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Original Article

Recombinant *Cryptosporidium parvum* p23 as a Candidate Vaccine for Cryptosporidiosis

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Abstract:

Background: *Cryptosporidium parvum* is a ubiquitous protozoan, which develops within the microvillous membrane of enterocytes in a wide variety of vertebrates, including man. Cryptosporidiosis is an important parasite causing severe diseases in the immunodeficient people especially AIDS patients. Cryptosporidiosis has been also reported as a com-mon serious primary cause of outbreaks of diarrhea in newborn calves. The aim of this study was to confirm that P23 was an immunogenic antigen in domestic isolates of *C. parvum*.

Methods: We isolated cryptosporidial oocysts from the naturally infected calves. The oocysts were then purified and characterized as *C. parvum* by nested PCR. To obtain the recombinant P23 protein, we isolated the mRNA from oocyst of *C. parvum*, and synthesized the cDNA. The cDNA was then amplified using specific primers for P23 gene.

Results: Sequencing of PCR product showed 100% homology to the known P23 sequences in GenBank. The double strand P23-cDNA was then cloned in pGEX-5X-2 expression vector and P23-recombinant protein was prepared. West-ern blot analysis of recombinant P23 showed that it could be recognized by the positive *C. parvum* serum. Furthermore, serum from immunized goat with the recombinant P23 protein also recognized a protein band with approximately 23 kDa in lysates prepared from the oocytes.

Conclusion: Since P23 is an immunodominant surface glycoprotein expressed in the early phase of infection and the immuno-genic epitopes are found in its residual chain of amino acid sequence, the recombinant P23 could be recom-mended as a favorable candidate for vaccination against *C. parvum* infection.

Keywords:

Cryptosporidium parvum , *Expression vector pGEX-5X-2* , *recombinant protein P23* , *PCR* , *RT-PCR* , *Western blot* , *Dot Blot*

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