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## Data Article

## Metagenomic data of free cyanide and thiocyanate degrading bacterial communities

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## ABSTRACT

The data presented in this article contains the bacterial community structure of the free cyanide (CN<sup>-</sup>) and thiocyanate (SCN<sup>-</sup>) degrading organisms that were isolated from electroplating wastewater and synthetic SCN<sup>-</sup> containing wastewater. PCR amplification of the 16S rRNA V1-V3 regions was undertaken using the 27F and 518R oligonucleotide primers following the meta-community DNA extraction procedure. The PCR amplicons were processed using the illumina<sup>®</sup> reaction kits as per manufacturer's instruction and sequenced using the illumina<sup>®</sup> MiSeq-2000, using the MiSeq V3 kit. The data was processed using bioinformatics tools such as QIIME and the raw sequence files are available via NCBI's Sequence Read Archive (SRA) database.

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## Specifications Table

|                            |  |
|----------------------------|--|
| Subject area               | Biology, Microbial ecology, Biodiversity |
| More specific subject area | Metagenomics                             |
| Type of data               | Table                                    |

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|                       |   |
|-----------------------|---|
| How data was acquired | Sequencing was conducted on an Illumina <sup>®</sup> MiSeq-2000, using a MiSeq V3 (600 cycle) kit following the procedures developed at Inqaba Biotech (Pretoria, South Africa) ( <a href="http://www.inqababiotec.co.za">www.inqababiotec.co.za</a> ).   |
| Data format           | Raw data  |
| Experimental factors  | The flanking regions of the 16S rRNA gene (V1-V3) were PCR amplified using the 27F and 518R oligonucleotide primers.  |
| Experimental features | Cyanide degrading organisms (CDOs) were isolated in electroplating wastewater. Since the CDOs were unable to degrade SCN <sup>-</sup> , a gravimetric technique was employed in synthetic wastewater containing SCN <sup>-</sup> outside the BioERG laboratory. Metacommunity DNA was extracted from both the CDOs and TDOs for sequencing. |
| Data source location  | BioERG laboratory, Cape Town, South Africa (33.9324°S, 18.6406°E)<br>Electroplating facility, Cape Town, South Africa (33.9708°S, 18.5780°E)  |
| Data accessibility    | The accession numbers of the sequence data are publicly available on a public repository ( <a href="http://hdl.handle.net/11189/5110">http://hdl.handle.net/11189/5110</a> ) and are also embedded within <a href="#">Supplementary Table 1 and 2</a> .   |

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### Value of the data

- This research data provides crucial information on the bacterial community structure and differences between the CDOs and TDOs post-CN<sup>-</sup> and SCN<sup>-</sup> exposure, respectively.
  - The presented data can be utilized by researchers for comparative studies related to CN<sup>-</sup> and SCN<sup>-</sup> biodegradation.
  - The bacterial organisms detected in both the CDOs and TDOs were mainly dominated by bacteria which have never been reported to possess CN<sup>-</sup> and SCN<sup>-</sup> degradation capabilities, and future research necessitates for the determination of the role that these organisms play in CN<sup>-</sup> and SCN<sup>-</sup> biodegradation processes.
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## 1. Data

The presented dataset contains the bacterial composition of free cyanide (CDO) and thiocyanate degrading (TDO) organisms from electroplating and synthetic SCN<sup>-</sup> containing wastewater, respectively. [Table 1](#) shows the comparative analysis of the bacterial compositions between the CDOs and TDOs.

## 2. Experimental design, materials and methods

### 2.1. Sample collection and isolation procedure

The CDOs were isolated from an electroplating facility wastewater. The wastewater was collected in sterile non-transparent 20 L polypropylene containers and the cyanide concentration was immediately quantified to be above 150 mg CN<sup>-</sup>/L, using the detection technique developed by [1]. The TDOs were isolated from synthetic SCN<sup>-</sup>-containing wastewater solution (500 mL) containing (g/L); K<sub>2</sub>HPO<sub>4</sub> (3.4), KH<sub>2</sub>PO<sub>4</sub> (4.3), Glucose (0.01), SCN<sup>-</sup> (0.2) and CN<sup>-</sup> (0.2), at a pH of 10 (± 0.05), using the gravimetric technique. Briefly, the solution was exposed for two months to allow airborne microorganisms to settle on the media outside the laboratory. A fraction (100 mL) of both the synthetic and electroplating wastewater solutions was filtered sterilized in a 0.22 μm Millipore membrane and the microbial cells were re-suspended in 5 mL of sterile Millipore water in preparation of DNA extraction procedures.

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