DRJAMM Is Involved in the Oxidative Resistance in *Deinococcus radiodurans*

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Proteins containing JAB1/MPN/MOV34 metalloenzyme (JAMM/MPN+) domains that have Zn2+-dependent deubiquitinase (DUB) activity are ubiquitous across among all domains of life. Recently, a homolog in *Deinococcus radiodurans*, DRJAMM, was reported to possess the ability to cleave DRMoaD-MoaE. However, the detailed biochemical characteristics of DRJAMM in vitro and its biological mechanism in vivo remain unclear. Here, we show that DRJAMM has an efficient in vitro catalytic activity in the presence of Mn2+, Ca2+, Mg2+, and Ni2+ in addition to the well-reported Zn2+, and strong adaptability at a wide range of temperatures. Disruption of drJAMM led to elevated sensitivity in response to H2O2 in vivo compared to the wild-type R1. In particular, the expression level of MoaE, a product of DRJAMM cleavage, was also increased under H2O2 stress, indicating that DRJAMM is needed in the antioxidant process. Moreover, DRJAMM was also demonstrated to be necessary for dimethyl sulfoxide respiratory system in *D. radiodurans*. These data suggest that DRJAMM plays key roles in the process of oxidative resistance in *D. radiodurans* with multiple-choice of metal ions and temperatures.

**Keywords:** Deinococcus, JAMM/MPN+, deubiquitinase, antioxidation, DMSO

**INTRODUCTION**

Proteins containing JAMM/MPN+ domain (JAMMs) have been found in prokaryotes, eukaryotes, and archaea. They play important roles in all kinds of cellular processes such as DNA repair (Zeqiraj et al., 2015), pre-mRNA-processing (Galej et al., 2014), and sulfur mobilization to form molybdenum cofactors (Cao et al., 2015). Generally, JAMMs cleave the ubiquitin-like small archaeal modifier proteins (SAMP1/2) or MoaD-MoaE in the presence of Zn2+, for instance, HvJAMM1 (Hepowit et al., 2012), PfHAMM1 (Cao et al., 2017), and CSN5 (Altmann et al., 2017). It is worth noting that the MPN domain super-family has two main subclasses: MPN+ and MPN−. The MPN+ domain-containing proteins are zinc-dependent isopeptidases with the conserved sequence (E-x[2]-H-S/T-H-x[7]-S-x[2]-D) (Cope et al., 2002; McCullough et al., 2004; Moretti et al., 2010). The functional activity of zinc-dependent isopetidases involves zinc bound to the proteins via two histidines and one aspartic acid residues, such as AMSH (Davies et al., 2011) and CSN5 (Echalier et al., 2013), which are JAMM/MPN+ proteins and have the similar organization and composition. The proteins of the MPN− family lack catalytic activity due to the absence of pivotal residues in the typical JAMM motif and are usually found in pairs in multi-protein complexes with JAMM+.
proteins. For example, the elf3 and COP9 complex has elf3f and CSN5 representatives of the JAMM/MPN family and MPN− family members, namely elf3h and CSN6 (Zhou et al., 2008; Sharon et al., 2009).

After decades of research, it has been found that the action of deubiquitinating enzymes or DUBs controls most ubiquitination events dynamically (Amerik and Hochstrasser, 2004). In addition, the deubiquitination process is achieved by hydrolyzing the last residue of the isopeptide bond after Gly76 or the peptide bond of the polyubiquitin chains connected to Met1 (Wilkinson, 1997; Love et al., 2007). According to the structural analysis of the active domain, DUBs can be divided into five subfamilies: the Ub C-terminal hydrolases (UCHLs), the Ub-specific proteases (UBPs), ovarian tumor proteases (OTUs), the Josephin domain proteases (JIDs), and JAB1/MPN/MOV34 (JAMMs) (Nijman et al., 2005). For example, in Haloferax volcanii, HvJAMM1 can cleave proteins attached to SAMP1 by linear and isopeptide bonds, and the C-terminal diglycine motif of SAMP1 is not required for HvJAMM1-mediated cleavage of linear protein fusions (Hepowit et al., 2012). In Pyrococcus furiosus, the PfJAMM1 can identify SAMP2 with accuracy, regardless of the target protein connected to the C-terminal Gly of the SAMP2 (Cao et al., 2017). In eukaryotes, AMSH is demonstrated to have DUB activity (Kyuuma et al., 2006).

Deinococcus radiodurans is well-known for its powerful capacity to endure extreme stresses such as ionizing radiation (IR), desiccation, and oxidation (Makarova et al., 2001; Daly, 2012). Studies demonstrated that oxidative stress is incurred by reactive oxygen species (ROS) (Goswami et al., 2006). The antioxidant defense mechanism of D. radiodurans is active against all three main ROS, including hydroxyl radicals (OH−), superoxide radicals (O2−), and hydrogen peroxide (H2O2). To remove the dangerous ROS and adapt to the oxygen-rich environment of Earth, D. radiodurans has evolved a variety of mechanisms to cope with stressful situations. For example, MnSOD (DR1279), a superoxide dismutase (SOD) of D. radiodurans, scavenges the superoxide more efficiently than its homologs in humans and Escherichia coli due to a more rapid protonation and release of H2O2 (Abreu et al., 2008). As reported, D. radiodurans contains a high concentration of manganese and keeps high intracellular total manganese to total iron ratio of 0.24 compared to that of radiation-sensitive bacteria (< 0.01 in E. coli) (Daly et al., 2004). And D. radiodurans contains three eukaryotic-type catalases, which are constitutively expressed in normal conditions (Lipton et al., 2002; Jeong et al., 2016). These findings give us insights into understanding the oxidative damage response mechanisms in D. radiodurans, while numerous genes related to oxidative resistance in D. radiodurans have not been clearly studied in detail (Yang et al., 2014).

Recently, dr_0402 has been found to encode the JAMM/MPN protein, and the product of its expression, DRJAMM, could cleave the MoaD-MoaE fusion protein (DR2607) and generate a C-terminal Gly residue (Yang et al., 2018). MoaD-MoaE is known as the MPT synthase that catalyzes the formation of MPT from cyclic pyranopterin monophosphate (cPMP) converted from 5'-GTP, while the two sulfur molecules on cPMP are carried as thiosulfates on the C-terminal glycine of MoaD (Leimkuhler et al., 2001; Zupok et al., 2019). During the formation of MPT, the substrate pocket of MoaE can bound the cPMP, MPT, and the C-terminal of MoaD. It has been shown that the utilitarian action of MoaD-MoaE as an MPT synthase must be cut-activated by JAMMs (Cao et al., 2015; Narrandes et al., 2015). In addition, MoaD and MoaE are essential for molybdenum cofactor (Moco)-dependent dimethyl sulfoxide (DMSO) reductase activity in archaea (Miranda et al., 2011). However, only one pair of DRMoaD-MoaE fusion protein is encoded in D. radiodurans, and the detailed catalytic activity of DRJAMM in vitro and its biological significance in vivo are still unknown.

In the present study, we found that DRJAMM could efficiently cleave DRMoaD-MoaE not only in the presence of Zn2+ but also in the presence of other metal ions in vitro under either low or high temperatures. Meanwhile, mutation of drJAMM led to a decreased survival rate and elevated transcriptional levels of DMSO reductase in response to H2O2 in vivo compared to the wild-type R1, indicating that DRJAMM plays an important role in the antioxidant process of the organism.

**RESULTS**

**DRJAMM Cleaving Activity Is Dependent on a Variety of Metal Irons**

In previous study, MoaD-MoaE has been shown to play an irreplaceable role in the transformation of cyclic pyranopterin monophosphate (cPMP) into molybdopterin (MPT) in E. coli (Neumann et al., 2009). To confirm the importance of DRJAMM to MPT synthase, cPMP was oxidized into its stable fluorescence derivatives, compound Z (Wuebbens and Rajagopalan, 1995; Dahl et al., 2013). When the amount of cPMP was set to 100% in drJAMM mutant strain, it was not detected in wild-type R1 strain and drJAMM complementary strain (Supplementary Figure 1), indicating that DRJAMM is essential for the activation of DRMoaD-MoaE.

It has been previously revealed that DRJAMM requires Zn2+ to cleave the MoaD-MoaE fusion protein (Yang et al., 2018). However, we found that other metal ions could also catalyze this cleavage activity in vitro, such as Mn2+, Mg2+, Ca2+, and Ni2+ (Figure 1A). The remaining DRMoaD-MoaE fragments were quantified using ImageJ software (National Institutes of Health, United States) to demonstrate the catalytic efficiency of DRJAMM in the presence of different metal ions, as shown in Figure 1B. Unexpectedly, DRJAMM displayed the highest catalytic efficiency in the presence of Ca2+, about threefold higher than that of Zn2+ (Figure 1B). In addition, high temperatures did not inhibit the activity of JAMMs. Since the catalytic function of PfJAMM1 was the best at 100°C in Pyrococcus furiosus (Cao et al., 2017), we set a series of temperature gradients and found that the DRJAMM exhibits catalytic activities at different temperatures. Surprisingly, DRMoaD-MoaE is stably degraded by DRJAMM even under high temperatures (Figure 1C). These results suggested that the enzyme activity of DRJAMM has strong adaptability to a wide range of temperatures, even above 100°C.
**drJAMM** Is Involved in the Antioxidative Process

To investigate the function of DRJAMM in *D. radiodurans*, a *drJAMM* (*Δdr_0402*) knockout mutant was constructed and the cell survival rate and cell growth curves were measured (Figures 2A–C). It was shown that the mutant *ΔdrJAMM* was not sensitive to UV radiation but declined significantly under H$_2$O$_2$ (0–80 mM) than the R1 (wild-type). The sensitivity is nearly disappeared after complementation with *drJAMM* (*Δdr_0402_Cwt*) in the mutant. However, the ΔdrJAMM growth curve displayed no change during a stationary phase of approximately 30 h (Figure 2C), indicating that the *drJAMM* mutation does not influence the growth rate of *D. radiodurans* but affects its response to oxidative stress.

Further analysis of oxidative stress survival with a spot-test method also demonstrated that the mutant is highly sensitive to H$_2$O$_2$, and could not endure 80 mM H$_2$O$_2$, but could be recovered after gene complementation (Figure 2D). Hence, the ROS level was measured to verify the role of drJAMM in the antioxidation process of *D. radiodurans*. As shown in Figure 2E, the ROS accumulation level in mutant was about 1.3-fold than that of R1 following 40 mM H$_2$O$_2$ treatment, while 1.5-fold higher after 80 mM H$_2$O$_2$ treatment. Furthermore, the ROS level in the mutant rises with the increase in H$_2$O$_2$ concentration, which was found to be restored to wild-type levels in the complementary strain. This suggests that the absence of drJAMM will cause the accumulation of ROS. Therefore, drJAMM is critical for the antioxidation process in *D. radiodurans*.

**Levels of drJAMM and DRMoaD-MoaE Increase Under Oxidative Stress**

To test the cleavage efficiency of DRJAMM (DR0402) to DRMoaD-MoaE (DR2607) during antioxidation processes, a Histag was fused to the C-terminal of DRJAMM and DRMoaD-MoaE in situ. The transcriptional and expressional levels of drJAMM and drmoaD-MoaE were analyzed using qRT-PCR and western blot assays following H$_2$O$_2$ treatment in the wild-type R1 strain. The mRNA levels of drJAMM and drmoaD-MoaE are increased under H$_2$O$_2$ treatment (Figure 3A), suggesting that they both may be involved in the oxidative resistance of *D. radiodurans*. Similarly, western blot assays showed that the expression of DRJAMM and DRMoaD-MoaE are both remarkably elevated about 1.5-fold and 3-fold, respectively, following H$_2$O$_2$ treatment, while the expression level of DRMoaE is also increased significantly about 1.5-fold (Figures 3B–E), indicating that the cleavage activity of DRJAMM might be necessary for oxidative resistance.

**DRJAMM Is Required for Dimethyl Sulfoxide Respiration System**

DMSO reductase activity is dependent on molybdenum cofactor (Moco) synthesis that requires JAMM/MPN
metalloprotease (Miranda et al., 2011). Sequence alignment suggested that \textit{dr}_0397 encode a molybdopterin oxidoreductase that has been shown to play a role in dimethyl sulfoxide respiration in \textit{Rhodobacter capsulatus} (Solomon et al., 2000), and is homologous to \textit{E. coli} DMSO reductase (Supplementary Figure 2).

To confirm whether the absence of \textit{drJAMM} will affect the DMSO respiration system in \textit{D. radiodurans}, the transcriptional levels of DMSO reductase were measured using qRT-PCR. Compared with wild-type R1, the mRNA levels of the DMSO reductase are more strongly elevated in \textit{ΔdrJAMM} following exposure to H$_2$O$_2$, though the levels are also induced in R1 (Figure 4A), suggesting deletion of \textit{drJAMM} causes a large demand for DMSO reductase in the antioxidant process. In the absence of DMSO, the growth of all strains was inhibited, while the addition of DMSO restarted growth. However, the growth of \textit{ΔdrJAMM} is still in stagnation after adding DMSO (Figure 4B), indicating that \textit{drJAMM} is necessary for the DMSO respiration system. In \textit{Haloflexx volcanii}, the JAMM/MPN$^+$ metalloprotease HvJAMM1 can activate MPT synthase, and anaerobic growth using DMSO as a terminal electron acceptor can be used as a method to monitor the activation of MPT synthase by HvJAMM1 (Hepowit et al., 2012).

To further investigate the function of DRJAMM in the DMSO respiration system, the wild-type R1, \textit{drJAMM} mutant strain, and \textit{drJAMM} complementary strain were grown to OD$_{600}$ = 1.0 under aerobic conditions, and then incubated with DMSO under anaerobic conditions. The DMSO reductase activity was not detected in the cell lysate of \textit{drJAMM} mutant strain, but could be readily detected in wild-type R1 and \textit{drJAMM} complementary strain (Figure 4C), suggesting that DRJAMM is important for the maturation of DMSO reductase protein.
FIGURE 3 | Analysis of transcriptional and expressional levels of drJAMM and drMoaD-MoaE under oxidative stress. (A) The mRNA levels of drJAMM and drMoaD-MoaE after exposure to 40 mM H$_2$O$_2$ for 15, 30, and 60 min. (B,C) The expression level of DRJAMM in the presence or absence of H$_2$O$_2$. GroEL was used as a control, and an anti-GroEL antibody was used for detection. The relative band strength was scanned and quantified from three independent experiments using ImageJ software. The expression level of each protein was normalized based on the expression level of GroEL. **$p$ < 0.01, ***$p$ < 0.001.

FIGURE 4 | Determination of levels of DMSO reductase under oxidative stress, growth of strains under anaerobic conditions, and the activity of DMSO reductase. (A) The mRNA levels of the DMSO reductase at different times following exposure to 40 mM H$_2$O$_2$ in R1 and ΔdrJAMM. Values were means of three independent assays (mean ± SD), ns, not significant, ***$p$ < 0.001. (B) Anaerobic growth of R1, ΔdrJAMM and ΔdrJAMM_Cwt in medium containing 0 or 100 mM DMSO for 4 days. Values were means of three independent assays (mean ± SD), **$p$ < 0.01, ***$p$ < 0.001. (C) The DMSO reductase activity was monitored with nitrogen at A600 nm. The U was defined as 1 μmol substrate consumed per minute at room temperature. Values were means of three independent assays (mean ± SD).
**DISCUSSION**

A broad spectrum of species encoding JAMM/MPN domain proteins are dependent on Zn$^{2+}$. For instance, the activity of HvJAMM1 can be activated by the addition of excess ZnCl$_2$ (Hepowit et al., 2012), and the loss of structural zinc leads to a significant reduction in the thermal stability of AMSH (Bueno et al., 2015). However, in the present study, the JAMM/MPN + protein DRJAMM could be activated by different metal ions such as Mn$^{2+}$, Mg$^{2+}$, Ca$^{2+}$, and Ni$^{2+}$ besides Zn$^{2+}$. Interestingly, more and more multi-metal-dependent nucleic acid enzymes (NAE) have been found to choose sulfophilic metal based on the characteristics of the reaction, or to perform the response through polymetallic collaboration (Zhou and Liu, 2018). Hence, as to how different metal particles control the action of DRJAMM amid the antioxidant handle, and whether there are numerous administrative components like multi-metal-dependent NAE needs further structural explanation.

Usually, proteins will be denatured and lose their function at high temperatures (Bischof and He, 2005). However, DRJAMM displayed stable protease activity even at 100°C. From the perspective of genome evolution, it is proposed that *D. radiodurans* has obtained many genes from *Thermus thermophilus* (Makarova et al., 2001), which may explain the resistance of high temperatures by DRJAMM.

A previous study revealed that HvJAMM1 regulates sumoylation and HvJAMM1-type proteins are thought to release SAMP (Hepowit et al., 2012). Meanwhile, DRJAMM contains a conserved motif similar to HvJAMM1 (Yang et al., 2018), and has the same reaction product MoaE (Hepowit et al., 2012). MoaE usually forms the MPT synthase with MoaD that shares a common globular $\beta$-grasp fold with Ub (Narrandes et al., 2015; Yang et al., 2018). For example, TtuB is a bacterial ubiquitin-like protein that has a similar globular $\beta$-grasp fold to the Ub of archaea (Shigi, 2012). In addition, DRMoaE is also associated with the function of ubiquitin-like (Ubl) proteins (Humbard et al., 2010). We speculated that Ubl protein modification system may exist in *D. radiodurans* and DRJAMM might play an important role.

Previous studies showed that BRCC36 is a JAMM (JAB1/MPN/Mov34 metalloenzyme) domain DUB enzyme and is involved in the DNA damage response (Patterson-Fortin et al., 2010). Although the survival rate of $\Delta$drJAMM is identical to R1 under UV stress (Supplementary Figure 3), the growth of this mutant is dramatically inhibited relative to the wild-type R1 under H$_2$O$_2$ stress. Simultaneously, the mRNA level of DRJAMM is significantly increased under oxidative stress, implying the importance of this protein in improving oxidation resistance.

According to previous studies, MoaD-MoaE in molybdenum cofactor (Moco) biosynthesis was able to catalyze some redox reactions in vivo (Zupok et al., 2019). In the present study, we found the expression levels of DRMoaE and DRJAMM are increased simultaneously under oxidative stress, indicating that DRJAMM is necessary for DRMoaE activation. A recent study revealed that HvJAMM1 plays an important role in releasing MoaE in Moco biosynthesis through deubiquitination (Cao et al., 2015). Therefore, DRJAMM might participate in the antioxidant process by cleaving the DRMoaD-MoaE fusion protein to release MoaE in *D. radiodurans*.

As an essential enzyme in the Moco biosynthesis pathway of bacteria, MoaE is located upstream of MobB, while MobB is responsible for forming molybdenum guanine dinucleotide commonly found in the DMSO reductase family (McLuskey et al., 2003; Miranda et al., 2011). The SAMPI-MoaE is ineffective in DMSO respiration, and this process requires metalloprotease HvJAMM1 (Cao et al., 2015). Under anaerobic conditions, wild-type R1 and complementary strains can remain in abnormal growth, while growth of the *drJAMM* knockout strain is almost completely stopped. After supplement with DMSO, the mutant still showed weak growth, while both wild-type R1 and the complementary strain recovered. Furthermore, the DMSO reductase activity is nearly completely lost in the *drJAMM* mutant strain. These results verified that *drJAMM* is necessary for the DMSO respiratory system in *D. radiodurans*.

In addition, a variety of microorganisms grow through the respiration of DMSO as an electron acceptor, and several DMSO respiratory systems with different compositions have been identified (Miralles-Robledillo et al., 2019). In a previous study, the thioredoxin (Trx) system, which is composed of NADPH, thioredoxin reductase (TrxR), and thioredoxin, provides the electrons to thiol-dependent peroxidases (peroxiredoxins) to remove ROS, and contributes to the resistance toward oxidative stress in *D. radiodurans* (Holmgren, 2000). Here, the levels of DMSO reductase in *D. radiodurans* are gradually increased under oxidative stress, especially in *drJAMM* knockout strains, implying that DMSO respiratory systems might be involved in the oxidation resistance similar to the Trx system, and *drJAMM* could play an important role in this process.

Taken together, DRJAMM is essential for resistance to oxidative stress and the DMSO respiration system in *D. radiodurans* (Figure 5). When oxidative damage is encountered, DRMoaD-MoaE is cleaved by DRJAMM to produce DRMoaE, which ultimately affects the DMSO reductase involved in the antioxidant process as demonstrated in *E. coli* (Leimkuhler, 2020). Overall, our findings provide new insights into the role of the JAMM/MPN domain proteins DRJAMM, which can accommodate a multiple-choice of metal ions and temperatures in *D. radiodurans* under oxidative stress.

**MATERIALS AND METHODS**

**Strains and Growth Conditions**

All strains, plasmids and primers used in this study are listed in Supplementary Tables 1, 2. *E. coli* strains were grown in Luria-Bertani (LB) liquid medium (1% tryptone, 0.5% yeast extract, and 1% sodium chloride) or on agar (1.5% Bacto-agar) plates supplemented at 37°C with appropriate antibiotics. All *D. radiodurans* strains were grown at 30°C in tryptone glucose yeast extract (TGY) liquid media or on agar plates (0.5% tryptone, 0.1% glucose, and 0.3% yeast extract) supplemented with appropriate antibiotics.
Expression and Purification of Proteins
The \textit{dr}_0402 and \textit{dr}_2607 genes were amplified and cloned into a modified pET28a expression vector at NdeI and BamHI site (Supplementary Table 2), respectively (Li et al., 2019). Then, the constructed plasmids were transformed into \textit{E. coli} BL21 (DE3), and induced in LB medium containing 50 \(\mu\)g/mL kanamycin and 0.4 mM isopropyl-\(\beta\)-D-thiogalactopyranoside (IPTG) at 30\(^\circ\)C for 5 h. Cells were collected and resuspended in lysis buffer (20 mM Tris–HCl, pH 8.0; 500 mM NaCl; 5\% (w/v) glycerol; 3 mM \(\beta\)-mercaptoethanol and 9 mM imidazole), followed by sonication. After centrifugation at 20,000 g for 30 min at 4\(^\circ\)C, the supernatant was purified using a Ni-NTA column (1 mL, GE Healthcare Biosciences, United States) equilibrated with buffer A (20 mM Tris–HCl, pH 8.0; 500 mM NaCl; 5\% (w/v) glycerol and 3 mM \(\beta\)-mercaptoethanol), and washed by buffer B (20 mM Tris–HCl, pH 8.0; 500 mM NaCl; 5\% (w/v) glycerol; 3 mM \(\beta\)-mercaptoethanol; 500 mM imidazole). Finally, the proteins were concentrated and purified using a Superdex75 column (GE Healthcare Biosciences, United States), DRJAMM was eluted with buffer C (20 mM Tris–HCl, pH 8.0; 200 mM KCl; and 1 mM EDTA), DRMoaD-MoaE was eluted with buffer D (20 mM Tris-HCl, pH 8.0; and 200 mM KCl).

DRJAMM Activity Assays
The DRJAMM (DR0402) activity assays were performed as described previously with minor modifications (Yang et al., 2018). The 40 \(\mu\)M of DRJAMM and 10 \(\mu\)M of DRMoaD-MoaE were added into the reaction buffer (100 mM KCl; 20 mM Tris-HCl, pH 8.0; and 1 mM dithiothreitol), and then 0.4 mM of different metal ions were added into samples. The reactions with 0.4 mM Ca\(^{2+}\) were incubated at 4, 16, 30, 37, 70, and 100\(^\circ\)C for 30 min, respectively, and quenched by the addition of SDS loading buffer followed by immediate boiling. The products were identified by Tricine-SDS-PAGE (12\%).

Construction of Mutant Strains
The mutant strains were constructed by a tripartite ligation method, as described previously (He et al., 2020). Briefly, the DNA fragment upstream of \textit{dr}_0402 was amplified by PCR using the primers \textit{1dr}_0402-p1 and \textit{1dr}_0402-p2, which was digested with BamHI (Supplementary Table 2). The DNA fragment downstream of \textit{dr}_0402 was amplified by PCR using the primers \textit{1dr}_0402-p3 and \textit{1dr}_0402-p4, which were digested with HindIII (downstream) (Supplementary Table 2). The digested fragment was connected to a streptomycin resistance gene. After the triplet ligation product was transformed into the \textit{D. radiodurans} wild-type R1 strain, the mutant colonies were then selected on TGY plates containing 10 \(\mu\)g/ml streptomycin, and confirmed by genomic PCR using primers \textit{1dr}_0402-p5 and \textit{1dr}_0402-p6, and DNA sequencing. For complementary strain construction, the wild type \textit{dr}_0402 was amplified by PCR using \textit{Adr}_0402-p5 and \textit{Adr}_0402-p6, and DNA sequencing. For complementary strain construction, the wild type \textit{dr}_0402 was amplified by PCR using \textit{Adr}_0402_Cwt-F and \textit{Adr}_0402_Cwt-R, and cloned into the plasmid pRADK containing the \textit{D. radiodurans} groEL promoter; and then transformed into the \textit{Adr}_0402 mutant strain to obtain the complementary strain \textit{Adr}_0402_Cwt.

Western Blot Analysis
Western blotting was used to confirm protein expression levels were performed as described previously (Dai et al., 2018). The
were grown to OD\textsubscript{600} = 1.0. For anaerobic growth, the strains were transferred to TGY medium containing 100 mM DMSO as a terminal electron acceptor at 30°C for 4 days.

**Dimethyl Sulfoxide Reductase Activity Assay**

DMSO reductase activity assay was performed as described previously (Miranda et al., 2011). A total of 250 mL of TGY cultures of each strain were grown to OD\textsubscript{600} = 1.0, harvested by centrifugation, washed in 15 mL buffer A (50 mM Tris-HCl, pH 7.5; 1 mM EDTA, pH 8.0; 2 M NaCl), resuspended in 20 mL buffer A, and lysed by ultrahigh pressure homogenizer (Shanghaiilitu, China), successively. Cell lysates were clarified by centrifugation (15,000 rpm, 30 min), and protein concentrations were measured using the Bradford assay kit (Beyotime Biotechnology, China). The DMSO reductase activity was monitored at A\textsubscript{600nm} (15 s intervals for 3.5 min). Assays (4 mL) included cell lysate (1–1.5 mg protein), 0.3 mM methyl viologen in buffer A, and the top filled with nitrogen. The mixture was titrated to 1–1.2 A\textsubscript{600nm} units with fresh 20 mM sodium dithionate (Na\textsubscript{2}S\textsubscript{2}O\textsubscript{4}) in 20 mM sodium bicarbonate (NaHCO\textsubscript{3}) prior to the addition of 10 mM DMSO. One unit (U) of enzyme activity was defined as 1-μm substrate consumed per minute at room temperature, with an extinction coefficient A\textsubscript{600nm} of 13.6 (mM\textsuperscript{-1}.cm\textsuperscript{-1}) for methyl viologen.

**DATA AVAILABILITY STATEMENT**

The original contributions presented in the study are included in the article/Supplementary Material, further inquiries can be directed to the corresponding author/s.

**AUTHOR CONTRIBUTIONS**

YH conceived the project. JC, CP, LW, and YH designed the experiments and drafted the manuscript. JC constructed the vectors and mutants and purified the proteins. CP was responsible for qRT-PCR, enzyme activity, and phenotype analysis. JC, BT, HX, and YZ participated in the data analysis. All authors reviewed the manuscript and approved the version to be published.

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**SUPPLEMENTARY MATERIAL**

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fmicb.2021.756867/full#supplementary-material
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