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Bioavailability of clay-adsorbed dioxin to *Sphingomonas wittichii* RW1 and its associated genome-wide shifts in gene expression

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1 **Abstract**

2   Polychlorinated dibenzo-*p*-dioxins and dibenzofurans are a group of chemically-related

3 pollutants categorically known as dioxins. Some of their chlorinated congeners are among the

4 most hazardous pollutants that persist in the environment. This persistence is due in part to the

5 limited number of bacteria capable of metabolizing these compounds, but also to their limited

6 bioavailability in soil. We used *Sphingomonas wittichii* strain RW1 (RW1), one of the few

7 strains able to grow on dioxin, to characterize its ability to respond to and degrade clay-bound

8 dioxin. We found that RW1 grew on and completely degraded dibenzo-*p*-dioxin (DD)

9 intercalated into the smectite clay saponite (SAP). To characterize the effects of DD sorption

10 on RW1 gene expression, we compared transcriptomes of RW1 grown with either free

11 crystalline DD or DD intercalated clay, i.e. sandwiched between the clay interlayers (DDSAP).

12 Free crystalline DD appeared to cause greater expression of toxicity and stress related

13 functions. Genes coding for heat shock proteins, chaperones, as well as genes involved in DNA

14 repair, and efflux were up-regulated during growth on crystalline dioxin compared to growth

15 on intercalated dioxin. In contrast, growth on intercalated dioxin up-regulated genes that might

16 be important in recognition and uptake mechanisms, as well as surface

17 interaction/attachment/biofilm formation such as extracellular solute-binding protein and

18 LuxR. These differences in gene expression may reflect the underlying adaptive mechanisms

19 by which RW1 cells sense and deploy pathways to access dioxin intercalated into clay. These

20 data show that intercalated DD remains bioavailable to the degrading bacterium with

21 implications for bioremediation alternatives.

22 **Key words**: dioxin, bioavailability, clay-adsorbed, *Sphingomonas wittichii*, toxicity,

23 transcriptome

Introduction

The ubiquitous occurrence of polychlorinated dibenzo-*p*-dioxins and dibenzofurans

(PCDD/Fs) is a result of their widespread formation and distribution as an unintentional

chemical byproduct of industrial and incineration processes as well as through natural

formation during forest fires and volcanic activity (US EPA, 2006). While the contribution of

anthropogenic PCDD/Fs sources has decreased markedly since the 1980's, they remain a

significant contaminant of concern based on their extreme environmental persistence and

toxicity at low exposure dosage (Alcock and Jones, 1996; Van den Berg et al., 1998). Efforts to

remediate highly contaminated sites, including many Superfund sites in the United States,

typically involve remediation technologies characterized by high cost and high environmental

impact. The remedies most commonly employed include excavation and dredging followed by

landfilling of contaminated soils and sediments (Kulkarni et al., 2008; Bridges et al., 2010).

For this reason, researchers have been motivated to develop new remediation technologies for

the cleanup of these sites that are less costly and minimize habitat destruction. Two of the most

promising include the application of sorbent amendments to sequester PCDD/Fs in forms that

reduce or eliminate their bioavailability, as well as microbiologically mediated biodegradation

of PCDD/Fs (Kulkarni et al., 2008; Ghosh et al., 2011; Chai et al., 2016).

A significant challenge in microbiologically mediated biodegradation has been the

isolation of bacteria and/or bacterial communities that can detoxify PCDD/Fs in the

environment (Moreno-Forero et al., 2015). While a number of isolated bacterial strains have

shown the ability to grow on dibenzofuran (DF) or co-metabolize specific PCDD/F congeners,

few isolates have been discovered with the ability to utilize dibenzo-*p*-dioxin (DD) as a sole

carbon source (Field and Sierra-Alvarez, 2008). *Sphingomonas wittichii* strain RW1 (RW1),

isolated from the Elbe River in Germany, is one of the few bacterial strains with the ability to grow on PCDD/Fs, specifically DD (Wittich et al., 1992), DF, and 4-chloro-dibenzofuran (Field and Sierra-Alvarez, 2008; Change, 2008; Wilkes et al., 1996). It is also able to co-metabolize PCDD/F congeners with up to six Cl substituents (Wilkes et al., 1996). Furthermore, RW1 is to our knowledge the only such strain with a sequenced genome.

Previous studies have investigated the specific metabolic pathways used by RW1 in the oxidation of DD and DF including the common upper pathway responsible for transforming DF into salicylate and DD into catechol followed by further catabolism into aliphatic compounds prior to entering the TCA cycle for complete oxidation (Chai et al., 2016). The RW1 genome consists of one chromosome and two mega plasmids (Wittich et al., 1992). Recent gene-knockout studies show that degradation of DD, but not DF, by RW1 requires at least one chromosomally-encoded upper pathway gene in addition to the plasmid-encoded upper pathway genes (Thamer and Zylstra, 2016). This indicates that conjugal plasmid transfer alone would be insufficient to confer DD degradation ability and leads to the question of whether RW1 contains other chromosomally-encoded features optimized for DD degradation.

Our previous work used transcriptomic analysis to outline the differences in RW1's transcriptional responses to DD and DF (Chai et al., 2016). The stress response was stronger with DD, suggesting higher toxicity compared to DF. Furthermore, it was found that either DD or clay resulted in the category-wide down-regulation of genes associated with cell motility and chemotaxis.

The effect of clay exposure on RW1 represents an environmentally significant finding for a number of reasons. First, clays are a major class of geosorbents in soils and as such are a major component of soil microbes' microenvironment. Smectite clays are also important

1   components of clay landfill liners and slurry walls. Smectite clays possess high sorptive

2   affinity for dioxins, and have been suggested as potential sorbent amendments aimed at

3   reducing contaminant bioavailability (Liu et al., 2009; US EPA, 2013). While the recently

4   demonstrated bioavailability of clay-intercalated PCDD/Fs to a mammalian (mouse) model has

5   raised concerns regarding its utility in remediation efforts (Boyd et al., 2011), the

6   bioavailability of clay-sorbed dioxin to bacteria remains unknown. Therefore, our objectives

7   were first to determine the bioavailability of intercalated DD to RW1, and after finding

8   intercalated DD bioavailable to RW1, to examine differences in RW1 gene expression during

9   growth on DD as sole carbon source with DD either in crystalline form or intercalated into the

10   smectite clay mineral saponite (SAP).

11   **Materials and Methods**

12   RW1 strain

13   *Sphingomonas wittichii* strain RW1 was kindly provided by Dr. R. Halden of Arizona

14   State University. Sequences of RW1 chromosome (NC_00911) and plasmids (NC_00907 and

15   NC_00908), and corresponding annotations were downloaded from Joint Genome Institute

16   (JGI)[1] where the genome was sequenced and assembled. Pathway information for RW1 was

17   obtained from Kyoto Encyclopedia of Genes and Genomes (KEGG)[2].

18   Analytical materials

19   DD with 97% purity was purchased from Wako Pure Chemical Industries Ltd (Osaka,

20   Japan). Cesium chloride and other inorganic salts with 99% purity were obtained from Fisher

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[1] http://genome.jgi-psf.org/sphwi/sphwi.home.html

[2] http://www.kegg.jp/kegg/

1 Scientific (Pittsburgh, Pa). All chemicals were used as received. SapCa-2, a saponite clay

2 mineral from Ballarat, CA was acquired from the Source Clays Repository of the Clay

3 Minerals Society (Purdue University, West Lafayette, IN). The physical/chemical properties

4 for the SapCa-2 are given in Table 1.

5 Preparation of the Cs saturated SapCa-2

6   Preconditioning of the SAP mineral was necessary to ensure intercalation of DD, i.e.

7 the positioning of dioxin molecules between the planar 2:1 (silicon tetrahedral: aluminum

8 octahedral: silicon tetrahedral) clay layers; dioxin was oriented parallel to the planar clay

9 layers. $Cs+$ was selected for use as the exchangeable cation as it is less hydrated than other

10 monovalent cations (Na, K) and far less hydrated than divalent cations. The reduced hydration

11 is important because it allows the negatively charged oxygen in DD to interact with the

12 exchangeable cation $Cs+$, and creates larger absorption domains between exchangeable cations

13 that are unobscured by water. Furthermore, the lower hydration resulting from the use of $Cs+$

14 results in limited water molecules in the clay interlayers creating a subaqueous environment

15 that is energetically favorable for the hydrophobic DD molecules. Previous studies have shown

16 the efficacy of DD intercalation by Cs-saponite (Liu et al., 2009; Rana et al., 2009).

17   The preparation of homoionic Cs^+-SapCa-2 followed the method of Arroyo et al.

18 (2004). Briefly, the clay suspension was first titrated with 0.5 M sodium acetate buffer (pH 5)

19 until a stable pH at 6.8 was reached to remove carbonate impurities. Clay-sized particles (<2

20 μm) were obtained by low-speed centrifugation and then re-suspended in 0.1 M CsCl solution

21 four times to ensure complete Cs^+ saturation. The resultant Cs- SapCa-2 was washed using

22 Milli-Q water until free of chloride as indicated by a negative test with $AgNO_3$, then quick

23 frozen, freeze-dried and stored.

1   Loading of dibenzo-*p*-dioxin on Cs- SapCa-2

2   Sorption of dibenzo-*p*-dioxin to Cs-SapCa-2 was performed using a method similar to

3   Liu et al. (2009) and Rana et al. (2009). Briefly, for each reaction a 28 ml aliquot of 1000 ppm

4   stock DD solution in methanol was added to 35 L of Milli-Q water containing 0.1 M cesium

5   for a DD concentration of 0.8 mg/L. Then Cs-saturated SAP clay (600 mg) was then added to

6   the DD solution, agitated to equilibrate for 24 h, after which the clay particles were allowed to

7   settle by gravity for 24-48 h. The supernatant was removed and the remaining clay suspension

8   was collected, concentrated by centrifugation and the pellets were quick-frozen, dried and

9   stored prior to use.

10   Confirmation of DD loading using TOC analysis

11   To measure the initial concentration of DD sorbed to the Cs-SAP, total organic carbon

12   (TOC) analysis was performed using a Shimadzu SSM-5000A analyzer following the

13   EPA/600/8-87/020 method. Briefly, total carbon content (TC) was measured by catalytically

14   aided combustion oxidation at 900°C and inorganic content was measured by pre-acidification

15   at 250°C. TOC was then calculated as the difference between TC and inorganic carbon (IC).

16   Analysis was performed on 1 g samples both pre and post loading of DD and the

17   concentrations of DD were determined by the increase in TOC between the two samples. Prior

18   to loading with DD, IC content and TOC content of the purified Cs-SAP clay, were determined

19   to be below their respective detection limits. After loading, the dioxin concentration on Cs-

20   SAP was determined by the increase in TOC to a value of 0.53±0.01% (w/w). DD consists of

21   78.26 % carbon, thus the percentage of DD in SAP is 0.67 % (6700 mg/Kg). This is in

22   agreement with values reported by Liu et al. (Liu et al., 2009). Furthermore, the intercalation of

23   DD by SAP from aqueous solution was supported by three lines of evidence (Liu et al., 2009;

Rana et al., 2009): First, the interlayer basal spacing between clay layers increased consistently

as DD sorption increased (Table 1). Second, Fourier-transform infrared (FTIR) spectroscopy of

rinsed and air-dried clay films showed a similar steady increase in FTIR intensities of DD

vibrational bands. Third, the vast majority of Cs^+ cations are in the interlayer, and several

cation-dependent FTIR vibrational shifts provided strong evidence for direct DD-Cs^+

coordination complexes. The dioxin-loaded clay was added to bacterial cultures to final

concentration of 1%, with a corresponding dioxin concentration of 365 μM (67.2 ppm).

Extraction and quantification of DD from bacteria plus clay culture

The amount of DD in the bacteria clay culture was measured by High Performance

Liquid Chromatography (HPLC). DD was extracted from cultures containing 1 ml of medium

and 10 mg of Cs-SAP by dual extractions using 0.5 ml dimethylsulfoxide (DMSO) in 25 ml

corex glass centrifuge tubes. Samples were mixed with DMSO for 30 min prior to

centrifugation at 3300g. Combined supernatants were subject to HPLC analysis using a Perkin-

Elmer series 200 (Norwalk, CT) with a UV detector set at a wavelength of 223 nm and a C18

HPLC column (Supelcosil Discovery, 15cm×4.6mm, 5μm) with a mobile phase mixture

consisting of 80% methanol and 20% water at a flow rate of 1.0 ml/min. Matrix matched

calibrations were prepared at 5, 10, 20, 40, and 60 ppm and a coefficient of correlation of

0.9994 was achieved. The percent recovery of the extraction and analytical method was 92.4%

determined by repeat measurements of DD from 10 mg fortified Cs-SAP with the known

concentration of DD in the SAP having been confirmed through TOC analysis described

above.

Culture conditions

1   RW1 was grown at 30°C using defined mineral DSMZ medium 457 (Brunner medium)

2   (Deutsche Sammlung von Mikroorganismen und Zellkulturen GmbH[3]). Carbon substrates were

3   added as follows: succinate (SUC) was added to a final 20 mM concentration, dibenzo-*p*-

4   dioxin (DD) (365 µM) was added to sterile flasks from acetone stock and the flasks were left

5   open in a sterile hood for 5 h to allow acetone to completely evaporate. Brunner medium was

6   then added and flasks were sonicated for 10 sec to dislodge and break up substrate crystals.

7   Clay-sorbed-dioxin (DDSAP) was added to 1% (w/v) final concentration (1% DDSAP

8   corresponded to 365 µM DD) in Brunner medium and sonicated to homogenize clay particles.

9   The DD concentration in both DD and DDSAP cultures were equivalent at 365 µM (67.2

10 ppm). Scanning electron microscopy was used to visualize RW1 growth under the two

11 different culture conditions (Figure 1). Cell density of cultures (CFU, colony forming units)

12 was determined by plating serial dilutions on nutrient agar for succinate-only or Brunner agar

13 with DF crystals (added onto Petri dish lid) for dioxin and DDSAP cultures.

14 RNA isolation for transcriptomes

15   Total RNA was isolated from early-to-mid log phase cultures (CFU of 1.0×10^9 to

16 1.5×10^9 cells/mL) using PureLink Mini Kit (Life Technologies, Carlsbad, CA) with lysis step

17 modified as per the RiboPure Kit instructions (Life Technologies, Carlsbad, CA). To ensure

18 complete removal of DNA, RNA samples underwent a two-step DNAse I treatment, first with

19 DNAse I (amplification grade, Life Technologies) followed by TURBO DNAse I (Life

20 Technologies). RNA concentration was measured using Qubit (Life Technologies) and its

21 quality was validated using BioAnalyzer (Agilent Technologies, Santa Clara, CA).

Transcriptomes were sampled in triplicate from cultures grown under three different conditions: succinate (SUC), dibenzo-*p*-dioxin (DD), and clay-sorbed dibenzo-*p*-dioxin (DDSAP).

Transcriptome sequencing and data processing

Enrichment of mRNA was performed by removing ribosomal RNA at the Michigan State University Research Technology Support Facility (RTSF) using Duplex Specific Normalization (DSN), in which cDNA libraries were treated with duplex-specific nuclease to deplete ribosomal RNAs and increase the abundance of mRNA-derived cDNAs. Library preparations and Illumina sequencing (GA II) were performed at RTSF. Sequence reads that passed the purity filter (Illumina chastity filter) were retained, processed and mapped to the RW1 genome as in our previous transcriptome study (Chai et al., 2016). Differentially expressed genes were called at a False Discovery Rate (FDR) of 5% using DESeq2 (Love et al., 2014) with R (R Core Team, 2014).

Gene set enrichment analysis

Enrichment analysis aims to identify differentially expressed groups of functionally related genes, such as genes whose products are involved in the same metabolic pathway. This was achieved by determining whether a functional gene set is statistically over-represented, i.e. enriched among all gene sets. Enrichment tests were performed using a Python script ("functional_enrichment.py") written by Yulia Mostovoy[4] and Biocyc[5].

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[4] https://cgrlucb.wikispaces.com/Functional+Enrichment+Analysis (this web site terminated its service in September, 2018 due to the funding problem.)

[5] https://biocyc.org/

1   Genomic comparisons

2   To find other organisms sharing genes in the Swit_0683 to Swit_0703 genomic region

3   regardless of synteny, we conducted tBLASTn searches on GenBank's nt database at cutoffs of

4   60% sequence identity and 60% of gene length aligned. *Sphingomonas sp*. strain DC-6 shared

5   all 21 genes with RW1, while the next closest genome (*Sphingobium yanoikuyae* strain S72)

6   shared only seven genes. The RW1 genome was compared to DC-6 genome using JGI's

7   neighborhood ortholog tool[6] while average amino acid identity (AAI) and average nucleotide

8   identity (ANI), along with percent shared genome and percent shared ORFs were calculated

9   using MIGA[7](Rodriguez-R et al., 2018).

10   **Results**

11   We found that RW1 was able to grow in DDSAP cultures with intercalated dioxin as

12   the sole carbon source (Table 1), at the same growth rate as for the cultures with free

13   (crystalline) dioxin (Fig.2). HPLC analysis of the culture extract showed that the reduction in

14   DD concentration corresponded with the growth of RW1 (Fig. 2).

15   We performed pairwise differential expression tests between transcriptomes from the

16   three culture conditions, i.e. dioxin-alone (DD), dioxin intercalated in clay (DDSAP), and

17   succinate control (SUC) and identified 1151 genes including 248 hypothetical protein-coding

18   genes (1000 for DDSAP vs SUC, 695 for DD vs SUC and 86 for DDSAP vs DD) as

19   differentially expressed between at least two different culture conditions (Table S1, S2),

20   representing 20 COG categories. Expression profiles of these genes are presented with log2 of

21   scaled normalized abundance values (Fig. 3). Based these three comparisons, we found:

[6] https://img.jgi.doe.gov/

[7] http://microbial-genomes.org/

(1). RW1's response to dioxin compared to SUC reconfirmed what we found in our previous study, including up-regulation of catabolic pathways for dioxin, its intermediate metabolite catechol, and the more general aromatic compounds (Table 2, Table S2) as well as down-regulation of flagellar assembly pathway and genes for chemotaxis (Table 3). With many more mapped reads in this study than our previous study, we were able to detect a higher number of differentially expressed genes (695; 462 up-regulated and 233 down-regulated vs 183 up-regulated and 137 down-regulated). Although 496 of the 695 genes found in this study did not reach our statistical criteria for differential expression in our previous study, the majority of these (410 out of 496) were positively correlated in expression pattern between the two studies. Only three genes (Swit_3144, Swit_3190, Swit_2651) showed statistically significant opposite expression patterns between these two studies, well within the 5% false discovery rate used in both studies. Gene set enrichment analysis indicated up-regulation of degradation pathways of aromatic compounds including dioxin and down-regulation of flagellar assembly and chemotaxis (Table 2, 3).

(2). Differences in expression between DDSAP and DD were found for a relatively small number (86) of differentially expressed genes (Table S1). The majority (19/21) of the genes up-regulated in DDSAP vs DD were also up-regulated in DD vs SUC, including 14 genes from the Swit_0683-Swit_0694 and Swit_0702-Swit_0703 gene clusters, which span six transcription units (according to our RNA-Seq assemblies, data not shown). Three TonB-dependent receptor genes were up-regulated including one (Swit_0687) in the gene clusters. One LysR transcriptional regulator gene (Swit_4320) was up-regulated but did not respond to DD in this and our previous study.

1   Sixty-five genes were up-regulated in DD compared to DDSAP. They include several

2   gene clusters, e.g. Swit_1152-Swit_1154, Swit_1712-Swit_1714, Swit_4786-Swit_4792, and

3   Swit_5364-Swit-5371.

4   **Discussion**

5   Our study demonstrated that dioxin intercalated into clay is bioavailable and sufficient

6   as the sole carbon source for RW1 growth. Further, the similar growth rates between the

7   sorbed and free dioxin suggests that bioavailability was not rate limiting. This is congruent

8   with studies involving biphenyl which showed that rates of mineralization could not be

9   explained by desorption into the liquid phase even if instantaneous desorption was assumed

10   (Feng et al. 2000). Desorption of DD into bulk culture solution is expected *a priori* to be

11   insignificant. The low hydration of Cs^+ provides subaqueous conditions in the Cs^+ saturated

12   clay interlayers, and FTIR data from a similarly prepared DD intercalated Sap (Liu et al, 2009;

13   Rana et al., 2009) indicates the formation of Cs^+-DD complexes. Both factors provide

14   favorable energetics for DD sorption, especially in the clay interlayers. The relative

15   concentrations of DD in the aqueous phase vs. clay-sorbed DD is defined by the sorption

16   isotherm (Liu et al., 2009), and by far most of the DD mass is in the clay-sorbed state. Also,

17   desorption hysteresis is common over the time scale of the experiments, which would further

18   suppress any dissolved DD.

19   The access to intercalated dioxin may be facilitated by hydration of the exchangeable

20   cation, which controls the spacing between clay interlayers. If the Cs^+ ion is replaced by Na^+

21   for instance, the layers expand substantially and this would make it easier to access dioxin.

22   However, Cs^+ sorption to smectite clays is highly selective (Maes et al., 1985) and desorption

23   has been shown to be very low even under aggressive acid dissolution treatments (Zachara et

al., 2002). The exact mechanisms microbes use to access intercalated substrates remains

unclear. In clays, the only known abiotic degradation processes involving dioxin required the

presence of a transition metal as an exchangeable cation and occurs only in the absence of

water and therefore not possible under our experimental conditions (Boyd, S.A. and M. M.

Mortland. 1985).

We detected 10 genes known to be directly involved in stress/toxicity response up-

regulated in DD vs DDSAP, most of which were also up-regulated in DD compared to SUC,

consistent with our previous study (Chai et al., 2016). These included high temperature stress

genes, e.g. heat shock sigma factors σ32 RpoH gene (Swit_0060) and two heat shock protein

genes: chaperon protein DnaK (Hsp70) (Swit_1250) and Hsp20 (Swit_0619), RND efflux

pump genes (Swi_1152, Swit_1153, Swit_1154) belonging to an outer-membrane protein

family in Gram-negative bacteria known for removing antimicrobials and toxins (Venter et al.,

2015), a gene (Swit_3457) from the Glutathione S-transferase superfamily best known for

detoxification of xenobiotics (Allocati et al., 2008), a gene (Swit_3139) from the a

Bacterioferritin family, which was reported to provide bacteria cells protection against acid,

oxidative and other environmental stresses (Choi et al., 2000; Calhoun and Kwon, 2011), a

Rhodanese family genes (Swit_3732) encoding a cyanide detoxification enzyme (Chaudhary

and Gupta, 2012) and a gene from the TetR transcriptional regulator family (Swit_4362),

known to interact with a wide diversity of other small molecule inducing ligands, including

toxic chemicals, and to regulate genes nearby or at different genome locations (Cuthbertson

and Nodwell, 2013). Three other stress response genes, i.e. phage shock proteins, PspA and

PspC (Swit_2937, Swit_2939), known to be induced by various membrane stresses in bacteria

1   (Kleerebezem et al., 1996), and a glutathione S-transferase family gene (Swit_0145) also

2   showed higher expression levels in DD than DDSAP, although short of statistical significance.

3   Another interesting finding is the strong up-regulation of some bacterial conjugative

4   type IV secretion system genes (Swit_5364-Swit_5369, Swit_5371 on plasmid pSWIT01) in

5   DD vs DDSAP. Their expression levels follow DD > DDSAP > SUC, sharing the same trend

6   with stress/toxicity genes. The type IV secretion system is generally associated with

7   pathogenicity of Gram-negative bacteria (Wallden et al., 2010), however it is possible that the

8   up-regulation of these genes is also linked to a stress response.

9   Regarding effects from clay, we showed in our previous study that RW1's response to

10   clay alone (SAP) was limited to down-regulation of genes in cell motility pathways and for

11   these genes there were no detectable differences between DDSAP and DD. This suggests that

12   it was the interaction of dioxin with clay in the intercalated state that mainly contributes to the

13   responses unique to DDSAP, while the simple additive effect from clay alone was negligible.

14   Since expression profiles of DDSAP and DD were in general very similar (Table 2, Fig. 3), the

15   effect of dioxin intercalation, and not the presence of clay, was likely the driving factor for the

16   observed expression differences.

17   We hypothesized that dioxin intercalated into clay creates a unique challenge for RW1

18   in accessing the carbon source, compared to the freely available dioxin in DD, and therefore

19   may require a different set of genes to be up-regulated. Hence, we sought to determine what

20   genes and pathways are important in enabling RW1 to utilize dioxin in this deemed less

21   accessible, clay-sorbed state. Genes up-regulated in DDSAP compared to DD are the logical

22   candidates. Of these 21 genes, the majority were those that responded to both DDSAP and DD

23   but significantly stronger to DDSAP than DD (Fig. 4). This created an apparent differential

gradient of DDSAP > DD > SUC and most belong to the gene cluster Swit_0683-Swit_0703.

Interestingly, nine genes from this cluster (Swit_0684, Swit_0687, Swit_0688, Swit_0689, Swit_0690, Swit_0692, Swit_0693, Swit_0702, Swit_0703) also showed very uniform expression patterns in our previous study, i.e. they were strongly up-regulated in DF vs DD (and DF vs SUC). This suggests that while these genes were induced by dioxin, the lower toxicity of DF and intercalated DD in DDSAP may be responsible for their stronger expression, and further suggests that the elevated expression of these genes may be required for unimpeded growth with dioxin (DD or DF). The fact that Swit_0683-Swit_0703 spans several confirmed transcription units (RNA-Seq assembly data not shown) suggests their up-regulation in DDSAP and, to a lesser level DD, is a response regulated at the transcriptional level. Considering that most of these genes are known for encoding membrane-associated proteins such as porin, TonB-dependent receptors, LuxR, amino acid permease, and extracellular solute-binding protein, it is possible that the form and intensity of cell surface level interactions, e.g. the possible increased access to dioxin through sensing (LuxR quorum sensor, Swit_0694), channeling (porin, Swit_0689), and binding (extracellular solute-binding protein, Swit_0692) between dioxin and RW1 constitutes what differentiates the RW1 response to dioxin in DDSAP compared to other states. Apart from this gene cluster, another LuxR gene (Swit_5012) was also more strongly up-regulated in DDSAP than in DD.

In a search for genomes that share a similar Swit_0683 to Swit_0703 genomic region with JGI's neighborhood ortholog tool, we found *Sphingomonas sp.* DC-6 (GenBank accession: CP021181.1) (DC-6) to be the only genome that shares with RW1 the complete gene synteny in this region, while the next closest organism (*Sphingobium yanoikuyae* strain S72) only shared seven genes with RW1. This synteny comprises 35 genes, well beyond

Swit_0679 and Swit_0713, in both directions (Fig. 5). In fact DC-6 is also the closest relative to RW1 at the whole genome level. Using MiGA (Rodriguez-R et al., 2018) we calculated 97.08% Average Amino Acid Identity (AAI) and 98.87% Average Nucleotide Identity (ANI) between their chromosomal genomes. These two genomes share 82.7% of open reading frames and 75.3% of the total genome. The next closest relative to RW1 was *S. sanxanigenens* with a considerably lower AAI of only 60.3% and sharing only 16 of the genes in the cluster. DC-6 shared with RW1 most lower pathway genes (16 of 24) for dioxin degradation. However, it lacks at least two key genes (3-oxoadipate CoA-transferase) in the catechol pathway and is missing the entire gene set for the upper pathway. *Sphingomonas sp.* DC-6 was isolated for its ability to degrade chloroacetanilide herbicide acetochlor (Chen et al., 2013) and, as far as we are aware, has not been tested for ability to metabolize DD or DF.

In related work, Thamer and Zylstra (2016) found from gene knockout-complementation experiments that in addition to plasmid-borne genes several chromosomal genes are required for dioxin upper pathways. Our transcriptomic data indicated that some of those genes (Swit_0910, Swit_3046, and Swit_3055) were expressed in all three growth conditions. Among these, Swit_0910, Swit_3055 were up-regulated by dioxin (DDSAP and DD compared to SUC). Additional analysis of RNA-seq reads mapped to Swit_0910 and its nearly identical homolog Swit_0886 confirmed that only Swit_0910 was highly expressed and up-regulated by dioxin while Swit_0886 was hardly active, consistent with the observation that Swit_0886 in its native form did not enable dioxin degradation unless fused to a strong promoter. These up-regulated genes are candidates for future knockout experiments.

In summary, RW1 was able to access and utilize dioxin intercalated between clay layers. While the exact mechanism used by RW1 to access intercalated DD is not known, the

up-regulation of genes associated with recognition, uptake, and surface interactions/attachment/biofilm formation suggest these processes facilitate the accessibility of RW1 to dioxin in the intercalated state. Further, the intercalation of DD into clay reduced toxicity and stress to the cells as compared with exposure to fully available crystalline DD. These results suggest that sequestration via intercalation by clay minerals do not eliminate the bioavailability of DD to the degrader RW1 and may support the efficacy of remedial efforts combining geosorbent amendments with microbial bioremediation. Our studies were with freshly prepared clay-DD but historically DD contaminated sites could have aged clay-DD and hence its biodegradability would need to be evaluated.

Data Availability

The data discussed in this publication have been deposited in NCBI's Gene Expression Omnibus (Edgar et al., 2002) and are accessible through GEO Series accession number GSE115658 (https://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE115658). (access token for editors and reviewers: mvepiagubhktjkx)

Author Contributions

TT, BT, SB, and JT designed the study. TT, CL, MB and JL carried out the experiment work. BC, BS, TT, HL, CJ, JC, and GZ carried out the data analysis and discussions. BC, BS, TT and JC, SB, and JT wrote the manuscript, with contributions from all co-authors.

Conflicts of Interest Statement

The authors declare that the research was conducted in the absence of any commercial

or financial relationships that could be construed as a potential conflict of interest.

Acknowledgements

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Supplementary Material: Table S2. Differentially expressed genes in DDSAP vs SUC (1000) and DD vs SUC (659).

Figure 1. Scanning Electron Microscope images of Sphingomonas wittichii Strain RW1 cultures growing on 365 µM dibenzo-p-dioxin introduced as crystals (A) or intercalated in saponite clay interlayers with bacteria cells identified (B & C).

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