile to the third quartile) while the whiskers indicate the values located within 1.5 times the interquartile range (note that sometimes the whiskers are not visible because the first or third quartile coincides with the minimum or maximum value, respectively). The median is represented as a black horizontal line (note that sometimes the box and whiskers are not visible because almost all values are equal to the median). Outliers are shown as empty dots.



**Fig. 5** *In vitro* phenotypes of the  $\Delta XC_{3388}$  mutant strain. (a) Motility assay showing swimming abilities after 48 h on soft 0.3% agar plates. The WT strain and the  $\Delta fliC$  nonmotile flagellin mutant are positive and negative controls, respectively. (b) Mean exopolysaccharide (EPS) production (ratio dry weight EPS/dry weight bacteria) after 24 h of growth in MOKA-rich medium. (c) Mean amylase activity measured after 24 h on MOKA plates supplemented with 0.125% potato starch. (b, c) Statistical significance of differences was assessed with the Wilcoxon test (\*\*,  $P < 0.01$ ). (d) Xcc biofilm architecture using spinning disk confocal microscopy after 24 h (upper panels) and 72 h (lower panels). Representative images are shown. Cells were grown in chambers with a coverslip containing MME minimal medium supplemented with 20 mM glucose. Propidium iodide staining was used to visualize dead cells (red) among live Xcc 8004::GUS-GFP cells (green). Square panels show the biofilm on the x and y planes, while left rectangles show the biofilm on the z plane. Side panel size varies depending on the thickness of the biofilm. Scale bar, 50  $\mu$ m. (e) Number of differentially expressed genes identified by RNAseq ( $|\log_2 FCI| \geq 1.5$  and  $FDR \leq 0.05$ ) during exponential growth in MOKA.

synthesis of c-di-GMP (Table S4). XC<sub>3388</sub> might thus be involved in the metabolism of this important intracellular messenger, providing a plausible explanation for the pleiotropic phenotypes of the  $\Delta XC_{3388}$  mutant. XC<sub>3388</sub> possesses a domain of unknown function, DUF1631, for which no information is available. Structure prediction and comparisons with ALPHAFOLD2 (Jumper *et al.*, 2021) or ROSETTAFOLD (Baek *et al.*, 2021) could not provide additional insights into the biological roles of XC<sub>3388</sub>. BLASTP analysis revealed that XC<sub>3388</sub> is highly conserved among the Xanthomonadales and plant-associated  $\gamma$ -proteobacteria (for conservation and structure predictions, see Methods S1). Together, these results suggest a c-di-GMP-mediated regulatory function for XC<sub>3388</sub>, which is probably conserved in a broad range of

$\gamma$ -proteobacteria. Accordingly, XC<sub>3388</sub> was renamed *pacR* (for pathogenicity and adaptation through c-di-GMP regulation).

## Discussion

### Colonization of hydathodes is constrained by postinvasive selection

By counting the number of barcoded strains entering individual hydathodes, we showed that hydathode pores do not constitute a significant bottleneck with our dip inoculation procedure. It is consistent with the observation that hydathode pores cannot fully close in response to biotic and abiotic stimuli (Cerutti *et al.*, 2017)

and that a single *Xcc* cell can be sufficient for successful hydathode infection (Robeson *et al.*, 1989). Therefore, hydathodes could represent easy infection routes for any microbe. Yet, very few pathogens are reported to infect hydathodes (Cerutti *et al.*, 2019). This paradox suggests that hydathode immunity supported by hostile growth conditions exists and that postinvasive *in planta* selection rather than infection bottlenecks drives the evolution of hydathode pathogens. The drivers of this infectious behavior are unknown but could include the microbial ability to sustain growth in low-nutrient conditions or to overcome hydathode-specific immune responses.

### Metabolic adaptation of *Xcc* to plant environments relies on high-affinity scavenging of mineral and organic matter from the xylem sap and guttation fluid

Our genetic screens highlighted the importance of several amino acid biosynthetic pathways for *Xcc* fitness in hydathodes and xylem sap. We hypothesize that the apoplastic plant tissue environment is limiting for such nutrients. Consistently, *P. syringae* mutants in the tryptophan, histidine and leucine/isoleucine/valine biosynthetic pathways have reduced growth in bean leaf mesophyll (Helmann *et al.*, 2019). Sulfur-containing amino acids methionine and cysteine biosynthetic pathways were important during *Dickeya dadantii* infection of chicory (Royet *et al.*, 2019) in contrast to our study and others (Duong *et al.*, 2018; Gonzalez-Mula *et al.*, 2019; Helmann *et al.*, 2019, 2020; Royet *et al.*, 2019; Torres *et al.*, 2022). These TnSeq results suggest that, even though amino acids are key nutrients, their requirements for fitness vary among pathogenic bacteria and are probably dependent on microbial metabolic features as well as individual amino acid availability within plant tissues.

Cabbage xylem sap contains amino acids, organic acids and sugars in the low millimolar range and most of these compounds are actively metabolized by *Xcc* (Dugé de Bernonville *et al.*, 2014). However, the abundance or the utilization of these amino acids present in xylem sap did not correlate with the gene fitness analysis in xylem sap, suggesting that *Xcc* feeds on multiple nutrients. Interestingly, *relA* (*XC\_1173*), which is important for (p)ppGpp biosynthesis, contributes to fitness in xylem sap. Since accumulation of the (p)ppGpp alarmone triggers the stringent response and allows bacterial cells to face nutrient starvation, it indicates that *Xcc* faces starvation in xylem sap (Bai *et al.*, 2022). Guttation fluid in hydathodes is derived from xylem sap and is presumably depleted from most mineral and organic components, as reported for a wide range of plants (Cerutti *et al.*, 2019). In support of this observation, *Xcc* expresses a diversity of nutrient-scavenging systems inside hydathodes (Luneau *et al.*, 2022a) including high-affinity transporters for phosphate, nitrate and sulfate. Future metabolomic analyses of guttation fluids and xylem sap will be needed to fully characterize the nutritional environment of *Xcc in planta*. These metabolomic and RB-TnSeq results will be instrumental to construct metabolic networks of vascular pathogens and understand their adaptation to a vascular lifestyle (Kim *et al.*, 2019; Gerlin *et al.*, 2020).

### RB-TnSeq identifies genes of unknown functions relevant for pathogenicity

RB-TnSeq allows high-throughput genome-wide screens and provides phenotypic information about individual gene contributions to bacterial fitness. Price *et al.* (2018) exploited this feature to investigate the adaptive value of genes from over 30 ecologically distant bacteria in a wide variety of conditions. This analysis identified condition-dependent cofitness relationships for more than 11 000 previously uncharacterized genes, showing that this approach is particularly powerful to unravel biological functions of hypothetical proteins. Based on our RB-TnSeq screening *in planta*, two proteins with unknown function, *XC\_3253* and *XC\_3388*, were studied in greater detail. In agreement with a preliminary report (Qian *et al.*, 2005), we showed that *XC\_3388*-PacR contributes to adaptation to environmental conditions and pathogenicity. Our study also showed that PacR could be a master regulator of *Xcc* physiology and virulence. The strong deregulation of c-di-GMP-related genes is consistent with the impact on biofilm formation and motility, which are known to be regulated by c-di-GMP (Ryan *et al.*, 2007). Hypothetically, PacR could be involved in c-di-GMP homeostasis, but direct evidence for this is missing. Further studies will be required to elucidate the functions of PacR- and DUF1631-containing proteins at the molecular and cellular levels.

### RB-TnSeq analysis reveals *Xcc* social behaviors inside hydathodes

A key result of our TnSeq study is the gain of fitness displayed by mutants in several major regulators of pathogenicity during hydathode infection. In TnSeq approaches, the fitness of individual cells is measured in a population context. Therefore, mutations associated with the production of public goods (i.e. secreted resources available for the whole population) can be cross-complemented in a population. *Xcc* mutants in those virulence regulators are not only cross-complemented but also have an increased fitness. These results are evidence of a fitness cost associated with the expression of virulence-related genes during hydathode infection. Because *Xcc* cells are confined within hydathodes, we suggest that hydathodes constitute a conducive environment for cheater-like behaviors and that RB-TnSeq can finely quantify the cost of virulence during infection.

There is a growing amount of evidence for spatio-temporal heterogeneity in the dynamics of microbial populations (Ackermann, 2015; Friesen, 2020). Infection strategies sometimes result in very few bacterial cells entering plant tissues, resulting in the direct challenge of their intrinsic fitness. For instance, *P. syringae* infection of tomato leaf mesophyll counter-selects cheaters at low inoculum density but promotes their fitness at high population density (Rufián *et al.*, 2016, 2018). Bistable phenotypic heterogeneity of *P. syringae* was also evidenced in a clonal population leading to the coexistence of two subpopulations that either express or do not express the T3SS (Rufián *et al.*, 2016). Such phenotypic heterogeneity was also reported in the population of phytopathogenic *Ralstonia pseudosolanacearum* during tomato infection (Poussier *et al.*, 2003; Perrier *et al.*, 2019). Based on



frequency-dependent gain of fitness for the *Xcc*  $\Delta hrpX$  mutant in competition assays, bistable T3SS expression could occur theoretically in *Xcc* during hydathode colonization. Further studies will be required to test this hypothesis.

Mechanistically, faster growth of the *hrpX* mutants could be explained by lower expression cost for the T3S regulon in an environment where immunosuppression is established by cooperators, as described for *Salmonella typhimurium* (Sturm *et al.*, 2011). In addition to dividing labor at the population scale, bistable expression of the T3SS in *Salmonella* maintains virulence traits fixed in the population and limits the emergence of T3SS-deficient mutants (Diard *et al.*, 2013). From an evolutionary standpoint, a high abundance of *Xcc* T3SS-deficient mutants could lead to the collapse of infection in proximal and distal tissues. Indeed, according to the Black Queen Hypothesis, emergence of T3SS-deficient mutants would be favored during hydathode infection because it confers local short-term fitness advantages (Morris *et al.*, 2012) allowing mutations occurring early in the infection cycle to be fixed in the population and drive the evolution of *Xcc*. This mechanism may explain how nonpathogenic *Xanthomonas* strains lacking a T3SS and/or T3E are routinely isolated in the phyllosphere (Vorholt, 2012; Merda *et al.*, 2017). Nevertheless, in the larger context of the *Xcc* life cycle, losing the ability to repress host immunity is a weakness compromising *Xcc* systemic spread from the hydathode, seed-mediated propagation of the disease and the infection of neighboring host plants.

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## Competing interests














None declared.

## Author contributions

AB, LDN, AMD and JDL conceived the project. AB supervised the project. JSL performed most of the experiments. MB built the *Xanthomonas* RB-TnSeq library. JF and CG constructed some mutant

strains. TQM and AB performed some competitive index experiments. EL performed *in vitro* and *in planta* bacterial growth assays. MA and BT performed phylogenetic analysis on some selected genes. SC conducted bioinformatic analyses. M-FJ conducted biostatistic analyses. OB contributed to the bottleneck analysis. JR performed the library mapping. The manuscript was written by JSL, LDN and AB. All authors read and approved the final manuscript.

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## Data availability

The fitness data are available from the Fitness Browser (<http://fit.genomics.lbl.gov>). Sequence read archive accession numbers for RNA-Seq libraries are openly available using accession no. SRP363574. Sequence read archive accession numbers for bottleneck analysis are openly available using accession no. SRP363580.

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## Supporting Information

Additional Supporting Information may be found online in the Supporting Information section at the end of the article.

**Fig. S1** Biological reproducibility of RB-TnSeq assays.

**Fig. S2** Growth of the RB-TnSeq library *in vitro* identifies genes contributing to multiplication in plant-independent conditions.

**Fig. S3** *Xanthomonas campestris* pathovar *campestris* (*Xcc*) fitness measured by RB-TnSeq for virulence-associated genes (as defined by Y-W. He *et al.*, 2007) in plant-associated conditions.

**Fig. S4** Correlation between RB-TnSeq fitness results, competitiveness and pathogenicity during hydathode infection with *Xanthomonas campestris* pathovar *campestris* (*Xcc*) mutant strains in genes identified by RB-TnSeq during infection of cauliflower hydathodes.

**Fig. S5** *In vitro* phenotypes of the  $\Delta XC\_3253$  mutant strain.

**Fig. S6** Importance of XC\_3388 for *Xanthomonas campestris* pathovar *campestris* (*Xcc*) metabolism.

**Methods S1** Supplementary material and methods for fitness screening *in vitro*; carbon and nitrogen substrate phenotyping; growth measurements *in vitro*.

**Table S1** Strains, plasmids and oligos used in this study.

**Table S2** List of essential genes of *Xanthomonas campestris* pathovar *campestris* (*Xcc*).

**Table S3** Genes affected in fitness under each tested condition.

**Table S4** RNAseq data.

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