Transcriptome Profiling of the Endophyte *Burkholderia phytofirmans* PsJN Indicates Sensing of the Plant Environment and Drought Stress

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**ABSTRACT** It is widely accepted that bacterial endophytes actively colonize plants, interact with their host, and frequently show beneficial effects on plant growth and health. However, the mechanisms of plant-endophyte communication and bacterial adaptation to the plant environment are still poorly understood. Here, whole-transcriptome sequencing of *B. phytofirmans* PsJN colonizing potato (*Solanum tuberosum* L.) plants was used to analyze *in planta* gene activity and the response of strain PsJN to plant stress. The transcriptome of PsJN colonizing *in vitro* potato plants showed a broad array of functionalities encoded in the genome of strain PsJN. Transcripts upregulated in response to plant drought stress were mainly involved in transcriptional regulation, cellular homeostasis, and the detoxification of reactive oxygen species, indicating an oxidative stress response in PsJN. Genes with modulated expression included genes for extracytoplasmic function (ECF) group IV sigma factors. These cell surface signaling elements allow bacteria to sense changing environmental conditions and to adjust their metabolism accordingly. TaqMan quantitative PCR (TaqMan-qPCR) was performed to identify ECF sigma factors in PsJN that were activated in response to plant stress. Six ECF sigma factor genes were expressed in PsJN colonizing potato plants. The expression of one ECF sigma factor was upregulated whereas that of another one was downregulated in a plant genotype-specific manner when the plants were stressed. Collectively, our study results indicate that endophytic *B. phytofirmans* PsJN cells are active inside plants. Moreover, the activity of strain PsJN is affected by plant drought stress; it senses plant stress signals and adjusts its gene expression accordingly.

**IMPORTANCE** In recent years, plant growth-promoting endophytes have received steadily growing interest as an inexpensive alternative to resource-consuming agrochemicals in sustainable agriculture. Even though promising effects are recurrently observed under controlled conditions, these are rarely reproducible in the field or show undesirably strong variations. Obviously, a better understanding of endophyte activities in plants and the influence of plant physiology on these activities is needed to develop more-successful application strategies. So far, research has focused mainly on analyzing the plant response to bacterial inoculants. This prompted us to study the gene expression of the endophyte *Burkholderia phytofirmans* PsJN in potato plants. We found that endophytic PsJN cells express a wide array of genes and pathways, pointing to high metabolic activity inside plants. Moreover, the strain senses changes in the plant physiology due to plant stress and adjusts its gene expression pattern to cope with and adapt to the altered conditions.

*Burkholderia phytofirmans* PsJN (1) is a naturally occurring endophyte isolated from onion roots (2) that is able to establish nonpathogenic relationships with a wide range of plant species, both monocotyledons and dicotyledons (3, 4). Numerous studies have reported beneficial effects of strain PsJN on host plants such as increased plant growth (for a review, see reference 5) and enhanced biotic and abiotic stress tolerance (4, 5).

There are increasing efforts to understand the plant physiological and genetic response to inoculation with *B. phytofirmans* PsJN. Bordic and colleagues (6) compared local defense reactions in grapevine cell cultures inoculated with either strain PsJN or the nonhost pathogen *Pseudomonas syringae* pv. *pisum*. The authors found that strain PsJN does not induce the defense events commonly found after pathogen attack in plants. Infection with the pathogen led to a two-phase oxidative burst and a hypersensitive response (HR)-like reaction, whereas no oxidative burst or cell death was observed in cells treated with strain PsJN (6).

Theocharis et al. (7) demonstrated that inoculation of Chardonnay grapevine plantlets with *B. phytofirmans* PsJN speeds up the plant response to chilling and plant adaption to cold temperatures. Numbers of cold-stress-related gene transcripts and metabolites increased earlier and faster and reached higher levels in bacterized plantlets than in control plants. Fernandez et al. (8) demonstrated that the higher tolerance of chilling of PsJN-
colonized grapevine plantlets could be related to changes in plant photosynthesis and sugar metabolism. More recently, Poupin and colleagues (9) studied the response of Arabidopsis thaliana to inoculation with B. phytofirmans PsJN. The bacterium affected the whole life cycle of Arabidopsis plants, increasing plant growth, especially at the early stage of ontogeny, and speeding up maturity. These physiological changes correlated with altered expression of plant growth regulator genes; i.e., genes involved in auxin and gibberellin pathways were induced in PsJN-inoculated plants, and flowering-control genes were induced earlier in PsJN-inoculated plants than in control plants.

Whereas the plant response to beneficial bacteria has been described in several studies, very little is known about bacterial physiology, response, and adaptation processes in planta. Efforts to characterize the effects of the plant environment on endophytic bacteria have been rare (10–12), and information on in planta gene expression of B. phytofirmans PsJN is missing. For example, how does B. phytofirmans PsJN recognize the plant environment? Does the bacterium respond to changing physiological conditions, e.g., due to plant stress, in plants?

Therefore, the aim of this study was to investigate gene expression patterns of B. phytofirmans PsJN cells colonizing potato (Solanum tuberosum L.) plants and, furthermore, to assess the effect of plant drought stress on the transcriptome of strain PsJN. In vitro-grown potato plants of two varieties (Russet Burbank and Bionta) were inoculated with B. phytofirmans PsJN, and drought stress was induced by application of polyethylene glycol (PEG). Bacterial transcriptomes of cells colonizing potato plants (cv. Bionta) were analyzed under nonstressed conditions (control) and at three different time points after drought stress induction by direct short-read deep sequencing (Illumina RNA-Seq). Differentially expressed genes included genes for extracytoplasmic function (ECF) group IV sigma factors. TaqMan quantitative PCR (TaqMan-qPCR) assays were performed to quantitatively assess ECF sigma factor activation in B. phytofirmans PsJN colonizing potato plants of two varieties (Russet Burbank and Bionta) in response to plant stress.

RESULTS

Detection of B. phytofirmans PsJN in plants. Six weeks after inoculation with B. phytofirmans PsJN, potato plants of two varieties (Bionta and Russet Burbank) showed increased shoot and root length in response to colonization by PsJN (see Fig. S1 in the supplemental material). Application of PEG caused visible symptoms of water deficiency in potato plants (see Fig. S2). PCR amplification with primers targeting bacterial 16S rRNA genes and plant 185 rRNA genes resulted in two bands in all inoculated plants, the mitochondrial band and a band of about 720 bp representing the bacterial 16S rRNA gene. No amplification of the bacterial fragment was found in control plants (see Fig. S3). Isolation and sequencing of the bacterial bands confirmed the presence of B. phytofirmans PsJN in all inoculated potato plants.

Transcriptome sequencing. Sequencing of cDNA samples yielded 35.7 to 42.2 million reads of cDNA, corresponding to over 2 billion nucleotides of cDNA per sample (Table 1). Around 10% of the reads were removed by initial quality filters to trim poor-quality data. After poly(A) tail removal and length trimming, more than 37% of the reads were removed, mainly because the bacterial mRNA molecules were poly(A) tailed during cDNA library preparation. Around 61% to 63% of total reads were considered for further analysis. By removal of RNA sequences, the data set was further reduced by 5%. Of the remaining sequences, 0.3% to 1.95% of the reads mapped to the genome of PsJN (Table 1). Normalization of transcript levels in control and stressed plants was done by RPKM (reads per kilobase per million mapped) normalization using NOISeq. For verification of this procedure, we used the expression data of selected housekeeping genes which were shown by qPCR to be stably expressed (see Fig. S4 in the supplemental material).

Transcriptomic profile of B. phytofirmans PsJN in potato plants. A total number of 4,591 different transcripts of B. phytofirmans PsJN were detected in PsJN-colonized potato plants. The expressed genes were evenly distributed on both chromosomes, and 76 of 167 genes carried on the plasmid were active in this experiment (Fig. 1). Among the latter, we found genes for type II B. phytofirmans 7353 (Bphyt_7353) and type IV (Bphyt_7341, Bphyt_7342, Bphyt_7347, Bphyt_7350, and Bphyt_7351) secretion system proteins.

Figure 2 shows a comparison of the genome of B. phytofirmans PsJN and the transcriptome of PsJN colonizing in vitro potato plants on the basis of the relative distributions of clusters of orthologous group (COG) categories. Overall, the functional categories of the in planta transcriptome and the genome of strain PsJN were highly similar. Few differences were found in the following COG categories: cell motility, defense mechanisms, and extracellular structures. The relative abundance of cell motility- and defense mechanism-related genes was reduced from 2% in the genome to 1% in the transcriptome. Genes encoding proteins in the COG category of “extracellular structures” covered 1% of the genes in the genome of PsJN but were not found in the transcriptome.

Expressed genes with an RPKM value of ≥5 were grouped in 74 clusters representing 354 functional classes by gene ontology (GO) analysis (see Table S1 in the supplemental material). The majority of functions were related to transcription regulation, general metabolism (sugars, amino acids, lipids, and nucleotides),

TABLE 1 Statistics of cDNA sequencing reads and their alignments on the B. phytofirmans PsJN genome

<table>
<thead>
<tr>
<th>Sample</th>
<th>Total no. of nucleotides</th>
<th>Total no. of RNA-seq reads</th>
<th>Total no. of trimmed reads</th>
<th>Total no. of rRNA reads</th>
<th>Total no. of reads</th>
<th>Total no. (%) of mapped PsJ reads</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>1,865,634,927</td>
<td>36,581,077</td>
<td>20,759,363</td>
<td>1,141,647</td>
<td>19,617,716</td>
<td>93,404 (0.48)</td>
</tr>
<tr>
<td>Stress 1</td>
<td>2,156,419,995</td>
<td>42,282,745</td>
<td>24,171,466</td>
<td>1,358,716</td>
<td>22,812,750</td>
<td>382,477 (1.70)</td>
</tr>
<tr>
<td>Stress 6</td>
<td>1,825,100,280</td>
<td>35,786,280</td>
<td>20,023,318</td>
<td>1,036,065</td>
<td>18,987,253</td>
<td>56,604 (0.30)</td>
</tr>
<tr>
<td>Stress 12</td>
<td>1,926,749,451</td>
<td>37,779,401</td>
<td>20,048,099</td>
<td>1,266,245</td>
<td>18,781,854</td>
<td>64,047 (0.34)</td>
</tr>
</tbody>
</table>

a Data indicate the total numbers of reads after quality assessment, trimming, and poly(A) removal.

b Data indicate the total numbers of reads after removal of the rRNA sequences which proceed to the mapping step.

c Data indicate the total numbers (percentages) of reads mapped on the B. phytofirmans PsJN (Ps) genome.
energy production, and cellular homeostasis. Furthermore, we found a high number of transcripts for signal transduction mechanisms such as two-component systems and extracytoplasmic (ECF) sigma factor genes (Bphyt_1327, Bphyt_1666, Bphyt_1784, Bphyt_2906, Bphyt_3189, Bphyt_4397, Bphyt_4980, Bphyt_5021, Bphyt_5131, and Bphyt_5142).

GO functions that were found in the PsJN transcriptome included functions generally considered important for endophytic plant colonization and plant growth promotion such as cell motility and chemotaxis, cellular iron homeostasis, and photosynthesis (see Table S1 in the supplemental material). Cellular iron homeostasis was represented mainly by ferritin (Bphyt_0714, Bphyt_2657, and Bphyt_5727) and bacterioferritin (Bphyt_1412 and Bphyt_2740) genes. By analyzing genes represented by the GO function "photosynthesis," we found NADH dehydrogenase subunit A, B, C, and D (Bphyt_1343 to Bphyt_1346) and polyprenyl synthetase (Bphyt_3450) genes, which do not clearly indicate putative photosynthetic activity. Furthermore, we found expression of an N-acyl homoserine lactone (AHL) synthase gene (bpI; Bphyt_4275), a quinolinate phosphoribosyl transferase gene (Bphyt_3152), indole-3-acetic acid (IAA) synthesis genes such as those encoding nitrile hydrolase (Bphyt_7182 and Bphyt_7181), and a gene encoding an IAM hydrolase of the indole-3-acetamide (IAM) pathway as well as aldehyde dehydrogenase (Bphyt_5803).

FIG 1 Circular maps representing the two chromosomes and the plasmid of Burkholderia phytofirmans PsJN. The following rings are included in each map: open reading frames (ORFs) on the plus (rings 1 and minus (rings 3) strands of the genome of strain PsJN (color by COG categories). Transcripts expressed on the plus (rings 2) and minus (rings 4) strands of the genome of strain PsJN during colonization of unstressed in vitro potato plants are indicated. The images were generated with a microbial genome context viewer (MgcV; http://mgcv.cmbi.ru.nl/) (53).
of the tryptophan side-chain oxidase pathway bypassing IPyA. Transcripts of IAA degradation (aromatic ring hydroxylating dioxygenase; Bphyt_2156) and 1-aminocyclopropane-1-carboxylate (ACC) deaminase (Bphyt_5397) genes were not detected.

Transcriptional response of \textit{B. phytofirmans} PsJN to plant stress. (i) One hour after stress induction. Analysis of differentially expressed genes in \textit{B. phytofirmans} PsJN colonizing non-stressed and drought-stressed potato plants identified 194 genes with modified expression 1 h after stress induction (see Table S2 in the supplemental material). Among these, 137 genes were upregulated and 57 genes were downregulated (Fig. 3).

For a better understanding of the genetic traits involved in the response of \textit{B. phytofirmans} PsJN to drought stress of its host plant, the differentially expressed genes were affiliated to biological processes using gene ontology (GO) analysis (see Table S3) such as those encoding various types of transcriptional regulators. The downregulated genes represented eight different GO biological processes at 1 h after stress induction. Genes that were upregulated in the transcriptome of \textit{B. phytofirmans} PsJN belonged mostly to the following functional categories: cellular homeostasis, homeostatic process, and cell redox homeostasis. Among them were genes such as those encoding bacterioferritins, glutaredoxin, redoxin domain protein, thioredoxin, and RNA polymerase factor sigma 70 (ECF sigma factor; Bphyt_1327) (see Table S3). The upregulated genes represented 52 different biological processes (see Table S3). The GO functions of these genes correspond to the functions with the highest enrichment score at all time points obtained using DAVID (Database for Annotation, Visualization, and Integrated Discovery) software. The first group consisted of two oxidoreductase activity-related genes (glutaredoxin and alkyl hydroperoxide reductase subunit) that are involved in cellular homeostasis and cell redox homeostasis. The second group represented genes involved in regulation of transcription.

(ii) Six hours after stress induction. A total number of 354 genes were differentially expressed compared to the control at this time point, with 229 genes being upregulated and 125 genes being downregulated (Fig. 3). The complete list of differentially expressed genes is available (see Table S2 in the supplemental material). Twelve hours after stress induction, 185 genes were upregulated and 81 genes were downregulated. These genes were grouped by GO analysis into 52 different biological processes (see Table S3). The GO functions showing highest enrichment at this time point were those corresponding to positive regulation of the cellular biosynthetic process, transcription, the macromolecule biosynthetic process, gene expression, the nitrogen compound metabolic process, and the macromolecule metabolic process. Peptidylprolyl isomerase FK306-binding protein (FKBP), UTP-glucose-1-phosphate uridylyltransferase, and glucose-6-phosphate dehydrogenase genes were upregulated at this time point under conditions of drought stress. Genes encoding histone family protein, transcriptional regulator GntR, transposable mutator type, peroxidasas, and catalase/peroxidas (HPI) were downregulated at this time point (see Table S3).

(iii) Twelve hours after stress induction. At this time point, potato plants treated with PEG showed severe wilting and 266 genes were differentially expressed in \textit{B. phytofirmans} PsJN (Fig. 3). The complete list of differentially expressed genes is available (see Table S2 in the supplemental material). Twelve hours after stress induction, 185 genes were upregulated and 81 genes were downregulated. These genes were grouped by GO analysis into 52 different biological processes (see Table S3). The GO functions showing highest enrichment at this time point were those corresponding to positive regulation of the cellular biosynthetic process, transcription, the macromolecule biosynthetic process, gene expression, the nitrogen compound metabolic process, and the macromolecule metabolic process. Peptidylprolyl isomerase FK306-binding protein (FKBP), UTP-glucose-1-phosphate uridylyltransferase, and glucose-6-phosphate dehydrogenase genes were upregulated at this time point under conditions of drought stress. Genes encoding histone family protein, transcriptional regulator GntR, transposable mutator type, peroxidasas, and catalase/peroxidas (HPI) were downregulated at this time point (see Table S3).

(iv) Transcriptional response of \textit{B. phytofirmans} PsJN to plant stress at all three time points. As shown by comparisons of the genes that are expressed in control plants and stressed plants, 47 genes were upregulated and 16 genes were downregulated in \textit{B. phytofirmans} PsJN in response to plant stress at all three time points (Fig. 3). These genes were subjected to further hierarchical clustered analysis and were classified into four groups by Cluster 3.0 (13). The functions of these genes correspond to the functions with the highest enrichment score at all time points obtained using DAVID (Database for Annotation, Visualization, and Integrated Discovery) software.

The first group consisted of two oxidoreductase activity-related genes (glutaredoxin and alkyl hydroperoxide reductase subunit) that are involved in cellular homeostasis and cell redox homeostasis. The second group represented genes involved in regulation of transcrip-

FIG 2 Relative distributions of functional COG categories in the \textit{in planta} transcriptome and genome of \textit{B. phytofirmans} PsJN.
tion activity and consisted of, among others, those encoding a GntR family transcriptional regulator and a cold-shock DNA-binding protein. The third group consisted of UTP-glucose-1-phosphate uridylyltransferase and glyceraldehyde-3-phosphate dehydrogenase, which showed similar (upregulated) expression patterns at all time points. The proteins encoded by those genes represent glucose metabolic processes. In the fourth group, we found an elongation factor, Tu, and ribosomal protein S7 (Fig. 3C and D).

FIG 3 (Left) Venn diagrams illustrating numbers of downregulated (A) and upregulated (B) differentially expressed (DE) genes either shared or unique at all time points after drought stress induction from pairwise comparisons of control samples at each of the drought stress time points (control at 1 h [1Hr], control at 6 h [6Hr], and control at 12 h [12Hr]). The total numbers of up- or downregulated DE genes in each time point are indicated in parentheses. (Right) Hierarchical clustering heat map of expression changes for downregulated (C) and upregulated (D) differentially expressed genes that are common across all time points (16 genes in downregulated genes and 47 genes in upregulated genes). The heat map was constructed based on RPKM expression values. Rows correspond to differentially expressed genes, and columns represent control and drought-stressed samples at the indicated time points (in hours) of drought stress. Green and red boxes represent differentially expressed genes that decreased and increased their expression levels at the indicated time points of drought stress, respectively. The identifiers (ID) and descriptions of differentially expressed genes are listed at the right in the table.
The roles of differentially expressed genes in cellular metabolic pathways were analyzed using the KEGG database. Seven pathways were found to be differentially expressed under conditions of plant stress. Genes corresponding to four pathways, namely, those corresponding to oxidative phosphorylation, sulfur metabolism, pentose and glucuronate interconversions, and aminosugar and nucleotide sugar metabolism, were upregulated. The genes corresponding to the KEGG pathway for glutathione metabolism, a two-component system, and the pentose phosphate pathway were downregulated.

Furthermore, KEGG analysis revealed that the oxidative phosphorylation pathway was the only metabolic pathway that was activated under conditions of drought stress at all three different time points (see Fig. S5 in the supplemental material). This pathway was activated at 1 h after drought stress induction with upregulation of ATP synthase subunit delta. After the organism had been maintained under conditions of drought stress for 6 h, the number of expressed genes reached 16 and included genes for cytochrome o ubiquinol oxidase subunit III, succinate dehydrogenase hydrophobic membrane anchor protein (SdhC), NADH-quinone oxidoreductase, and several ATP synthase subunits and NADH dehydrogenase subunits. This pathway was still active after 12 h, and the genes expressed included those encoding protoheme IX farnesyltransferase, succinate dehydrogenase, cytochrome b556 subunit (SdhD), NADH-quinone oxidoreductase, and several ATP synthase subunits and NADH dehydrogenase subunits.

Expression of ECF sigma factor genes in \textit{B. phytofirmans} PsJN colonizing potato plants. Expression of extracytoplasmatic function (ECF) sigma factor genes in \textit{B. phytofirmans} PsJN colonizing potato plants (cv. Bionta and Russet Burbank) with and without drought stress was tested by real-time qPCR with cDNA of four biological replicates per treatment. Transcripts of six ECF sigma factor genes (ECF_164, ECF_886, ECF_429, ECF_718, and ECF_474) were detected in PsJN-colonized plants (see Fig. S6 in the supplemental material). The number of expressed ECF sigma factor genes in stressed plants was different from the number in control plants, and the numbers varied over time in stressed plants. ECF_164, ECF_429, and ECF_886 were expressed in control plants of both cultivars and were also active in stressed plants at all time points. The number of expressed ECF sigma factor genes increased under conditions of drought stress to a maximum of five genes after 1 h in Bionta and after 6 h in Russet Burbank (see Fig. S6). Transcripts of ECF_718 were detected in both cultivars, whereas ECF_474 was found in Russet Burbank and ECF_230 in Bionta only.

**DISCUSSION**

The genome of \textit{B. phytofirmans} PsJN encodes 7,405 genes, 4,591 (62%) of which were expressed in PsJN colonizing cv. Bionta potato plants \textit{in vitro}. The active genes were evenly distributed across both chromosomes and the plasmid. Interestingly, about 46% of...
the coding sequences (CDS) located on the plasmid were expressed by PsJN inside potato plants. The genome of _B. phytofirmans_ PsJN shows a high degree of similarity to that of _B. xenovorans_ LB400, but the pBPHYT01 plasmid is different from the megaplasmid of _B. xenovorans_. In _B. xenovorans_ LB400, the small chromosome is the “lifestyle-determining” replicon, whereas strain-specific functions are encoded on the megaplasmid (14). In _B. phytofirmans_ PsJN, only 29% of the CDS on pBPHYT01 could be functionally described (15); consequently, the majority of plasmid-carried genes that are active in PsJN colonizing cv. Bionta potato plants are of unknown function. Several reports of studies have proposed that plasmids are genetic determinants of functional diversification and niche adaptation (16, 17). We can only speculate on the role of pBPHYT01 in the endophytic lifestyle of _B. phytofirmans_ PsJN. Experiments designed to test plasmid-free PsJN for the ability to colonize plants and to establish endophytic population could give further insights.

The transcriptome profile of _B. phytofirmans_ PsJN colonizing potato plants indicates that the bacterium is metabolically active in plants. The majority of expressed traits were related to transcription regulation, general metabolism (sugars, amino acids, lipids, and nucleotides), energy production, and cellular homeostasis. The overall patterns of functions encoded on the genome of _B. phytofirmans_ PsJN and expressed in _in vitro_ potato plants were highly similar. We conclude from this that the plant interior as a habitat for bacteria does not require very selected and specialized functionalities. Only a few differences in the COG patterns were found, with cell motility and defense mechanisms being less represented in the transcriptome than in the genome of _B. phytofirmans_ PsJN. Cell motility might be important in plant invasion and in the spreading of endophytic microorganisms throughout plant organs and tissues (18). Recently, Balsanelli and colleagues showed that motility-related functions such as chemotaxis and type VI pilus functions play an important role in the initial contact with plants and the epiphytic colonization of maize roots by _Herbaspirillum seropedicae_ (19). Our data indicate that active movement is less important once a bacterial population is successfully established inside plants. Interestingly, defense-related traits seem to play a minor role in the endophytic life of _B. phytofirmans_ PsJN also. A possible explanation is that the plant endosphere is a protected niche allowing endophytes to escape the high competitive pressure in the rhizosphere and soil.

_B. phytofirmans_ PsJN stimulates plant growth in many of its host plants. Metabolic properties suggested to be involved in this activity include the production and degradation of auxin phytohormone indole-3-acetic acid (IAA) (20), ACC deaminase activity (21), quinolinate phosphoribosyl transferase (QPRTase) or nicotinate-nucleotide pyrophosphorylase activity (22), and sidrophore production (1). In our experiment, _in vitro_ potato plants colonized by _B. phytofirmans_ PsJN showed increased growth compared to an untreated control. By showing the expression of quinolinate phosphoribosyl transferase and indole-3-acetic acid (IAA) synthesis genes of strain PsJN colonizing potato plantlets, the results of the present study support previous reports of the importance of these functions for the beneficial interaction between PsJN and plants.

Plants colonized by _B. phytofirmans_ PsJN often show increased tolerance of abiotic stress such as chilling (23) and drought (4). One of the main ambitions of this study was to elucidate whether and how endophytic _B. phytofirmans_ PsJN is affected by host plant drought stress. Analysis of differentially expressed genes in _B. phytofirmans_ PsJN colonizing nonstressed and drought-stressed potato plants showed that 137, 229, and 185 genes were upregulated and 57, 125, and 81 genes were downregulated in response to host plant drought stress at 1, 6, and 12 h after PEG application. This indicates that _B. phytofirmans_ PsJN adjusts gene expression to physiological conditions that have been altered in host plants due to plant stress responses. Genes that were significantly upregulated in _B. phytofirmans_ PsJN in response to host plant stress are mostly involved in transcription regulation, cellular homeostasis, and cell redox homeostasis, indicating a stress response in _B. phytofirmans_ PsJN. Drought stress provokes an oxidative burst in plants as a primary immune response. This increase in the production of reactive oxygen species (ROS) serves on the one hand as an alarm signal that triggers acclimation and defense reactions and is kept in tight balance by the antioxidant system in plants (24). If, on the other hand, the drought stress continues for a certain period of time, the oxidative burst may lead to extensive cellular damage and finally to cell death (24). Upregulation of genes involved in detoxification of ROS in strain PsJN colonizing potato plants suffering from drought stress led us to the assumption that endophytic _B. phytofirmans_ PsJN senses and is affected by plant-produced ROS. We propose the following scenario. Water limitation leads to ROS burst in plants. Endophytic PsJN cells respond with the expression of genes involved in the defense against oxidative stress in order to prevent cell damage. ROS scavenging by endophytes is also very important during the early stage of plant colonization, as previously shown for _Glucanacetobacter diazotrophicus_ PAL5 (11). Whether bacterial antioxidant activity may help to maintain the redox balance in plants and thus reduce the effects of drought stress on plants remains elusive and merits further investigation.

Oxidative phosphorylation was found to be activated in _B. phytofirmans_ PsJN over time during plant drought stress. In the process of cellular respiration, aerobic bacteria pass electrons from oxidizable substances to molecular oxygen via the so-called electron transport chain. The released energy is used to produce energy-rich ATP from ADP by phosphorylation. Oxidative phosphorylation generates the energy needed for almost all vital processes (25). Apart from this, pentose and glucuronate interconversions and amino sugar and nucleotide sugar metabolism were also activated. Upregulation of genes involved in these processes in _B. phytofirmans_ PsJN colonizing drought-stressed potato plants could indicate activation of bacterial growth. At least for _Epichloë_ endophytes, it is well documented that the mutualistic interaction of fungi and host plant is tightly regulated. Perturbations of this balance result in a switch from restricted to proliferative growth of the endophyte inside the plant and a breakdown from mutualistic to pathogenic behavior (26). Furthermore, anarchic proliferation of otherwise asymptomatic bacterial endophytes is a common phenomenon in _in vitro_ plant propagation when cultures are under stress (27). We therefore quantified PsJN in the control and stressed potato plants by performing qPCR with the selected housekeeping genes used for data normalization but did not find a significant increase in copy numbers over time under conditions of drought stress (data not shown). It is possible that the time span (12 h) was too short to observe a significant increase in cell numbers, but it is also likely that the increase in metabolic activity in _B. phytofirmans_ PsJN under conditions of host plant drought stress was not coupled with proliferated growth.
One way that bacteria sense and react to the extracellular environment is by the so-called cell surface signaling-employed extracytoplasmatic function (ECF) sigma factors (28). This signal transduction system consists of an outer membrane receptor, an inner membrane-bound sigma factor regulator (anti-sigma factor), and, bound to that, an ECF sigma factor. In the absence of a signal, the anti-sigma factor tightly binds the ECF sigma factor, thereby keeping it in its inactive state. The anti-sigma factor is proteolytically degraded in the presence of a stimulus. As a result, the sigma factor is released and activates expression of its target genes (28). The ECF subfamily is the largest group in the sigma 70 family, and its members are involved in a wide range of environmental responses, such as metal homeostasis, starvation, and resistance to antimicrobial peptides, and are also required for pathogenesis in some cases (28). ECF sigma factors may also play a role in the establishment of plant-microbe interactions. Gourion and colleagues (29) showed that an extracytoplasmatic sigma factor is involved in symbiotic efficiency in the plant symbiont Bradyrhizobium japonicum USDA110. The genome of B. phytofirmans PsJN harbors eighteen different CDS putatively coding for extracytoplasmatic function (ECF) sigma factors. Analysis of the in planta transcriptome of B. phytofirmans PsJN revealed the expression of eleven extracytoplasmatic sigma factor genes, and the expression of one of these genes was upregulated upon plant stress induction. TaqMan-quantitative PCR experiments were performed to quantitatively assess ECF sigma factor activation in B. phytofirmans PsJN in response to plant stress. Six ECF sigma factor genes were expressed in B. phytofirmans PsJN colonizing drought-stressed potato plants. One of these genes (Bphyt_1327; EcfD) was found to be involved in the response to chlorhexidine (31). The biological role of Bphyt_1327 in B. phytofirmans PsJN remains unclear and requires further investigation.

Another ECF sigma factor (Bphyt_4063; ECF_164) was down-regulated in B. phytofirmans PsJN. It is orthologous to EcfI in Burkholderia cenocepacia and can be found in 31 other Burkholderia genomes in similar genetic neighborhoods. Little is known of the function of this ECF sigma factor in Burkholderiaeae; only the ortholog in Burkholderia cenocepacia, Ecfl, was found to be involved in the response to chlorhexidine (31). The biological role of Bphyt_1327 in B. phytofirmans PsJN remains unclear and requires further investigation.

In this study, the plant growth-promoting rhizosphere bacterium and endophyte Burkholderia phytofirmans PsJN (= LMG 22146 T = CCUG 49060 T) was used (1). The bacterial strain was grown by loop inoculation of a single colony in LB broth. The bacterial culture was incubated at 28 ± 2°C for 2 days at 180 rpm in a shaking incubator.

**Materials and Methods**

**Bacterial strains and growth conditions.** In this study, the plant growth-promoting rhizosphere bacterium and endophyte Burkholderia phytofirmans PsJN (L = LMG 22146 T = CCUG 49060 T) was used (1). The bacterial strain was grown by loop inoculation of a single colony in LB broth. The bacterial culture was incubated at 28 ± 2°C for 2 days at 180 rpm in a shaking incubator.

**Plant experiment.** Two potato varieties (Solanum tuberosum cv. Russet Burbank and Bionta) were grown in vitro in a growth chamber at 20°C with a 16-h-light/8-h-dark-photoperiod cycle. Four-week-old elongated apical shoots (10 cm in length) were used directly as explants. They were inoculated with B. phytofirmans PsJN by dipping for some seconds in a bacterial culture (1.3 × 10^7 CFU/ml). Inoculated plants were grown in 10 ml solid Murashige-Skoog (MS) medium containing 8% Duxhefa Daishin agar (pH 5.8) and 20% saccharose for 4 weeks in glass tubes (2.5-cm diameter). Rooted potato plantlets were transferred into custom-tailored glass tubes with a narrow neck allowing hydroponic plant culturing. The plantlets were grown in 15 ml liquid MS medium containing 20% saccharose for two more weeks. Drought stress was induced by adding polyethylene glycol (PEG) (molecular weight [MW], 6,000) to reach a final concentration of 45%. Control plants were not treated. Shoots of stressed and control plantlets of each variety (6 replicates per treatment) were harvested at 1, 6, and 12 h after PEG application, immediately frozen in liquid nitrogen, and stored at −80°C for further analysis. To evaluate the presence of B. phytofirmans PsJN in inoculated potato plantlets, 16S rRNA gene PCR was performed using universal primers 799F [5'-AAC (AC)GGATTGATAACC(GT)-3'] (37) and 1520R [5'-AAGGAGGTGA TCCAGCGCA-3'] (38). Amplification with this primer pair allows ex-
clclusion of chloroplast 16S rRNA gene-based amplicons but results in co- amplification of plant mitochondrial small-subunit RNA gene fragments and the bacterial 16S rRNA gene (see Fig. S1 in the supplemental mate- rial). Each PCR (50 µl) contained 30 to 50 ng/µl bacterial or potato DNA as the template, 1× PCR buffer, 2.5 mM MgCl₂, 200 µM deoxy nucleoside triphosphate (dNTP) mix (Thermo Scientific), a 150 nM concentration of each forward and reverse primers, 2.5 U of Firepol DNA polymerase (Solis Biodyne, Estonia), and PCR-grade water. Cycling conditions were as fol- lows: initial denaturation for 5 min at 95°C; 34 cycles of 30 s at 95°C, 1 min at 60°C, 2 min at 72°C; and final elongation for 4 min at 72°C. Amplified PCR products (5 µl) were separated by electrophoresis (80 V) on 1% (wt/vol) agarose gels. Agarose gels were stained with ethidium bromide. Bac- terial PCR amplicons were sequenced, making use of the sequencing service at LGC Genomics (Germany). Retrieved sequences were visualized and aligned with ClustalW as implemented in BioEdit v7.1.3 (39). For identification, sequences were subjected to BLAST analysis with the NCBI database.

Total RNA isolation and cDNA synthesis from plant tissue and bac- terial cells. Frozen plant material (100 mg) was prechilled with liquid nitrogen in 2-ml Safe-Lock tubes (Greiner Bio-One, Germany) and ho- mogenized by the use of a ball mill MM301 mixer (Retsch GmbH & Co., Germany) at 30 Hz for 2 min using a single steel ball (5-mm diameter). Afterward, the material was immediately subjected to RNA isolation as described by Chang et al. (40). Extraction of total RNA from pure bacterial cultures was done using TRIZol reagent (Invitrogen, USA) according to the manufacturer’s instructions, and the RNA was used as a control in qPCR experiments. RNA samples were treated with DNase (Turbo DNA- free kit; Ambion, USA) according to the manufacturer’s protocol for pu- rification from DNA contamination. RNA samples were tested for co- containing DNA by 16S rRNA gene PCR. RNA was analyzed at a 260-nm/ 280-nm ratio using a NanoDrop 1000 spectrophotometer (Thermo Scientific, USA), and the integrity was checked by electrophoresis in a 1% agarose gel. Purified RNA samples were reverse transcribed to cDNA with an iScript cDNA synthesis kit (BioRad Inc., USA) using random hexamers according to the manufacturer’s instructions.

Transcriptome sequencing. In order to get a comprehensive image of the total transcriptome of B. phytofirmans PsJN colonizing potato plants (cv. Bionta), the total RNAs of four biological replicates for each treat- ment and control samples were pooled at equal concentrations to obtain approximately 48 µg RNA per treatment. rRNA depletion, cDNA library preparation, and sequencing on an Illumina HiSeq 2000 system with a 50-bp single-end read length were outsourced to Vertis Biotechnologie (Germany). PCR amplification was per- formed in a T-gradient PCR thermocycler (Biometra, Germany). PCR-amplicon sequencing as well as sequence analysis and identifi- cation were done as described above. Three primer sets (Bphyt_5131, Bphyt_7017, and Bphyt_1666) were not specific for their target ECF sigma cation were done as described above. The analysis of PCR amplification, sequencing, and sequence analysis and identifi- cation was performed by making use of the ARB software package with its subfunc- tion “Probe design,” version ARBuntu 2.0 (49). Probes were labeled with the reporter dye 6-carboxyfluorescein (FAM) at the 5’ end and with black hole quencher 1 (BHQ-1) fluorophore at the 3’ end.

Preprocessing of sequencing data and mapping of reads. RNA-seq reads were subjected to quality filtering using Prinseq (41) on the basis of the criterion of a minimum read length of 40, and good-quality reads were ob- tained with Q20 (sequencing error rate lower than 1%) for all reads. The poly(A) tails with a minimum length of 10 bp were removed by the use of Prinseq. Reads longer than 30 bp were considered for further analysis. Before read alignment, rRNA fragments were filtered from the transcriptomics data by the use of SortMeRNA software and the default rRNA database included in the software package (42). Reads were aligned to the genome of B. phytofir- mans PsJN, and gene annotations were obtained from the NCBI database of bacterial genomes (ftp://ftp.ncbi.nlm.nih.gov/genomes/Bacteria/Burkhold- eria_phytofirmans_PsJN_uid58729/).

Transcriptome analysis. RNA-seq reads were aligned to the genome of B. phytofirmans PsJN using the Burrows-Wheeler Alignment Tool (BWA v0.6.2) (43). Transcript abundance was calculated using in-house Python scripts. Differential gene expression levels were analyzed and vi- sualized with NOIseq (44). Gene expression levels were normalized using the number of reads per kilobase of coding sequence per million mapped reads (RPKM) in B. phytofirmans PsJN. As there was no replicate available for our data set, NOIseq-sim was used with the highest threshold (q = 0.9) to compute the probability of differential expression of genes under differ- ent conditions. To determine the variations of the differentially ex- pressed genes which were expressed across all the time points from con- trol compared to drought-stressed samples in B. phytofirmans PsJN, hierarchical cluster analysis was performed using Cluster 3.0 (13). The clustering results were visualized using TreeView (http://jtreeview. sourceforge.net/). Functional annotation of target genes based on Gene Ontology terms was performed using David v6.7 software (45) and all PsJN genome data available at NCBI Refseq database (46) as the back- ground. Venn diagrams were drawn using BioVenn software (47). Func- tional COG (clusters of orthologous groups) categories of the transcripts expressed in the control were listed using the COG database (48) and compared to the COG categories of the B. phytofirmans PsJN genome obtained from Integrated Microbial Genomes (IMG) systems (http://img.jgi.doe.gov).

Design of primers and probes for amplification of ECF sigma factor genes. Oligonucleotides and probes for amplification of extracytoplasmatic function (ECF) sigma factor genes (see Table S4 in the supplemental mate- rial) were designed on the basis of the genome sequence of B. phyto- firmans PsJN (GenBank project accession no. CP001052, CP001053, and CP001054) by making use of the ARB software package with its subfunc- tion “Probe design,” version ARBuntu 2.0 (49). Probes were labeled with the reporter dye 6-carboxyfluorescein (FAM) at the 5’ end and with black hole quencher 1 (BHQ-1) fluorophore at the 3’ end.

Test for specificity of primers and probes for ECF sigma factor genes. Primer specificity was checked by PCR amplification using genomic DNA of B. phytofirmans PsJN. DNA was extracted from bacterial cell pellets using a FastDNA Spin kit for soil (MP Biomedicals, LLC). DNA concentrations were measured with a NanoDrop 1000 spectrophotometer (Thermo Scientific, Wilmington, DE). PCR amplification was per- formed in a T-gradient PCR thermocycler (Biotera, Germany). PCR and PCR-amplicon sequencing as well as sequence analysis and identifi- cation were done as described above. Three primer sets (Bphyt_5131, Bphyt_7017, and Bphyt_1666) were not specific for their target ECF sigma factor genes and were removed from further analysis.

Selection of housekeeping genes for normalization of expression data. Primers and probes were designed for 12 (see Table S4 in the supplemen- tary material) gene candidates using the ARB software package (49), and probes were labeled with the 6-FAM reporter dye at the 5’ end and with black hole quencher 1 (BHQ-1) at the 3’ end. The primers were designed to be specific for B. phytofirmans PsJN and to avoid amplification of host plant material. The specificity of primers was tested by PCR, using genomic DNA of B. phytofirmans PsJN and potato plants as the template in separate reactions. The analysis of PCR amplification, sequencing, and sequences was performed as described above. Expression of all house- keeping genes was checked by TaqMan-qPCR using cDNA synthesized from total RNA isolated from stressed and control plants. The expression stability of selected housekeeping genes under conditions of drought stress and the variation in quantitative PCR efficiency in B. phytofirmans PsJN were calculated using qBasePLUS software with the geNormPLUS algorithm implemented (50, 51). Glutamine synthetase (Bphyt_2615) was the candidate housekeeping gene product that showed the most stable expression at different time points of drought stress and in both potato varieties (see Fig. S4).
TaqMan real-time PCR assays. Quantitative PCR was carried out using the TaqMan-qPCR assay and a Bio-Rad CFX-96 real-time detection system (Bio-Rad, Hercules, CA). Triplicate qPCR reactions were performed with 1 μl of cDNA as the template, 1× Bio-Rad SsoFast probe mix (BioRad Inc., Hercules, CA), 10 μM of forward and reverse primers, and 5 μM probe in a final volume of 10 μl. Cycling conditions were as follows: a hot start at 95°C for 2 min, 69 cycles of denaturation at 95°C for 5 s, and 20 s of elongation at 60°C. In each run, 2 negative controls were used, one as a no-template control performed with PCR-grade water instead of cDNA and another that included cDNA from non-PsJN-inoculated control plant samples. Four biological replicates of both potato varieties were tested. Data were analyzed with Bio-Rad CFX Manager software (version 3.0). Based on the Pfaffl equation (52) implemented in this software, normalized relative quantity (NRQ) values of ECF sigma factor genes in comparison to the most stably expressed reference gene (Bphyt_2615) were determined.

SUPPLEMENTAL MATERIAL


REFERENCES


