



Inhibition of *Leishmania major* PTR1 Gene Expression by Antisense in *Escherichia coli*

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Abstract

Background: Protozoa related to Trypanosome family including *Leishmania*, synthesize enzymes to escape from drug therapy. One of them is PTR1 that its enzymatic activity is similar to dihydrofolate reductase (DHFR). Dihydrofolate reductase - thymidylate synthase has a major role in DNA synthesis, if it is inhibited, the result would be the death of parasite. Since PTR1 activity is similar to DHFR, causes the decrease of inhibition effect of drug. The aim of this study was inhibition of Iranian *L. major* PTR1 expression with mRNA antisense in prokaryotic system as an approach to appear of the drugs therapeutic effects more.

Methods: PTR1 gene was ligated to pACYCDuet-1 and pcDNA3 plasmids as sense and antisense plasmids, respectively. Simultaneously transfer of sense and antisense plasmids was done in *E. coli* strain M15. SDS-PAGE and western blot analysis were carried out to analyze the expression.

Results: Sense and antisense plasmids were prepared and confirmed by restriction analysis and PCR then simultaneously transfer of them was done. SDS-PAGE and western blot analysis showed PTR1 gene was inhibited by mRNA antisense in bacterial cells.

Conclusion: Expression of PTR1 gene in sense plasmid was inhibited successfully by antisense plasmid.

Keywords: PTR1, *Leishmania major*, Antisense plasmid, Dihydrofolate reductase, Expression

Introduction

Leishmania major protozoa are the causative agent of human zoonotic cutaneous leishmaniasis. Studies have shown that the growth of protozoa related to Trypanosome family depends on (folate-pteridine) (1). Protozoa related to Trypanosome family including *Leishmania* synthesizes enzymes to escape from drug therapy. One of these enzymes is PTR1 (1-9). The enzymatic activity of PTR1 is similar to dihydrofolate reductase (DHFR) (10, 11). DHFR

plays a role in activating the enzyme of thymidylate synthase (TS) that catalyzes the conversion of deoxyuridylate in to deoxythymidylate and another role of DHFR is the reduction of dihydrofolate (H₂-folate) to tetrahydrofolate (H₄-folate) (9); the recent product is co-factor of thymidylate synthase (9-11). Thus DHFR-TS has a major role in DNA synthesis, if it is inhibited, according to the mentioned reactions of DNA synthesis, the result would be the death of

parasite (1). Genesis drugs on metabolism in mediating cell proliferation are important affect; one of the mechanisms of drug is inhibition of DHFR-TS (1). Since PTR1 activity is similar to DHFR causes the decrease of inhibition effect of drug (10, 11). On the other hand, PTR1 causes synthesis and reduction of pteridine that increases the resistance of parasite against oxidative stress-related drug or reactions of the host (1, 2, 12). Because of production of enzymes by *Leishmania* including PTR1, the drug efficacy has been decreased and no response to treatment is observed, therefore, it is suggested that in addition of prescription of anti-parasite drugs, these enzymes inhibit in such way that the therapeutic effects of drugs appear more (1).

The aim of this study was inhibition of Iranian *L. major* PTR1 expression with antisense RNA in prokaryotic system.

Materials And Methods

Sense and antisense plasmids construction

In order to transfer of sense and antisense plasmids in one bacterial cell, two different plasmids with two different origins of replication, and two different antibiotic resistance genes were needed. The position of restriction enzyme cut sites used on the plasmids was in two different directions.

BamH I and Hind III were considered on the sense plasmid, as the expression, PTR I protein was produced. In addition, BamH I and KpnI were considered on antisense plasmid, in this case gene order placement is in the reverse mode.

For this purpose, plasmids pACYCDuet-1 and pcDNA3 were used for the sense and antisense plasmids, respectively.

Plasmid pQE-ptr that has been described previously (13), was digested by Hind III and BamH I to release the PTR1 gene. The fragment was cloned into Hind III and BamH I digested pACYCDuet-1. This vector was transformed in *E. coli* strain M15 then extracted and confirmed by restriction analysis and PCR.

For preparing of antisense plasmid, pQE-ptr (13) was digested by kpn I and BamH I to release the PTR1 gene. The fragment was cloned into pcDNA3, which was lineared by kpn I and

BamH1. Recombinant plasmid was transformed in *E. coli* strain M15.

Recombinant plasmids were extracted and confirmed by restriction analysis and PCR.

Simultaneously transformation of sense and antisense plasmid in one bacterial cell

The *E. coli* strain M15 was transformed with pACYCDuet-ptr and selected on Luria Bertani agar containing 30 µg/ml chloramphenicol. These cells were transformed again with pcDNA-Rptr and selected on Luria Bertani agar containing 50 µg/ml ampicillin and 30 µg/ml chloramphenicol.

Transformation of pACYCDuet-ptr and pcDNA-Rptr in the *E. coli* strain M15 was done separately as control and selected on Luria Bertani agar containing selective antibiotics.

E. coli strain M15 was susceptible to used selective antibiotics.

Gene expression

Gene expression was done as previously described (13, 14) by a little modification. Briefly, the transformant *E. coli* strain M15 with both of sense and antisense plasmids was inoculated into 3 ml culture tube containing X medium (1.2% bacato trypton , 2.4% yeast extract , 0.04% glycerol , 1% M9 salts) (M9 salt containing 6.4% Na₂HPO₄ - 7H₂O, 1.5% KH₂PO₄ , 0.025% NaCl , 0.05% NH₄Cl) and allowed to grow at 37°C in a shaker incubator at 200 rpm overnight. The day after, it was inoculated in to 50 ml flask and allowed at 37°C in a shaker incubator at 200 rpm. The culture in the logarithmic phase (at OD 600 = 0.6) was induced with 0.8mM isopropyl-B-D-thiogalactopyranosid (IPTG) for 5 hours. After induction, cells were withdrawn and analyzed by 10% SDS-PAGE (15). Induced and uninduced the transformant bacterial cells with pcDNA-Rptr and pACYC Duet-ptr were analyzed in parallel.

Western blot analysis

For immunoblotting, proteins resolved by SDS-PAGE were electrophoretically transferred onto nitrocellulose membrane (13). Then UV cross linker was used for protein fixation. The membrane was blocked with 3% BSA (Bovine Serum

Albumin) at room temperature and washed twice with TBS (Tris-Buffered Saline) after 1 h. After that incubated for 1 h at 37 °C with the rabbit anti-PTR1 antiserum at a 1:1000 dilution as the primary antibody. The membrane was washed three times with TBS-T (TBS-TWEEN 20) and incubated for 1 h at 37 °C with the goat anti-rabbit IgG HRP conjugate solution at a 1:5000 dilution as secondary antibody (13, 16). Antibody binding was visualized using calorimetry with diaminobenzidine (DAB) and H₂O₂.

Results

Preparation of sense and antisense plasmids:

Plasmid pQE-PTR1 that was the source of the *L. major* PTR1 gene (13) was digested with restriction enzymes and released the PTR1 gene. The PTR1 gene was ligated to pACYCDuet-1 and pcDNA3 then confirmed by restriction analysis and PCR (Fig. 1, Fig. 2, Fig. 3, Fig. 4) and named pACYC Duet-ptr and pcDNA-Rptr, respectively.

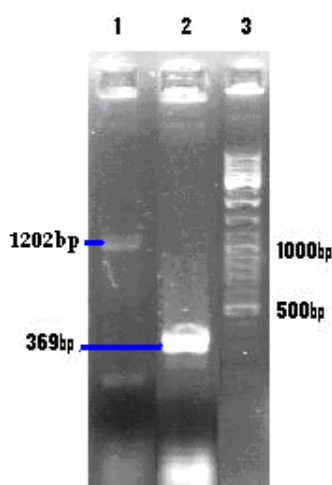


Fig. 1: Electrophoresis of PCR product on 1% agarose gel

Lane 1: The 1202 bp as PCR product of pACYCDuet-ptr

Lane 2: The 369 bp as PCR product of pACYC-Duet-1

Lane 3: 100 bp DNA ladder marker

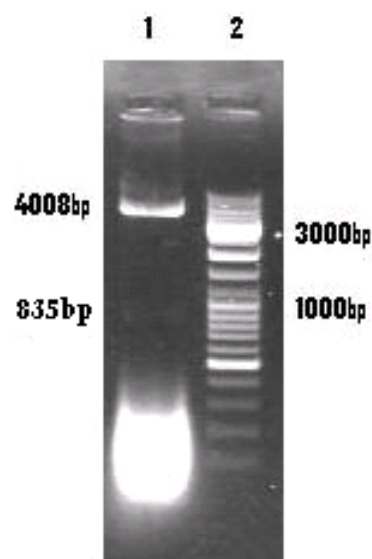


Fig. 2: 1.5% Agarose gel electrophoresis

Lane 1: Digested pACYCDuet-ptr by BamH I and Hind III

Lane 2: 100 bp DNA ladder marker

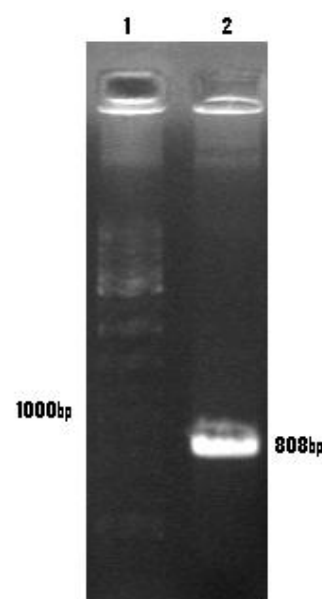


Fig. 3: Electrophoresis of PCR product on 1% agarose gel

Lane 1: 100 bp DNA ladder marker

Lane 2: The 808 bp as PCR product of pcDNA-Rptr

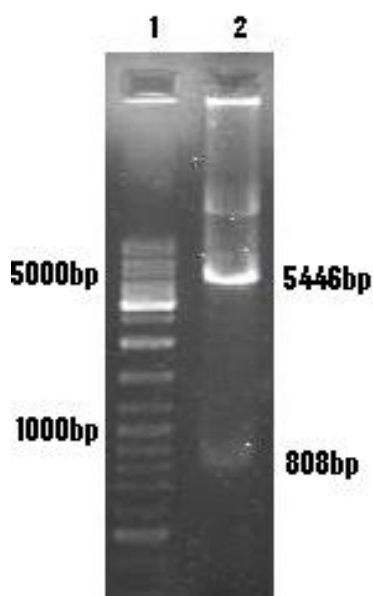


Fig. 4: 1.5% Agarose gel electrophoresis
Lane 1: 100 bp DNA ladder marker
Lane 2: Digested pcDNA-Rptr by BamH I and KpnI

The results of the simultaneously transfer of sense and antisense plasmids:

Plate from simultaneously transfer of pACYC-Duet-ptr and pcDNA-Rptr plasmids contained colonies. Since the plate contained selective antibiotics (ampicillin, chloramphenicol) in every plasmid, the presence of colonies in such plate indicated the simultaneous presence of both plasmids in one bacterial cell.

Inhibition of gene expression in the sense plasmid by antisense plasmid:

SDS-PAGE and Western blot analysis:

SDS-PAGE on lysate collected transformed bacteria with both of pACYCDuet-ptr and pcDNA-Rptr plasmids before and 5 hours after induction was performed. Induced bacteria containing pACYCDuet-ptr and induced bacteria containing pcDNA-Rptr also were analyzed in parallel. If the cloned PTR1 gene expressed in sense plasmid (pACYCDuet-ptr), the protein with molecular weight of about 30 kDa was expected. While the lysate bacteria containing both sense

and antisense plasmids also lysate bacteria containing antisense plasmid, after induction any trace of protein was observed in this range that this results indicate the inhibition of gene expression in sense plasmid by antisense plasmid (Fig. 5). Final confirmation was performed with western blotting (Fig. 6).

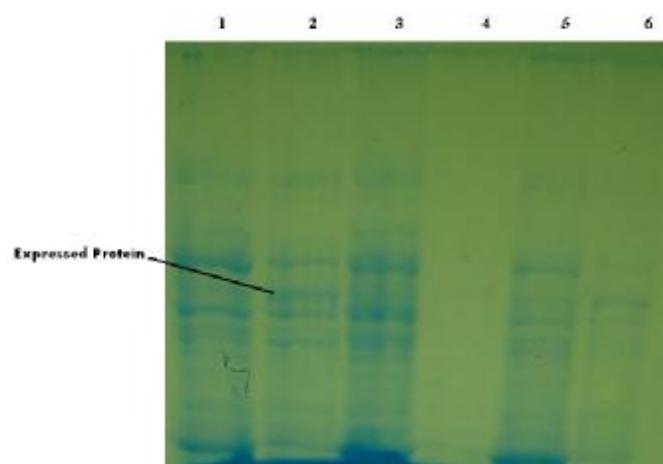


Fig. 5: 10% SDS-PAGE of bacterial lysate
Lane 1: Lysate of M15 cell containing pcDNA-Rptr collected 5hrs after induction
Lane 2: Lysate of M15 cell containing pACYC-Duet-ptr collected 5hrs after induction
Lane 3: Lysate of M15 cell containing pACYC-Duet-ptr and pcDNA-Rptr collected 5hrs after induction
Lane 4: Lysate of M15 cell containing pcDNA-Rptr collected before induction
Lane 5: Lysate of M15 cell containing pACYC-Duet-ptr collected before induction
Lane 6: Lysate of M15 cell containing pACYC-Duet-ptr and pcDNA-Rptr collected before induction



Fig. 6: Western blot analysis
Lane 1: Lysate of M15 cell containing pcDNA-Rptr collected 5hrs after induction

Lane 2: Lysate of M15 cell containing pACYC-Duet-ptr collected 5hrs after induction

Lane 3: Lysate of M15 cell containing pACYC-Duet-ptr and pcDNA-Rptr collected 5hrs after induction

Lane 4: Lysate of M15 cell containing pcDNA-Rptr collected before induction

Lane 5: Lysate of M15 cell containing pACYC-Duet-ptr collected before induction

Lane 6: Lysate of M15 cell containing pACYC-Duet-ptr and pcDNA-Rptr collected before induction

Discussion

Leishmaniasis is one of the parasitic diseases in most parts of the world. Out of the 30 provinces of Iran in the 15 provinces, leishmaniasis is endemic (17, 18). In addition to reports in the world about *Leishmania* which is resistant to drugs (19, 20) in Iran 10 to 15 percent of people infected with cutaneous leishmaniasis have been treated with Glucantime; they have no response to treatment that one of the reasons is drug-resistant parasites (21). Studies concerning methods of treatment of cutaneous leishmaniasis was conducted but none has been quite favorable (22, 23). According to the mentioned subjects, there is no completely effective drug therapy for treatment of cutaneous leishmaniasis and some failures are seen in the treatment that the lack of proper medical response is due to drug resistance, which needs more research. As was mentioned DHFR in parasitic DNA synthesis has an important role and inhibit this enzyme is causing the death. Methotrexate, one of anti-metabolite drugs, is an antagonist for folic acid, which competitively will inhibit DHFR, and its result is the death of parasite (24-25). PTR1 enzyme of *Leishmania* with similar enzymatic activity to DHFR reduced parasite susceptibility to anti-metabolite drugs such as Methotrexate (26) and was shown that deletion of PTR1 gene in *Leishmania*, which are treated with Methotrexate causes the death of parasite (27).

To inhibit gene expression, different several ways have been reported such as, oligodeoxy nucleo-

tides (ODNs) (28), mRNA antisense (29) and small interfering RNA (siRNA) molecules (30). The purpose of this study is to identify inhibiting power of sense plasmid by complete antisense plasmid in vitro. After preparing the sense and antisense plasmids, since origin of replication of both plasmids was different and were in compatibility group, transformation of both in a bacterial cell was done successfully and then expression was induced. The results of SDS-PAGE and western blot analysis showed inhibition of sense plasmid by antisense plasmid. In addition, it was shown that the inhibition occurs in the cytosol, as was reported by Dumas et al (31).

In conclusion, expression of PTR1 gene in sense plasmid was inhibited successfully by antisense plasmid. Because the mRNA antisense can be used in gene therapy, the results of this study showed that this method could be successful.

Ethical considerations

Ethical issues (Including plagiarism, Informed Consent, misconduct, data fabrication and/or falsification, double publication and/or submission, redundancy, etc) have been completely observed by the authors.

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