A Cytoplasmic NAD(P)H-Dependent Polysulfide Reductase with Thiosulfate Reductase Activity from the Hyperthermophilic Bacterium Thermotoga maritima

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ABSTRACT Thermotoga maritima is an anaerobic hyperthermophilic bacterium that efficiently produces H2 by fermenting carbohydrates. High concentration of H2 inhibits the growth of T. maritima, and S0 could eliminate the inhibition and stimulate the growth through its reduction. The mechanism of T. maritima sulfur reduction, however, has not been fully understood. Herein, based on its similarity with archaeal NAD(P)H-dependent sulfur reductases (NSR), the ORF THEMA_RS02810 was identified and expressed in Escherichia coli, and the recombinant protein was characterized. The purified flavoprotein possessed NAD(P)H-dependent S0 reductase activity (1.3 U/mg for NADH and 0.8 U/mg for NADPH), polysulfide reductase activity (0.32 U/mg for NADH and 0.35 U/mg for NADPH), and thiosulfate reductase activity (2.3 U/mg for NADH and 2.5 U/mg for NADPH), which increased 3-4 folds by coenzyme A stimulation. Quantitative RT-PCR analysis showed that nsr was upregulated together with the mbx, yeeE, and rnf genes when the strain grew in S0- or thiosulfate-containing medium. The mechanism for sulfur reduction in T. maritima was discussed, which may affect the redox balance and energy metabolism of T. maritima. Genome search revealed that NSR homolog is widely distributed in thermophilic bacteria and archaea, implying its important role in the sulfur cycle of geothermal environments.

IMPORTANCE The reduction of S0 and thiosulfate is essential in the sulfur cycle of geothermal environments, in which thermophiles play an important role. Despite previous research on some sulfur reductases of thermophilic archaea, the mechanism of sulfur reduction in thermophilic bacteria is still not clearly understood. Herein, we confirmed the presence of a cytoplasmic NAD(P)H-dependent polysulfide reductase (NSR) from the hyperthermophile T. maritima, with S0, polysulfide, and thiosulfate reduction activities, in contrast to other sulfur reductases. When grown in S0- or thiosulfate-containing medium, its expression was upregulated. And the putative membrane-bound MBX and Rnf may also play a role in the metabolism, which might influence the redox balance and energy metabolism of T. maritima. This is distinct from the mechanism of sulfur reduction in mesophiles such as Wolinella succinogenes. NSR homologs are widely distributed among heterotrophic thermophiles, suggesting that they may be vital in the sulfur cycle in geothermal environments.

KEYWORDS elemental sulfur reduction, thiosulfate reduction, sulfur reductase, NAD(P)H-dependent polysulfide reductase, Thermotoga maritima, thermophile

Thermotoga maritima is a strict anaerobic hyperthermophilic bacterium that was originally isolated from geothermally heated marine sediments (1). It is unique in that it belongs to a deep-branching lineage of domain Bacteria and possesses numerous genes found only in Archaea (2). T. maritima has drawn considerable attention for
its ability to utilize diverse carbohydrates and for its high hydrogen production (1–4). It can ferment 1 mol glucose to about 2 mol acetate, 2 mol CO₂, and 4 mol H₂, which is close to the Thauer limit of hydrogen production (4 mol H₂ per mole glucose) (1, 5). Previous research found that T. maritima growth was significantly inhibited by high H₂ concentration, whereas in the presence of elemental sulfur (S⁰) and thiosulfate, production of H₂ decreased and H₂S was produced instead (1, 5, 6), suggesting that sulfur reduction might have important effects on the metabolism of T. maritima.

Sulfur reduction is a common physiological feature shared by many bacteria and archaea (7–9). S⁰ can be used as the terminal electron acceptor of the respiratory system in several sulfur-reducing bacteria (7, 8, 10). In Wolinella succinogenes, for example, sulfur reduction is linked to H₂ oxidation. H₂ is oxidized by a membrane-bound hydrogenase, and the electrons are transferred to S⁰ by cytochromes and the polysulfide reductase PsrABC. Protons are pumped into the periplasm from the cytoplasm during this process, and ATP is further generated by the electrical H⁺ potential (11, 12). The mechanism of S⁰ reduction in W. succinogenes has been known as a model of bacterial sulfur reduction.

T. maritima reportedly reduced S⁰ to hydrogen sulfide (H₂S) after its isolation (1, 5). In the presence of sulfur, T. maritima growth was stimulated, however, the production of acetate, lactate, and CO₂ remained unaltered (1). Thus, sulfur reduction was assumed to be irrelevant to energy metabolism, and S⁰ was thought to act as an electron sink to prevent H₂ accumulation (1, 5, 6). Besides, neither has typical membrane-bound polysulfide reductase been discovered in the genome of T. maritima nor have cytochrome or quinone been found to be required for the electron transport chain in T. maritima (2, 3). Unlike W. succinogenes, T. maritima cannot reduce S⁰ through traditional sulfur respiration; rather, S⁰ reduction occurs through a different mechanism.

Besides sulfur-reducing bacteria, many archaea can also reduce S⁰, such as Pyrococcus furiosus, a hyperthermophilic archaea (13). Like T. maritima, P. furiosus is strictly heterotrophic; it produces H₂ fermentatively, and its growth is also inhibited by H₂. Sulfur can also stimulate the growth of P. furiosus under H₂ pressure. Considering these similar properties, T. maritima and P. furiosus may have similar mechanisms of sulfur metabolism. A coenzyme A (CoA)-dependent NAD(P)H:sulfur oxidoreductase (NSR) and a membrane-bound sulfane reductase (MBS) have been discovered and characterized in P. furiosus (14, 15), representing a novel mechanism of sulfur reduction. However, no sulfur reductase in T. maritima has been reported until now.

In this study, we investigated the sulfur reduction activities in cell extracts of T. maritima; we predicted through bioinformatics analysis that an ORF (THEMA_RS02810, the locus tag in GenBank; its old locus tag in GenBank is TM0379) in its genome can encode a protein homologous to the NSR from P. furiosus. The predicted NSR from T. maritima was characterized to catalyze the NAD(P)H-dependent reduction of S⁰, polysulfide, and thiosulfate through heterologous expression, purification, and biochemical analysis. Reverse transcription PCR (RT-PCR) analysis further confirmed that the NSR, together with a putative membrane-bound sulfur oxidoreductase (MBX), may function in the sulfur reduction of T. maritima. Our genome search shows that the NSR homolog is widely distributed in thermophilic bacteria and archaea, suggesting that it might play an important role in the biological sulfur cycle of geothermal environments.

RESULTS

Effects of S⁰ on T. maritima growth. Unlike other sulfur-reducing bacteria, T. maritima is a heterotrophic bacterium and its growth was not dependent on sulfur respiratory systems. T. maritima grew slowly in the absence of S⁰, reaching an OD₆₀₀ of 0.6 after around 30 h in a starch medium at 80°C (Fig. 1). When cultivated in the starch medium with S⁰, the growth of T. maritima significantly accelerated compared to the growth without S⁰, reaching an OD₆₀₀ of 0.8 after 20 h. Meanwhile, when S⁰ was added to the medium during the growth of T. maritima, H₂S formation dramatically increased 1 h after sulfur addition (Fig. S1 in the supplemental material).
Sulfur reductase activity in *T. maritima* cell extracts. *T. maritima* cells cultured in both starch and sulfur-containing media were harvested at the late exponential growth phase to prepare cell extracts by sonication. After a 100,000 × g centrifugation, cytoplasmic and membrane fractions were used for sulfur reductase activity assays. No H$_2$S production was detected with the membrane fraction, no matter whether NAD(P)H or reduced ferredoxin was used as the electron donor. No ferredoxin-dependent sulfur reductase activity was detected in the cytoplasmic fractions, either. Significant H$_2$S production was detected with the cytoplasmic fraction when NAD(P)H was used as the electron donor in the presence of S$^0$ (Table 1). NAD(P)H oxidation was also detected by monitoring the absorption decrease at 340 nm (Table 1). The results indicate that the sulfur reduction of *T. maritima* is an NAD(P)H-dependent reaction and occurs mainly in the cytoplasm. The results also revealed that the NAD(P)H-dependent sulfur reductase activity was dramatically stimulated by CoA (Table 1). CoA-dependent sulfur reductase activity was also observed in thermophilic archaea, such as *P. furiosus* (14), suggesting that *T. maritima* may share a mechanism of sulfur reduction similar to *P. furiosus*. In addition, cell extracts from the S$^0$-containing medium presented higher sulfur reductase activities than those from starch medium (Table 1), indicating that S$^0$ may promote the expression of the presumed cytoplasmic sulfur reductase.

**Heterologous expression and characterization of the cytoplasmic sulfur reductase NSR of *T. maritima*.** Attempts to purify the sulfur reductase from *T. maritima* cell extracts failed because of their poor stability. However, we observed that the sulfur reductase reaction of *T. maritima* was stimulated by CoA, which is similar with that of the reported cytoplasmic NSR in hyperthermophilic archaea *P. furiosus* and *Thermococcus kodakarenensis*, implying that the sulfur reductase of *T. maritima* may be similar to NSR from the two archaea. After searching the *T. maritima* genome, we found that the protein encoded by the ORF THEMA_RS02810 (its old locus tag is TM0379) contains similar conserved domains (a FAD/NAD binding domain and a FAD/NAD-linked reductase dimerization domain) with NSRs from *P. furiosus* and *T. kodakarenensis* (Fig. S2 in the supplemental material). It shares 57% identity in protein sequence with the NSR from

**TABLE 1** H$_2$S production and NADH oxidation activities in the cytoplasmic fraction of *T. maritima* cell extracts

| Cell extracts          | H$_2$S production (U/mg) | NADH oxidation (U/mg) |
|------------------------|---------------------------|-----------------------|
|                        | NADH + S$^0$ | NADH + S$^0$ + CoA   | NADH + S$^0$ | NADH + S$^0$ + CoA   |
| Starch medium (no S$^0$) | 0.01 ± 0.00 | 0.12 ± 0.10           | 0.09 ± 0.04 | 0.21 ± 0.11           |
| S$^0$ medium           | 0.14 ± 0.08 | 0.48 ± 0.10           | 0.40 ± 0.16 | 0.81 ± 0.25           |

**FIG 1** Effects of S$^0$ and thiosulfate on *T. maritima* growth. *T. maritima* was cultivated in starch medium (black circles), starch medium containing S$^0$ (5 g S$^0$ powder per L of medium) (red squares), and starch medium containing thiosulfate (30 mM Na$_2$S$_2$O$_3$) (blue triangles).
**T. kodakarensis** and 34% identity with the NSR from *P. furiosus*. Besides, sequence analysis indicated that this protein does not have a transmembrane domain or signal peptide. This suggests that THEMA_RS02810 may encode a cytoplasmic enzyme involved in sulfur reduction in *T. maritima*.

THEMA_RS02810 was then heterologously expressed in *Escherichia coli* strain BL21, and the corresponding His-tagged protein was purified anaerobically. The purified recombinant enzyme had a molecular weight of around 50 kDa, as shown by SDS-PAGE analysis (Fig. 2A). This is consistent with the deduced molecular mass according to its protein sequence. The enzyme is yellow in color and exhibits an UV–visible (UV–vis) spectrum with two peaks at around 320 and 430 nm (Fig. 2B). Thin-layer chromatography analysis showed that it binds FAD as its cofactor (Fig. S3 in the supplemental material). The purified recombinant enzyme presented a specific NAD(P)H: S0 reductase activity (Table 2). With S0 as the electron acceptor, the recombinant protein had a NADH oxidation activity of 0.8 U/mg and a NADPH oxidation activity of 1.3 U/mg at 45°C. (One unit (1 U) is defined as the oxidation of 1 μmol NAD(P)H per minute or the formation of 1 μmol H2S per minute.) Moreover, when CoA was added to the reaction mixture, the activities increased to 3.2 U/mg (NADH, Fig. 2C) and 4.2 U/mg (NADPH), which is consistent with the observations of *T. maritima* cell extracts. The apparent K_m values of NADH and NADPH were also comparable (around 0.05 mM for NADH and 0.08 mM for NADPH) (Table 2). The protein also presented H2S production activities from S0 reduction with NAD(P)H. No ferredoxin-related activity was detected with the recombinant enzyme. The results indicate that the recombinant enzyme is an NAD(P)H-dependent sulfur reductase, and CoA can significantly stimulate the reaction.

Several CoA-dependent NAD(P)H:sulfur oxidoreductases have been characterized in thermophilic archaea, in which the TK1481 from *T. kodakarensis* has the highest (57%)

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**TABLE 2 Activities of recombinant NSR of *T. maritima***

| Substrates       | NAD(P)H oxidation (U/mg) | H2S production (U/mg) | Apparent K_m (μM) |
|------------------|--------------------------|-----------------------|-------------------|
| NADH + S0        | 0.8 ± 0.08               | 0.39 ± 0.27           | 50 ± 6.3 (NADH)   |
| NADH + S0 + CoA  | 3.2 ± 0.10               | 1.38 ± 0.76           |                   |
| NADPH + S0       | 1.3 ± 0.02               | 0.45 ± 0.20           | 80 ± 21.2 (NADPH) |
| NADPH + S0 + CoA | 4.2 ± 0.05               | 1.21 ± 0.32           |                   |
| NADH + Na2S2O3   | 2.5 ± 0.01               | 1.35 ± 0.49           | 10 ± 1.2 (Na2S2O3) |
| NADH + Na2S2O3 + CoA | 6.8 ± 0.04 | 2.79 ± 1.20           | 60 ± 10.3 (NADH)  |
| NADPH + Na2S2O3  | 2.3 ± 0.02               | 1.56 ± 0.31           | 10 ± 3.1 (Na2S2O3) |
| NADPH + Na2S2O3 + CoA | 6.7 ± 0.08 | 2.93 ± 0.82           | 80 ± 2.5 (NADPH)  |

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*The NAD(P)H oxidation activities were determined by monitoring absorption at 340 nm at 45°C.*

*The apparent K_m was determined and calculated by using NAD(P)H oxidation assays.*
identity in protein sequence to THEMA_RS02810 of *T. maritima*. Other characterized NSR homologs include PF1186 from *P. furiosus* (34% identity) and TK1299 from *T. kodakarensis* (34% identity). The enzyme TK1481 from *T. kodakarensis* was identified as a NAD(P)H oxidase and a NAD(P)H:polysulfide oxidoreductase, as well as a CoA-independent NAD(P)H:S0 oxidoreductase (16, 17). Both PF1186 and TK1299 were identified as NAD (P)H:sulfur oxidoreductases with strictly CoA-dependent activities; additionally, TK1299 had NAD(P)H oxidation activity with O2 as electron acceptor, whereas PF1186 had no oxygen-related activities despite the fact that it is oxygen insensitive. The recombinant THEMA_RS02810 in this research presented some properties different from its archaeal homologs. Unlike TK1299 and TK1481, the NSR of *T. maritima* (THEMA_RS02810) did not show any NAD(P)H oxidation activity with O2 as electron acceptor. Besides, the recombinant THEMA_RS02810 was sensitive to O2. After exposure to air for 30 min at room temperature, it lost its sulfur-reducing activity. Besides, the NSR of *T. maritima* was not strictly CoA-dependent. The recombinant NSR showed a lower sulfur reduction activity without CoA, and the activity was enhanced about four folds in the presence of CoA.

When polysulfide was used as electron acceptor, the recombinant NSR showed a NAD(P)H oxidation activity (0.32 ± 0.11 U/mg for NADH and 0.35 ± 0.08 U/mg for NADPH), and this activity was also enhanced by CoA (1.19 ± 0.22 U/mg for NADH and 1.22 ± 0.32 U/mg for NADPH). The activity was only detected at a high concentration of polysulfide (> 1 mM), where 1 M phosphate buffer (pH 7.0) was used in the reaction to balance pH change. And in the same buffer, the recombinant NSR could hardly reduce S0. Considering that S0 is insoluble in water, and NSR is located in the cytoplasm of *T. maritima*, S0 is not likely a substrate *in vivo*, polysulfide might be a more possible substrate. So, NSR most likely functions as NAD(P)H-dependent polysulfide reductase.

CoA showed a significant stimulation on sulfur reductase activity of NSR. A possible reason is that the thiol group may react with S0 or polysulfide. A NSR homolog from *P. horikoshii* was reported with CoA persulfide and CoA polysulfide reductase activities (18). The NSR of *T. maritima* is likely a sulfane sulfur reductase, and CoA could enhance its access to substrate. However, other compounds with thiols, such as cysteine and glutathione, showed no influence on its activities. The mechanism of CoA enhancement on NSR still needs further study.

**Thiosulfate reductase activity of the sulfur reductase NSR from *T. maritima***. Members of the genus *Thermotoga* can also reduce thiosulfate in addition to S0 (19). Thiosulfate enhanced the growth of *T. maritima*. In the presence of thiosulfate, *T. maritima* grew faster and reached an OD600 of around 1.0 in 16 h (Fig. 1), whereas in the absence of thiosulfate, it only reached an OD600 of around 0.6 in 30 h. Meanwhile, the addition of thiosulfate during the growth of *T. maritima* led to an obvious increase in H2S production (Fig. S3). With NADH as electron donors, thiosulfate reductase activity was observed in *T. maritima* cytoplasmic fractions (Table S1 in the supplemental material), but no thiosulfate reductase activity was detected in the membrane fractions, indicating that thiosulfate reduction occurs in the cytoplasm. Interestingly, NAD(P)H-dependent thiosulfate reductase activities were detected with the purified recombinant NSR (Table 2), and the apparent Km of thiosulfate was low (around 10 μM). Similar to the S0 reductase activity, the thiosulfate reductase activities were enhanced 3-fold by CoA. These results suggest that in contrast to its *P. furiosus* and *T. kodakarensis* homologs, the NSR participates in both S0 reduction and thiosulfate reduction. Previous studies have shown that *P. furiosus* and *T. kodakarensis* could not reduce thiosulfate to H2S (20, 21).

**Transcription analysis of sulfur-reduction-related genes**. Quantitative RT-PCR (qRT-PCR) analysis was used to examine the transcription of the *T. maritima nsr* gene before and after *T. maritima* was exposed to S0 or thiosulfate. We discovered that 20 min after adding S0 to the culture, the transcription level of the nsr gene increased 2.74-fold, and 1 h later, it increased 6.99-fold. These results are consistent with the finding that the sulfur reductase activity of the S0-containing culture was significantly higher than that of normal culture (Table 1), and that H2S generation dramatically increased following sulfur addition (Fig. S3 in the supplemental material). The results
indicate that the *nsr* gene is upregulated in the presence of S\(^0\), and that NSR might be involved in the sulfur-reduction metabolism. Furthermore, the addition of sodium thiosulfate also boosted the transcription level of the *nsr* gene, which increased 3.18-fold after 20 min and 7.50-fold after 1 h.

The transcription levels of other genes were also analyzed by qRT-PCR. Besides the *nsr* gene, *mbxA* (THEMA_RS08165), *yeeE* (THEMA_RS09360), and *rnfC* (THEMA_RS03485) genes were upregulated in the presence of S\(^0\) or thiosulfate. There were more significant transcription level changes in these genes in the presence of thiosulfate (Fig. 3). The *mbxA* gene is located in a transcriptional unit that encodes a 13-subunit membrane-bound oxidoreductase (MBX), which is a homolog of the MBS (membrane-bound sulfane reductase) complex from *P. furiosus*. The MBS complex of *P. furiosus* was recently reported to reduce sulfane to H\(_2\)S and to pump protons to the periplasm using reduced ferredoxin as an electron donor (15). The transcription level of the *mbxA* gene increased 4.36-fold after the addition of S\(^0\). These results suggest that the MBX complex may also participate in sulfur reduction in *T. maritima*.

The *yeeE* gene is annotated to encode a membrane protein YeeE, which reportedly mediates thiosulfate uptake in *E. coli* and *Spirochaeta thermophila* (22). Interestingly, several *dsrE/tusA* homologs are also located in the same transcription unit, which are involved in thiosulfate transport (23). After exposure to S\(^0\), the transcription level of *yeeE* increased slightly; when exposed to thiosulfate, however, the transcription level of *yeeE* increased 42.6-fold. Given that *T. maritima* could reduce thiosulfate in its cytoplasm, the *yeeE/dsrE/tusA* gene cluster of *T. maritima* may play an important role in thiosulfate uptake.

*RnfC* is a subunit of the Rnf complex, a membrane-bound Na\(^+\)-translocating respiratory enzyme that oxidizes reduced ferredoxin and reduces NAD\(^+\). Rnf has been found to function in the energy metabolism of *T. maritima* (24). In this study, the transcriptional change of *rnfC* was insignificant after S\(^0\) addition, whereas in the presence of thiosulfate, the transcription level of *rnfC* was upregulated 15.7-fold, indicating that the Rnf complex plays a key role in energy conservation and balancing reducing equivalents in the presence of thiosulfate. In the presence of S\(^0\), however, the contribution of the MBX complex might be more important than Rnf for metabolism.

**DISCUSSION**

Sulfur is an important element in the lithosphere and biosphere, and microorganisms play a vital role in the sulfur cycle. Sulfate, S\(^0\), and sulfide are the three most important forms in the sulfur cycle. Sulfate reduction is thought to be a key process in the biological sulfur cycle (7, 25). However, in various geothermal environments where
S\textsuperscript{0} is widely distributed, such as deep-sea hydrothermal vents, marine sediments, sulfidic springs, etc., the reduction of S\textsuperscript{0} (vs. sulfate) may be more crucial in the sulfur cycle. Diverse S\textsuperscript{0}-reducing microorganisms have frequently been isolated from these hydrothermal habitats (7, 8, 26). Sulfur-reducing prokaryotes are widespread within the phylogenetic tree of life, especially the thermophilic bacteria and archaea, although the metabolism of S\textsuperscript{0} reduction is still not completely understood. The mesophilic bacterium \textit{W. succinogenes} has been studied as a model organism for sulfur metabolism. The membrane of \textit{W. succinogenes} has an electron transport chain, including a hydrogenase and a molybdenum-bound polysulfide reductase (PsrABC), which oxidizes H\textsubscript{2} and ultimately transfers electrons to polysulfide. During this process, protons are pumped into the periplasm and subsequently utilized in ATP synthesis via chemiosmosis (27, 28). In some thermophiles, like \textit{Acidianus ambivalens} and \textit{Aquifex aeolicus}, membrane sulfur reductase Sre is used for sulfur reduction (29, 30). A group of heterotrophic thermophiles, such as \textit{Thermococcus}, \textit{Pyrococcus}, \textit{Thermotoga}, and \textit{Thermoanaerobacter} species, also reduce S\textsuperscript{0} to H\textsubscript{2}S. These thermophiles are widely distributed and are evident in the sulfur cycle of geothermal environments, such as hot springs, hot vents, and marine sediments. However, unlike autotrophic sulfur reducers, no typical sulfur reductases like Psr or Sre, have been found in these heterotrophic microbes; neither have any cytochromes nor quinones been found. Therefore, sulfur reduction in these species is not likely accomplished by a respiratory system analogous to \textit{W. succinogenes} or \textit{A. ambivalens}. A different mechanism may contribute to the sulfur reduction in these microbes.

In this study, we report a cytoplasmic NSR from \textit{T. maritima} that is similar to the NSR of \textit{P. furiosus}. The NSR of \textit{T. maritima} catalyzed the NAD(P)H-dependent reduction of S\textsuperscript{0}, and transcription of the \textit{nsr} gene increased in the presence of sulfur, suggesting that \textit{nsr} could participate in sulfur reduction. Besides \textit{nsr}, the transcription of \textit{mbxA} increased, suggesting that the MBX complex may be involved in sulfur metabolism. The MBX complex of \textit{T. maritima} was a homolog of the \textit{P. furiosus} MBS complex. The MBS complex is thought to be a key enzyme in the sulfur reduction in \textit{P. furiosus}, which reduces S\textsuperscript{0} with reduced ferredoxin and pumps out H\textsuperscript{+} (15, 31, 32). Thus, we predicted that the MBX complex of \textit{T. maritima} would also function as a sulfur reductase.

Based on these results and on existing knowledge of \textit{T. maritima} metabolism, we proposed a simplified mechanism of sulfur reduction in \textit{T. maritima} (Fig. 4), namely, that only glucose is utilized through the EMP and ED pathways, during which NADH, NADPH, and reduced ferredoxin are generated. NADPH can also reduce NAD\textsuperscript{+} and ferredoxin by the electron-bifurcating Nfn (24, 33). Electrons of these reduced equivalents can be further transferred via several pathways: (i) NADH and reduced ferredoxin are oxidized by the electron-bifurcating [FeFe]-hydrogenase (Hyd) to produce H\textsubscript{2} (34); (ii) NADH or NADPH is used for sulfur reduction with NSR to produce H\textsubscript{2}S; (iii) reduced ferredoxin is used for sulfur reduction with the MBX complex, which is predicted to pump out Na\textsuperscript{+} and generate a Na\textsuperscript{+} potential (Rnf and F\textsubscript{1}F\textsubscript{0}ATPase of \textit{T. maritima} is Na\textsuperscript{+}-dependent (24)); (iv) NADH is used for lactate production (34); and (v) reduced ferredoxin is used to regenerate NADH through the membrane-bound Rnf complex (the Rnf complex in \textit{T. maritima} is a Na\textsuperscript{+}-coupled respiratory reduced ferredoxin:NAD\textsuperscript{+} oxidoreductase) (24).

In the absence of sulfur, the reductants are released mainly by electron-bifurcating Hyd to produce H\textsubscript{2}, as only a small amount of lactate is produced (Fig. S4 in the supplemental material). In the presence of sulfur, the electrons are mainly released through sulfur reduction catalyzed by NSR and MBX instead of H\textsubscript{2} production (Fig. S5). The transcriptional analysis showed that the increase of \textit{rnfC} gene transcription was not evident after S\textsuperscript{0} addition, whereas \textit{mbxA} and \textit{nsr} were significantly upregulated, indicating that MBX rather than Rnf may play a key role in the presence of S\textsuperscript{0}. As more electrons are transferred through the MBX in the process of H\textsubscript{2}S formation than H\textsubscript{2} formation, more ATP is formed by the Na\textsuperscript{+} potential; therefore, \textit{T. maritima} grows better in the presence of sulfur. If all the electrons are released by H\textsubscript{2} production, 0.038 ATP molecules are generated by
chemiosmosis per molecule of glucose (Fig. S4), whereas if all the electrons are released by sulfur reduction, 0.96 molecules of ATP per glucose molecule are generated (Fig. S5) (assume that only glucose is utilized). Therefore, *T. maritima* grows better in the presence of sulfur, especially when the H$_2$ pressure is high, and electrons are not released by hydrogenase.

Interestingly, the NSR of *T. maritima* showed a specific activity of thiosulfate reduction with NAD(P)H that is not found in its archaeal homologs. Thiosulfate is also an important intermediate in the sulfur cycle of anoxic sediments (35), and is involved in sulfur reduction, oxidation, and disproportionation. Thiosulfate reduction is thought to be primarily carried out by sulfate-reducing bacteria in sediments, where thiosulfate functions as an alternative electron acceptor for sulfate respiration. Other bacteria also exhibit thiosulfate reductase activity. The thiosulfate respiratory system has been studied in *Salmonella enterica* and *Pyrobaculum aerophilum*, among other species, wherein thiosulfate was reduced by a membrane-bound thiosulfate reductase complex (36, 37). In geothermal environments, thermophiles with thiosulfate reductase activity are widely distributed. For example, researchers view thiosulfate reduction as an important physiological feature of the order Thermotogales (19). However, neither thiosulfate nor a sulfate respiratory system has been found in *T. maritima*. In this study, the cytoplasmic polysulfide reductase NSR of *T. maritima* showed significant thiosulfate reductase activity. After the addition of thiosulfate, the *nsr* gene and the putative thiosulfate transferase gene *yeeE* were upregulated; H$_2$S was formed dramatically after these genes were expressed. This indicates that thiosulfate reduction may occur in the cytoplasm instead of the membrane, and that NSR may play a role in this metabolism. In the presence of thiosulfate, the *rnfC* gene was significantly upregulated, indicating the involvement of the Rnf complex in the thiosulfate metabolism. Thus, reduced ferredoxin may be oxidized by the Rnf complex for NADH regeneration, and the NADH/NADPH is consumed by NSR to reduce thiosulfate to H$_2$S. Similar to the case of S$^0$, 0.96 molecules of ATP per glucose molecule is expected to be generated by chemiosmosis

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**FIG 4** The proposed model for metabolism of *T. maritima* in the presence of S$^0$ or thiosulfate. Nfn, the electron-bifurcating NAD$^+$-dependent NADPH:ferredoxin oxidoreductase; Hyd, the electron-bifurcating [FeFe]-hydrogenase; Rnf, reduced ferredoxin:NAD$^+$ oxidoreductase; MBX, the putative membrane-bound sulfur oxidoreductase; NSR, NAD(P)H:sulfur oxidoreductase.
if all the electrons are released by thiosulfate reduction (Fig. 36 in the supplemental material) (assume that only glucose is utilized).

A BLAST search was performed to investigate the distribution of NSR in Bacteria and Archaea. Search results revealed that NSR homologs are widely distributed in some groups of bacteria and archaea (Fig. 57 in the supplemental material). Most of these microbes, originally isolated from geothermal environments, are obligate heterotrophic thermophiles with physiological properties similar to *T. maritima* belonging to three groups: Clostridia, Thermotogales, and Thermococcales. Members of these groups have been frequently reported as sulfur-reducing microbes (19); further investigations confirmed that many of them reduce S\(^{0}\) to H\(_{2}\)S. Some can also reduce thiosulfate (Fig. 57) (7, 38). *Caldanaerobacter subterraneus* subsp. *tengcongensis* (previously named *Thermoanaerobacter tengchongensis*), for example, is a heterotrophic thermophile that can ferment carbohydrates to acetate and reduces S\(^{0}\) and thiosulfate; its growth is stimulated by S\(^{0}\) and thiosulfate (39). *C. subterraneus* possesses an NSR homolog (48% identity with the NSR of *T. maritima*) and contains no typical sulfur respiratory system or MBS complex in its genome, suggesting that NSR may play a key role in the strain’s ability to reduce S\(^{0}\) and thiosulfate.

In sum, this study found that the NSR from *T. maritima* functions as a reductase for S\(^{0}\), polysulfide and thiosulfate. The NSR-catalyzed reduction of S\(^{0}\) or thiosulfate may significantly affect the strain’s redox balance and energy metabolism, which also involves membrane-bound Rnf and MBX complexes and electron-bifurcating Hyd and Nfn. Considering the wide distribution of the NSR homolog in thermophilic bacteria and archaea and the ubiquitous S\(^{0}\)/thiosulfate reduction in geothermal environments, NSR may be important in the thermophile-driven sulfur cycle.

**MATERIALS AND METHODS**

**Strains and cultivation.** *T. maritima* (strain DSM 3109) was obtained from DSMZ (German Collection of Microorganisms and Cell Cultures GmbH, Braunschweig, Germany). It was cultured strictly anaerobically at 80°C with 100% gas-phase N\(_{2}\) in modified DSMZ medium 1232. The medium was composed of the following (per L): 0.5 g NH\(_{4}\)Cl; 1.9 g MgSO\(_{4}\) × 7H\(_{2}\)O; 1.6 g KH\(_{2}\)PO\(_{4}\); 1.0 g NaH\(_{2}\)PO\(_{4}\); 26 g NaCl; 10 mL trace element solution; 2.0 g yeast extract; 2.0 g tryptone; 5.0 g soluble starch; 10 mL vitamin solution; and 0.6 g L-cysteine-HCl. Resazurin (0.5 mg/L) was added to the medium as a redox-sensitive indicator.

The trace element solution contained the following: 1.5 g nitritotriacetate acid; 3.0 g MgSO\(_{4}\) × 7H\(_{2}\)O; 0.5 g MnSO\(_{4}\) × H\(_{2}\)O; 1.0 g NaCl; 0.1 g FeSO\(_{4}\) × 7H\(_{2}\)O; 0.18 g CoSO\(_{4}\) × 7H\(_{2}\)O; 0.10 g CaCl\(_{2}\); 0.18 g ZnSO\(_{4}\) × 7H\(_{2}\)O; 0.01 g CuSO\(_{4}\) × 5H\(_{2}\)O; 0.01 g H\(_{3}\)BO\(_{3}\); 0.01 g Na\(_{2}\)MoO\(_{4}\) × 2H\(_{2}\)O; 0.03 g NiCl\(_{2}\) × 6H\(_{2}\)O; 0.3 mg Na\(_{2}\)SeO\(_{3}\) × 5H\(_{2}\)O; and 0.4 mg Na\(_{2}\)WO\(_{4}\) × 2H\(_{2}\)O in 1 L distilled water. The vitamin solution contained the following: 2.0 mg biotin; 2.0 mg folic acid; 10.0 mg pyridoxine-HCl; 5.0 mg thiamine-HCl; 5.0 mg riboflavin; 5.0 mg nicotinic acid; 5.0 mg Ca-pantothenate; 0.1 mg vitamin B12; 5.0 mg p-aminobenzoic acid; and 5.0 mg lipoid acid in 1 L distilled water. The *T. maritima* cells were harvested by centrifugation at the late exponential growth phase (OD\(_{600}\) ≈ 0.4–0.6) and stored at −80°C until use.

*E. coli* strains were cultivated in LB or TB medium for heterologous expression of specific proteins. The LB medium contained 5 g yeast extract, 10 g tryptone, and 10 g NaCl per L. The TB medium contained 12 g tryptone, 24 g yeast extract, 2.3 g KH\(_{2}\)PO\(_{4}\); 12.5 g K\(_{2}\)HPO\(_{4}\); and 4 mL glycerol.

**Preparation of *T. maritima* extracts.** *T. maritima* cells (1 g wet weight) were resuspended with 5 mL of anaerobic 50 mM phosphate buffer (pH 7.4), and 20 mg lysozyme was added. After incubating at 37°C for 6 h, the cells were disrupted by ultrasonication in a vinyl anaerobic chamber (Coy Laboratory Products, Inc.) filled with 95% N\(_{2}\) and 5% H\(_{2}\). After centrifugation at 10,000 × g to remove cell debris, the cell extracts were centrifuged at 100,000 × g, and the supernatant and the membrane fraction were used for enzyme assays.

**Heterologous expression and purification of *T. maritima* sulfur reductase.** A recombinant *E. coli* BL21 strain was constructed to express the putative sulfur reductase gene (THEMA_RS02810, old_locus_tag: TM0379) of *T. maritima* (strain DSM 3109, old_locus_tag: TM0379). The gene was amplified via PCR with the genomic DNA of *T. maritima* as a template. The following primers were used: 5′-CGGATCCATGGAGTACGATGTTGTAG-3′ (forward primer, BamHI restriction site underlined) and 5′-CCCTCGAGTAATACCATTTCAGCCG-3′ (reverse primer, XhoI restriction site underlined). After being digested by the restriction enzymes, the PCR product was ligated to the pET-24b (+) plasmid with T4 ligase. The recombinant plasmid was then transferred into *E. coli* BL21 for heterologous expression.

*E. coli* BL21-TM0379 was cultivated in LB medium (with 50 mg/L kanamycin); 0.3 mM isopropyl β-D-1-thiogalactopyranoside (IPTG) was added when the OD\(_{600}\) reached 0.8. After incubation at 25°C for 16–18 h, the cells were harvested by centrifugation and stored at −20°C for use.

The *E. coli* BL21-TM0379 cells were disrupted anaerobically in a vinyl anaerobic chamber (Coy Laboratory Products, Inc.) by ultrasonication, and the extracts were centrifuged at 35,000 × g for 1 h to remove the debris. The following purification steps were performed anaerobically. The supernatant was
water-bathed at 80°C for 30 min and centrifuged at 35,000 x g; the supernatant was subsequently pumped into a HisTrap HP column (GE Healthcare) for purification. The column was then washed by 25-, 50-, 75-, 100-, 150-, 250-, and 500-mM imidazole solutions (20 mL for each concentration), and the His-tagged protein was eluted at both 100- and 150-mM imidazole solutions. The purified recombinant protein was concentrated by 30K Amicon Ultra-4 centrifugal filters (Merck Millipore, Ireland), resuspended in 50 mM Tris–HCl (pH 7.4), and stored anaerobically at −80°C for subsequent studies.

Heterologous expression and purification of *T. maritima* ferredoxin. The gene encoding ferredoxin (TM0927) was chemically synthesized and cloned to the pETDuet-1 vector. The recombinant plasmid was transferred to *E. coli* C41 (with a pRKISC plasmid to enhance iron–sulfur cluster synthesis). The *E. coli* C41–TM0927 strain was cultured in TB medium (with the addition of 0.2 g/L FeSO₄, 0.2 g/L ferric citrate, 8 mg/L chloramphenicol, 3 mg/L tetracycline, and 8 mg/L carbenicillin). The expression of ferredoxin was induced by the addition of 0.5 mM IPTG. After incubation at 30°C for 8 h, cells were harvested by centrifugation and stored at −20°C.

*E. coli* C41–TM0927 cells were disrupted anaerobically by ultrasonication, and debris was removed by 35,000 x g centrifugation. The supernatant was water-bathed at 80°C for 30 min to denature the *E. coli* proteins. After centrifugation at 100,000 x g for 1 h, the supernatant was collected. The purity of the heterologously expressed ferredoxin was confirmed by SDS-PAGE and UV-visible absorption analysis (Fig. S8). The purified ferredoxin was stored anaerobically at −80°C until use.

**S⁰ reductase activity assays.** The standard sulfur reductase activity assay mixture (200 μL) contained 50 mM phosphate buffer (pH 7.0), 10 mM NAD(P)H, and 2 mg S⁰. When indicated, 100 μM CoA was added to the mixture. For ferredoxin-related activity, a reduced ferredoxin regeneration system was added to the mixture instead of NAD(P)H. The reduced ferredoxin regeneration system contained 30 μM ferredoxin, 10 U *Clostridium pasteurianum* hydrogenase (Cpl), and 100% gas-phase Hz (40). To measure the formation of H₂S, a modified double-vial technique (14) was used. Gas-phase Nz was used, and a 1.5-mL Eppendorf tube was placed in a 10-mL serum bottle and surrounded by 2 mL 1 M NaOH (100% gas-phase Hz was used when the reduced ferredoxin regeneration system was used). The reaction mixture was added into the inner tube, and purified enzymes or cell extracts were injected into the inner tube to start the reaction. After incubating at 80°C (a temperature of 40°C was set when the reduced ferredoxin regeneration system was used) for 5 min; the reaction was stopped, and H₂S was released by the addition of 400 μL sulfuric acid (2 M). The H₂S was captured by NaOH and its amount measured by methylene blue assay, as described previously (41).

**Thiosulfate reductase activity assays.** The thiosulfate reductase activity assay followed the same procedure as the S⁰ reductase activity assay, except that 40 mM thiosulfate was added as an electron acceptor instead of S⁰. The same double-vial system was used to measure the production of H₂S.

**NAD(P)H oxidation with S⁰, polysulfide, or thiosulfate.** The oxidation of NAD(P)H was measured in a cuvette (sealed with a rubber stopper), the assay (400 μL) contained 50 mM (1 M was used to stabilize pH when polysulfide as substrate (42)) phosphate buffer (pH 7.0), 0.24 mM NAD(P)H, and 2 mg S⁰, 10 mM sodium polysulfide (prepared as previously described (42)) or 4 mM thiosulfate. When indicated, 100 μM CoA, glutathione or cysteine was added to the mixture, and gas-phase Nz was used. Purified enzymes or cell extracts were added to start the reaction. The oxidation of NAD(P)H was monitored by a UV–vis spectrophotometer at 340 nm (εS⁰ₐ₅₅ = 6.2 mM⁻¹·cm⁻¹). The assay was performed at 45°C because it was easier to handle the cuvettes at 45°C than at 80°C (the optimum temperature for *T. maritima* growth). Most enzymes increase their activity by a factor of 2 for every 10°C rise in the temperature (the Q₁₀ rule).

All solutions, vials, and cuvettes used were prepared or treated anaerobically. One unit (1 U) was defined as the oxidation of 1 μmol NAD(P)H per minute or the formation of 1 μmol H₂S per minute.

**Quantitative RT-PCR analyses.** *T. maritima* was cultivated in 100 mL normal starch medium, and 0.5 g S⁰ or 30 mM thiosulfate was added at the middle of its exponential growth phase (OD₆₀₀ ≈ 0.2–0.25). Cells were collected for RNA extraction both before and after (20 min and 1 h) S⁰ or thiosulfate addition. Total RNA of *T. maritima* was extracted by the EasyPure RNA Kit (TransGen Biotech Co., Ltd., Beijing, China) and stored at −80°C until use. cDNA was prepared with an All-In-One 5 μL RT MasterMix (Applied Biological Materials Inc., Canada). The 16S rRNA gene of *T. maritima* was selected as the reference. The genes *nrn*, *mbxA*, *yeeC*, and *mfeC* were selected for analyses. Primers for these genes in qPCR were designed by Primer-BLAST (https://www.ncbi.nlm.nih.gov/tools/primer-blast/) (Table S2 in the supplemental material). All qPCR experiments were performed with the LightCycler 480 system (F. Hoffmann-La Roche Ltd., Switzerland) using TransStart Top Green qPCR SuperMix (TransGen Biotech Co., Ltd., Beijing, China). The comparative Ct method was used to analyze the results (43).

**Bioinformatic analyses.** Nucleotide and protein sequences were retrieved from the National Center for Biotechnology Information (NCBI) (www.ncbi.nlm.nih.gov/). The *T. maritima* NSR protein sequence (AAD35465.1) was applied as the query sequence to search for NSR homologs using BLASTp of the NCBI. Protein sequences with identity higher than 40% were selected for further analyses. The neighbor-joining method was used to conduct the multiple sequence alignment and phylogenetic analyses with MEGA 7 (www.megasoftware.net).

**SUPPLEMENTAL MATERIAL**
Supplemental material is available online only.

**SUPPLEMENTAL FILE 1** PDF file, 2 MB.
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