Loss of virulence in *Leishmania donovani* deficient in an amastigote-specific protein, A2

WEN-WEI ZHANG AND GREG MATLASHEWSKI*

Institute of Parasitology, McGill University, 21 111 Lakeshore Road, Ste-Anne de Bellevue, Québec, Canada H9X 3V9

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**ABSTRACT** *Leishmania donovani* is the etiologic agent of fatal visceral leishmaniasis in man. During its life cycle, *Leishmania* exist as flagellated promastigotes within the sandfly vector and as nonflagellated amastigotes in the macrophage phagolysosomal compartment of the mammalian host. The transformation from promastigotes to amastigotes is a critical step for the establishment of infection, and the molecular basis for this transformation is poorly understood. To define the molecular basis for amastigote survival in the mammalian host, we previously identified an amastigote stage-specific gene family termed “A2.” In the present study, we have inhibited the expression of A2 mRNA and A2 protein in amastigotes using antisense RNA and show that the resulting A2-deficient amastigotes are severely compromised with respect to virulence in mice. Amastigotes that did survive in the mice had restored A2 protein expression. These data demonstrate that A2 protein is required for *L. donovani* survival in a mammalian host, and this represents the first identified amastigote-specific virulence factor identified in *Leishmania*. This study also reveals that it is possible to study gene function in *Leishmania* through the expression of antisense RNA.

*Leishmania donovani* protozoa are the causative agents of human visceral leishmaniasis, which is a potentially fatal disseminating infectious disease that effects over 80 countries worldwide (1, 2). To define the molecular basis for amastigote survival in the mammalian host, we previously identified an amastigote stage-specific gene family termed “A2,” whose corresponding mRNA and protein are abundant in amastigotes but are largely absent in promastigotes (3–5). There are at least seven members of the A2 gene family that encode a family of proteins ranging from 45 to 100 kDa and that are specific to the amastigote stage (5). A2 proteins are mainly comprised of a repetitive amino acid sequence; each repeat encodes a stretch of 10 amino acids (3, 5) that shares partial identity with the S antigens of the *Plasmodium falciparum* V1 strain, which is responsible for malaria in man (3). Like A2, the S antigens are stage-specific to the mammalian host and the function has yet to be defined.

To determine whether the A2 proteins are required for the survival of amastigotes in *in vitro* in macrophages and *in vivo* in a model mammalian host, we undertook to develop A2-deficient *L. donovani* amastigotes. Although it is possible to block gene expression in *Leishmania* through creating null mutants by homologues recombination (6–8), we explored the use of antisense RNA to inhibit A2 gene expression and protein synthesis for the following reason. The A2 genes are present as a multigene family alternating with another gene, termed “A2rel,” that is constitutively expressed in promastigotes and amastigotes (4), making it difficult to specifically target the A2 genes without effecting A2rel genes. Although it is possible to make A2 gene knockouts, it would be difficult. Transfecting and selecting cells with an antisense-expressing plasmid construct is technically simpler, and the information gained through this approach may provide justification for subsequently developing the gene knockouts. Finally, if the antisense approach was successful, this would represent a significant advance in studying gene function in *Leishmania*.

From this study, we present data showing that it is possible to develop A2-deficient amastigotes using the antisense RNA approach. A2 antisense RNA containing amastigotes displayed an absence of both A2 mRNA and corresponding A2 proteins. The A2-deficient amastigotes were viable in culture and proliferated as well as control plasmid-containing amastigotes. A2-deficient amastigotes, however, displayed a reduced ability to multiply in cultured macrophages and were severely compromised with respect to survival in mice. Of interest, the A2-deficient amastigotes that did survive in mice had lost the ability to express A2 antisense RNA and consequently had restored the ability to express A2 protein. These data reveal that A2 can be considered an amastigote-specific virulence factor that is required for *L. donovani* survival in a mammalian host.

**MATERIALS AND METHODS**

**Plasmid Construction.** The A2 antisense plasmid (pKSneoA2–1R) was constructed by reversing the 2.4-kb BamHI fragment containing a 1.6-kb A2–1 encoding sequence and partial A2 3′ untranslated region (UTR) sequence that was originally derived from the A2 sense plasmid (pKSneoA2–1), which has been described in detail (5).

**Cell Culture and Transfection.** The *L. donovani* 1S/C12D cell line was kindly provided by Dennis Dwyer of the National Institutes of Health, Bethesda, MD. These cells can be cycled between promastigotes at 26°C and amastigotes at 37°C *in vitro* under defined conditions in which both promastigotes and amastigotes remain virulent. The details concerning the formulation of these culture procedures were previously described (9). The medium for promastigotes was 199 plus 10% fetal bovine serum (pH 6.8) and for amastigotes was RPMI 1640 medium plus 20% fetal bovine serum (pH 5.5). The A2 antisense plasmid was electroporated into *L. donovani* promastigotes, and transfected cells were selected with 20 μg/ml G418 as described (4, 5). After establishment of transfected promastigotes, these transfected cells were routinely cycled between 26°C promastigotes and 37°C amastigotes in the above described media containing 100 μg/ml G418 on a weekly basis.

**Northern Blot Analysis.** Total RNAs were isolated from *L. donovani* promastigotes and *in vitro* amastigotes with TRIZol reagent (GIBCO). Northern blot analyses were performed as described (5).

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*Abbreviation: UTR, untranslated region.*

*To whom reprint requests should be addressed. e-mail: greg.matlashewski@maclan.mcgill.ca.*
described (3) using 10 μg of total RNA. The hybridizations were carried out for 12 h at 42°C in a solution containing 1% SDS, 1M NaCl, 10% dextran sulfate, and 50 μg/ml salmon sperm DNA. The antisense oligonucleotides [CGGGG(A)CG(A)GACG(A)GA] specific for A2 mRNA and the sense oligonucleotides [G(C)TCC(T)GTGT(G)GGC(T)CCGC] specific for A2 antisense RNA were derived from the repeat region of A2 encoding sequence (3) and labeled with 32p by T4 polynucleotide kinase.

**Western Blot Analysis.** Protein samples of *L. donovani* cells were prepared as described (5) and separated on 10% SDS/PAGE, transferred to nitrocellulose, and analyzed by immunoblotting for A2 proteins with anti-A2 mAb C9 (5). Western blot was performed with an enhanced chemiluminescence kit (Amersham) following the manufacturer’s instruction except 10% skim dry milk was used for blocking the membrane and 5% skim dry milk was included in the primary antibody and the second antibody solutions.

**Infection of Macrophages in Culture.** Mice bone marrow cell preparation and infection with *L. donovani* cells were performed as described (10) with some modification. In brief, macrophages were infected in suspension for 24 h at an amastigote-to-cell ratio of 20:1. Noningested amastigotes were removed by centrifugation and three washes in RPMI complete medium, and the infected cells were incubated at 37°C. Amastigote infection level and growth rate in macrophages were evaluated daily by cytopsin and Giemsa staining.

**Visceral Infection in Mice.** Female BALB/c mice (Charles River Breeding Laboratories) weighing 20–25 g were injected via the tail vein with 1.5 × 10⁸ wild-type amastigotes or A2-deficient amastigotes or amastigotes containing the control vector; 4 weeks after infection, mice were examined for *L. donovani* parasite burden by counting the number of amastigotes in the Giemsa-stained imprints of the liver. Liver parasite burdens, expressed as Leishman–Donovan Units, were calculated as follows: number of amastigotes per 1000 cell nuclei × liver weight (g) (11). Amastigotes were recovered from the infected mice liver and spleen as described (12), and the recovered amastigotes were cultured in promastigote or amastigote culture medium described above either with or without 20 μg/ml G418. Cells were collected when they reached a density of 3 × 10⁷ cells/ml and subjected to Western blot analysis.

**RESULTS**

The plasmid construct developed to express A2 antisense RNA and the control plasmid are shown in Fig. 1A. This antisense RNA-expressing construct was designed to express a chimeric A2 antisense RNA (complementary to A2 coding region and part A2 3′ UTR sequence) flanked by A2 5′ UTR and 3′ UTR sequences. We previously have demonstrated that A2 3′ UTR was essential for A2 mRNA accumulation and stabilization in amastigotes (4). Thus, the construct was designed to have the chimeric A2 antisense RNA accumulate at higher levels in *Leishmania* amastigotes than promastigotes. Plasmids were electroporated into the promastigotes of the *L. donovani* 1S/CL2D strain, and the recombinant promastigotes were selected with G418 as a pooled population of resistant cells. Norther blot was then performed on RNA isolated from promastigotes and amastigote populations to identify antisense and sense A2 transcripts (Fig. 1B). As predicted, the A2 antisense plasmid-transfected cells contained higher levels of A2 antisense RNA in amastigotes than in promastigotes, and no antisense RNA was present in the wild-type cells (Fig. 1B, Upper). Although A2 mRNA was clearly abundant in the wild-type amastigotes, it was strikingly absent from the amastigotes expressing the A2 antisense transcripts (Fig. 1B, Lower). However, very low amounts of A2 mRNA could be detected when the film was overexposed (data not shown).

These data demonstrate that the A2 antisense RNA either inhibited the expression of the A2 genes or impaired the processing and maturation of the A2 transcripts.

We next examined the level of A2 proteins in the different *L. donovani* cells by Western blot analysis using previously developed anti-A2 mAbs (5). As expected, the A2 protein family was present in wild-type and control vector transfected amastigotes but not in promastigotes (Fig. 2, lanes 2 and 6, respectively), and this is consistent with our previous observation that A2 proteins are specific to amastigotes (5). However, the A2 proteins were absent in the amastigotes containing the A2 antisense RNA (Fig. 2A, lane 4), and this is consistent with the data showing that these cells contained dramatically reduced levels of A2 mRNA. To further examine the extent of A2 protein suppression in the A2 antisense RNA containing amastigotes, we compared the level of A2 protein...
Amastigotes within macrophages were subsequently monitored containing, and wild-type amastigotes, and the growth rates of infected for 24 h with A2-deficient, control plasmid-containing, and wild-type amastigotes. 

Bone marrow-derived macrophages from BALB/c mice were infected with A2-deficient cells as indicated. Female BALB/c mice were injected with amastigotes via the tail vein (1.5 × 10⁸ amastigotes/mouse, three mice per group). Four weeks after infection, mice were examined for hepatic parasite burdens by counting amastigotes from liver impression. This experiment was repeated four times with consistent results. As shown in Fig. 3A, A2-deficient amastigotes grew slower in macrophages than wild-type cells and much slower than amastigotes containing the control plasmid. This revealed that A2-deficient amastigotes were compromised in their ability to proliferate in cultured macrophages.

The most stringent assay for virulence is the ability of the amastigotes to establish an infection in vivo in an animal model. Therefore, BALB/c mice were infected with A2-deficient, wild-type, and control plasmid-containing amastigotes. Mice were infected via the tail vein (1.5 × 10⁸ amastigotes/mouse, three mice per group), and, 4 weeks after infection, mice were examined for hepatic parasite burdens by counting amastigotes from liver impression. This experiment was repeated four times with consistent results. As shown in Fig. 3A, the A2 antisense RNA can efficiently block A2 protein expression in L. donovani amastigotes.

We next determined whether inhibition of A2 expression resulted in altered virulence in vitro in cultured macrophages. Bone marrow-derived macrophages from BALB/C mice were infected for 24 h with A2-deficient, control plasmid-containing, and wild-type amastigotes, and the growth rates of amastigotes within macrophages were subsequently monitored over a 5-day period by cytopsin analysis. There was no difference in the ability to invade macrophages during the 24-h infection period. The percentage of infected macrophages for wild-type, A2-deficient, and control plasmid-containing amastigotes was 74%, 80%, and 86%, respectively, and the initial numbers of amastigotes within infected macrophages were similar. However, as shown in Fig. 3A, A2-deficient amastigotes grew slower in macrophages than wild-type cells and much slower than amastigotes containing the control plasmid. This revealed that A2-deficient amastigotes were compromised in their ability to proliferate in cultured macrophages.

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Fig. 3B, the parasite burden of mice infected with A2-deficient amastigotes was low compared with the parasite burdens of mice infected with wild-type or control plasmid-containing amastigotes. Similar results were obtained when mice were infected with wild-type or A2-deficient promastigotes (data not shown). It is unclear why amastigotes containing the control plasmid were more virulent than wild-type cells in mice and in cultured macrophages. We do not believe that this was caused by an homologous recombination event resulting in the overexpression of A2 from the plasmid episome because this would have resulted in overexpression of one, but not all, A2 protein species, and there is no evidence of this, as shown in Fig. 2A. The control plasmid-containing cells did, however, undergo a prior selection for the plasmid using G418, which the wild-type cells did not, and this may have somehow selected for a more virulent population. Nevertheless, it is clear that the presence of the plasmid itself did not reduce virulence in these amastigotes. Taken together, these data show that loss of A2 expression was concomitant with reduced virulence in vivo.

As revealed in Fig. 3B, there was a low but consistent level of infection with the A2-deficient amastigotes. It was therefore of interest to determine the status of the A2 protein in these amastigotes that had survived the 4-week period and had established infection in the liver of these mice. Amastigotes were isolated from the liver and spleen of mice whose A2 protein was not expressed before infection. These amastigotes were unable to survive in BALB/c mice infected with A2-deficient amastigotes and these were placed in promastigote or amastigote culture medium both in the presence and absence of G418. Only amastigotes retaining the transfected plasmid conferring Neo resistance would have survived in the G418-containing media. Both promastigotes and amastigotes grew equally well in the presence or absence of G418. When cells reached a density of \(3 \times 10^7\) cells/ml, they were harvested, lysed, and subjected to Western blot analysis with the anti-A2 mAb. As shown in Fig. 4A, A2 protein expression was restored in the surviving amastigotes that were originally A2-deficient before infection. This demonstrated that during the infection period, there was a selection for A2 protein-containing amastigotes and against A2-deficient amastigotes. Northern blot analysis was performed to determine whether the recovered amastigotes lost the ability to express the antisense RNA. As shown in Fig. 4B, the recovered amastigotes had lost the ability to express the A2 antisense RNA but did retain the ability to express the neo mRNA. Therefore, the surviving amastigotes retained the transfected plasmid but the plasmid selectively lost the ability to express the A2 antisense RNA. Taken together, these data demonstrate that the A2-deficient amastigotes were unable to survive in these mice whereas the amastigotes with restored A2 protein did survive. It is possible that the A2-deficient amastigotes were either unable to establish an infection in the mice or that they were unable to proliferate after infection. In either case, this establishes the A2 protein as an amastigote-specific virulence factor required for survival in the mouse model.

**DISCUSSION**

Previous studies (3–5) have shown that the expression of the *L. donovani* A2 transcripts and protein family are amastigote-specific. In the present study, we examined the role of the A2 protein family in amastigote survival in axenic culture, in infectivity and survival in macrophages cultures in vitro, and in infectivity and survival in BALB/c mice in vivo. The principal observations were that amastigotes deficient in A2 protein were compromised with respect to proliferation in cultured macrophages and were unable to survive in BALB/C mice. Of particular interest, amastigotes that were originally A2-deficient and did survive in BALB/C mice had regained the ability to express the A2 protein. This provides a compelling argument that A2 protein is required for amastigotes to survive in a mammalian host and thus can be considered an amastigote-specific virulence factor. We have also revealed that antisense RNA can be successfully used to study gene function in *Leishmania*. Antisense RNA can therefore be considered as an additional genetic tool to study the biology of this important parasitic protozoan.

The amastigote-specific expression of A2 is suggestive of a role in the parasite’s survival in macrophages, and this hypothesis is supported by the *in vitro* and *in vivo* infection experiments reported within. It has been reported that nitrogen oxides (NO) are the major mediators of macrophage leishmanicidal activity (13). We know that A2-deficient amastigotes and wild-type amastigotes were equally capable of suppressing NO release in infected macrophages (data not shown). This suggests that A2 is not involved in altering host macrophage function, and this is consistent with our previous observation that A2 is located predominantly in the cytoplasm of amastigotes (5). A2 represents a multiprotein family con-
taining repeated subunits, so we are pursuing the possibility that A2 proteins form a complex, which may have a structural role in the amastigote.

We show that antisense RNA can be used in place of gene targeting to examine gene function in *Leishmania*. Because of its simplicity and effectiveness, the antisense RNA approach could be widely applied to examine the function of a variety of genes in *Leishmania*. This could be particularly useful in situations like that of the A2 gene, which is part of a multigene family. Nevertheless, our data also reveal that antisense RNA cannot replace gene knockouts in situations in which a stable phenotype is needed, such as in the development of live, attenuated vaccine strains. For example, it has been shown that gene replacement in the dihydrofolate reductase locus has provided an attenuated strain of *L. major*, which can induce protection against wild-type *L. major* in mice (14). Targeting the cysteine protease genes in *L. mexicana* resulted in parasites retaining the ability to produce s.c. lesions in mice but at a slower rate than the wild-type parasites (15). Based on the data obtained in this study, there is clear justification for targeting the A2 gene in *L. donovani*, and we are currently undertaking this. With regard to a live, attenuated vaccine strain, it may be advisable to knockout several relevant genes, including the A2 gene, to ensure a stable but viable attenuated phenotype. We also have shown recently that *L. donovani* can be used to express high levels of recombinant proteins (16); therefore, live attenuated strains can be further engineered to express a variety of protective antigens or immunostimulatory cytokines.

In conclusion, the present study has established the use of antisense RNA to study gene function in *Leishmania* and has identified the amastigote-specific A2 protein as a virulence factor required for parasite survival in a mammalian host. A2 was not, however, required for parasite survival in axenic culture, and these observations therefore may have implications for the development of a live, attenuated vaccine strain of *L. donovani*.

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