A MATE Transporter is Involved in Pathogenicity and IAA Homeostasis in the Hyperplastic Plant Pathogen *Pseudomonas savastanoi* pv. *nerii*

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**Abstract:** During the last years, many evidences have been accumulating about the phytohormone indole-3-acetic acid (IAA) as a multifaceted compound in the microbial world, with IAA playing a role as a bacterial intra and intercellular signaling molecule or as an effector during pathogenic or beneficial plant–bacteria interactions. However, pretty much nothing is known on the mechanisms that bacteria use to modulate IAA homeostasis, in particular on IAA active transport systems. Here, by an approach combining *in silico* three-dimensional (3D) structural modeling and docking, mutagenesis, quantitative gene expression analysis, and HPLC FLD auxin quantitative detection, for the first time a bacterial multidrug and toxic compound extrusion (MATE) transporter was demonstrated to be involved in the efflux of IAA, as well as of its conjugate IAA–Lysine, in the plant pathogenic hyperplastic bacterium *Pseudomonas savastanoi* pv. *nerii* strain Psn23. Furthermore, according to the role proved to be played by Psn23 MatE in the development of plant disease, and to the presence of Psn23 MatE homologs in all the genomospecies of the *P. syringae* complex, this membrane transporter could likely represent a promising target for the design of novel and selective anti-infective molecules for plant disease control.

**Keywords:** *Pseudomonas savastanoi*; indole-3-acetic acid (IAA); IAA–Lysine; multidrug and toxic compound extrusion transporter; MATE; Type Three Secretion System (TTSS)

1. Introduction

Auxin are plant hormones whose correct homeostasis is pivotal for proper plant growth and development, as well as for plant defense [1]. Indole-3-acetic acid (IAA) is the main and most abundant naturally occurring auxin in plants, as well as the best studied, whose *de novo* biosynthesis is mainly through four interlinked pathways having L-tryptophan (Trp) as a precursor. Generally, the Trp-dependent pathways are two-step reactions, named accordingly to their specific key intermediate molecule, specifically, indole-3-pyruvic acid (IPyA), indole-3-acetamide (IAM), tryptamine (TAM), or indole-3-acetaldoxime (IAOX). Less information is definitely available for Trp-independent IAA biosynthesis, where indole-3-glycerol phosphate or indole are considered the main precursors. The IPyA and IAM pathways are considered the most conserved and used routes for IAA biosynthesis in plants. However, many other important aspects still remain to be fully elucidated,
such as which pathways are used in the different plant species and if they are likely to play alternative roles [2,3].

Firstly discovered in human urine and structurally similar to melatonin in animals [4], in addition to plants IAA is also produced by microalgae, archaea, bacteria, fungi, and yeasts [5]. Although the ability to synthesize IAA in bacteria and fungi is not restricted to those associated to plants, the role of microbial IAA in the interactions between plants and phytopathogenic or beneficial bacteria and fungi is the most studied [6].

Microbial IAA biosynthesis is strictly Trp-dependent, according to at least five different routes, including the IPyA and TAM pathways, as well as the tryptophan side-chain oxidase (TSO) pathway [7,8]. In gall- and tumor-forming bacteria and fungi, IAA has been shown pivotal for the development of hyperplastic symptoms, and its biosynthesis is generally through the IAM pathway. Conversely, the IPyA pathway is mainly represented in beneficial bacteria and fungi. Interestingly, the hyperplastic plant pathogenic bacterium Pantoea agglomerans has both the IPyA and IAM pathways, that are preferentially expressed during epiphytic colonization and the pathogenetic process, respectively [9,10].

Phylogenetic analysis carried out on key genes for IAA biosynthesis in organisms and microorganisms indicates that an independent but convergent evolution was occurred [5]. This finding strongly suggests a universal role of IAA as a signal molecule, both for the producers and during their biotic interactions at different taxonomic levels (e.g., intra and interspecies and even interkingdom) [11]. Plant pathogens have been demonstrated to produce IAA to hijack plant immunity, by subverting plant auxin signaling to increase host susceptibility to infection [6,12–14]. In addition, microbial IAA is also essential as signal molecules within the producer populations, and in plant pathogenic bacteria, IAA was demonstrated to affect the expression of genes of their virulence network [7,15–17].

However, the multiple effects triggered or dynamically modulated by IAA do not exclusively depend on its de novo biosynthesis. In plants, significant and coordinated changes occur over time for local IAA concentrations, as well as for its bioactive forms, also as a consequence of the IAA active polar transport throughout the whole plant and of other processes, such as its catabolism, conjugation, oxidation, storage, and even its signal transduction [18]. A similar fine and dynamic control of IAA homeostasis seems to occur also in bacteria, such as clearly demonstrated for the hyperplastic plant pathogen Pseudomonas savastanoi pv. nerii. Its ability to cause “knots” on its hosts relies on a functional Type Three Secretion System (TTSS), as well as on the bacterial IAA biosynthesis by the IAM pathway [19–21]. In addition, during the pathogenetic process, P. savastanoi pv. nerii regulates free IAA levels in the infected tissues by its conversion to the conjugate IAA–Lysine (hereafter indicated as IAA-Lys), supposed to be less biologically active than the IAA free form. This reaction is mediated by the enzyme IAA-Lys synthase, encoded by the iaaL gene [17]. Interestingly, most of the P. syringae pathogens and strains possess the iaaL gene in their genomes, even if they do not cause hyperplastic symptoms, and this gene appears to be very well conserved and present independently from the genes for IAA biosynthesis [22,23]. It is worth pointing out that the conversion of IAA to IAA-Lys is an exclusive trait of bacteria belonging to the P. syringae complex, and plants neither produce IAA-Lys nor are able to degrade it. Overall, these findings suggest for the bacterial conversion of IAA to IAA-Lys a widely conserved role in the dynamic regulation of the IAA content at and near the infection site. In this frame, it is thus not surprising that in P. savastanoi pv. nerii both the expression of the operon for IAA biosynthesis and that of iaaL gene are also under the control of TTSS, in addition to being auxin-regulated, to further stress the involvement of bacterial IAA and IAA-Lys in the plant–pathogen dialogue since the very first steps of their interaction [17].

Obviously, if a similar dialogue has to take place, it is reasonable to assume that bacterial IAA, and perhaps also its IAA-Lys conjugate, needs to be transported some way out of the bacterial cell into the apoplast. In the P. savastanoi pv. nerii strain Psn23 genome, a gene coding a putative multidrug and toxic compound extrusion (MATE) efflux transporter (hereafter named matE) was found upstream to the iaaL gene, and whose expression was demonstrated to be TTSS-regulated [17]. MATE transporters are widely distributed in Gram-positive and Gram-negative bacteria, where they
are usually associated with the efflux of organic cations for multidrug resistance. Conversely, MATE pumps found so far in plants have been demonstrated to be involved in the transport of a broader range of substrates than in bacteria, and having many other roles beyond detoxification, including the efflux of plant hormones and the regulation of plant disease resistance to pathogens, respectively [24,25].

The aim of this study was to analyze the structure of the putative MATE transporter identified in Psn23 (hereafter named Psn23 MatE) through the application of bioinformatics tools and to evaluate the role played by Psn23 MatE in the development of plant disease and its relationship with the IAA efflux and homeostasis.

2. Materials and Methods

2.1. Bacterial Strains and Growth Conditions

The bacterial strains used in this study are listed in Table S1. Pseudomonas savastanoi pv. nerii strain Psn23 and its mutants were routinely grown at 26 °C on King’s B (KB) [26] or hrp-inducing minimal medium (MM) [27], while Escherichia coli strains TOP10 and ER2925 were grown on Luria-Bertani (LB) [28], as liquid or agarized cultures. Bacterial growth in liquid media was monitored by measuring optical density (OD) at 600 nm (OD600) with a spectrophotometer (Infinite® M200 PRO Multimode Reader, Tecan Group Ltd., Männedorf, Switzerland), while the concentration of viable bacteria was evaluated by plate counts and expressed as colony forming units per milliliter (CFU/mL). For long-term storage, bacteria were maintained at −80 °C on 40% (v/v) glycerol, and P. savastanoi cultures were periodically monitored by using specific PCR-based assays to exclude any bacterial contamination [29,30]. Antibiotics were added to growth medium if needed, and used at the following final concentrations: 20 μg/mL streptomycin, 50 μg/mL nitrofurantoin, 10 μg/mL gentamicin, and 50 μg/mL kanamycin.

2.2. Molecular Techniques

Unless otherwise stated, routine DNA manipulations and PCR were carried out using standard procedures [31] or according to manufacturers’ instructions. The plasmids used in this study are reported in Table S1. Genomic DNA from P. savastanoi strains was extracted from single bacterial colonies using thermal lysis [29], or from bacterial cultures (OD600 = 0.8), using Puregene® Genomic DNA Purification Kit (Qiagen, Hilden, Germany). DNA concentration was evaluated both spectrophotometrically with NanoDrop™ ND-1000 (NanoDrop Technologies Inc., DE, USA) and visually by standard agarose gel electrophoresis on 1% agarose (w/v) in TBE 1× [31]. For plasmid DNA extraction, NucleoSpin® Plasmid (Macherey-Nagel GmbH & Co. KG, Düren, Germany) was used according to the manufacturer’s protocol. Amplicons were purified from agarose gel using NucleoSpin® Gel and PCR clean-up (Macherey-Nagel GmbH & Co. KG) and then double-strand sequenced at Eurofins Genomics (Ebersberg, Germany). Primers were designed using Beacon Designer 7.7 software (Premier Biosoft International, Palo Alto, CA, USA), and their sequences and features are reported in Table S2.

2.3. Construction of Psn23 Mutants for matE Gene

Five mutants were here produced from the wild-type strain Psn23 [30] for the matE gene. The primers were designed according to the matE nucleotidic sequence of Psn23 strain (GenBank Accession Number KU351686), and here used to generate and analyze the mutants listed in Table S2. The suicide vector for P. syringae sensu lato pkI8-AhrpA [17,32] was used to clone the mutated matE constructs into E. coli cells, and then for their transfer into electrocompetent Psn23 cells by using Gene Pulser XCell™ (Bio-Rad Laboratories Inc., Hercules, CA, USA) to replace the native matE gene by marker exchange [17]. A preliminary PCR screening of the putative matE mutants was carried out on transformed Psn23 Suc®/Kan® colonies, and then the marked mutations were confirmed by DNA sequencing. A stable knockout ΔmatE mutant was constructed by an in-frame deletion of matE gene from the Psn23 genome. Three alanine-substituted mutants for the putative Psn23 MatE were also
generated. The matE gene from Psn23 was cloned into the PstI and EcoRI sites of pK18ΔhrpA to produce the pK18-matE recombinant vector (Table S1). On this plasmid, the alanine substitutions D182A, Y200A, and T17035A were introduced into matE by using mutagenic PCR primers and QuickChange II Site-Directed Mutagenesis Kit (Agilent Technologies, La Jolla, CA, USA). The overexpressing mutant Psn23_pT3-matE was also constructed. The recombinant plasmid pT3-matE (Table S1) was produced starting from the vector pLPVM-T3A, which contains the native promoter of hrpA gene for Psn23 [33]. The matE gene, amplified from pK18-matE using the primers pT3_matE_BamHI_For/pT3_matE_KpnI_Rev, was cloned into the KpnI and BamHI restriction sites of pLPVM-T3A and then electroporated into Psn23 cells.

2.4. In Planta Phenotypic Characterization of Psn23 matE Mutants

Hypersensitive Response (HR) assay was carried out on 2 months old Nicotiana tabacum plants (var. Burley White), grown at 24 °C and at 75% relative humidity with a photoperiod of 16/8 h of light/dark. Bacterial cultures, grown overnight in KB medium at 26 °C, were washed twice in sterile physiological solution (SPS, 0.85% NaCl in distilled water) and then resuspended to reach an OD600 = 0.5. Bacteria were then injected into the mesophyll of fully expanded leaves of Tobacco plants, using a 2 mL blunt-end syringe pressed against the abaxial surface (approximately 100 μL/spot) [34]. The appearance of macroscopic symptoms associated to the development of HR was monitored in the next 48 h post-infiltration, taking photographic records of the results obtained. Pathogenicity trials with Psn23 and its mutants were carried out on in vitro micropropagated oleander plants (var. Hardy Red), grown on Murashige–Skoog medium (MS) [35] without addition of any phytohormone for 3 weeks, at 26 °C and with a photoperiod of 16 h/light and 8 h/dark, as described previously [30,36]. Plants were periodically monitored for symptoms appearance, and the bacterial growth was estimated at 7, 14, and 21 days post inoculation (dpi). Three independent experiments were performed, and three plants for each P. savastanoi strain were used in each run.

2.5. Reverse Transcription-Quantitative PCR (qRT-PCR) and Gene Expression Analysis

Liquid cultures of the wild-type strain Psn23 and its mutants were grown overnight at 26 °C in KB, on an orbital shaker (100 rpm), then washed twice with SPS and transferred into MM medium supplemented with L-Trp (0.25mM) to reach an OD600 = 0.5. After 24 h at 26 °C and under shaking condition, bacterial cultures were collected and used for RNA extraction, performed with NucleoSpin® RNA Plus (Macherey-Nagel GmbH & Co. KG), after a treatment with NucleoSpin® gDNA Removal Column (Macherey-Nagel GmbH & Co. KG) to eliminate any genomic DNA. Then, RNA Reverse transcription was carried out on about 5 μg of total RNA per sample, by using iScript™ Advanced cDNA Synthesis kit (Bio-Rad Laboratories Inc.). Diluted cDNA was analyzed with SsoFast™ EvaGreen® Supermix (Bio-Rad Laboratories Inc.) according to manufacturer’s protocols, using the CFX96 Real-Time PCR Detection System and CFX Manager software v1.6 (Bio-Rad Laboratories Inc.). The specific primers pairs here designed and used are listed in Table S2. The expression of each monitored gene was quantified using the 2−ΔΔCT method, and the 16S rRNA gene for normalization. For each sample, three biological replicates were processed in each of the three independent qRT-PCR experiments here carried out.

2.6. Quantification of Bacterial IAA Synthesis

The amount of IAA produced in vitro by the wild-type strain Psn23 and its mutants was assessed both by the colorimetric Salkowski’s method [37] and by high-performance liquid chromatography coupled with fluorescence detection (HPLC FD). Bacterial cultures were grown overnight at 26 °C in KB on an orbital shaker (100 rpm), then washed twice with SPS and transferred into MM supplemented with L-Trp (0.25 mM). At 24 and 48 dpi, bacterial growth was recorded as OD600. The bacterial cultures were centrifuged in a microcentrifuge at 5,000 rpm for 10 min. The supernatants were collected, filter-sterilized on 0.2 μm pore size membranes (Sarstedt, Nümbrecht, German), and then used as such for IAA determination both by Salkowski’s assay and HPLC FD. Standards for
IAA, L-Trp, and IAM at high purity grade (98%) for analytical applications were purchased from Sigma-Aldrich Co. (St. Louis, MO, USA), while IAA–Lysine was kindly synthesized by Department of Chemistry of the University of Padova, Italy. Standard curves were prepared by five ten-fold dilutions of each molecule, starting from 100 ppm in 35% MeOH. The HPLC analyses were performed on a HP 1100 Series chromatograph (Agilent Technologies, Waldbronn, Germany), equipped with diode array (DAD) and fluorescence (FLD) detectors. Chromatographic separations were carried out using a reverse-phase HPLC column ZORBAX ODS (4.6 mm x 250 mm; 5μm) (Agilent Technologies), whose temperature was set up at 40 °C. A 50 μl injection volume and flow rate at 0.9 mL/min were selected. Analytes were separated with an isocratic elution in 35% MeOH. The detection was performed in absorbance at 273 nm and in fluorescence using λex 280 nm and λem 340 nm. The HPLC–DAD/FLD system control, as well as data acquisition and analysis were performed using the ChemStation A.10.01 software (Agilent Technologies).

2.7. Bioinformatic Analysis

Multiple sequence alignments and comparisons were performed by using the computer package Clustal Omega (https://www.ebi.ac.uk/Tools/msa/clustalo/) [38] and basic local alignment search tool (BLAST) (http://www.ncbi.nlm.nih.gov/blast) [39]. Phylogenetic analyses were carried out using neighbor-joining statistical method [40], and phylogenetic trees were generated in MEGA, version 7.0.18 [41]. Bootstrap analysis used 500 replications, Poisson model for substitutions and pairwise deletion method for data treatments (gaps). The cut-off value for condensed tree was 90%. According to the putative MATE protein from Psn23, three-dimensional (3D) structural models were produced by Phyre2 (http://www.sbg.bio.ic.ac.uk/phyre2/html/page.cgi?id=index) [42] and RaptorX (http://raptorx.uchicago.edu/) under default settings [43]. Molecular models visualization was performed using the software USCF Chimera (https://www.cgl.ucsf.edu/chimera/) [44]. The in silico prediction of ligand-binding sites for the interaction with target molecules was made by molecular docking, using the GEMDOCK software (BioXGEM) [45] and AutoDock Vina [46], and the molecular structure of L-Trp, IAM, IAA, and IAA–lysine were assembled in digital format using MarvinSketch 17.6 software (ChemAxon, Budapest, Hungary) (http://www.chemaxon.com).

2.8. Data Collection and Statistical Analysis

The experiments reported in this study were always carried out in triplicate, and at least three independent experiments were performed. The results are reported as means values ± standard deviation (SD). PAST software version 3.11 was used [47] to perform one-way ANOVA followed by Tukey–Kramer’s post-test analysis, and p values ≤0.05 were considered to be statistically significant.

3. Results

3.1. In Vitro IAA Production by Psn23 Depends from a Functional matE Gene

Upstream to the gene iaal, an ORF encoding a putative MATE efflux transporter (GenBank A0R51355) was found near the iaam/iaah operon in P. savastanoi pv. neri Psn23, having a TTSS-dependent expression and hypothesized to mediated IAA efflux, in addition to confer resistance to drugs such as 8 hydroxyquinoline [17]. To test this hypothesis, the in-frame deleted mutant AmatE was produced. As a control, the overexpressing mutant pT3-matE also was generated. Here, the matE gene was under the control of an inducible promoter, that is, the promoter driving the expression of hrpA in the TTSS of Psn23, hereafter named pT3. This promoter can be switched on in vitro when Psn23 is grown on minimal medium (MM), mimicking the apoplast condition [27]. No significant differences were observed between the in vitro growth of Psn23 and its mutants AmatE and pT3-matE, when incubated at 26 °C on MM or on KB, during the first 72 h in shaking condition (100 rpm) (data not shown). Similarly, no differences were found in the ability of the wild-type strain Psn23 and of its mutants AmatE and pT3-matE to cause HR after their infiltration into the mesophyll of Tobacco leaves (Figure 1).
Hypersensitive Response assay on *N. tabacum*. Tobacco leaves were infiltrated with bacterial suspensions of ΔmatE and pT3-matE mutants. For comparison, the wild-type *Psn23* (WT) was also tested. As negative control, sterile physiological solution was used (Neg.Ctrl). Picture was taken 48 h post-infiltration.

Conversely, strongly statistically significant differences were found between *Psn23* and its mutants as far as the *in vitro* production of IAA is concerned. After 24 and 48 h of incubation at 26°C on MM supplemented with Trp, the concentration of IAA in the bacterial free culture supernatants was evaluated by the colorimetric assay based on Salkowski’s reagent [17,48]. As shown in Figure 2, after 24 h the amounts of IAA released into the culture medium by the mutants ΔmatE and pT3-matE were significantly lower and higher, respectively, than that of the wild-type *Psn23*. The exceptionally increased value in IAA production obtained for the mutant pT3-matE was then confirmed as statistically significant also after 48 h.

IAA *in vitro* production by *Psn23* and its ΔmatE and pT3-matE mutants. Bacteria were grown on minimal medium (MM) supplemented with L-tryptophan (Trp, 0.25mM), at 26 °C in shaking condition. Salkowski’s assay was carried out on bacterial supernatants collected after 24 h (blue) and 48 h (red) of growth. The data represent the average of three independent experiments, each with
replicates ± standard deviation (SD). Different letters indicate significant differences among means of mutants at p < 0.05, according to Tukey’s test.

3.2. Expression of matE Gene Influences Expression of Genes for IAA Production and Pathogenicity

The expression of genes related to IAA biosynthesis (iaaM and iaaH) and metabolism (iaaL), to the activation of the master pathogenicity system TTSS (hrpRS and hrpA) as well as to the putative Psn23 MatE transporter here studied (matE), was then evaluated by real-time PCR on Psn23 and its matE mutants, grown on MM supplemented with Trp. The mutants ΔiaaM and ΔiaaL were also included for comparison. The expression of iaaM and iaaH was statistically significantly reduced in the mutant ΔmatE in comparison with the wild-type Psn23, while it was upregulated in the overexpressing mutants pT3-matE (Figure 3). These results are in accordance with the data obtained on the in vitro IAA production for the same bacteria, and with the feedback inhibition of IAA on its own biosynthesis, as known for Psn23 [17].

![Gene expression analysis](image)

**Figure 3.** In vitro gene expression analysis of the Psn23 mutants ΔmatE, pT3-matE, ΔiaaL, and ΔiaaM. Bacteria were grown in vitro on MM for 24 h, and their gene expression was compared with that of the wild-type Psn23, for the genes *hrpA* (light blue), *hrpRS* (orange), *iaaH* (grey), *iaaM* (yellow), *iaaL* (blue), and *matE* (green). Data are averages of triplicates ± standard deviation (SD). Asterisks indicate significant differences compared with the untreated sample at p < 0.05.

Interestingly, in the mutant ΔmatE, also the genes related to the TTSS were downregulated, and just the gene *iaaL* appeared to be overexpressed in comparison with the wild-type Psn23. The gene expression profile of the ΔmatE mutant resulted here quite close to that of ΔiaaM mutant, with the exception of the gene *iaaL*.

Similarly to the pT3-matE mutant, the genes for IAA biosynthesis and matE were statistically significantly overexpressed in the ΔiaaL mutant. Overall, these findings strongly suggest the involvement of the putative Psn23 MatE membrane protein also in the IAA homeostasis of this bacterium, in particular to mediate IAA efflux.

3.3. Virtual 3D Modelling of Psn23 MatE, and Prediction of IAA and IAA-Lys as Putative Substrates

Concerning microbial IAA, up to now just the fungal MATE transporter Mte1 of *Tricholoma vaccinum* has been demonstrated to have a role in IAA efflux, by an indirect approach based on the use of the IAA transport inhibitor 2,3,5-triiodobenzoic acid [49]. Unfortunately, the crystal structure
of MteI is not yet available to perform the most appropriate structure–activity studies to unequivocally demonstrate its involvement in IAA active transport. Conversely, the crystal structures of some bacterial MATE transporters are already available, and all of them are membrane proteins characterized by the presence of twelve transmembrane helices (TM 1–12), forming an internal cavity. In this pocket, several quite specific and conserved residues provide the binding sites for the substrates and for those ions (H⁺ or Na⁺) whose gradient across the membrane serves as energy source [50]. As shown in Figure 4, also the putative Psn23 MatE transporter consists of twelve TMs arranged as two bundles (TM 1–6 and TM 7–12), one at the N- and the other at the C-terminal domain, forming the above-mentioned pocket with a typical V-shaped conformation.

![Diagram](image)

**Figure 4.** Predicted membrane topology and structure of the Psn23 MatE transporter. A) Twelve putative transmembrane (TM) domains were predicted for Psn23 MatE, on the basis of amino acid hydrophobicity; B) Ribbon three-dimensional (3D) model of Psn23 MatE, viewed parallel to the membrane (left) and along the membrane normal from the periplasmic side (right), with the twelve TM helices numbered starting from the first N amino acid (methionine).

According to this 3D model, in the TM1 and TM5 of the N lobe of Psn23 MatE two negative amino acids (E35 and D182) were found to be located, as occurring in the same TMs (D41 and D184) of the H⁺-driven MATE transporter from *Pyrococcus furiosus* (known as PfMATE) to give the cation-binding site [51,52]. In contrast, no negative-charged amino acids were found in the TM7 and TM10 of Psn23 MatE as well as of PfMATE, as occurring in the NorM-type MATE transporters which usually use the Na⁺ motive force to drive the substrate transport across the membrane [52,53].

Overall, these findings support the hypothesis of Psn23 MatE transporter belonging to the DinF subfamily of the prokaryotic MATE transporters. On these bases, a virtual structure-based ligand analysis was then performed to assess if IAA and its conjugate IAA-Lys could be substrates for Psn23 MatE, by using the GEMDOCK software (BioXGEM) and PfMATE as comparative protein structure model [45]. The binding free energy calculations showed that free IAA and IAA-Lys could be realistically considered among the potential substrates transported by Psn23 MatE across the bacterial membrane, and that the Y200, T170, T173, and T175 residues are important in substrate binding (Table 1).
Table 1. Energy values of interaction between selected Psn23 MatE amino acids and several putative ligands. The value for each amino acid residue represents the energy of the single bond (expressed as kcal/mol). H = hydrogen bonding; V = van der Waals forces.

| Ligand    | Energy | H T175 | H Y200 | V T170 | V T173 | V T175 | V Y200 |
|-----------|--------|--------|--------|--------|--------|--------|--------|
| L-Trp     | −72.1  | 0      | 0      | 0      | 0      | 0      | −11.1  |
| IAM       | −71.2  | −2.9   | −1.4   | 0      | 0      | −0.9   | −10.6  |
| IAA-free  | −102.5 | 0      | 0      | 0      | −1.5   | −28.5  |
| IAA-lysine| −95.1  | −5.9   | 0      | −2.1   | −3.5   | −3.9   | −16.8  |

While Y200 has to be considered potentially involved in the binding of both free IAA and of its conjugate with Lysine, the residues T170, T173, and T175 showed to have a stronger interaction with IAA-Lys than with IAA, probably because of a specific affinity of polar uncharged amino acids for lysine, such as occurring for threonine.

3.4. Site-Directed Mutagenesis of Psn23 MatE to Confirm Its Involvement in Pathogenicity and IAA Secretion

According to the data from virtual 3D modelling and docking analysis performed on Psn23 MatE, site-directed mutagenesis was then carried out on those residues of its N-lobe supposed to be involved in cation-binding (i.e., D182) or in substrate recognition and interaction (i.e., T170, T173, T175, and Y200). Therefore, the alanine-substituted mutants D182A and Y200A were obtained by single substitution events, while by a triple substitution the mutant T17035A was generated. As previously reported for the Psn23 mutants ΔmatE and pT3-matE, the in vitro growth of the alanine-substituted mutants D182A and Y200A, and T17035A was not impaired in comparison with the wild-type Psn23, when incubated at 26 °C on MM or on KB in shaking condition (data not shown).

Pathogenicity tests were then carried out on these mutants, as well as on the ΔmatE and pT3-matE mutants, by using in vitro micropropagated oleander plants (var. Hardy Red) [17]. For comparison, the wild-type Psn23 was also used. Plants were periodically monitored for symptoms appearance, and the development of the typical hyperplastic knot was firstly visible at naked eyes at about 7 dpi. The in planta bacterial growth was also periodically evaluated at 7, 14, and 21 dpi.

The results obtained at 21 dpi are shown in Figure 5, where it is obvious the significant reduction in the size of the knots occurring in the ΔmatE-inoculated plants, as well as on those infected by the alanine-substituted mutants D182A and Y200A.
Figure 5. Hyperplastic symptoms obtained at 21 dpi in pathogenicity tests carried out on micropropagated oleander plants with Psn23 wild type and its mutants ΔmatE, pT3-matE, D182A, Y200A, and T17035A. Control: sterile physiological solution (SPS, NaCl 0.85%) inoculated plants. Complemented mutant for ΔmatE: ΔmatE(matE).

Their hypovirulent phenotype was further confirmed by the data on their growth in planta, which was statistically significantly reduced in comparison with the wild-type Psn23 (Figure 6). A reduction in their in planta growth ability was also observed at 21 dpi for the mutants T17035A and pT3-matE, although the hyperplastic galls they generated on infected plants were slightly bigger or comparable in size to those for the wild-type Psn23, for pT3-matE and T17035A, respectively (Figure 6).
These findings undoubtedly demonstrated that the putative MATE transporter coded in *Psn*23 by the *matE* gene is definitely involved in the virulence of this plant pathogen, expressed as ability to cause symptoms and to grow inside the infected plant host tissues. Most importantly, this role appears to depend on *Psn*23 MatE ability to transport bacterial biosynthesized IAA using the H+ motive force, similarly to *Pf*MATE [52]. It was therefore essential to increase the resolution of the analysis for the quantification of IAA and its conjugate IAA-Lys, as well as its intermediate IAM, synthesized in vitro by *Psn*23 and its MatE mutants by using high-performance liquid chromatography coupled with a fluorescent detector (HPLC–FLD). In Figure 7, the data related to the in vitro production of IAA and IAA-Lys are evaluated by HPLC–FLD on the cell-free filtrates of *Psn*23 and the mutants *matE* and pT3-*matE*, after 24 and 48 h of growth on MM supplemented with Trp. For comparison, the hypovirulent ΔiaaM and the hypervirulent ΔiaaL mutants were also tested, thus to confirm what was already known and expected, that is the ΔiaaM inability to synthesize IAA as well as the hyperproduction of free IAA obtained by ΔiaaL. The mutants *matE* and pT3-*matE* have been here shown to have a behavior similar to that of ΔiaaM and ΔiaaL, respectively, for both IAA production and virulence.
Figure 7. IAA in vitro production by the wild-type Psn23 (WT) and its mutants ΔmatE, pT3-matE, ΔiaaM, and ΔiaaL. After 24 h (A) and 48 h (B) of in vitro growth on MM supplemented with Trp, free IAA and IAA-Lys were quantified in the bacterial supernatants by HPLC–FLD. Data are averages of triplicates ± standard deviation (SD). Different letters indicate significant differences among means of mutants at p < 0.05, according to Tukey’s test.

As far as the Psn23 MatE alanine-substituted mutants are concerned, in Figure 8 the results obtained are reported. As expected, the hypovirulent mutants D182A and Y200A showed a reduced production of IAA and increased levels of IAA-Lys, with a behavior coherent to that of ΔmatE, that is the other hypovirulent mutant. Accordingly, no particular differences were found in the biosynthesis of IAA and its metabolite for the T17035 mutant in comparison with the Psn23 wild type, as occurring also for its ability to cause symptoms on the host plant.

Figure 8. IAA in vitro production by the wild-type Psn23 (WT) and its MatE alanine-substituted mutants D182A, Y200A, and T17035A. For comparison, the mutant ΔmatE was also used. After 24 h of in vitro growth on Minimal Medium MM supplemented with Trp, free IAA and IAA-Lys were
quantified in the bacterial supernatants by HPLC–FLD. Data are averages of triplicates ± standard deviation (SD). Different letters indicate significant differences among means of mutants at \( p < 0.05 \), according to Tukey’s test.

4. Discussion

In plants, phytohormones are known to finely regulate plant morphogenesis and development, and their involvement in plant–microbe interactions has been demonstrated as well [1]. In this frame, a pivotal role is played by the auxin IAA, whose levels in planta can be modulated by phytopathogens to promote susceptibility in their potential hosts. Gram-negative phytopathogenic bacteria belonging to the so-called *P. syringae* complex have been shown to hijack IAA accumulation or auxin signaling by specific virulence factors, such as several TTSS effectors. In addition, most *P. syringae* bacteria produce IAA, even if they do not cause any hyperplastic symptoms, and its role as a signaling molecule able to regulate bacterial gene expression as well as virulence has been ascertained [9,10]. The hyperplastic activity of *P. savastanoi* pv. *nerii* strain *Psn23* on oleander was shown to depend on the balance between free IAA and its conjugate IAA-Lys in the infected tissues, and this process was demonstrated to be under the control of TTSS [17]. In this auxin-based dialogue with plants, up to now the aspect definitely less investigated has been how IAA-producing phytopathogenic bacteria secrete this phytohormone into the apoplast.

Here, for the first time, a bacterial MATE transporter was demonstrated to mediate IAA efflux in the plant pathogen *P. savastanoi* pv. *nerii* strain *Psn23*, as already known for some plant MATE membrane proteins which mediate transport of several phytohormones, including auxins [24]. The same role has been demonstrated just for a microbial MATE, that is Mte1 from *T. vaccinum* [49]. By targeted mutagenesis, several amino acid residues involved in *Psn23* MatE functionality have been identified. According to these data and to its putative structure, obtained by in silico 3D modeling, *Psn23* MatE appears to belong to the DinF subfamily of the prokaryotic MATE transporters, such as the H⁺-driven MATE transporter from *P. furiosus* [51].

Just one other bacterial MATE transporter has been identified so far in plant pathogenic bacteria, particularly in *Erwinia amylovora*. The norM gene from *E. amylovora* codes for a protein highly homologous to the NorM MATE transporter of *E. coli* and *Vibrio parahaemolyticus*, and it was demonstrated to confer resistance to toxins produced by several epiphytic bacteria colonizing the same habitat in addition to the canonical resistance to some hydrophobic cationic antibiotics [54].

Here, for the first time, a bacterial MATE transporter was demonstrated to be involved in the molecular dialogue between a phytopathogenic bacterium and its potential host plant, by modulating IAA homeostasis in *Psn23* through the MATE-mediated auxin transport. It is well established that in *P. savastanoi* the expression of several TTSS genes is downregulated by IAA. Therefore, in the first step of the infection process induced by *P. savastanoi* it is reasonable to hypothesize that the intracellular IAA levels have to be carefully modulated by some homeostatic mechanisms, for the most part unknown but which certainly include IAA conjugation, to give auxin metabolites having a lower biological activity than IAA, such as IAA-Lys. In the *ΔiaAL* mutant, unable to conjugate IAA with Lysine, IAA secretion is strongly increased in association with a hypervirulent phenotype [17]. The same phenotype was here found for the overexpressing mutant pT3-*matE*. Conversely, a hypovirulent phenotype was scored for the *Psn23* mutants having their MATE transporter somehow impaired, as occurring for the deleted mutant *ΔmatE* and for the alanine-substituted mutants D182A and Y200A. In addition, these mutants also showed a reduced *in vitro* IAA production in comparison with the wild-type *Psn23*. Overall, these data demonstrated that *Psn23* MatE mediates IAA efflux, thus to contribute to maintain the most appropriate IAA intracellular concentrations in each step of the whole infective process, as also suggested by its TTSS-dependent expression, in order to maximize the chances of success. This regulation of IAA homeostasis is played together with IAA-Lysine synthase. The *iaaL* expression is also TTSS-dependent, as well as coordinated with that of *matE* and IAA-inducible [17]. According to the experimental data obtained by HPLC–MS on the IAA *in vitro* production by *Psn23* and its MATE-mutants, *Psn23* MatE can be fairly hypothesized to transport
IAA but not its conjugate IAA-Lys, for which additional secretion mechanisms have to be taken into consideration.

However, in addition to the virtual structure-based ligand analysis here carried out, the crystal structure determination and analysis of Psn23 MatE is the next key step to definitely elucidate those substrates specifically recognized and transported by this membrane protein, as well as the actual antiporter cation (H+ or Na+). A deeper understanding of mechanisms underlying the physiology and the activity of these evolutionarily conserved transport proteins also in bacterial plant pathogens is advisable. For its role in the efflux of a virulence factor such as IAA and also in the resistance to several antimicrobials [17], Psn23 MatE has indeed to be considered an important promising target for the development of innovative and ecofriendly strategies for the control of Psn23 as well as of other pathogenic bacteria, possibly by the use of natural indole-mimicking competitors.

Supplementary Materials: The following are available online at www.mdpi.com/xxx/s1, Table S1: Bacterial strains, mutants, and plasmids used in this study; Table S2: Primers used in this study.

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