Cloning and characterization of a gene encoding a protein disulfide isomerase from Neospora caninum

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A gene encoding a protein disulfide isomerase (PDI) was isolated from a *Neospora caninum* cDNA expression library. The nucleotide sequence of the cDNA clone revealed the presence of an ORF of 1,416 bp, which encoded 461 amino acids, showing a high degree of homology to *Toxoplasma gondii* PDI. The gene was cloned into a pGEX vector and expressed in *E. coli* as a GST fusion protein. The *Nc*PDI was detected in *N. caninum* tachyzoite lysate and ES products with a molecular weight of 50 kDa. IgA antibody in 58.0% of individual cattle tear samples recognized both the recombinant and native *Nc*PDI, which suggests that the PDI-specific antibody may be involved in defense against parasites. In addition, PDI-specific inhibitors showed significant inhibitory effect on the growth of *N. caninum* tachyzoites. The purified recombinant *Nc*PDI demonstrated biological activity in vitro by catalysis and refolding of reduced RNase, and assisted in the recovery of native from denatured lysozyme. These findings indicate that the *Nc*PDI possesses specific-PDI enzymatic activity and it offers a putative target for prevention and chemotherapy of neosporosis.