

Two Nitrate/Nitrite Transporters Are Encoded within the Mobilizable Plasmid for Nitrate Respiration of *Thermus thermophilus* HB8

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***Thermus thermophilus* HB8 can grow anaerobically by using a membrane-bound nitrate reductase to catalyze the reduction of nitrate as a final electron acceptor in respiration. In contrast to other denitrifiers, the nitrite produced does not continue the reduction pathway but accumulates in the growth medium after its active extrusion from the cell. We describe the presence of two genes, *nark1* and *nark2*, downstream of the nitrate reductase-encoding gene cluster (*nar*) that code for two homologues to the major facilitator superfamily of transporters. The sequences of NarK1 and NarK2 are 30% identical to each other, but whereas NarK1 clusters in an average-distance tree with putative nitrate transporters, NarK2 does so with putative nitrite exporters. To analyze whether this differential clustering was actually related to functional differences, we isolated derivatives with mutations of one or both genes. Analysis revealed that single mutations had minor effects on growth by nitrate respiration, whereas a double *nark1 nark2* mutation abolished this capability. Further analysis allowed us to confirm that the double mutant is completely unable to excrete nitrite, while single mutants have a limitation in the excretion rates compared with the wild type. These data allow us to propose that both proteins are implicated in the transport of nitrate and nitrite, probably acting as nitrate/nitrite antiporters. The possible differential roles of these proteins in vivo are discussed.**

Nitrate can be used as an alternative to oxygen in respiration by many bacteria and by the mitochondria of certain fungi (1, 17). In most cases, nitrate reduction is the first step in a pathway catalyzed by a series of membrane-bound reductases whose final products are dinitrogen and ammonia (1, 17).

We have recently described the presence of a nitrate reductase operon in the extreme thermophile *Thermus thermophilus* HB8, a bacterium formerly considered a strict aerobe (11). Interestingly, the absence of a nitrite reductase in this bacterium results in the long-term accumulation of nitrite in the medium of anaerobically grown cultures. Accordingly, a very active nitrite export system should function in this bacterium to escape from its high toxicity.

Genes encoding polytopic membrane proteins belonging to the major facilitator superfamily of transporters (9) have been found close to the *nar* gene cluster for all such genes so far sequenced (1, 17). The role of such proteins is still controversial as nitrate/nitrite or H⁺/nitrite antiporters and nitrate/H⁺ symporters (17). There is experimental evidence that the NarK protein, encoded upstream of the *narGHII* operon of *Escherichia coli*, is essentially implicated in nitrite export by using the electrochemical gradient as the energy source (an H⁺/nitrite antiporter) (14). On the other hand, and despite their similarity to the *E. coli* NarK, the NarT and NasA proteins from *Staphylococcus carnosus* and *Bacillus subtilis*, respectively, have been proposed to function as nitrate/H⁺ symporters (5, 10).

Thus, the role of such membrane transporters in different bacteria is still unclear due to the intrinsic difficulties in measuring the transport of these anions and also because of the ability of these bacteria to overcome mutations in the corresponding genes through secondary transporters (17). Moreover, the ability of most denitrifiers to use alternative anaerobic pathways for growth increases the difficulties in analyzing the effects of mutations in the corresponding nitrate/nitrite transporters.

Since *T. thermophilus* HB8 does not have any alternative way for anaerobic growth than nitrate respiration, and keeping in mind that this bacterium lacks a nitrite reductase, a requirement for nitrite extrusion proteins seems crucial for its viability. Thus, any mutation in the putative nitrite transporters should have a strong phenotypic consequence in this bacterium. Here we describe two NarK homologues (named NarK1 and NarK2) encoded by genes downstream of the *narGHII* operon of *T. thermophilus* HB8, one of which (*nark2*) overlaps the replicative origin of the *nar*-carrying conjugative plasmid (13). We demonstrate that mutations in each of the coding genes (*nark1* and *nark2*) have minor effects on the ability to grow anaerobically, whereas double *nark1 nark2* mutants are unable to grow under these conditions. This, and the analysis of nitrite excretion, led us to conclude that both proteins are implicated in nitrate import and nitrite export during anaerobic growth.

MATERIALS AND METHODS

Bacterial strains, plasmids, and growth conditions. *T. thermophilus* HB8 (ATCC 27634) was obtained from the American Type Culture Collection (Rockville, Md.). Its nitrate reductase (NR) deletion derivative (*narGH::kat*) was described previously (12). *E. coli* strains TG1 [*supE* Δ (*nsdM mcrB*) 5(*r_K⁻ m_K⁻ mcrB*) *thi* Δ (*lac-proAB*) F' (*traD36 proAB⁺ laqI^qZ Δ M15*)] and DH5 α F' [F' *supE44* Δ (*lacZYA-argF*)U169 Φ 80*lacZ* Δ M15 *hsdR17 recA1 endA1 gyrA96 thi-1 relA1*] (Bethesda Research Laboratories, Gaithersburg, Md.) were used as hosts

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for plasmid construction. Plasmids pNIT5 and pNIT9 (12) were pUC119 derivatives containing the *narK1* and *narK2* genes, respectively. Plasmid pKT1 was the source of the *kat* gene, encoding thermostable resistance to kanamycin (7).

Aerobic growth conditions for *T. thermophilus* were attained by incubation at 70°C in a nitrate-free rich medium (12) with vigorous shaking. For NR induction, cells were grown in the same medium containing 40 mM KNO₃ without stirring. For anaerobic growth, cells grown under aerobic conditions up to an optical density at 550 nm (OD₅₅₀) of 0.05 were divided into 10-ml aliquots inside 15-ml tubes filled to the top with mineral oil (12). Plates were incubated at 70°C in a water-saturated atmosphere. For liquid and solid selection, kanamycin (30 µg/ml) was added. *E. coli* strains were grown at 37°C in Luria-Bertani (LB) medium (8) or in LB agar plates. Ampicillin (100 µg/ml) or kanamycin (30 µg/ml) was added when needed.

Mutant isolation. Insertional mutagenesis was done through electroporation of *T. thermophilus* HB8 (3) with *Hind*III-linearized forms of defined plasmids, followed by selection on kanamycin-containing plates.

Plasmid pNIT9Kkat1 was used for the inactivation of *narK1*. It was obtained by insertion of the *kat* cassette (7) into the *Kpn*I site of pNIT9 (12), interrupting the sequence of the *narK1* gene at a position corresponding to amino acid 41 of the encoded protein. Isolation of the *narK2::kat* mutant was achieved by transformation with plasmid pNIT5kat1. In this plasmid, the *kat* gene replaced a *Sma*I fragment internal to the *narK2* gene, resulting in a truncated protein of 200 amino acids. To obtain double *narK1 narK2::kat* mutants, plasmid pNIT9Kkat1 was digested with *Eco*RI. After treatment with the Klenow fragment of DNA polymerase I from *E. coli*, the linearized plasmid was partially digested with *Sma*I and religated. The new plasmid, pNIT9Kkat2, was used to produce a deletion of positions corresponding to encoded amino acids 41 of NarK1 to 304 of NarK2.

Genetic analysis. General methods were used for DNA manipulation (15). Southern blots of total DNA from putative mutants digested with the appropriate enzyme(s) were hybridized with fluorescein-labeled oligonucleotides O-70 (5'CGGAGAGGAAGATGCCG3'), which hybridized to the sequence of *narK2*, and Okat-2 (5'GAAACTTCTGGAATCGC3'), directed against the 3' region of the *kat* gene, and revealed with the ECL kit (Amersham Ibérica SA). DNA was sequenced by automatic methods (Applied Biosystems) with synthetic primers (Isogen Bioscience, Maarsse, The Netherlands). Partial sequences were assembled with the software of the University of Wisconsin Genetics Computer Group (4). DNA amplification from colonies of the putative mutants was developed in an MJ Research minicycler (MJ Research Inc., Watertown, Mass.). Oligonucleotides O-28 (5'CACCCTCATGTTCCGCC3'), O-54 (5'CCACCCTCTCCTCCTC3'), O-65 (5'CGGGCCGATGAACTTGG3') and O-70 (see above) were used for the amplification of specific fragments. Their approximate hybridization sites are labeled in Fig. 2A.

NR activity and nitrite analysis. For induction of the NR, cells of *T. thermophilus* HB8 were grown at 70°C in a shaker bath to an OD₅₅₀ of 0.5. After addition of potassium nitrate (40 mM), the cultures were incubated at 70°C without stirring for an additional 2 h. Cells were then recovered by centrifugation, washed twice with phosphate buffer (50 mM, pH 7) by centrifugation (10,000 × g, 5 min, room temperature), and disrupted by sonication at a cell density of ~24 OD₅₅₀ units/ml. Intracellular concentrations of nitrite were measured in the above cell extracts by assuming a cell volume and a ratio of OD₅₅₀ to cell mass similar to those of *E. coli* (10⁻¹² ml/cell and 10⁹ cells per OD₅₅₀ unit per ml). The NR activity was measured as described before (12) after 5 min of incubation at 80°C with methyl viologen as the electron donor (16) and nitrate as the electron acceptor. The nitrite excreted was measured in cell-free samples of the growth medium.

RESULTS

Sequence of the region downstream of the *narGHJI* operon.

Analysis of the sequence downstream of the *narGHJI* operon from *T. thermophilus* HB8 (accession number AJ237974) revealed the presence of two open reading frames encoding 435 and 443 amino acids, both of them preceded by putative Shine-Dalgarno sequences. Due to their similarities to the sequences for nitrite transporters found close to *nar* operons from other bacteria (see below), the corresponding encoding genes were named *narK1* and *narK2*, respectively.

The putative translation ATG start codon of *narK1* overlaps the last codon of *narI*, the gene encoding the cytochrome *b* (γ subunit) component of the NR. Thus, a translational coupling from a common mRNA seems very likely. By contrast, 22 bp separate the last codon of *narK1* and the ATG start codon of *narK2*, keeping the possibility of differential expression between the proteins open. However, the absence of putative transcription terminator sequences between the genes suggests that *narK2* could be cotranscribed within the same mRNA.

Secondary-structure predictions support a polytopic integral

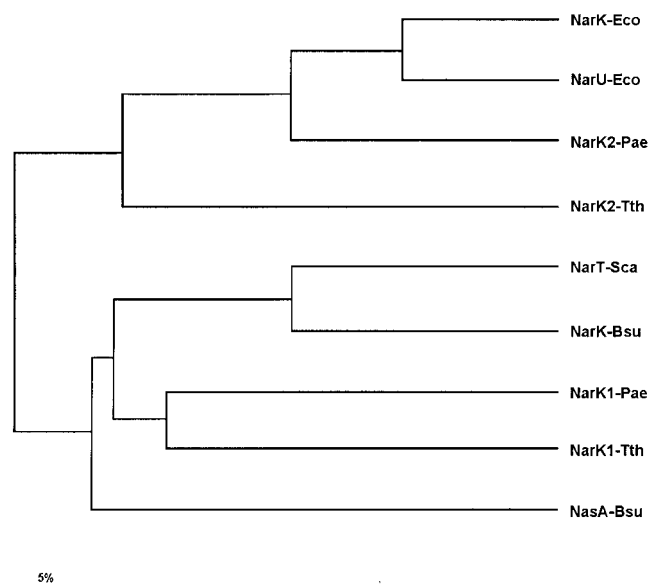


FIG. 1. Average-distance tree for NarK proteins. The sequences of the putative nitrate and nitrite transporters were aligned through the CLUSTAL program, and the average distance was represented through the JALview editor. In addition to the NarK1 and NarK2 proteins from *T. thermophilus* HB8 (Tth), the proteins used in the alignment included NarK (P10903) and NarU (P37958) from *E. coli* (Eco), NarK1 and NarK2 from *P. aeruginosa* (accession no. Y15252) (Pae), NarT from *S. carnosus* (accession no. U40014) (Sca), and NarK (P46907) and NasA (P42432) from *B. subtilis* (Bsu).

membrane nature for NarK1 and NarK2, with 12 putative α -helix domains spanning the cytoplasmic membrane. No typical general secretory pathway-dependent signal peptide was present in any of them. Comparison of their sequences with those in the data banks revealed similarities to membrane transporters belonging to the major facilitator superfamily of proteins (9). High scores were found when NarK1 and NarK2 were compared with proteins implicated in the extrusion of nitrite and/or nitrate transport from different bacteria. As shown in Fig. 1, the calculation of the average-distance tree from the alignment of the NarK1 and NarK2 sequences with those from proteins implicated in the transport of nitrate and/or nitrite in different bacteria revealed a differential grouping of both proteins. Whereas NarK1 groups with NarT and NasA, two proteins suggested to participate in nitrate transport in *S. carnosus* (5) and *B. subtilis* (10), respectively, NarK2 groups with the nitrite extrusion proteins NarK and NarU from *E. coli*.

Consequently, a different role for these proteins is suggested from these comparisons, in which NarK1 could be responsible for nitrate transport and NarK2 for the extrusion of nitrite. This putative difference in roles is also supported by the low identity (30%) found between the two proteins.

The presence of two in-tandem genes encoding NarK homologues (*narK1* and *narK2*) has been identified close to the *nar* operon within the genome sequence of *Pseudomonas aeruginosa* (accession number Y15252) and, more recently, in *Pseudomonas stutzeri*, where they have been named *narK* and *narC* (6). Interestingly, inclusion of the sequences of NarK1 and NarK2 from *P. aeruginosa* in the above comparison revealed a clustering similar to that of the NarK1 and NarK2 proteins from *T. thermophilus* HB8. Moreover, the NarK1 proteins from both organisms were 43% identical and 54.5% similar, whereas their corresponding NarK2 proteins were 41%

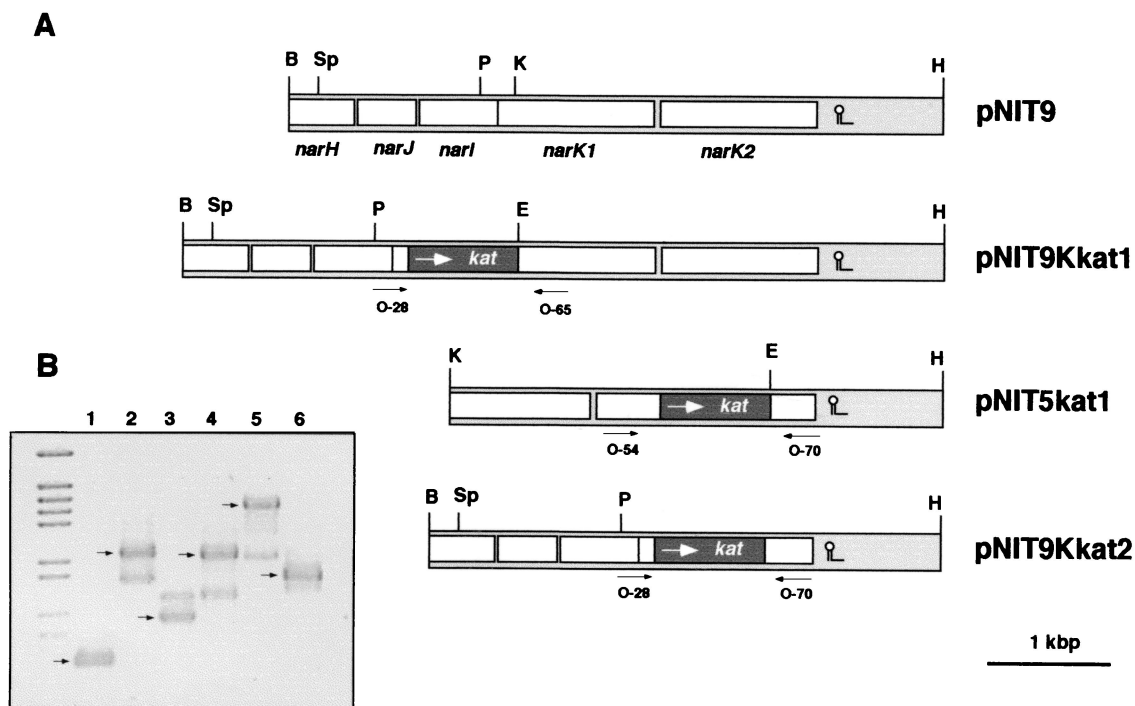


FIG. 2. Isolation and analysis of *narK::kat* mutants. (A) Restriction maps of the plasmids used to get single and double *narK::kat* mutants. Enzymes: B, *Bam*HI; E, *Eco*RI; H, *Hind*III; K, *Kpn*I; P, *Pst*I; Sp, *Sph*I. (B) PCR analysis of *narK::kat* mutants. PCR amplification products from the DNA of single *narK1* (lane 2) and *narK2* (lane 4) and double *narK1 narK2* (lane 6) mutants were compared with those obtained from the DNA of the parental strain (lanes 1, 3, and 5). Primers used for amplification: lanes 1 and 2, O-28 and O-65; lanes 3 and 4, O-54 and O-70; lanes 5 and 6, O-28 and O-70. Arrows indicate the product expected from the corresponding amplifications. The approximate positions at which these primers hybridize are shown in panel A.

identical and 51.3% similar. Thus, although in *P. aeruginosa* the *narK1* and *narK2* genes are located upstream of the corresponding *narGHJ* cluster, the high similarity found between these proteins supports similar roles for them in both organisms.

Insertional inactivation of *narK1* and *narK2*. The analysis described above suggested a role for NarK1 and NarK2 in nitrate transport and nitrite extrusion, respectively, suggesting that both proteins were required for nitrate respiration. To analyze this hypothesis, we isolated individual *narK1::kat* and *narK2::kat* mutants as well as a double *narK1K2::kat* mutant by using the plasmids shown in Fig. 2A (see Materials and Methods for construction details). After transformation and selection for kanamycin resistance, the presence of the expected mutations was confirmed by DNA amplification with primer pairs that hybridized at the approximate positions shown in Fig. 2A. As shown in Fig. 2B, the use of primers O-28 and O-65 allowed the amplification of a 0.4-kbp fragment in the wild-type strain (lane 1), whereas in the *narK1::kat* mutant, this fragment was replaced by a 1.3-kbp fragment as a result of insertion of the *kat* gene (lane 2). In the *narK2::kat* mutant, the use of primers O-54 and O-70 resulted in amplification of the expected 1.29-kbp fragment (Fig. 2B, lane 4) instead of the 0.69-kbp DNA obtained from the wild type (lane 3). Finally, the 2.2-kbp fragment amplified from the wild type with primers O-28 and O-70 (Fig. 2B, lane 5) was replaced by a 1-kbp fragment in the double mutant as a consequence of replacement with the *kat* cassette of most of the sequence from both genes (lane 6). These results were further confirmed by Southern blot analysis (not shown).

Phenotypic analysis of *narK::kat* mutants. As could be expected, the growth of single and double mutants under aerobic

conditions was indistinguishable from that of the wild-type strain (not shown), indicating that neither of these genes is required for the aerobic metabolism of the bacterium.

Under anaerobic conditions, however, single mutants grew at slightly lower rates than the wild type, reaching 5 to 10% lower cell mass densities than the wild type (Fig. 3A). By contrast, the double *narK1K2::kat* mutant was unable to grow anaerobically, a behavior indistinguishable from that of the *narGH::kat* mutant used as a negative growth control (Fig. 3A).

In order to analyze whether the lack of anaerobic growth in the double mutant was due to a defect in nitrate transport or to the absence of an active NR, induction experiments of this enzyme were undertaken. The results showed that the NR activity in single and double mutants was similar to that in the wild type (from 300 to 600 nmol of nitrite produced per min per OD₅₅₀ unit). Thus, the inability to grow anaerobically had nothing to do with a defect in NR induction, suggesting instead a defect in nitrate and/or nitrite transport.

To analyze this, cells were induced as above, washed twice by centrifugation, resuspended to an OD₅₅₀ of 0.5 in nitrate-free medium, and separated into 2-ml aliquots. These aliquots were subsequently incubated at 70°C with increasing concentrations of nitrate, and the nitrite excreted was measured at different times. In Fig. 3B, the amount of nitrite excreted by each mutant after 10 min of incubation is plotted against the amount of nitrate added. As can be observed, the amount of nitrite excreted by the single mutants reached a plateau far below that of the wild type. By contrast, nitrite was not detected in the medium when the double mutant was assayed under these conditions.

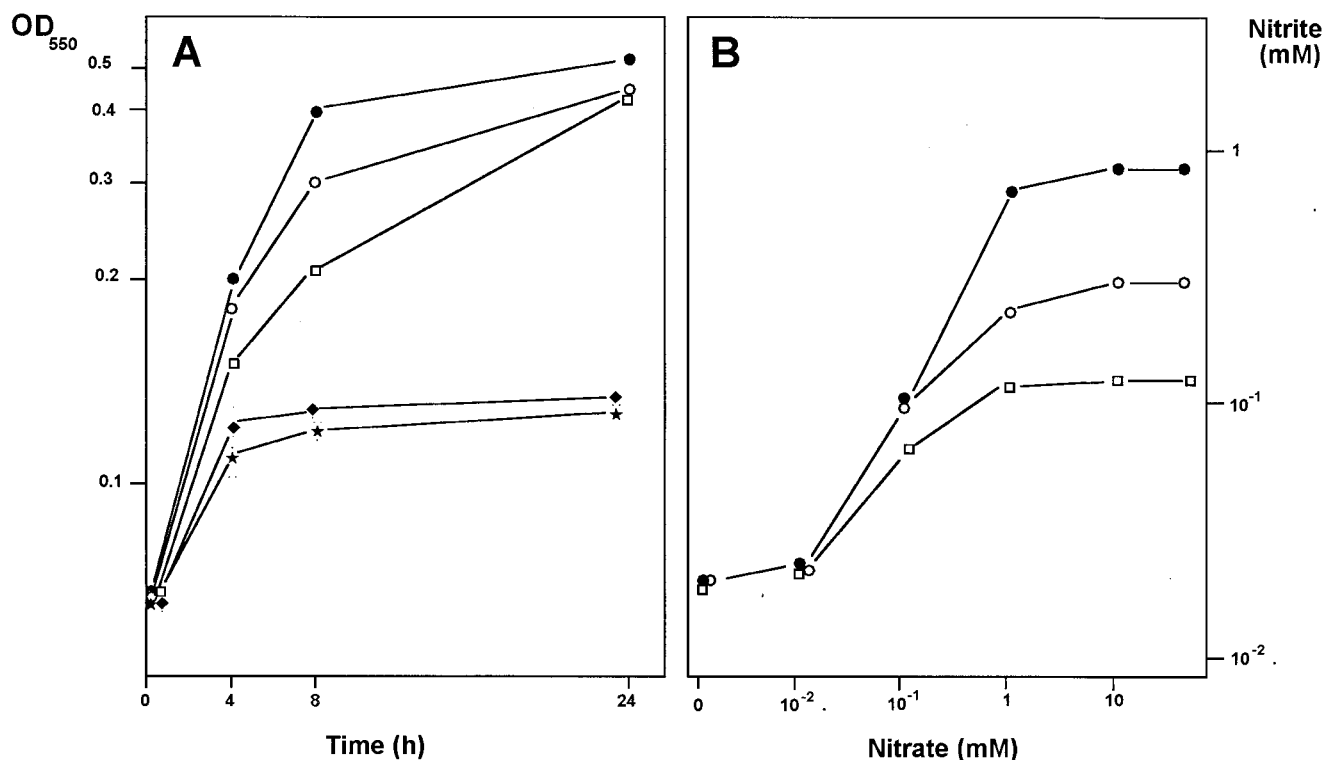


FIG. 3. Anaerobic growth and nitrite export of *narK::kat* mutants. (A) Anaerobic growth. Growth curves under anaerobic conditions were performed as described in Materials and Methods. Tubes were taken at the indicated times to measure the OD₅₅₀. (B) External concentration of nitrite at different nitrate concentrations. Samples from induced cultures of single and double *narK* mutants were incubated in preheated culture medium containing increasing concentrations of nitrate. After 10 min, the nitrite produced was measured and plotted against the external concentration of nitrate (see the text for details). Symbols: ●, wild type; ○, *narK1::kat*; □, *narK2::kat*; ◆, *narK1K2::kat*; ★, *narGH::kat*.

DISCUSSION

The *narK1* and *narK2* genes described in this study are predicted to encode proteins belonging to the major facilitator superfamily of transporters. Proteins from this family have a broad substrate spectrum that includes H⁺, sugars, and antibiotics, and its members share 12 membrane-spanning α -helix and diverse sequence motifs (9). This fact, and the location of the genes downstream of the *narGHJI* cluster, suggested that NarK1 and NarK2 were involved in the transport of nitrate and/or nitrite. The average-distance tree shown in Fig. 1, in which each protein clusters in a different group, suggested a role in nitrate transport for NarK1 and a more specific role in nitrite extrusion for NarK2.

The results shown in Fig. 3A demonstrate that a double mutation in *narK1* and *narK2* abolishes the ability of *T. thermophilus* HB8 to grow anaerobically. Thus, either the substrate (nitrate) or the product (nitrite), or both, has no alternative way to cross the membrane than by using the NarK1 or NarK2 protein. On the other hand, as the presence of either of these proteins suppresses this defect in anaerobic growth, it follows that they share at least one of these functions. Whether this function is nitrate transport or nitrite excretion cannot be deduced from this experiment. However, a detailed analysis of our present results led us to support a nitrate/nitrite antiporter capability for both proteins.

The arguments in favor of this interpretation are various. First, if both proteins shared only the ability to extrude nitrite, the presence of an alternative way to bring nitrate into the cell should result in a dramatic intracellular accumulation of nitrite because of the presence of a normal level of NR. Most prob-

ably, such levels of intracellular nitrite would be lethal to the cell. In fact, in the double mutant, intracellular nitrite remains at concentrations similar to or even lower than that of the wild type (0.5 to 1 mM) in all the experiments in which it was checked. Thus, we concluded that there is not an alternative way to transport nitrate into *T. thermophilus* HB8 than the NarK1/NarK2 proteins. In consequence, both proteins should have the ability to act as nitrate transporters to the cell.

On the other hand, the inability of the double mutant to excrete a detectable amount of nitrite at the highest nitrate concentration used (40 mM) in the experiment shown in Fig. 3B could be expected from the absence of nitrate transporters. In addition, the plateau of nitrite secretion reached by each single mutant could reflect the existence of a limiting step only in nitrate transport. Alternatively, it could be related to a limitation in the excretion of nitrite that could be expected from the absence of nitrite transporters outside the *nar* cluster. In this sense, the fact that the *nar* cluster of *T. thermophilus* HB8 is located within a self-mobilizable element, along with the small size of its chromosome, strongly argues against the existence of nitrite extrusion transporters outside this genetic element. In fact, no homologous genes could be detected either with a labeled probe of *narK1* and *narK2* in Southern blot analysis or by comparison with the unfinished genome sequence of *T. thermophilus* HB27, a closely related aerobic strain to which the *nar* cluster can be transferred and expressed. Thus, although the inability of the double mutant to excrete a detectable amount of nitrite (Fig. 3B) does not exclude it, the putative existence of an alternative nitrite extruder other than the NarK1/NarK2 proteins seems unlikely.

As noted above, the *T. thermophilus* HB8 chromosome is quite small (1.8 Mbp) (2), and consequently, genetic redundancy is rare (e.g., there are only two copies of DNA encoding rRNA). Thus, a good but yet-unknown reason should justify the presence of two genes encoding functionally redundant proteins in the *nar* cluster for anaerobic respiration of *T. thermophilus*. Despite the above discussion, the low similarity between the two proteins suggests different roles for each protein in the natural environment in which these microorganisms live. For instance, the extrusion of nitrite and the transport of nitrate would require the proton motive force at low nitrate concentrations. In this scenario, one of the proteins could have the ability to function as an H⁺/nitrate symporter and the other as an H⁺/nitrite antiporter, and only at higher concentrations of nitrate could they function as nitrate/nitrite antiporters. Alternatively, the presence of two enzymes with redundant enzymatic activities could be related to putative differential expression between them. In this sense, the distance between the genes (20 bp) and the existence of a T-rich sequence overlapping the C-encoding region of *narK1* could be related to differential expression of each protein.

Interestingly, the two *narK* genes identified upstream of the nitrate respiration cluster of *P. aeruginosa* encode proteins with high similarity to those described here from *T. thermophilus* (Fig. 1). Moreover, this similarity is conserved at the DNA sequence level, not only between the *narK* genes but also along many stretches of the whole *nar* cluster. Keeping in mind the plasmidic nature of the genetic element that encodes anaerobic respiration in *T. thermophilus* HB8 and the ubiquity, respiratory character, and similarity in codon usage of *Thermus* spp. and *Pseudomonas* spp., it is tempting to speculate about a common origin for both groups of genes. Meanwhile, future work with *P. aeruginosa* would confirm the functional relationship between the tandemly organized *narK* genes of these bacteria.

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