Effects of genetically modified potatoes with increased zeaxanthin content on the abundance and diversity of rhizobacteria with in vitro antagonistic activity do not exceed natural variability among cultivars

Nicole Weinert · Remo Meincke ·
Christine Gottwald · Viviane Radl · Xia Dong ·
Michael Schloter · Gabriele Berg · Kornelia Smalla

Abstract To assess potential effects of genetically modified (GM) potatoes on the abundance and diversity of rhizobacteria with in vitro antagonistic activity in relation to natural variability among cultivars, two GM potato lines accumulating the carotenoid zeaxanthin in their tubers, the parental cultivar and four additional commercial cultivars were planted at two field sites in Germany. Rhizosphere samples were taken at three developmental stages of the plants. A total of 3,985 bacteria isolated from the rhizosphere were screened for their in vitro antagonistic activity towards *Rhizoctonia solani*, *Verticillium dahliae* and *Phytophthora infestans* using a dual-culture assay. Genotypic characterisation, 16S rRNA gene sequencing and antifungal metabolite analysis was performed to characterize the 595 antagonists obtained. The 16S rRNA gene-based identification of in vitro antagonists revealed strong site-dependent differences in their taxonomic composition. This study showed that the site was the overriding factor determining the proportion and diversity of antagonists from the rhizosphere of potato while the effect of the genetic modification on the proportion of antagonists obtained did not exceed natural variability among the five commercial cultivars tested.

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N. Weinert · C. Gottwald · K. Smalla (✉)
Julius Kühn-Institute - Federal Research Centre for
Cultivated Plants (JKI),
Institute for Epidemiology and Pathogen Diagnostics,
Messeweg 11–12, 38104 Braunschweig, Germany
e-mail: kornelia.smalla@jki.bund.de

R. Meincke · G. Berg
Environmental Biotechnology,
Graz University of Technology,
Petersgasse 12,
A-8010 Graz, Austria

V. Radl · M. Schloter
Department for Terrestrial Ecogenetics,
Helmholtz Zentrum München,
Ingolstädter Landstr. 1,
85758 Oberschleissheim, Germany

X. Dong
Technical University of Munich,
Chair for Plant Breeding,
Am Hochanger 4,
85350 Freising, Germany

Introduction

The rhizosphere, which is defined as part of the soil influenced by the plant, has a central role as interface

between the soil and the plant (Hiltner 1904; Brimecombe et al. 2001). In the rhizosphere the abundance and metabolic activity of some soil microorganisms is enhanced in response to root exudates and deposits. In turn rhizosphere microorganisms contribute to plant nutrition, health and growth and can induce systemic resistance (Van Loon et al. 1998; Whipps 2001). Cultivation-independent studies (16S rRNA gene fingerprints) have shown that the rhizosphere often harbors a reduced number of dominant bacterial populations whose relative abundance clearly differs from bulk soil (Costa et al. 2006; Marilley et al. 1998; Smalla et al. 2001). Differences in exudate level and composition, together with rhizosphere microsite location, root-mass and structure, but also the nutritional status of the plant are assumed to contribute to individual microbial communities of distinct plant species (Gyamfi et al. 2002; Marschner et al. 2001; Yang and Crowley 2000). Some studies reported on differences in the rhizosphere as well as endophyte bacterial community composition among cultivars of the same plant species (Adams and Kloepper 2002; Dunfield and Germida 2001; Siciliano and Germida 1999). But not only the plant species itself is of vital importance, also the plant developmental stage, soil properties, climatic conditions and agricultural management practice are assumed to have a major influence on the composition of rhizosphere microbial communities (Bossio et al. 1998; Crecchio et al. 2004; Kennedy et al. 2005; Rasche et al. 2006).

Antagonistic microorganisms that reside in the rhizosphere are able to defend the plants against pathogens. Several traits are assumed to play an important role for antagonistic activity: (i) antibiosis, implying the inhibition of a pathogen through the release of antibiotics, toxins or biosurfactants (Fravel 1988), (ii) competitive root colonization (Lugtenberg et al. 2001), and (iii) production of extra-cellular cell wall degrading enzymes like ß-1,3 glucanase, chitinase or cellulase (Chernin and Chet 2002).

Although the application of GM plants in agriculture and forestry is of increasing importance (James 2007) the overall impacts of GM crops on soil quality are, however, still poorly understood. Despite their importance for soil and plant health, the response of soil microbes to the application of GM crops has not been addressed adequately. Unintended side-effects due to altered characteristics of the modified plants

cannot be *a priori* excluded and thus their impact has to be evaluated case-by-case. Genetic modifications of plants often result in phenotypic changes and may also result in alteration of the root exudation which in turn could cause changes in the structural and functional diversity of plant-associated microorganisms. Potential effects of GM crops on beneficial rhizobacteria might be of particular interest as their relative abundance and diversity could affect plant health and ecosystem sustainability. Only few studies attempted to assess potential impacts of GM plants on antagonistic species (Lottmann et al. 1999; Lottmann et al. 2000; Lottmann and Berg 2001). However, these studies compared only the two GM potato lines to a GM control and the parental cultivar.

In the present study two GM potato lines which accumulate the carotenoid zeaxanthin in their tubers (Römer et al. 2002) were investigated under field conditions. Although the two constructs used in this study do not harbor new genes and the modifications are based on antisense respectively co-suppression of an existing pathway (zeaxanthin epoxidase) and should only be active in the potato tuber, it could not be excluded that root exudation might be changed as a result of the genetic modification. Indeed the root length, carbohydrate pattern and pH in the rhizosphere of both GM lines were changed compared to the wild type. Interestingly, also differences between both GM lines were observed (Neumann et al. personal communication). These data clearly indicate that a modification of a single gene can influence the plant phenotype in many ways. The two GM potato lines with increased zeaxanthin level in their tubers were grown together with their parental cultivar and four additional commercial cultivars in a randomised field trial with six replicated plots per cultivar and GM line at two sites in Germany. Rhizosphere samples were taken at three developmental stages. In this study we aimed to assess potential effects of GM potatoes on the abundance and composition of bacteria with in vitro antagonistic activity towards three potato pathogens (*Rhizoctonia solani*, *Verticillium dahliae*, *Phytophthora infestans*) and relate these to natural variability among the potato cultivars investigated. Furthermore, this study provides insights into the phenotypic and genotypic diversity of bacterial isolates with in vitro antagonistic activity.

Materials and methods

Potato cultivars and GM lines

For genetic modifications *Solanum tuberosum* L. cultivar 'Baltica' was used. The construction of the two GM zeaxanthin-accumulating potato lines through co-suppression (SR47/00#18, further referred to as SR47) and antisense (SR48/00#17, further referred to as SR48) was described in detail by Römer et al. (2002). The promoter used in both GM lines were tuber specific and thus no increased zeaxanthin levels in the above ground parts of the plants were expected. This could be confirmed by studying the expression of the zeaxanthin epoxidase in different plant compartments (Wenzel et al. personal communication). The accumulation of zeaxanthin in the tuber reached up to 40 µg/g dry weight (dw) (SR47) and 17 µg/g dw (SR48) compared to 0.2 µg/g dw of the wildtype. In addition to the GM lines and the parental cultivar 'Baltica', the potato cultivars 'Selma', 'Désirée', 'Ditta' (all used for food production) and 'Sibu' (used for industrial purposes) were grown in the field.

Field design

The two sites used for this study are located in southern Germany. The Roggenstein soil was characterized by 26.1% sand, 44.0% silt, 28.1% clay, C_{org}: 1.1%, N_t: 0.1% and pH 6.6. The Oberviehhausen soil contained 54.6% sand, 31.3% silt, 14.1% clay, C_{org}: 1.9%, N_t: 0.2%, and had a pH of 6.5. The climatic conditions in 2005 (Roggenstein) and 2006 (Oberviehhausen) are available from the weather station located nearby both field sites. Plant protection measures were performed according to agricultural practice.

The experimental setup was designed as a randomized field trial: Each plot had a size of 9×3 m, resulting in four rows with a total of 40 potatoes. The cultivars and GM lines were grown in six replicated plots, of which four plots were sampled. The data presented in this study are based on the field trials in 2005 (Roggenstein) and 2006 (Oberviehhausen).

Isolation of root-associated bacteria

Sampling of the plants was carried out at three developmental stages, young plants (EC30), flowering plants (EC60) and senescent plants (EC90) according to Hack et al. (1993). The GM lines showed a delayed emergence but at EC60 no obvious differences between the GM lines and commercial cultivars were observed. Therefore, we decided to sample not completely randomized but to select plants in a comparable growth stage at EC30. From each of the plots the roots of five plants were combined as a composite sample and transported to the laboratory. To extract the root-associated bacteria, 10 g of the root material of each plot was transferred into sterile stomacher bags. Thirty ml Milli-Q water was added and samples were homogenized for 60 s in a Stomacher laboratory blender (Seward, West Sussex, UK) at high speed. This step was repeated three times. The resulting suspensions were serially diluted with 0.85% NaCl and plated onto R2A (Merck, Darmstadt, Germany) supplemented with cycloheximide (100 µg/ ml) to avoid the growth of fungi. After 5 days of incubation at 28°C colony forming units (CFU) were determined and 100 visually different colonies were picked from each cultivar and GM line, resulting in a total of 700 isolates per sampling time and site. The isolates were purified and stored at −70°C.

Screening for bacteria with in vitro antagonistic potential

The bacterial isolates obtained were screened in a dual-culture in vitro assay for their antagonistic potential towards three major potato pathogens: *Rhizoctonia solani* Kühn AG3 (*Basidiomycete* with a chitin-glucan cell wall), *Verticillium dahliae* Kleb. ELV25 (*Ascomycete* with a chitin-glucan cell wall) and *Phytophthora infestans* [Mont.] De Bary strain 20/01 (*Oomycete* with a cellulose-glucan cell wall). Activity of the isolates against *R. solani* and *V. dahliae* was determined on Waksman agar (WA), containing 5 g of bactopeptone (Merck, Darmstadt, Germany), 10 g glucose (Merck), 3 g meat extract (Chemex, München, Germany), 5 g NaCl (Merck) and 20 g bacto-agar (Becton Dickinson, NJ, USA) per l distilled water. Agar disks with mycelia were directly cut out from WA plates with *R. solani* (grown for 7 days) and placed between the streaks of four isolates. *V. dahliae* was grown in liquid culture in Czapek Dox medium (Difco, Detroit, MI, USA). 150 µl of the suspension containing hyphal fragments

were plated onto the agar and after surface drying the bacteria were streaked on the same plate. Inhibition zones were measured after 5 days of incubation at room temperature as described by Berg et al. (2002). The dual-culture assay against *P. infestans* was carried out on pea-agar (modified according to Hollomon 1965). For this purpose, 125 g frozen peas were cooked in 500 ml distilled water for 20 min. The broth obtained after filtration was adjusted with distilled water to 1 l after adding 15 g bacto-agar. Agar disks from pea-agar plates with *P. infestans* grown for 10–12 days were cut out and pre-grown on the pea-agar test plates for 1–2 days before streaking the bacterial isolates. Inhibition zones were determined after 7 days at room temperature. Every bacterial strain was tested twice independently.

Extraction of genomic DNA

Genomic DNA from the in vitro antagonistic strains was extracted using the Qiagen Genomic DNA Extraction Kit (Hilden, Germany) for cell lysis and the MoBio UltraClean™ 15 DNA Purification Kit (Carlsbad, CA, USA) for DNA-preparation, as described by the manufacturers.

Genotypic characterization using ARDRA and BOX-PCR genomic fingerprinting

Amplified Ribosomal DNA Restriction Analysis (ARDRA) was performed by amplification of the 16S rRNA gene with primers U8-U27 (5'-AGAGTTT GATC(A/C)TGGCTCAG-3') and 1492–1514 (5'-TACGG(C/T)TACCTTGTTACGACTT-3') (Weissburg et al. 1991). Amplicons obtained were digested with the enzymes *Hin*6I and *Bsh*1236I (Fermentas) for 3 h at 37°C and 300 rpm in a thermomixer (Eppendorf, Thermomixer comfort). Fifteen µl of the resulting fragments were separated in a 4% agarose gel (NuSieve 3:1) in 0.5 × TBE buffer for 4 h, stained with ethidium bromide and visualized using a UV transilluminator. Antagonists displaying similar ARDRA-patterns (cut-off level 85%) were further analyzed using BOX-PCR genomic fingerprinting. BOX-PCR fingerprints were performed using the BOX_A1R primer as described by Rademaker and De Bruijn (1997). Eight µl of the PCR-product were separated in a 1.5% agarose gel in 0.5 × TBE buffer

for 4 h, stained with ethidium bromide and photographed under UV.

Identification of bacterial in vitro antagonists

Antagonists with either individual ARDRA patterns or different BOX patterns (cut-off level 85%) were identified by partial sequencing (approx. 700 bp) of the 16S rRNA gene. The sequences obtained were aligned with reference 16S rRNA gene sequences using the nucleotide basic local alignment and search tool (BLASTN) from NCBI databases.

Enzymatic activity, production of siderophores and N-acyl-homoserine-lactone (AHL)-production

Protease activity (casein degradation) was determined from clearing zones in skim milk agar (400 ml sterilised skim milk mixed with 1/10 TSB and 16 g bacto-agar at 55°C) after 5 days of incubation at 28°C. β-glucanase and cellulase activity were tested using chromogenic AZCL and reamazolbrilliant blue R (AZO) substrates, respectively (Megazyme International, Ireland). Formation of blue and white haloes was determined after 5 days at 28°C. Chitinolytic activity was tested in chitin minimal medium as described by Berg et al. (2001). Clearance haloes indicating β-1,4-glucosamine polymer degradation were determined after 7 days of incubation at room temperature. Production of siderophores under Fe^{3+}-limited conditions was analyzed using the plate assay developed by Schwyn and Neilands (1987). Orange haloes, indicating a reduction of iron(III) to iron(II) were measured after 3 days of incubation at room temperature. AHL-production was investigated in a cross-streak assay using the bioluminescent sensor plasmid pSB403 in *E. coli* to detect 3-oxo-C6-HSL molecules produced by the bacterial antagonists (Winson et al. 1998). The sensor strain *Chromobacterium violaceum* CV026 (McClean et al. 1997) changes its color to purple in the presence of short chain AHLs. *Serratia plymuthica* HROC48 (Liu et al. 2007) served as a positive control in both assays.

Statistical analysis

Significance tests for CFU values were performed using ANOVA and Tukey Test. Significance tests for

numbers of in vitro antagonists (Chi square exact) were performed with Statistical Product and Service Solutions for Windows v. 15.0. (SPSS Inc., Chicago, IL, USA). Computer-assisted evaluation of both ARDRA and BOX-PCR genomic fingerprints was performed using the GelCompar II program version 4.5. (Applied maths, Kortrijk, Belgium). Cluster analysis was performed using the Dice correlation matrix and the unweighted pair group method using average linkages.

Nucleotide sequence accession numbers

The nucleotide sequences determined in this study were deposited in the GenBank database under accession numbers EU834335-EU834701.

Results

Colony forming units of bacterial isolates from the potato rhizosphere

Significant differences in the CFU counts (p≤0.05) were revealed at EC60 in Roggenstein and at EC30 and EC60 in Oberviehhausen. These differences were detected for the GM lines as well as for some of the commercial cultivars (Table 1).

In vitro antagonistic activity of bacterial isolates towards *R. solani* AG3, *V. dahliae* ELV25 and *P. infestans*

A total of 1,946 isolates originating from the rhizosphere of potatoes grown at the site in Roggenstein and 2,039 isolates from the potato rhizosphere in Oberviehhausen were tested in dual culture assays for their in vitro antagonistic activity. The total numbers of antagonists obtained from the Roggenstein site revealed significant differences both between GM lines and commercial cultivars, as well as between different commercial cultivars for all three pathogens tested (Table 2). The majority of in vitro antagonists towards all three pathogens was obtained from the cultivar 'Désirée'. For the Oberviehhausen site significant differences in the total numbers of antagonists were observed for *R. solani* and *P. infestans*, whereas the numbers of antagonists towards *V. dahliae* revealed no significant differences between the cultivars and GM lines. Significant differences could be observed both between GM lines and commercial cultivars and between different commercial cultivars (Table 2).

At both sites significantly more isolates showed in vitro antagonistic activity towards *P. infestans* than towards *R. solani* and *V. dahliae*. Moreover, the

Table 1 Mean values (Log$_{10}$) of colony forming units (CFU) per gram of root fresh weight of the potato rhizospheres for the field sites in Roggenstein (A) and Oberviehhausen (B) for the three sampling times. Values followed by different letters are significantly different (p≤0.05) as determined with the Tukey-Test

Cultivar/Line	EC30 Log$_{10}$ CFU (mean)	EC60 Log$_{10}$ CFU (mean)	EC90 Log$_{10}$ CFU (mean)
(A) Roggenstein			
'Baltica'	6.99 ± 0.34 (a)	7.44 ± 0.09 (cb)	6.98 ± 0.44 (a)
SR47	6.94 ± 0.24 (a)	6.95 ± 0.15 (a)	7.22 ± 0.54 (a)
SR48	6.99 ± 0.13 (a)	7.26 ± 0.09 (b)	7.24 ± 0.17 (a)
'Selma'	6.51 ± 0.25 (a)	7.25 ± 0.06 (b)	6.77 ± 0.06 (a)
'Désirée'	6.79 ± 0.39 (a)	7.56 ± 0.12 (c)	7.19 ± 0.32 (a)
'Ditta'	6.40 ± 0.28 (a)	7.57 ± 0.10 (c)	6.68 ± 0.32 (a)
'Sibu'	7.00 ± 0.21 (a)	7.57 ± 0.10 (c)	6.81 ± 0.11 (a)
(B) Oberviehhausen			
'Baltica'	7.53 ± 0.09 (abc)	7.40 ± 0.02 (abc)	7.52 ± 0.21 (a)
SR47	7.62 ± 0.15 (abc)	7.28 ± 0.09 (b)	7.38 ± 0.19 (a)
SR48	7.47 ± 0.09 (b)	7.37 ± 0.14 (abc)	7.32 ± 0.08 (a)
'Selma'	7.77 ± 0.07 (c)	7.51 ± 0.11 (abc)	7.51 ± 0.06 (a)
'Désirée'	7.55 ± 0.09 (abc)	7.52 ± 0.13 (ac)	7.38 ± 0.15 (a)
'Ditta'	7.53 ± 0.17 (abc)	7.41 ± 0.08 (abc)	7.53 ± 0.15 (a)
'Sibu'	7.39 ± 0.15 (ab)	7.52 ± 0.09 (ac)	7.33 ± 0.09 (a)

Table 2 Proportions of in vitro antagonistic isolates in the potato rhizospheres of the sampling site in Roggenstein (A) and Oberviehhausen (B) towards *Rhizoctonia solani* AG3, *Verticillium dahliae* V25 and *Phytophthora infestans* as determined by dual-culture assays

	Isolates tested	Total no. of antagonists EC30-EC90[a]		
		R. solani	*V. dahliae*	*P. infestans*
(A) Roggenstein				
'Baltica'	283	17 (a)	11 (a)	37 (a)
SR47	288	26 (abd)	24 (bd)	44 (a)
SR48	286	14 (a)	16 (ab)	39 (a)
'Selma'	256	30 (bcd)	12 (ab)	34 (a)
'Désirée'	286	50 (c)	43 (c)	69 (b)
'Ditta'	272	20 (abd)	12 (ab)	41 (a)
'Sibu'	275	30 (d)	29 (cd)	52 (ab)
total no.	1946	187	147	316
(B) Oberviehhausen				
'Baltica'	297	12 (a)	18 (a)	65 (ab)
SR47	297	15 (ab)	21 (a)	60 (a)
SR48	298	24 (bc)	20 (a)	64 (ab)
'Selma'	299	35 (c)	24 (a)	80 (ab)
'Désirée'	250	19 (abc)	24 (a)	69 (ab)
'Ditta'	298	37 (c)	25 (a)	82 (b)
'Sibu'	300	31 (c)	22 (a)	67 (ab)
total no.	2039	173	154	487

[a] EC30 = young plants, EC60 = flowering plants, EC90 = senescent plants

numbers of antagonists towards *P. infestans* were significantly higher in Oberviehhausen than in Roggenstein (data not shown).

Genotypic characterization of the antagonists using ARDRA and BOX-PCR

A total of 243 and 339 ARDRA-patterns were generated from rhizosphere isolates of EC60 and EC90 potato plants grown in Roggenstein and Oberviehhausen, respectively. EC30 isolates of both sites were excluded from further identification, because the roots of all cultivars and GM lines were very heterogeneous in size at that sampling time. Furthermore, only the in vitro antagonists displaying either activity towards the *Oomycete P. infestans* alone or additionally towards one or both of the tested fungi were further analyzed since they accounted for the vast majority of antagonists (243 of 253 in Roggenstein and 352 of 368 in Oberviehhausen, respectively, data not shown). In vitro antagonists displaying identical ARDRA-patterns (cut-off level 85%) were further characterized by BOX-PCR, which offers a higher resolution below the species. Isolates displaying unique ARDRA-patterns or one representative isolate of a

group sharing the same BOX-pattern were identified by 16S rRNA gene sequencing.

Identification of bacterial in vitro antagonists from both sampling sites

Based on the characterization by ARDRA and BOX patterns, the 16S rRNA genes of 192 and 295 in vitro antagonistic isolates from Roggenstein and Oberviehhausen, respectively were selected for 16S rRNA gene sequencing (approx. 700 bp). Although both sampling sites revealed the highest proportion of in vitro antagonists towards *P. infestans*, the composition of antagonistic bacteria revealed strong differences between the sites (Table 3). Most of the 229 isolates identified from the Roggenstein site (66.4%) were affiliated to the class of *Gammaproteobacteria* (n=152 isolates), the majority of them showing the highest similarity to the genus *Pseudomonas* (n=83 isolates), belonging to 17 different species (see Table S2, supplementary material). Fifty antagonists from Roggenstein (21.8%) were assigned to the *High G+C Gram-positives*, 40 isolates of which belonged to the genus *Streptomyces* (11 different species). In contrast to the Roggenstein site, only 69 (21.6%) of

Table 3 Phylogenetic affiliation of identified in vitro antagonists from both sampling sites as determined by partial 16S rRNA gene sequence analysis

	Roggenstein (n)	Oberviehhausen (n)
Gammaproteobacteria	**152 (66.4%)**	**69 (21.6%)**
Pseudomonas	83	36
Dickeya/Pectobacterium	34	14
Lysobacter	12	17
Acinetobacter	13	0
Stenotrophomonas	0	1
Serratia	4	0
Pantoea	4	0
Citrobacter	1	0
Enterobacter	1	1
High GC Gram-positives	**50 (21.8%)**	**149 (46.7%)**
Streptomyces	40	135
Arthrobacter	7	3
Microbacterium	3	1
Amycolatopsis	0	3
Lentzea	0	3
Saccharothrix	0	1
Micromonospora	0	1
Promicromonospora	0	1
Nocardioides	0	1
Firmicutes	**24 (10.5%)**	**93 (29.2%)**
Bacillus	13	87
Paenibacillus	11	0
Brevibacillus	0	3
Staphylococcus	0	3
Betaproteobacteria	**1 (0.4%)**	**7 (2.2%)**
Delftia	1	3
Variovorax	0	2
Massilia	0	1
Janthinobacterium	0	1
CFB group bacteria	**2 (0.9%)**	**0**
Flavobacterium	2	0
Alphaproteobacteria	**0**	**1 (0.3%)**
Phyllobacterium	0	1
Total number of isolates identified	**229**	**319**

the 319 antagonists from the Oberviehhausen site were affiliated to the *Gammaproteobacteria*, with 36 isolates belonging to the genus *Pseudomonas* (9 different species). In Oberviehhausen the majority of antagonists from the rhizosphere (46.7%) were assigned to the High G+C Gram-positives (n=149 isolates) with most sequences displaying the highest similarity to the genus *Streptomyces* (135 isolates belonging to 43 different species). The second largest group of antagonists from the potato rhizosphere in Oberviehhausen (29.2%) was assigned to the phylum *Firmicutes* (n=93 isolates) with 87 isolates belonging to the genus *Bacillus* (4 different species). In contrast, only 24 antagonists (10.5%) from Roggenstein were assigned to the phylum *Firmicutes*, with 12 isolates belonging to *Bacillus*, all of them assigned to the species *Bacillus pumilus* (see Table S2, supplementary material).

Cell wall degrading enzymatic activity,
siderophore- and AHL-production

A large proportion of the antagonists displayed
glucanase-, cellulase- and protease-activity, while
isolates with chitinase-activity were less frequently
detected. Strong site-dependent differences were found
mainly for glucanase and siderophore production,
reflecting the proportion of dominant bacterial genera
(*Pseudomonas*, *Streptomyces* and *Bacillus*) in the
collection of rhizobacteria with in vitro antagonistic
activity from both sites. As revealed by principal
component analysis, these three genera separated due
to their enzymatic properties (Fig. 1). The proportion
of antagonists producing short chain AHLs also
exhibited remarkable differences between both sites,
with the numbers in the rhizosphere of Roggenstein
exceeding those of Oberviehhausen three to four
times (data not shown). The majority of antagonists
producing AHLs belonged to the genus *Pseudomonas*.
AHL production was detected for antagonists
identified as *Ps. thivervalensis*, *Ps. corrugata*, *Ps.
brassicacearum* and *Ps. stutzeri* with both indicator
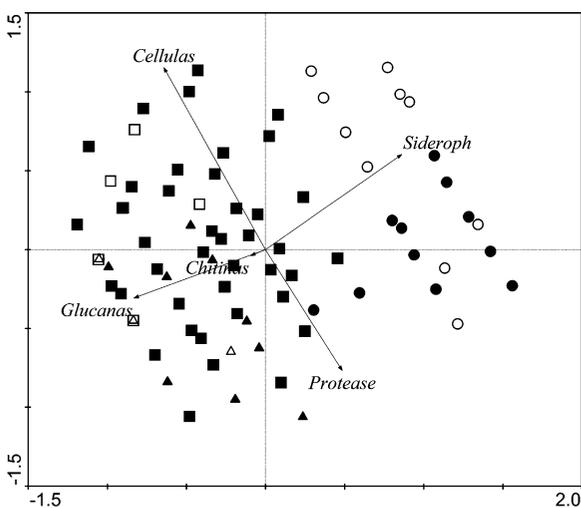
Fig. 1 Principal components analysis (PCA) ordination biplot
of in vitro antagonistic *Pseudomonas* (*circle*), *Streptomyces*
(*square*) and *Bacillus* (*triangle*) isolates and their respective
enzymatic activity and siderophore production properties
(*arrows*). Antagonists were grouped in the ordination diagram
according to Euclidian distances calculated on the basis of their
enzymatic and siderophore traits. Arrows pointing in the same
direction indicate a positive correlation between enzymatic
properties and phylogenetic assignment of the antagonists. Blue
symbols: antagonists isolated from the Roggenstein site, black
symbols: antagonists isolated from the Oberviehhausen site

strains while only *E. coli* pSB403 responded to the
AHL molecules produced by antagonists assigned to
Ps. chlororaphis, *Ps. putida* and *Ps. jessenii*. In
Oberviehhausen, only antagonists identified as *Ps.
brassicacearum* produced AHLs in vitro. Antagonists
identified as *Dickeya chrysanthemi* were the only
AHL producers which were obtained from both sites.
While AHL producing antagonists identified as
Pectobacterium carotovorum were found only in the
Roggenstein collection, *P. atrosepticum* was retrieved
from the rhizosphere of Oberviehhausen.

Comparison of BOX-profiles of the same species
from different sites

Only for a few species sufficiently high numbers of
antagonists were obtained from both sites to allow a
comparison of their phenotypic and genotypic diver-
sity which was influenced by sites, cultivars and plots.

Almost the same numbers of antagonists from both
sites were identified as *Ps. fluorescens* (Roggenstein:
n=20; Oberviehhausen: n=18). The genetic diversity
of all *Ps. fluorescens* antagonists was compared by
BOX PCR fingerprints. The results revealed both a
high intraspecific diversity for each site as well as
strong site-dependent differences between the profiles
(Fig. 2). Within one site profiles with more than 80%
similarity were found not only for isolates from the
same replicate plots, but also for different plots of the
same cultivar, yet rarely for different cultivars.

A considerable number of antagonists were identified
as *B. pumilus* with 13 and 80 isolates from the potato
rhizospheres of Roggenstein and Oberviehhausen,
respectively. The comparison of the BOX-profiles
revealed clear site-specific differences, with a very
stable profile for the majority of isolates from the
Oberviehhausen site (68/80), whereas the profiles of
the remaining isolates were clearly distinct from these.
Some representative isolates displaying the stable
BOX-profile are shown in Fig. 3. *B. pumilus* isolates
with identical BOX-patterns were obtained from all
four analyzed plots of the cultivars 'Selma', 'Désirée',
'Ditta' and 'Sibu', as well as from one plot of the
parental cultivar 'Baltica' and the GM line SR47, and
from two plots of the GM line SR48. Compared to the
Oberviehhausen site, isolates obtained from Roggen-
stein showed completely different profiles, except for
one isolate which displayed the same BOX-profile as
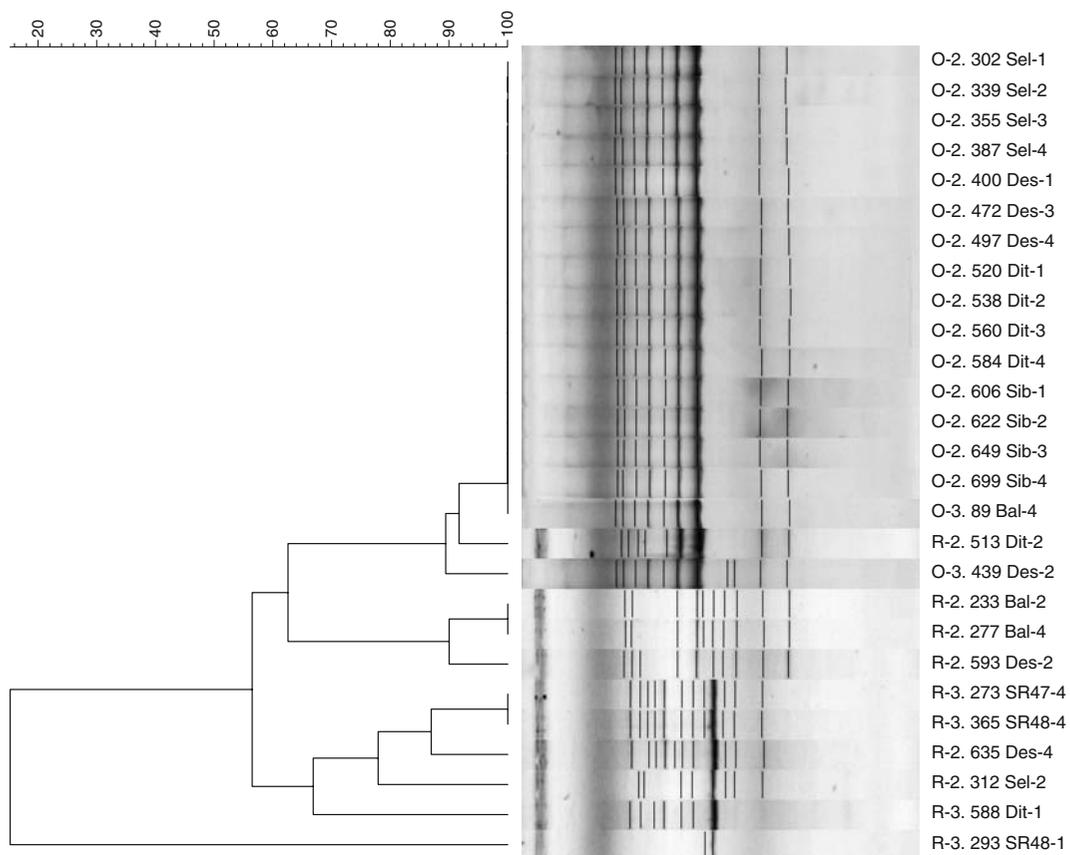the majority of isolates from Oberviehhausen (Fig. 3).
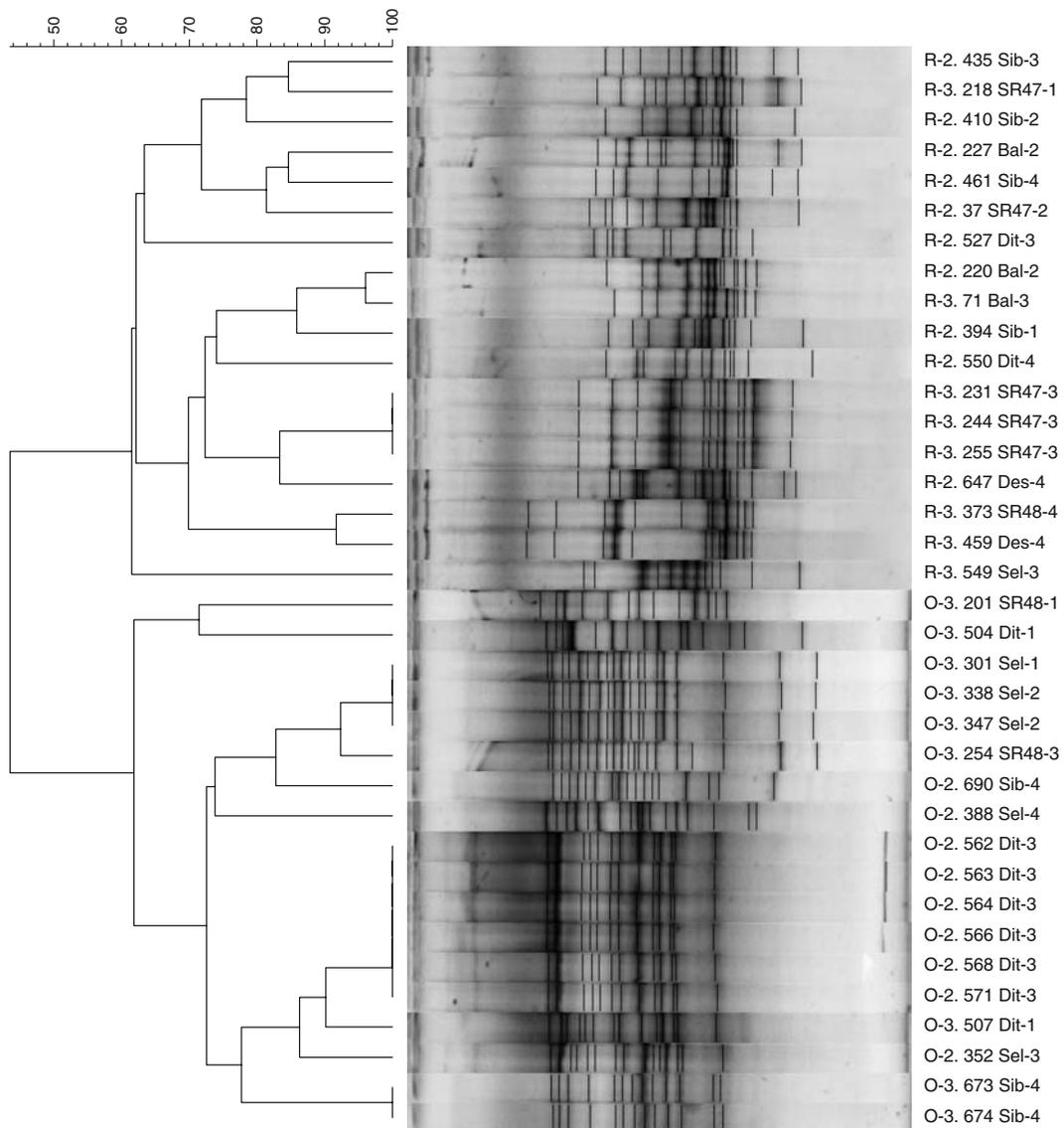
Fig. 2 Dendrogram based on BOX-PCR fingerprints of in vitro antagonistic isolates from the potato rhizospheres in Roggenstein (R) and Oberviehhausen (O), identified by 16S rRNA gene-sequencing as *Pseudomonas fluorescens*. The UPGMA algorithm was applied to the similarity matrix generated with the Dice coefficient. Strain code: site, sampling time (2 = EC60, 3 = EC90), strain number, cultivar/GM line (Bal = 'Baltica', SR47 = 'Baltica' co-suppresion, SR48 = 'Baltica' antisense, Sel = 'Selma', Des = 'Désirée', Dit = 'Ditta', Sib = 'Sibu'), plot (1 to 4) (original gel see Fig. S1)

Twenty-six and 11 antagonists from the rhizosphere of potato plants grown in Roggenstein and Oberviehhausen, respectively, were assigned to *Dickeya chrysanthemi*. Eight antagonists obtained exclusively from Roggenstein were identified as *Pectobacterium carotovorum*. Two antagonists assigned to *P. atrosepticum* were found in the collection of antagonists from Oberviehhausen. Interestingly, all *D. chrysanthemi* strains from the Roggenstein site were isolated from all four plots planted with the cultivar 'Désirée' (EC60: n=3; EC90: n=23), while most *D. chrysanthemi* isolates from the Oberviehhausen site originated from the cultivar 'Ditta' (EC60 n=2; EC90 n=7) and only two from 'Désirée'. All antagonists identified as *P. carotovorum* originated from one plot of the GM line SR48. For all 26 *D. chrysanthemi* strains of the cultivar 'Désirée' from Roggenstein the same BOX-patterns were obtained (Fig. 4). Seven of nine

Fig. 3 Dendrogram based on BOX-PCR fingerprints of in vitro antagonistic isolates from the potato rhizospheres in Roggenstein (R) and Oberviehhausen (O), identified by 16S rRNA gene-sequencing as *Bacillus pumilus*. The UPGMA algorithm was applied to the similarity matrix generated with the Dice coefficient. Strain code as described in Fig. 2 (original gel see Fig. S2)

D. chrysanthemi strains obtained from the cultivar 'Ditta' from the Oberviehhausen site displayed the same profile as the strains in Roggenstein and two showed a distinct pattern. The two *D. chrysanthemi* isolates obtained from 'Désirée' in Oberviehhausen shared the same pattern with the strains from the same cultivar in Roggenstein. The eight strains identified as *P. carotovorum* formed two separate clusters with five and three strains, respectively. The two strains identified as *P. atrosepticum* formed a tight cluster which was very distant from the other *Dickeya* and *Pectobacterium* isolates (data not shown).

Discussion

The rhizosphere was recently described as an important microenvironment for potentially antagonistic bacteria (Berg et al. 2002; Berg et al. 2006) because

bacterial populations growing in response to the easily available root exudates are often r-strategists which frequently also produce cell wall degrading extracellular enzymes to improve their competitive behavior in soil (Raaijmakers et al. 2008). Although the present study was mainly aiming to assess potential effects of GM potato lines on the group of beneficial bacteria and to relate these effects to natural variability among cultivars, its experimental design also allowed insights into the diversity of antagonistic rhizobacteria in dependence on the site. However, only for a few species (*Ps. fluorescens*, *B. pumilus* and *P. chrysanthemi*) the number of isolates was large enough to compare the diversity in relation to these factors and to assess their field distribution.

The CFU-counts determined at three developmental plant stages were comparable to those reported by Lottmann et al. (1999) for the rhizosphere of genetically modified, T4-lysozyme producing potato

Fig. 4 Dendrogram based on BOX-PCR fingerprints of in vitro antagonistic isolates from the potato rhizospheres in Roggenstein (R) and Oberviehhausen (O), identified by 16S rRNA gene-sequencing as *D. chrysanthemi*. The UPGMA algorithm was applied to the similarity matrix generated with the Dice coefficient. Strain code as described in Fig. 2 (original gel see Fig. S3)

plants, respective control lines and their parental cultivar. In the present study significant differences in the CFU-values were always transient and not stable throughout the whole season.

To assess whether the proportion and composition of plant beneficial bacteria in the rhizosphere of GM potatoes was changed compared to the parental cultivar and the four commercial cultivars, similar numbers of bacterial isolates were retrieved and tested for their in vitro antagonistic activity towards three major potato pathogens. The average proportion of in vitro antagonists towards *R. solani* and *V. dahliae* was comparable to results reported by other authors (Berg et al. 2002; Krechel et al. 2002; Lottmann et al. 1999). For the Roggenstein site the total numbers of antagonists obtained revealed significant differences both between GM lines and commercial cultivars, as well as between different commercial cultivars for all three pathogens tested. This was mainly due to the 26 *D. chrysanthemi* strains with strong antagonistic activity that were recovered from the cultivar 'Désirée' at EC90, thus leading to exceptionally higher numbers of antagonists for this cultivar. In Oberviehhausen, significant differences were observed both among commercial cultivars and GM lines and among different commercial cultivars towards *R. solani* and *P. infestans*. However, in all cases significant differences found were transient (see Table S1, supplementary material). This finding is consistent with results from other studies which indicated that changes due to the genetic modification of the plants were either transient (Donegan et al.

1995; Dunfield and Germida 2003; Griffiths et al. 2000) or negligible compared to natural factors like field site (Dunfield and Germida 2001) or season (Gyamfi et al. 2002; Heuer et al. 2002). Also altered plant characteristics, independent from the genetic modification were suggested as a possible reason for observed differences (Donegan et al. 1995; Siciliano et al. 1998). Siciliano and Germida (1999) found significant differences in the rhizosphere communities between cultivars of GM *Brassica napus* cv. Quest to cv. Excel and *Brassica rapa* cv. Parkland. However, their analysis focussed only on the flowering stage of the plants and the finding could not be generalized for other GM canola varieties (Dunfield and Germida 2001). In the present study the individual soil characteristics and cropping histories of both sites had the strongest influence on the composition of their inhabiting bacterial communities as reflected in the composition of in vitro antagonists. To date several studies showed that the soil type has a strong influence on the composition of microbial communities in the rhizosphere (Costa et al. 2006; Girvan et al. 2003; Latour et al. 1999; Marschner et al. 2001; Sessitsch et al. 2001). A remarkably high proportion of the rhizobacterial isolates from both sites displayed an in vitro antagonistic potential against the *Oomycete P. infestans*. However, the in vitro assay only evaluates the activity towards the mycelium of the *Oomycete* and not towards its zoospores which represent the infectious part (Lifshitz et al. 1985). The results of the dual-culture assays towards *R. solani* and *V. dahliae* also have to be interpreted with

care as in vitro tests do not necessarily correlate with the in situ antagonistic activity (Sari et al. 2006). Although comparable numbers of in vitro antagonists were retrieved from the rhizosphere of potatoes grown at both sites, their taxonomic composition and the proportion of antagonists producing siderophores, glucanase or AHLs showed strong site-dependent differences. One main difference was the proportion of in vitro antagonists affiliated to the *Gammaproteobacteria*, especially the genus *Pseudomonas* which was much higher in Roggenstein than in Oberviehhausen (Table 3). *Pseudomonas* species are supposed to benefit directly from root exudates provided by the plant and are therefore more often found in the rhizosphere than in other microenvironments (Costa et al. 2006; Raaijmakers et al. 2008). Also in the studies by Berg et al. (2002, 2006) the largest proportion of *Verticillium dahliae* antagonists was affiliated to the genus *Pseudomonas*. The diversity of isolated *Pseudomonas* species from the Roggenstein site was quite remarkable and several species such as *Ps. thivervalensis*, *Ps. putida* and *Ps. corrugata* were only obtained from this site. Moreover, this genus accounted for the high number of isolates from the Roggenstein site producing siderophores. A huge diversity of siderophores formed by *Pseudomonas* species has been reported (e.g. Cornelis and Matthijs 2002; Meyer et al. 2002). In a habitat like soil that is poor in disposable iron, siderophores can be important in rhizosphere colonization (Raaijmakers et al. 1995) and in competing with plant pathogens (De Boer et al. 2003; Lemanceau et al. 1992; Leong 1986). The comparison of BOX-fingerprints generated for *Ps. fluorescens* isolates revealed both a high genotypic diversity within each site but also between both sites. As reviewed by Picard and Bosco (2008) such variability is important in rhizosphere competence and biocontrol ability. Only few isolates from different plots of the same cultivar or of different cultivars at one field site displayed identical BOX-patterns. The endemism revealed by the use of BOX-PCR was also reported by Cho and Tiedje (2000) who argued that geographic isolation is important in bacterial diversification. Although the same cultivars and GM lines were grown in Oberviehhausen, antagonistic isolates belonged most frequently to the *High G + C Gram-positives* and *Firmicutes*. The BOX-patterns of *B. pumilus* showed that a genetically highly similar population colonized the rhizosphere of

the five different potato cultivars and of the two GM lines in Oberviehhausen (Fig. 3). As *B. pumilus* antagonists were retrieved from the rhizosphere of plants belonging to different cultivars grown in different plots this indicates that this population shows a homogeneous distribution at this site and obviously successfully colonized the rhizosphere of potatoes independent from the cultivar or the genetic modification. Interestingly, one *B. pumilus* isolate of 'Ditta' from the Roggenstein site shared the same BOX-pattern as the *B. pumilus* antagonists from Oberviehhausen.

The affiliation of the majority of in vitro antagonists in this study is similar to those reported for antagonistic or rhizocompetent bacteria from other studies (Berg et al. 2002, 2006; Juhnke et al. 1987). The presence of antagonists among the dominant cultured bacteria might also mirror their ability to form colonies on plates. The use of members of the genus *Pseudomonas* for biological control of plant pathogens as well as for plant growth promotion has been reported in several studies (Bloemberg and Lugtenberg 2001; Haas and Défago 2005; Mercado-Blanco and Bakker 2007; O'Sullivan and O'Gara 1992; Weller 1988). The genus *Bacillus* contains many species that were reported to have in vitro antagonistic activity. *B. pumilus* was recently described as an antagonist of *Gaeumannomyces graminis* var. *tritici*, causing take-all disease of wheat (Sari et al. 2007), but also as a plant growth promoting rhizobacterium inducing systemic resistance against *P. infestans* on tomato plants (Yan et al. 2002). Kloepper et al. (2004) reviewed several studies reporting on disease reduction by both *B. pumilus* and *B. subtilis* species in different hosts. Strains of both species are also available as commercial biocontrol products (Schisler et al. 2004). Also *Streptomyces* species have been described to reduce plant diseases caused by *R. solani* (Yuan and Crawford 1995), *V. dahliae* (Berg et al. 2000) and *P. infestans* (Malajczuk 1983) and many other plant pathogens.

Furthermore, it is interesting to note that among the in vitro antagonists obtained from both sites several isolates belonged to known potato pathogens, namely the species *Dickeya chrysanthemi* and *Pectobacterium carotovorum* as well as *S. scabies*, *S. turgidiscabies* and *S. acidiscabies*. The 26 strains identified as *D. chrysanthemi* from the potato rhizosphere of Roggenstein were obtained from all four replicates of the cultivar 'Désirée' indicating either that infected tubers

were planted or a root exudate dependent enrichment in the rhizosphere of 'Désirée'. Strains identified as *P. carotovorum* were obtained from one replicate of the GM line SR48. Although both strains are known as opportunistic pathogens, causing tuber soft rot and blackleg in the field but also tuber decay in storage (Pérombelon 2002), no infection symptoms were observed in the respective plant material retrieved from the field. Because all the strains obtained for SR48 originated from the same plot, this might either be the result of a local infection or a consequence of the pooling of the root material. Pathogenic strains of the genera *Dickeya* and *Pectobacterium* typically produce a broad range of extracellular enzymes which were described as major pathogenicity factors and are expressed in a cell-density dependent manner (Hélias et al. 2000; Kotoujansky 1987). The production of extracellular enzymes by the isolates might have contributed to their antagonistic activity. To prove whether the in vitro antagonistic *Dickeya* and *Pectobacterium* isolates showed pathogenicity in plants, they were tested in a potato slice assay and indeed the strains caused the typical symptoms (data not shown). Interestingly, even though the vast majority of the different *Dickeya* and *Pectobacterium* species produced extracellular enzymes and all strains produced AHLs, isolates belonging to *P. carotovorum* did not antagonize *R. solani* and *V. dahliae* in vitro (data not shown). Only *D. chrysanthemi* showed antagonistic activity towards all three pathogens tested, suggesting additional traits responsible for this antagonism.

Overall, in this study the effects of GM potatoes on rhizobacteria with in vitro antagonistic activity did not exceed natural variability among cultivars. The site was found to be the overriding factor determining the composition and diversity of in vitro antagonists in the rhizosphere of field grown potato plants.

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