Serum Resistance-Associated Protein Blocks Lysosomal Targeting of Trypanosome Lytic Factor in *Trypanosoma brucei*

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*Trypanosoma brucei* is the causative agent of nagana in cattle and can infect a wide range of mammals but is unable to infect humans because it is susceptible to the innate cytotoxic activity of normal human serum. A minor subfraction of human high-density lipoprotein (HDL) containing apolipoprotein A-I (apoA-I), apolipoprotein L-I (apoL-I), and haptoglobin-related protein (Hpr) provides this innate protection against *T. b. brucei* infection. This HDL subfraction, called trypanosome lytic factor (TLF), kills *T. b. brucei* following receptor binding, endocytosis, and lysosomal localization. *Trypanosoma brucei rhodesiense*, which is morphologically and physiologically indistinguishable from *T. b. brucei*, is resistant to TLF-mediated killing and causes human African sleeping sickness. Human infectivity by *T. b. rhodesiense* correlates with the evolution of a resistance-associated protein (SRA) that is able to ablate TLF killing. To examine the mechanism of TLF resistance, we transfected *T. b. brucei* with an epitope-tagged SRA gene. Transfected *T. b. brucei* expressed SRA mRNA at levels comparable to those in *T. b. rhodesiense* and was highly resistant to TLF. In the SRA-transfected cells, intracellular trafficking of TLF was altered, with TLF being mainly localized to a subset of SRA-containing cytoplasmic vesicles but not to the lysosome. These results indicate that the cellular distribution of TLF is influenced by SRA expression and may directly determine the organism’s susceptibility to TLF.

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(SRA) (8, 9, 10, 31, 40, 51, 37, 49). SRA is a member of the variable surface glycoprotein (VSG) family of proteins in African trypanosomes, and despite having low sequence homology (<25%) with VSGs, it shares several sequence and structural features with VSGs (6, 9, 11). SRA is an expression site-associated gene in T. b. rhodesiense and is located upstream of the VSG in the active telomeric expression site (51).

The role of SRA in resistance to human serum was conclusively shown in transfection studies of T. b. brucei with SRA (50, 51). These studies were extended to show that recombinant apol-I and SRA bind in vitro by a coiled-coil interaction between the two proteins, and this has been proposed to directly inhibit trypanosome killing by apol-I (47). Immunofluorescence microscopy analysis of T. b. rhodesiense indicated that apol-I and SRA colocalized to the lysosome in trypanosomes treated with apol-I. However, other studies have shown that TLF uptake and cellular localization differed in resistant and susceptible lines of T. b. rhodesiense (19). TLF accumulation was reduced approximately sixfold in resistant trypanosomes, and the cell-associated TLF was excluded from the lysosome (19). This indicated that differences in cellular trafficking of TLF might contribute to trypanosome susceptibility to TLF killing.

Here we show that transfection of SRA into three different bloodstream-stage T. b. brucei isolates expressing different VSGs was sufficient to confer high levels of resistance to TLF and human serum. Epitope-tagged SRA also conferred TLF resistance and allowed subcellular localization of SRA to non-lysosomal vesicles predominantly located between the nucleus and the kinetoplast. In SRA-expressing cells, TLF was not routed to the lysosome, and colocalization of TLF and SRA was observed in small cytoplasmic vesicles. Based on these observations, we conclude that the association of SRA with TLF-containing endosomes results in rerouting of TLF, thus preventing lysosomal localization and trypanosome death.

MATERIALS AND METHODS
Trypanosome isolates and culture methods. Culture-adapted monomorphic isolates of T. b. brucei MiTat 1.2 (427/221), pleomorphic BiTat 1.1 (KETRI 667), and GuTat 10.1 (TREU 927/4) cells were used throughout these studies. Bloodstream-stage cultures were grown in HMI-9 medium supplemented with 10% fetal bovine serum (FBS; HyClone, Logan, UT), and fresh serum was added at 24 h. The cell-associated TLF was observed in small cytoplasmic vesicles. Based on these observations, we conclude that the association of SRA with TLF-containing endosomes results in rerouting of TLF, thus preventing lysosomal localization and trypanosome death.

Trypanosome resistance to TLF. Susceptible and resistant bloodstream-stage T. b. brucei isolates expressing different VSGs were grown in HMI-9 medium supplemented with 10% human serum. Epitope-tagged SRA also conferred TLF resistance and allowed subcellular localization of SRA to non-lysosomal vesicles predominantly located between the nucleus and the kinetoplast. In SRA-expressing cells, TLF was not routed to the lysosome, and colocalization of TLF and SRA was observed in small cytoplasmic vesicles. Based on these observations, we conclude that the association of SRA with TLF-containing endosomes results in rerouting of TLF, thus preventing lysosomal localization and trypanosome death.

Transfection studies. SRA was transfected into bloodstream T. b. brucei cells. The DNA sequence encoding the Ty epitope, containing the coding information for a 10-amino-acid sequence of the major structural protein of the TLF-containing endosomes results in rerouting of TLF, thus preventing lysosomal localization and trypanosome death.

DNA and RNA analysis. Genomic DNA was isolated as described previously (30). Twenty nanograms of genomic DNA was used for PCRs to examine the presence of SRA (5’ primer, CACACCTCTAAAGATCACCATAG; 3’ primer, AACCTCATGAAAAATGGTAAAG) and tubulin (5’ primer, CGGTTGGCAT ATGCGAA; 3’ primer, GGGGGTGCACTTTGTC) gene sequences. Southern blots were performed according to standard protocols (42). Total RNA was isolated with Trizol reagent (Roche Biochemicals, Indianapolis, IN), and 5 μg of RNA was separated in 1% formaldehyde gels and electrophoresed in 1X TBE buffer for 1.5 h, followed by 50 V for 1.5 h, onto a positively charged nylon membrane (Roche Biochemicals, Indianapolis, IN). DNA probes were labeled by the random primer method (Life Technologies-Invitrogen, Carlsbad, CA) with [32P]dCTP. The blots were hybridized in 0.9 M NaCl, 5 × Denhardt's solution, 10% dextran sulfate, 5 mM EDTA, 0.1% sodium dodecyl sulfate [SDS], 40% formamide, 0.1 mg/ml salmon sperm DNA overnight at 42°C and subsequently washed at 60°C for 2× SSC (1× SSC, 0.15 M NaCl plus 0.015 M sodium citrate), 1% SDS, and finally 0.1× SSC, 1% SDS at room temperature. Alternatively, the probes were labeled with AlkPhos Direct (Amersham Pharmacia, Piscataway, NJ), hybridized, and developed according to the manufacturer's description.

Analysis of SRA by SDS-PAGE and Western blotting. Cultured cells were washed twice with phosphate-buffered saline (PBS; 137 mM NaCl, 2.7 mM KCl, 10 mM Na2HPO4, 1.8 mM KH2PO4, and 0.1 mM EDTA) and then resuspended in lysis buffer (100 mM Tris, pH 8; 10 mM EDTA; 0.5% SDS) containing a protease inhibitor cocktail (Complete Mini; Roche Biochemicals, Indianapolis, IN). Freshly prepared cell lysates (equivalent to 3 × 107 cells per lane) were separated in 10% SDS-polyacrylamide gel electrophoresis (SDS-PAGE) gels (24). Proteins were transferred to nitrocellulose membranes (0.2-μm Protran; Schleicher and Schuell, Dassel, Germany) for 45 min at 57 V. Polyclonal antibodies against VSG 221 were added at a 1:2,000 dilution, incubated for 1 h at room temperature, and subsequently incubated with an HRP-conjugated secondary antibody (1:5,000 dilution) for 1 h at room temperature. Slides were then washed with PBS, and Alexa Fluor 488-labeled goat anti-mouse immunoglobulin G (Molecular Probes, Eugene, OR) was added at a 1:1,000 dilution in PBS–1% FBS for 1 h at room temperature. 4,6-Diamidino-2-phenylindole (DAPI, 2-μg/ml final concentration) was added dur-
ing the secondary antibody incubation. Slides were washed two times in PBS and microscopically analyzed.

To determine whether SRA colocalized with the endoplasmic reticulum (ER), fixed cells were incubated with monoclonal anti-Ty (1:50) and polyclonal anti-βtubulin antibodies. For colocalization of SRA with TL or TLF, cells were stained with Alexa fluorochromes. Cells were incubated with 4 μg/ml TLF for 30 min at 37°C. After incubation, the cells were put on ice, washed, resuspended in PBSG at 4°C, smeared onto a slide, and fixed with cold methanol. For colocalization of SRA with the lysosomal marker p67, cells were stained with the anti-Ty antibody as described above and then incubated with a monoclonal antibody to the lysosomal marker p67 (1:1,000) conjugated to Alexa Fluor 594-labeled antibodies were added, and slides were processed as described above. To determine SRA localization with the lysosome, cells were stained with the anti-Ty antibody as described above and then incubated with a monoclonal antibody to the lysosomal marker p67 (1:1,000) conjugated to Alexa Fluor 594 (23).

To further localize SRA with the endocytic marker tomato lectin (TL; Vector Laboratories, Burlingame, CA) (34) and TLF, we directly conjugated TL and TLF to Alexa fluorochromes. Cells were incubated with 4 μg/ml TL and/or 10 to 20 μg/ml TLF for 30 min at 37°C. After incubation, the cells were put on ice, washed, resuspended in PBSG at 4°C, smeared onto a slide, and fixed with cold methanol. For colocalization of SRA with TL or TLF, cells were stained with anti-Ty antibody as described above. For localization of TLF and p67, cells were incubated with 50 μM chloroquine for 30 min before adding Alexa-conjugated TLF to a final concentration of 20 μg/ml. Cells were incubated for an additional 90 min before being processed and stained for p67 as described above. Trypanosomes were analyzed at a magnification of ×100 with a Zeiss fluorescence microscope, and digital images were captured with a Zeiss AxiosCam video camera. The contrast and brightness of some images were adjusted and overlaid with Adobe Photoshop software.

**RESULTS**

**Genotypic and phenotypic analysis of SRA-transfected** *T. b. brucei*. Previous studies have shown that transfection of procyclic *T. b. brucei* with SRA and subsequent transmission through tsetse flies resulted in bloodstream-stage *T. b. brucei* organisms that were resistant to normal human serum (51). The cyclical transmission of trypanosomes through tsetse flies is difficult due to the low transmission efficiency, variability in the ability of *T. b. brucei* isolates to infect tsetse flies, and specific developmental changes in gene expression influencing the VSG expression sites and other genes. To circumvent these problems and to directly determine if SRA and Ty epitope-tagged SRA were sufficient to confer resistance to normal human serum, three different isolates of bloodstream-stage *T. b. brucei* were transfected with SRA and SRA-Ty. The location of the Ty epitope within the SRA gene was not expected to disrupt the proposed N-terminal alpha-helical domains implicated in having a role in SRA resistance (Fig. 1A) (6, 47). Similar transfections of bloodstream-stage *T. b. brucei* with the hemagglutinin (HA) epitope fused to the amino or carboxyl terminus of SRA were reported previously (50). Genomic DNAs from wild-type 427, transfected 427-SRA and 427-SRA-Ty, wild-type 667, transfected 667-SRA and 667-SRA-Ty, wild-type 927, and transfected 927-SRA and 927-SRA-Ty organisms were analyzed for the presence of the SRA gene by PCR with primers complementary to the 5’ and 3’ coding sequences of SRA. A 1.4-kb SRA product was only detected in SRA- and SRA-Ty-transfected *T. b. brucei* lines and not in wild-type cells (Fig. 1B). Southern blotting confirmed single-copy integration of SRA in the SRA and SRA-Ty transfecants (data not shown). Control PCRs using primers specific to the α-tubulin gene produced the expected 0.75-kb product (Fig. 1B).

By light microscopy, no morphological differences were observed between SRA-transfected and wild-type cells. In addition, cells with and without SRA reached approximately the same cell densities and grew at the same rates, indicating that the expression of SRA did not influence the growth characteristics of bloodstream-stage *T. b. brucei* (data not shown).

**Susceptibility of SRA-transfected T. b. brucei to TLF killing.** In order to determine whether transfection with SRA was sufficient to confer the serum resistance phenotype, transfected cells were incubated with increasing concentrations of TLF (Fig. 2) or normal human serum (data not shown). The three wild-type lines of *T. b. brucei* were highly susceptible to lysis by TLF. Following SRA transfection, the three lines of *T. b. brucei* showed high levels of resistance to TLF (Fig. 2). The level of serum resistance in SRA-transfected *T. b. brucei* was similar to that observed for human infectious *T. b. rhodesiense* (data not shown). No difference in the level of TLF resistance was detected between cells transfected with SRA alone and those transfected with SRA-Ty (data not shown). To confirm whether
bromide staining of the agarose gel. These results show that nucleotides). Equal RNA loading was confirmed by ethidium bromide staining of the agarose gel and Western blot analysis show results for protein extracts from T. b. brucei transfected with vector alone (427) or with SRA (427-SRA-Ty). The Western blot was probed with antibodies to the Ty epitope, and proteins of approximately 59 and 65 kDa were detected in extracts from T. b. brucei 427-SRA-Ty (arrows).

FIG. 2. Transfection of three T. b. brucei lines with SRA confers resistance to TLF. Lysis assays were conducted with T. b. brucei transfected with empty vector (427 [■], 667 [●], and 927 [▲]) and with SRA-Ty vector (427-SRA [▲], 667-SRA [●], and 927-SRA [▲]). The percentages of trypanosomes lysed by increasing amounts of TLF under standard assay conditions were determined following a 4-h incubation at 37°C.

SRA-induced TLF resistance was comparable to that seen for T. b. rhodesiense, the survival of SRA-transfected T. b. brucei was examined at concentrations of TLF similar to those found in human serum (1,000 U/ml) and by incubation for 24 h under the same growth conditions in the presence of 50% human serum. The SRA-transfected T. b. brucei lines remained viable under these conditions (data not shown).

Early studies suggested that the VSG expressed on the surfaces of trypanosomes might influence their susceptibility to normal human serum. Each of the T. b. brucei lines used in the current studies expresses different VSG genes producing serologically distinct VSG coats. Based on the T. b. brucei isolates tested, it appears that resistance to TLF is highly dependent on SRA expression but independent of the VSG expressed on the trypanosome surface. The transfection of bloodstream-stage lines of T. b. brucei directly demonstrated that expression of the SRA gene is sufficient to confer resistance to TLF and may lead to human infectivity.

Expression of SRA mRNA in transfected cells. SRA mRNA is an abundant transcript in T. b. rhodesiense, representing as much as 10% of the total cellular RNA (10, 18, 31). In order to determine the level of SRA mRNA in transfected cells and to compare this level with that in human infectious T. b. rhodesiense, total cellular RNA was isolated from T. b. brucei transfected with vector alone (427) or SRA (427-SRA) and from a human-serum-resistant line of T. b. rhodesiense. Blots were hybridized with probes specific for the VSG expressed by T. b. rhodesiense (VSG-R) and the VSG-221 expressed by the T. b. brucei 427 cell line (Fig. 3A). The amount of SRA mRNA in human-serum-resistant T. b. rhodesiense was similar to the level of SRA mRNA in T. b. brucei 427-SRA (Fig. 3A). The SRA mRNA in T. b. brucei 427-SRA is approximately 460 nucleotides larger than that in T. b. rhodesiense because of additional 5′ and 3′ untranslated region sequences added to the SRA sequence in the expression vector (about 430 nucleotides) and the addition of the epitope tag within the gene (30 nucleotides). Equal RNA loading was confirmed by ethidium bromide staining of the agarose gel. These results show that while the transgenic SRA gene is expressed from the ribosomal locus in T. b. brucei 427-SRA, the expression level of the gene is similar to the endogenous expression level of SRA in human-serum-resistant T. b. rhodesiensi strains.

Expression of SRA in transfected T. b. brucei. Despite high levels of SRA mRNA, the detection of SRA protein in T. b. rhodesiense and in SRA-transfected T. b. brucei has been problematic in some cases (8, 9, 51). However, visualization of SRA has been reported for T. b. rhodesiense cells by use of a polyclonal mouse antiserum raised against recombinant SRA (31). In addition, endogenous SRA in T. b. rhodesiense as well as recombinant SRA (rSRA) expressed in T. b. brucei was detected using antibodies raised against an SRA polypeptide (47). To examine SRA expression in transfected T. b. brucei, total cell lysates from 3 × 10⁵ trypanosomes were separated by 10% SDS-PAGE and analyzed by Coomassie staining and Western blotting with monoclonal antibodies specific to the Ty tag inserted into the SRA gene. Epitope-tagged proteins, migrating at approximately 59 and 65 kDa, were detected in Western blots of cell lysates of both 427-SRA and 667-SRA but not in lysates of wild-type 427 and 667 (Fig. 3B; data for 667 and 667-SRA are not shown). The predicted size of mature SRA is approximately 38,000 Da. While the nature of the doublet is unknown, it may be the result of differences in posttranslational modification of SRA. The staining patterns observed are similar to previous Western blot results obtained with T. b. rhodesiense extracts analyzed with anti-rSRA and SRA-HA-transfected T. b. brucei probed with anti-HA (31, 50). As reported by others, we found that the detection of SRA
within the ER or targeting to the lysosome for degradation.

To determine whether SRA was located in endosomes or the ER, we performed double labeling with the endocytic marker TL (red) and the ER marker BiP (green; Fig. 4A, B, C, and D). We found that SRA-Ty staining did not overlap with TL staining (Fig. 4F), and to a limited extent, in endosomes located nearer to the flagellar pocket (green; Fig. 5C and G). While SRA-Ty localized to the region between the kinetoplast and the nucleus, SRA-Ty staining did not overlap with TL staining (Fig. 4E to H). To directly address lysosomal localization, we stained cells with an antibody to p67 (red) and found that SRA-containing vesicles (green) did not significantly overlap with localization of the lysosomal membrane protein p67 (Fig. 4I to L). This pattern suggests that SRA-Ty is predominately nonlysosomal.

TLF killing of T. brucei requires lysosomal localization and subsequent acidification (20, 26). T. brucei 427 was incubated with TL conjugated with Alexa Fluor 594 and TL conjugated with Alexa Fluor 488 for 30 min at 37°C. Under these conditions, both TL and TLF colocalized mainly to the lysosomes of wild-type cells (Fig. 5A to D). These results are consistent with previous studies that localized TLF to lysosomes of other isolates of T. brucei (20, 44). In contrast, TLF did not colocalize with the endocytic marker TL in T. brucei 427-SRA-Ty cells (Fig. 5E to H). After 30 min, small TLF-containing vesicles were seen close to the flagellar pocket (Fig. 5F).

Once we saw that TLF did not seem to be trafficking by its normal endocytic pathway in the SRA-expressing cells, we asked whether TLF could be concentrated in the lysosome in the presence of chloroquine. Previous studies have shown that chloroquine blocks lysosome acidification and prevents the degradation of endocytosed proteins (44). Cells were treated with chloroquine for 30 min before incubation with TLF for 90 min in an attempt to force TLF into the lysosome. As expected, in wild-type T. brucei, TLF accumulated and colocalized with...
the lysosomal marker p67 (Fig. 6A to D). However, in the T. b. brucei 427-SRA-Ty cells, TLF localization was quite different. TLF did not localize with p67 and accumulated in nonlysosomal vesicles (Fig. 6E to H). Similar results were obtained when non-chloroquine-treated cells were examined (data not shown). Based on these results, we conclude that the intracellular trafficking of TLF is altered in T. b. brucei 427-SRA-Ty and that the expression of SRA may prevent TLF from reaching the lysosome, the site of trypanolytic activity.

**SRA and TLF colocalize in T. b. brucei 427-SRA-Ty.** The essential nature of SRA expression in human-serum-resistant T. b. brucei suggests that SRA and TLF may interact with one another directly. In fact, studies have shown that SRA directly interacts with apol-I, one of the trypanolytic components of TLF (47). To examine whether SRA and TLF are contained in the same cellular compartments, we incubated T. b. brucei 427-SRA-Ty with Alexa Fluor 594-conjugated TLF and subsequently stained the cells for SRA using the Ty antibody. Fluorescence microscopy showed that TLF (red) and SRA-Ty (green) overlap in their subcellular distributions (Fig. 7A to H). TLF and SRA colocalized to a subpopulation of small cytoplasmic vesicles in SRA-transfected T. b. brucei, but neither was found in the lysosome. We observed many SRA-containing vesicles without TLF, but few vesicles containing TLF alone, without SRA, were detected (Fig. 7D and H). These findings are consistent with a proposed mechanism of resistance to TLF resulting from reduced targeting of TLF to the lysosome and with the idea that the association of SRA and TLF is required for resistance (19).

**DISCUSSION**

Initial studies using subtractive hybridization methods suggested that human-serum-resistant and -sensitive African trypanosomes might differ by the expression of a single gene (27). This possibility was tested in an elegant series of transfection experiments, in which the SRA gene from T. b. rhodesiense was introduced into a human-serum-susceptible line of T. b. brucei and was found to be sufficient to confer resistance (51). These studies relied on transfection of an insect-borne developmental stage of T. b. brucei and subsequent transmission through the tsetse fly prior to analysis of transfecteds. This strategy, though certainly successful, is both cumbersome and potentially prone to artifacts due to changes in gene expression unrelated to the transgene. In this paper and elsewhere, direct transfection of bloodstream-stage T. b. brucei with SRA was shown to render the parasites resistant to both TLF and normal human serum (50). Transfected T. b. brucei cells express the SRA transgene at high levels, resulting in steady-state amounts of the SRA mRNA that are similar to those detected in human infectious T. b. rhodesiense. Using antibodies against the Ty epitope, we examined the intracellular distribution of SRA in T. b. brucei 427-SRA-Ty and found it to be largely localized to small cytoplasmic vesicles between the flagellar pocket and the nucleus. The incubation of T. b. brucei 427-SRA-Ty with Alexa-conjugated TLF (red) and SRA-Ty (green) showed a high degree of overlap in their subcellular distributions, indicating that SRA and TLF may interact with one another directly. In fact, studies have shown that SRA directly interacts with apol-I, one of the trypanolytic components of TLF (47). To examine whether SRA and TLF are contained in the same cellular compartments, we incubated T. b. brucei 427-SRA-Ty with Alexa Fluor 594-conjugated TLF and subsequently stained the cells for SRA using the Ty antibody. Fluorescence microscopy showed that TLF (red) and SRA-Ty (green) overlap in their subcellular distributions (Fig. 7A to H). TLF and SRA colocalized to a subpopulation of small cytoplasmic vesicles in SRA-transfected T. b. brucei, but neither was found in the lysosome. We observed many SRA-containing vesicles without TLF, but few vesicles containing TLF alone, without SRA, were detected (Fig. 7D and H). These findings are consistent with a proposed mechanism of resistance to TLF resulting from reduced targeting of TLF to the lysosome and with the idea that the association of SRA and TLF is required for resistance (19).

**FIG. 6.** Cellular localization of TLF relative to p67 in T. b. brucei 427 (A to D) and T. b. brucei 427-SRA-Ty (E to H) in the presence of chloroquine. The positions of the nucleus and the kinetoplast were visualized by staining with DAPI (blue). (A and E) Cells visualized by bright-field (BF) microscopy overlaid with DAPI staining. (B and F) Cells incubated with Alexa-conjugated TLF (red). (C and G) Cells stained with anti-p67 (green). (D and H) Overlays of images of cells incubated with Alexa-conjugated TLF and stained for DNA and p67.

**FIG. 7.** Colocalization of TLF-1 and SRA in T. b. brucei 427-SRA-Ty. The positions of the nucleus and the kinetoplast were visualized by staining with DAPI (blue). (A to H). (A and E) Cells visualized by bright-field (BF) microscopy overlaid with DAPI staining of the nucleus and kinetoplast. (B and F) Cells incubated with Alexa-conjugated TLF (red). (C and G) Cells stained with anti-Ty (green) to visualize SRA. (D and H) Overlaid images of cells treated with Alexa-conjugated TLF and stained for DNA and SRA.
trypanosomes. It is possible that the cell surface reactivity was simply a consequence of cross-reactivity of the polyclonal mouse antiserum against SRA with the VSG. This is a particular concern since SRA is a member of the VSG gene family and shares characteristics with other trypanosome cell surface proteins (6, 31). Alternatively, the amounts of SRA present on the cell surface in different trypanosome lines may vary. Regardless, the significance of the cell surface-associated SRA is questionable, since little or no SRA was found on the surfaces of *T. b. rhodesiense* 427-SRA-Ty cells yet they were highly resistant to both TLF-1 and human serum. Other labs have also shown that the localization of endogenous SRA in *T. b. rhodesiense* is primarily intracellular (47). In this paper, we showed that SRA-Ty localizes to an intracellular, nonlysosomal, vesicular location in transfected *T. b. brucei*.

We were concerned that high-level expression of SRA might result in aberrant intracellular localization. Transgenes are often expressed at abnormal levels due to the use of heterologous promoters. In our construct, SRA expression is driven from a constitutively active rRNA polymerase I promoter known to mediate high levels of mRNA synthesis in African trypanosomes (25, 41). The 5′ and 3′ untranslated regions were derived from tubulin sequences, which contribute posttranscriptionally to the stability of mRNAs in both bloodstream and procyclic forms. Previous studies have shown that SRA mRNA is an abundant transcript in both wild-type *T. b. rhodesiense* and transfected *T. b. brucei*, representing as much as 5 to 10% of the total mRNA (31, 51). This is comparable to the levels of VSG mRNA in African trypanosomes. We found that the amount of SRA mRNA in our transfecants was similar to the level of SRA mRNA expression in *T. b. rhodesiense*. Since SRA in *T. b. brucei* 427-SRA-Ty confers resistance to human serum and the amount of SRA mRNA is similar to that in wild-type *T. b. rhodesiense*, it seems likely that the SRA-containing cytosolic vesicles revealed by immunofluorescence microscopy are the primary sites of SRA activity.

The subcellular localization of SRA and how it prevents TLF from reaching the lysosome may provide a better understanding of both the function of SRA and the endocytic pathways of trypanosomes. This is in contrast to the proposed site of SRA activity being at the lysosome, where it interacts with apoL-1 (47). However, an important difference between these studies is the use of purified human HDLs containing both Hpr and apoL-1 in our studies rather than purified and recombinant apoL-1. The uptake and trafficking of native TLF and its interaction with SRA may differ from that of recombinant apoL-1. Endocytosis in trypanosomes shares many characteristics with endocytosis in other eukaryotic organisms but also exhibits several unique features, including developmental regulation, selective trafficking of proteins from the flagellar pocket, and the potential routing of resident lysosomal proteins through the flagellar pocket prior to localization to the lysosome (35). Vesicular trafficking between organelles occurs through the fusion of donor and specific acceptor membranes. This process is highly regulated and ensures proper directionality in protein sorting and packaging. Monomeric GTPases of the Rab family play a pivotal role in the control of membrane fusion and vesicle trafficking. Several *T. brucei* Rab proteins have now been identified that localize to specific subcellular compartments (14–17, 28, 35). The availability of antibodies to the *T. b. brucei* Rab proteins makes it possible to determine whether the cytosolic vesicles containing SRA and TLF are part of a vesicle recycling pathway.

Although we have shown that SRA confers human serum resistance when transfected into bloodstream-stage *T. b. brucei* and that SRA and TLF colocalize within cytoplasmic vesicles, the mechanism of SRA function is still unknown. One proposed mechanism is that inhibition may depend on interactions between SRA and apoL-1 in the lysosome (47, 48). However, we previously showed that TLF was not targeted to the lysosome in *T. b. rhodesiense*, whereas a naturally occurring human-serum-sensitive variant of *T. b. rhodesiense* transported TLF to the lysosome prior to cell lysis (19). The distributions of TLF in *T. b. rhodesiense* and in *T. b. brucei* 427-SRA-Ty appear to be somewhat different. In contrast to the localization of TLF to small cytoplasmic vesicles in *T. b. brucei* 427-SRA-Ty, TLF was largely localized at or near the flagellar pocket of *T. b. rhodesiense* (19). Therefore, it remains possible that SRA can influence TLF uptake and lysosomal trafficking at either the flagellar pocket, where receptor-mediated endocytosis initiates, or at later steps in the endocytic pathway.

Regardless of the precise role of SRA, we have shown that SRA expression is sufficient to confer resistance to TLF and normal human serum in *T. b. brucei* bloodstream-stage trypanosomes. Furthermore, we have shown that SRA is responsible for the rerouting of TLF within the trypanosome endocytic pathway, with the majority of the TLF becoming associated with nonlysosomal, SRA-containing vesicles. Future experiments will address whether SRA alters TLF localization by either recycling it out of the cell or directing it to a cellular compartment where TLF degradation is accelerated. The elucidation of mechanisms of TLF resistance could lead to the development of inhibitors of the SRA-mediated resistance pathway, thereby increasing the susceptibility of *T. b. rhodesiense*-mediated human sleeping sickness trypanosomes to TLF and thus leading to novel treatment of the disease.

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