

Respiratory nitrate reductase metabolic pathway in *Corynebacterium pseudotuberculosis* biovar *Equi*

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Abstract

Corynebacterium pseudotuberculosis can be classified in two biovars, based on their ability to convert nitrate to nitrite. The nitrate-positive biovar is *Equi*, which causes ulcerative lymphangitis in equines, while the nitrate-negative biovar is known as *Ovis*, which is the etiologic agent of caseous lymphadenitis in small ruminants. Both diseases are globally distributed and cause large economic losses to goat, sheep, horse and cattle farmers. The nitrate reduction is associated with the bacterium's ability to breathe in the absence of oxygen and having two different metabolic pathways, (1) respiratory nitrate reductase and (2) dissimilatory nitrate reduction. In the first pathway, the denitrification process takes place where the nitrate is sequentially reduced to nitrite, nitric oxide, nitrous oxide, and finally to dinitrogen. In the second pathway nitrate is directed converted into ammonia, which is secreted from the cell, this process can be performed by organisms with the *nrf* gene. This is a less common method of nitrate reduction than denitrification in most ecosystems. Prokaryotic nitrate reductases include a class of assimilatory enzymes and two classes of respiratory enzymes, all contain a guanylate molybdenum cofactor, but differ in their substructures, cellular location, and requirement for cofactor. Variability among enzyme is also found into the classes. Aiming to discover the molecular mechanisms to related the ability bacteria nitrate reduction, 19 complete genomes of *C. pseudotuberculosis* were analysed. To identify the nitrate pathways, genes of these pathways were analyzed using databases such as BioCyc, ENZYME, KEGG. For the analysis of metabolic pathways Pathway tools were used. Were done in the blast database Uniprot and protein domain analysis through INTERPROSCAN. Genome analysis revealed that *C. pseudotuberculosis* biovar *Equi* possess *narKGHJI* gene clusters that are similar to the *narK* gene and *narGHJI* operon of *Escherichia coli*. The gene encodes a nitrate/nitrite transporter, whereas the operon encodes a respiratory nitrate reductase (*NarGH*) and one specific chaperone (*NarJ*) required for insertion of Mo-bisMGD cofactor in *NarG*. The enzymes that are involved in electron transport chain are also identified by in silico methods. Findings about pathogen metabolism can contribute to the identification of relationship between nitrate reductase and the *C. pseudotuberculosis* pathogenicity, virulence factors and discovery of drug targets.

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