*Burkholderia ferrariae* sp. nov., isolated from an iron ore in Brazil

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A Gram-negative, non-spore-forming bacterial strain with the ability to solubilize highly insoluble phosphatic minerals was isolated from a high-phosphorous iron ore from Minas Gerais State, Brazil. This strain, designated FeGl01T, was subjected to a polyphasic taxonomic investigation. Comparative 16S rRNA gene sequence analysis indicated that it formed a distinct phylogenetic lineage within the genus *Burkholderia* together with several other species of the genus, e.g. *Burkholderia sacchari*, *Burkholderia tropica* and *Burkholderia unamae*. Partial nucleotide sequencing and analysis of the recA gene roughly corroborated the phylogenetic position of strain FeGl01T within the genus *Burkholderia*. The chemotaxonomic properties of strain FeGl01T, such as ubiquinone Q-8 as the predominant quinone system and C₁₆:₀, C₁₇:₀ cyclo, C₁₈:₁ω7c and C₁₉:₀ω8c cyclo as the major fatty acids, were also consistent with its classification within the genus *Burkholderia*. DNA–DNA hybridization experiments between strain FeGl01T and the type strains of *B. unamae*, *B. sacchari* and *B. tropica* yielded reassociation values of 40% or lower, which, together with qualitative and quantitative differences in fatty acid composition and with differences in several phenotypic traits, support the separation of the new isolate from the phylogenetically most closely related species. Therefore, it is suggested that strain FeGl01T represents a novel species of the genus *Burkholderia*, for which the name *Burkholderia ferrariae* sp. nov. is proposed. The type strain is FeGl01T (=LMG 23612T = CECT 7171T = DSM 18251T).

Since Yabuuchi et al. (1992) proposed the genus *Burkholderia* to include the former rRNA group II pseudomonads, many other bacterial species have been described as belonging to this genus, which at the time of writing includes more than 40 species. Members of the genus *Burkholderia* have been found within many different ecological niches, but predominantly within the soil and the rhizosphere, from which some of the more recently described species have been isolated, such as *Burkholderia sacchari* (Brämer et al., 2001), *Burkholderia tropica* (Reis et al., 2004) and *Burkholderia unamae* (Caballero-Mellado et al., 2004).

Functionally, *Burkholderia* is a remarkably diverse genus that includes plant symbionts and both plant and animal pathogens. Some species of the genus are also known as opportunistic pathogens in humans. Certain species of *Burkholderia* have proved to be very efficient in biocontrol, bioremediation and plant growth promotion (Coenye & Vandamme, 2003; O’ Sullivan & Mahenthiralingam, 2005).
Many strains of *Burkholderia* species have, among other properties, the ability to solubilize highly insoluble phosphatic minerals and, therefore, are of significant interest to the agricultural sector with regard to their applicability in biofertilization (Artursson et al., 2006; Igal et al., 2001; Peix et al., 2001; Purnomo et al., 2005). Moreover, this property could also be economically useful for emerging industries such as biomining. Many of the current world iron ore resources contain over 0.08 % (w/w) phosphorus, a level above the accepted standard for the manufacture of metallic iron and steel (Cheng et al., 1999). Although there are chemical processes to reduce the phosphorus content of iron ores, the historically low prices of this raw material make them non-viable economically. In this context, biotechnology may have a role in overcoming this problem in a cost-effective and environmentally friendly way. In the course of isolating phosphate-solubilizing micro-organisms (PSMs) from a high-phosphorous iron ore in Minas Gerais State, Brazil, we isolated a bacterial strain, designated FeGl01T, that, based on its genotypic and phenotypic characterization, should be classified within a novel species of the genus *Burkholderia*.

Strain FeGl01T was isolated from a suspension of the ore material in sterile distilled water maintained under agitation for 24 h at ambient temperature. The suspension was serially diluted and spread on NBRIP agar plates. The medium NBRIP was described by Nautiyal (1999) for the detection of PSMs, and contains glucose as a carbon source and insoluble tricalcium phosphate as the sole source of phosphorus, allowing the detection of PSMs based on the formation of haloes around their colonies. Cultures used in further studies were purified from a single colony after 12 days incubation at 30°C on NBRIP medium, and subsequently cultivated on YED-P agar plates. On YED-P, colonies of strain FeGl01T were cream-coloured, circular, smooth and convex with diameters of 1–3 mm.

Genomic DNA was extracted as described by Rivas et al. (2001). The 16S rRNA gene of strain FeGl01T was analysed as described by Rivas et al. (2002). The sequence obtained was compared with those from GenBank using the FASTA program (Pearson & Lipman, 1988). Sequences were aligned using CLUSTAL X software (Thompson et al., 1997). Distances were calculated according to Kimura’s two-parameter method (Kimura, 1980). The phylogenetic tree was inferred using the neighbour-joining method (Saitou & Nei, 1987), and bootstrap analysis was based on 1000 resamplings. The MEGA2.1 package (Kumar et al., 2001) was used for all analyses. A neighbour-joining tree showing the phylogenetic position of strain FeGl01T based on its 16S rRNA gene sequence is presented in Fig. 1 (an extended tree is shown in Supplementary Fig. S1 available in IJSEM Online). The results of the phylogenetic analysis indicate that strain FeGl01T is related to members of the genus *Burkholderia*. The closest relatives to strain FeGl01T among recognized species of the genus *Burkholderia* are *B. sacchari* LMG 19450T, *B. tropica* Ppe8T and *B. unanae* MTI-641T showing, respectively, 16S rRNA gene sequence similarities of 97-6, 97-3 and 97-0 %. Two recently described *Burkholderia* species are also very closely related to strain FeGl01T: *Burkholderia silvatlantica* SRMrh-20T and *Burkholderia mimosarum* PAS44T show 16S rRNA gene sequence similarities of 97-4 and 97-6 %, respectively, to strain FeGl01T. The low similarities found between strain FeGl01T and its closest relatives suggest that it represents a novel species of the genus *Burkholderia*.

According to the results of Payne et al. (2005), *Burkholderia* species can be differentiated by analysis of an internal 385-bp sequence of the recA gene (spanning bases 76 to 461 relative to the *Burkholderia cenocepacia* J2315 genome recA gene). Moreover, Payne et al. (2005) also reported that analysis of this partial recA sequence, obtained with the *Burkholderia*-specific primers BUR3 and BUR4, produced phylogenetic trees with the same topology and discrimination as those derived from analysis of nearly full-length recA gene sequences. Although the recA analysis does not exactly match the phylogeny obtained with 16S rRNA gene sequences, it provides a greater degree of resolution among closely related species within the genus (Payne et al., 2005). Thus, to confirm the phylogenetic position of strain FeGl01T, we amplified and sequenced this partial recA region for strain FeGl01T, *B. tropica* Ppe8T and *B. unanae* MTI-641T as described by Payne et al. (2005), and these sequences were compared with those from GenBank and analysed as described above for the 16S RNA gene. A phylogenetic tree constructed with these partial recA sequences is shown in Fig. 2. The results roughly confirm the phylogenetic position of strain FeGl01T within the genus *Burkholderia* obtained by analysis of 16S rRNA gene sequences. Although *B. tropica* Ppe8T grouped in a cluster different from that containing strain FeGl01T, *B. sacchari* LMG 19450T and *B. unanae* MTI-641T, a pairwise analysis of the partial recA sequences showed that these three recognized species are the closest relatives to strain FeGl01T, with similarity values of 94-9 % (*B. sacchari* LMG 19450T), 93-5 % (*B. unanae* MTI-641T) and 92-0 % (*B. tropica* LMG 2129T).
Ppe8\textsuperscript{T}). These \textit{recA} sequence similarity values suggest that strain FeGl01\textsuperscript{T} may belong to a novel species.

For base composition analysis, DNA was prepared according to the method of Chun & Goodfellow (1995). The G + C content of the DNA was determined using the thermal denaturation method (Mandel & Marmur, 1968). The G + C content of strain FeGl01\textsuperscript{T} was 62.7 mol%. DNA–DNA hybridization was performed according to the method of Ezaki \textit{et al.} (1989), following the recommendations of Willems \textit{et al.} (2001). Mean levels of DNA–DNA relatedness of 40% were found between strain FeGl01\textsuperscript{T} and both \textit{B. sacchari} LMG 19450\textsuperscript{T} and \textit{B. tropica} Ppe8\textsuperscript{T}, and of 24% between strain FeGl01\textsuperscript{T} and \textit{B. unamae} MTI-641\textsuperscript{T} (mean of four replications). These results indicate that strain FeGl01\textsuperscript{T} does not belong to any of the recognized species of \textit{Burkholderia} based on the recommended minimum threshold value of 70% DNA–DNA relatedness for the definition of genomic species (Wayne \textit{et al.}, 1987).

Analyses of quinones and of the cellular fatty acid profile of strain FeGl01\textsuperscript{T} were performed at the DSMZ. As in all other species of the genus \textit{Burkholderia}, ubiquinone Q-8 was detected as the predominant quinone system. The fatty acid profile of strain FeGl01\textsuperscript{T} consisted of (only components comprising >1% of the total are given): \textit{C}_{14:0} (4.9%), \textit{C}_{16:0} (18.0%), \textit{C}_{17:0} cyclo (18.9%), \textit{C}_{16:1} \text{c9} (1.5%), \textit{C}_{16:2} \text{c3} (5.0%), \textit{C}_{16:0} \text{c3} (3.4%), \textit{C}_{18:1} \text{c7c} (16.7%), \textit{C}_{19:0} \text{c8c} cyclo (18.8%), \textit{C}_{18:1} \text{c5c} (1.5%) and summed features 2 (6.0%) and 3 (1.9%). Summed feature 2 corresponds to \textit{C}_{14:0} \text{c3} \text{OH}, iso-\textit{C}_{16:1} \text{c9} I, an unknown fatty acid with equivalent chain length of 10-928, \textit{C}_{12:0} \text{ALDE} or any combination of these fatty acids, and summed feature 3 corresponds to \textit{C}_{16:1} \text{c7c} and/or iso-\textit{C}_{15:0} \text{c9c} 2-\text{OH}. The components included in summed features 2 and 3 are similar to those reported in other \textit{Burkholderia} species (Caballero-Mellado \textit{et al.}, 2004; Chen \textit{et al.}, 2006; Coenye \textit{et al.}, 2001; Vandamme \textit{et al.}, 1997). The fatty acid profile of strain FeGl01\textsuperscript{T} shows significant differences from those of the phylogenetically most closely related species, \textit{B. sacchari} (Bra¨mer \textit{et al.}, 2001), \textit{B. unamae} (Caballero-Mellado \textit{et al.}, 2004) and \textit{B. mimosarum} (Chen \textit{et al.}, 2006); the proportions of \textit{C}_{17:0} \text{c7c} and \textit{C}_{19:0} \text{c8c} cyclo are considerably higher and the proportions of \textit{C}_{18:1} \text{c7c} and summed feature 3 are considerably lower in strain FeGl01\textsuperscript{T} than in these other three \textit{Burkholderia} species. In comparison with \textit{B. silvatlantica} (Perin \textit{et al.}, 2006), strain FeGl01\textsuperscript{T} contains relatively high proportions of both \textit{C}_{19:0} \text{c8c} cyclo and \textit{C}_{17:0} Cyclo (Table 1).

Phenotypic traits of strain FeGl01\textsuperscript{T} were analysed by using the API 20NE gallery (bioMérieux) as recommended by the manufacturer, and by using the API 50CH galleries (bioMérieux) inoculated with a suspension of cells in 0.7% (w/v) YNB minimal growth medium (Difco) adjusted to pH 7.0. Results are given in the species description below. Strain FeGl01\textsuperscript{T} can be differentiated from \textit{B. sacchari}, \textit{B. tropica}, \textit{B. unamae}, \textit{B. silvatlantica} and \textit{B. mimosarum} by its inability to assimilate sorbitol and D-arabinose and from the first four of these species by its ability to assimilate dulcitol.
Table 1. Differential phenotypic characteristics of strain FeGl01T and phylogenetically closely related Burkholderia species

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>1</th>
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<td>Growth on MacConkey medium</td>
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<td>D-Arabinose</td>
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<td>Rhamnose</td>
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<td>Fatty acid content (%):</td>
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<tr>
<td>C17:0 cyclo</td>
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<td>Summed feature 3*</td>
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<td>15-6</td>
<td>ND</td>
<td>7-5</td>
<td>12-7</td>
</tr>
</tbody>
</table>

*Summed feature 3 comprises C18:1ω7c and/or iso-C15:0 2-OH for strain FeGl01T (B. ferrariae sp. nov.). B. unamae (Caballero-Mellado et al., 2004) and B. mimosarum (Chen et al., 2006), C16:1ω7c for B. sacchari (Brämer et al., 2001) and C16:1ω6c and/or C16:1ω7c for B. silvatlantica (Perin et al., 2006).

and D-tagatose. Other differences in the assimilation of carbon sources are given in Table 1.

Strain FeGl01T can be differentiated genotypically and phenotypically from recognized species of the genus Burkholderia and we therefore suggest that it represents a novel species, for which the name Burkholderia ferrariae sp. nov. is proposed.

**Description of Burkholderia ferrariae sp. nov.**

Burkholderia ferrariae (fer.‘ra’ri.æ. L. gen. n. ferrariae of an iron mine).

Cells are Gram-negative, non-sporulating rods. Catalase- and oxidase-positive. Colonies on YED-P medium are cream-coloured, round, smooth and convex with diameters of approximately 1–3 mm. Nitrate is reduced to nitrite. In the API 20NE system, it produces β-galactosidase but does not produce indole, urease, arginine dihydrolase or gelatinase; it does not hydrolyse aesculin. The following substrates are assimilated as carbon sources in the API 20NE and API 50CH systems: glycerol, L-arabinose, ribose, D-xylose, adonitol, galactose, D-glucose, D-fructose, D-mannose, dulcitol, inositol, mannitol, N-acetylgalcosamine, cellobiose, trehalose, D-tagatose, L-fucose, D-arabitol, gluconate, 2-ketogluconate, malate, citrate, caprate, adipate and phenylacetate. It does not use erythritol, D-arabinose, L-xylose, methyl β-xyloside, L-sorbose, rhamnose, sorbitol, methyl α-D-mannoside, methyl α-D-glucoside, amygdalin, arbutin, salicin, maltose, lactose, melibiose, sucrose, inulin, melezitose, D-raffinose, starch, glycogen, xylitol, β-gentiobiose, D-turanose, D-lyxose, D-fucose, L-arabitol or 5-ketogluconate as carbon sources. The G+C content is 62-7 mol%.

The type strain, FeGl01T (=LMG 23612T=CECT 7171T=DSM 18251T), was isolated from ore material from the Jangada mine, Minas Gerais State, Brazil.

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**References**


Burkholderia ferrariae sp. nov.


