Characterization of the recombination activities of the *Entamoeba histolytica*

Rad51 recombinase

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Graphical abstract

Abstract

The protozoan parasite responsible for human amoebiasis is *Entamoeba histolytica*. An important facet of the life cycle of *E. histolytica* involves the conversion of the mature trophozoite to a cyst. This transition is thought to involve homologous recombination (HR), which is dependent upon the Rad51 recombinase. Here, a biochemical characterization of highly purified *ehRad51* protein is presented. The *ehRad51* protein preferentially binds ssDNA, forms a presynaptic filament and possesses ATP hydrolysis activity that is stimulated by the presence of DNA. Evidence is provided that *ehRad51* catalyzes robust DNA strand exchange over at least 5.4 kilobase pairs. Although the homologous DNA pairing activity of *ehRad51* is weak, it is strongly enhanced by the presence of two HR accessory cofactors, calcium and Hop2-Mnd1. The biochemical system described herein was used to demonstrate the potential for targeting *ehRad51* with two small molecule inhibitors of human RAD51. We show that 4,4'-diisothiocyanostilbene-2,2'-disulfonic acid (DIDS) inhibited *ehRad51* by interfering with DNA binding and attenuated encystation in *Entamoeba invadens*, while B02 had no effect on *ehRad51* strand exchange activity. These results provide insight into the underlying mechanism of homology-directed DNA repair in *E. histolytica*.

Keywords: homologous recombination, *Entamoeba histolytica*, RAD51, DNA repair, recombinase inhibitors

1. Introduction

Homologous recombination (HR) is a critical pathway involved in the repair of chromosomes with deleterious DNA double-strand breaks (DSBs) and interstrand crosslinks that may occur upon exposure to chemical reagents and ionizing radiation. HR is also important for the repair of blocked or stalled replication forks and is crucial for the proper segregation of homologous chromosomes in Meiosis I [1-7]. When the cell encounters a DSB, the ends of the break are nucleolytically processed to yield 3' single-strand DNA (ssDNA) overhangs. The exposed 3' overhang serves as the nucleation site for a DNA recombinase to form a nucleoprotein filament known as the presynaptic filament, which is responsible for the homology search in the sister chromatid. Once homology is found, the nucleoprotein filament invades the homologous duplex DNA resulting in the displacement of the homologous strand to form a displacement loop (D-loop). Following the formation of a D-loop, the recombinase facilitates strand exchange [8], and the invading strand primes DNA synthesis to restore the damaged DNA. In most eukaryotes, there are two orthologs of the *Escherichia coli* RecA recombinase, Rad51 and Dmc1, which are responsible for the homology search and strand exchange of HR [9]. Dmc1 is a meiosis-specific recombinase, and Rad51 is ubiquitously expressed, yet plays a supporting role in meiosis [10]. The activities of the recombinases during HR are regulated by a variety of different accessory cofactors, including replication protein A (RPA) [11], calcium [12-16], and Hop2-Mnd1 [1, 13, 17-20].

Entamoeba histolytica is a protozoan parasite responsible for amoebic dysentery and amoebic liver abscess. Potential hosts are most commonly exposed to *E. histolytica* through contaminated food and water sources, which puts 50 million humans at risk for amoebic dysentery and liver abscesses yearly, with approximately 70,000 deaths worldwide [21-23]. The life cycle

69 of *E. histolytica* involves host ingestion of tetra-nucleated cysts that undergo several divisions to
70 produce eight trophic amoebae through a process known as excystation. The mature trophozoite
71 (containing a single nucleus) can colonize the colon [24], where it divides by binary fission and
72 feeds on bacteria, host cells, and host cell debris. As a result of unknown cues, the pathogen
73 completes the life cycle by duplicating its genome and forming tetra-nucleated, environmentally-
74 stable cysts in a process known as encystation. This process facilitates host-to-host spread. It is
75 unknown if there is genetic exchange among the four nuclei in the *E. histolytica* cyst; however,
76 *Giardia intestinalis*, another cyst-forming enteric pathogen, possesses two nuclei that may
77 exchange genetic information [25]. It is conceivable that the ability of *E. histolytica* to go through
78 diverse life stages, including genome duplications and spontaneous genome amplification, is likely
79 mediated through HR [26-28]. In support of this argument, the *E. histolytica* genome contains
80 genes homologous to the majority of the RAD52 epistasis group of genes [29-31]. The RAD52
81 epistasis group of genes accounts for the core DNA repair genes of HR (*RAD50*, *RAD51*, *RAD52*,
82 *RAD54*, *RAD55*, *RAD57*, *RAD59*, *RDH54*, *MRE11*, and *XRS2*), which are highly conserved
83 among eukaryotes [32]. In recent years, *E. histolytica* was shown to differentially express this
84 group of genes in response to DNA damage [29, 30], and HR was demonstrated *in vivo* [33]. Other
85 protozoan parasites are reported to utilize HR as a means for adaptation and survival in host
86 environments. *Trypanosoma brucei* employs the Rad51 recombinase to recombine variant surface
87 glycoproteins to evade the host immune response. Specifically, *T. brucei* parasites that contain
88 defective Rad51 show a decreased ability to undergo variant surface glycoproteins switching [34].
89 *Leishmania major* use HR to amplify antibiotic resistance genes and up-regulate the expression of
90 Rad51 in response to DNA damage [35]. The role of the Rad51 recombinase in *E. histolytica*
91 (*ehRad51*) is largely unknown. In this study, we purified recombinant *ehRad51* and biochemically

characterized its recombinase activities. Our results will be valuable for defining the mechanism of *ehRad51*-mediated recombination in *E. histolytica*.

2. Materials and methods

2.1 DNA substrates and small molecules

All oligonucleotides were purchased from Integrated DNA Technologies, and the sequences can be found in Table 1. H3 was used as ssDNA in the DNA binding assay. H3c was annealed to H3 for dsDNA in the DNA binding assay. OL83-1 (ssDNA) and the complex OL83-1/OL83-2 (dsDNA) were used in the oligonucleotide strand exchange assay. OL90 (ssDNA) was used in the nuclease protection assay and D-loop assay. All oligonucleotides indicated as radiolabeled were done so using T4 polynucleotide kinase and [³²P-γ]-ATP as described [15]. φX174 (+) virion ssDNA and φX174 replicative form I dsDNA were purchased from New England BioLabs; the φX174 replicative form I dsDNA was linearized with *Apa*LI (New England BioLabs). pBluescript was purified from *E. coli* using a Plasmid Giga kit (Qiagen). 4,4'-diisothiocyanostilbene-2,2'-disulfonic acid, DIDS was purchased from Sigma-Aldrich. (E)-2-(2-(pyridin-3-yl)vinyl)quinazolin-4(3H)-one, B02 was purchased from Ryan Scientific Inc. [36].

2.2 Isolation and modification of the gene encoding *E. histolytica* Rad51

The isolation and modification of the *E. histolytica* Rad51 gene (XM_648984) to create the pET-*ehRAD51*-(HIS)₆ bacterial expression plasmid was previously described [37].

2.3 Growth of *E. invadens*

E. invadens (strain IP-1) (1-2 X 10⁶ cells/ml) were seeded into LYI-S-2 [38] axenic medium in 15 ml glass screw cap tubes at 25°C in the absence or presence of DIDS (30 μM or 150 μM). After 48 hr, cell viability was assessed by hemocytometer-based cell counting with Trypan blue exclusion.

2.4 Encystation of *E. invadens*

To induce encystation, *E. invadens* trophozoites (5×10^6) were transferred to encystation medium, LYI-LG [39], in the absence or presence of DIDS (30 μ M or 150 μ M). After 48 hr, encystation efficiency was estimated by incubation in 0.1% (v/v) sarkosyl in PBS on ice for 40 min, which lyses trophozoites. This allowed the percentage of mature detergent-resistant cysts in a population to be calculated.

2.5 Expression and purification of *ehRad51*

The purification of *ehRad51* was performed as previously described [37] with modifications. Briefly, *ehRad51* was expressed in the BL21 (Novagen) strain of *E. coli* cells. The cells were grown at 37°C until an $OD_{600} = 1.0$ and induced with 0.4 mM IPTG at 16°C for 20 hr. The cells were harvested by centrifugation. All successive steps in the purification were performed at 4°C. The cell paste (40 g) was resuspended in Buffer A (50 mM Tris-HCl pH 7.5, 1 mM EDTA, 10% sucrose, 250 mM KCl, 1 mM 2-mercaptoethanol, 0.01% Igepal, 0.1 mg/ml lysozyme, 5 mM benzamide, 1 mM PMSF, and a combination of protease inhibitors at 5 μ g/ml: aprotinin, chymostatin, leupeptin, and pepstatin A) followed by sonication. The lysate was clarified using ultracentrifugation at 100,000 $\times g$ (Beckman Ti-45 rotor) for 90 min. The clarified lysate was incubated with 3 ml of Ni-NTA Sepharose (GE Healthcare) overnight. The lysate containing the Ni-NTA matrix was poured into a glass column with an internal diameter of 1 cm (BioRad). After a wash with Buffer B (20 mM KH_2PO_4 pH 7.5, 10% glycerol, 1 mM EDTA) containing 150 mM KCl, the bound protein was eluted from the Ni-NTA beads in Buffer B containing 150 mM KCl and 500 mM imidazole. The eluate containing *ehRad51* was loaded onto an 8 ml hydroxyapatite column (BioRad) equilibrated with Buffer B containing 100 mM KCl. The column was developed using a linear gradient of Buffer B containing 0 – 300 mM KH_2PO_4 . The peak fractions containing

ehRad51 were pooled and loaded onto a 2 ml Source 15Q column (GE Healthcare) equilibrated with Buffer B containing 100 mM KCl. The column was developed using a linear gradient of Buffer B containing 100 - 450 mM KCl. The fractions containing *ehRad51* were pooled, concentrated to 8 mg/ml, and frozen in aliquots at -80°C. The yield was 12 mg of purified *ehRad51*.

2.6 Expression and purification of other proteins

Existing protocols were used to express and purify *E. histolytica* Dmc1 (*ehDmc1*) [37], murine Hop2-Mnd1 [40], human RAD51, and human RPA [41].

2.7 ATP hydrolysis

ehRad51 (10 µM) was incubated with 1.5 mM [³²P-γ]-ATP in Buffer C (25 mM Tris-HCl pH 7.5, 0.1 mg/ml BSA, 1 mM DTT, 2.4 mM MgCl₂, 20 mM KCl) in the absence or presence of ϕX174 (+) virion ssDNA (90 µM nucleotides) or linearized ϕX174 replicative form I dsDNA (90 µM base pairs) at 37°C. At the indicated times, the reactions were stopped by the addition of an equal volume of 0.5 mM EDTA and subjected to thin-layer chromatography using polyethyleneimine-cellulose plates (Sigma Aldrich). The amount of ATP hydrolysis was determined using a phosphorimager (Typhoon FLA 7000, GE Healthcare). When testing the effects of DIDS on ATP hydrolysis activity, 67 µM of DIDS was incubated with *ehRad51* in the absence or presence of ϕX174 ssDNA or ϕX174 dsDNA. The reactions were stopped at the indicated times and treated as described above.

2.8 DNA binding

Increasing concentrations of *ehRad51* (as indicated) were incubated in Buffer D (25 mM Tris-HCl pH 7.4, 0.1 mg/ml BSA, 1 mM DTT, 2.4 mM MgCl₂, 2 mM ATP) in the presence of ³²P-radiolabeled H3 (ssDNA) or ³²P-radiolabeled H3/H3c (dsDNA) for 20 min at 37°C. Reaction

products were subjected to 12% non-denaturing gel electrophoresis. The gels were dried on Whatman cellulose chromatography paper (Sigma-Aldrich), and analyzed using a phosphorimager (GE Healthcare Life Sciences Typhoon FLA 7000). When testing the effect of DIDS on ssDNA binding, *ehRad51* (7 μ M) was incubated with 32 P-radiolabeled H3 (ssDNA) in the presence of increasing amounts of DIDS (5, 10, 15, 20, 30, and 40 μ M). Also, increasing concentrations of DIDS (20, 40, 80, 100, 150, and 200 μ M) were incubated with *ehRad51* (35 μ M) in the presence of 32 P-radiolabeled H3/H3c (dsDNA). The reactions products were processed and analyzed as described above.

2.9 Nuclease protection assay

To determine the ability of *ehRad51* to form a stable nucleoprotein filament, *ehRad51* (2.5 μ M) was incubated with 32 P-radiolabeled OL90 (3 μ M nucleotides) at 37°C in Buffer C containing 2 mM ATP for 1, 2.5, 5, 7.5, and 10 min (as indicated) at which point DNase I (2 units; Promega) was added to the reaction. The reaction was incubated for 15 min, and the reaction products were deproteinized with the addition of SDS (0.5%) and Proteinase K (1.5 μ g/ml). The products were separated using 12% native polyacrylamide gel electrophoresis. The gels were dried on Whatman cellulose chromatography paper (Sigma-Aldrich), and analyzed using a phosphorimager (GE Healthcare Life Sciences Typhoon FLA 7000). When testing the nucleotide dependence of filament stability, *ehRad51* (2.5 μ M) was incubated with 32 P-radiolabeled OL90 (3 μ M nucleotides) at 37°C in Buffer C that was nucleotide free or contained ATP, ADP, ATP- γ -S, and AMP-PNP (2 mM) for 10 min at which point DNase I was added to the reaction. The reactions were processed and analyzed as described above. When testing the filament stability in the presence of DIDS, we used the same amount of DNA and *ehRad51* used in the DNA binding.

Here, *ehRad51* (7 μ M) incubated with increasing concentrations of DIDS (5, 10, 15, 20, 30, and 40 μ M). The reactions were processed and analyzed as previously described.

2.10 Strand exchange with plasmid-length DNA substrates

Increasing concentrations of *ehRad51* (4, 6, 8, 10, 12, 14, and 16 μ M) were incubated with ϕ X174 virion ssDNA (30 μ M nucleotides) in Buffer D containing 16 mM creatine phosphate and 36 μ g/ml creatine kinase at 37°C. After a 10 min incubation, RPA (3.8 μ M) and 150 mM KCl were added to the reactions, followed by an additional 8 min incubation at 37°C. Addition of linearized double-stranded ϕ X174 DNA (30 μ M base pairs) and spermidine (4 mM) initiated the reaction. The reactions were deproteinized after 90 min by the addition of SDS (0.5%) and Proteinase K (0.5 μ g/ml). The reaction products were separated on a 0.9% agarose gel, stained with ethidium bromide, and visualized using Image Lab software (BioRad).

2.11 Oligonucleotide-based strand exchange

Increasing concentrations of *ehRad51* (0.25, 0.5, 0.75, and 1 μ M) were incubated with unlabeled OL83-1 (3 μ M nucleotides) in Buffer C containing 2 mM ATP for 10 min at 37°C. Duplex DNA (³²P-radiolabeled OL83-1/OL83-2; 3 μ M base pairs) and spermidine (4 mM) were added to the reactions followed by a 60 min incubation at 37°C before being deproteinized with the addition of SDS (0.5%) and Proteinase K (0.5 μ g/ml). When testing the nucleotide dependence, *ehRad51* (1 μ M) was incubated with unlabeled OL83-1 (3 μ M nucleotides) in Buffer C containing either no nucleotide, ATP, ADP, ATP- γ -S, or AMP-PNP (all at 2 mM) for 10 min at 37°C. The reactions were deproteinized as described above, and the reaction products were separated using 10% native polyacrylamide gel electrophoresis and analyzed using a phosphorimager (GE Healthcare Life Sciences Typhoon FLA 7000). To test the stimulatory effect of calcium or mHop2-Mnd1 on *ehRad51* DNA strand exchange activity, a time course analysis

was performed. The experiment was processed in the same manner as above, except for the incubation of *ehRad51* (1 μ M) in the absence or presence of CaCl_2 (2 mM), mHop2-Mnd1 (0.6 μ M; H2M1), or both CaCl_2 and mHop2-Mnd1. After the addition of duplex DNA and spermidine, the reaction products were deproteinized after 2.5, 5, 7.5, 10, 12.5, 15, 30, 45, and 60 min. The reaction products were separated on 12% polyacrylamide gels and analyzed using a phosphorimager. When testing *ehRad51* strand exchange activity in the presence of DIDS, *ehRad51* (1 μ M) was incubated with increasing concentrations of DIDS (2.5, 5, 10, 15, 20, 30, and 40 μ M) under the experimental conditions described above. The reaction products were analyzed as described above. When testing B02, the reactions were processed similar to a previous report [36]. Briefly, human RAD51 (1 μ M), *ehRad51* (1 μ M), or *ehDmc1* (1 μ M) was incubated with OL83-1 in Buffer C containing ATP (2 mM) and CaCl_2 (1 mM). Increasing concentrations (100, 200, and 300 μ M) of B02 (diluted in DMSO—reactions contained 6% DMSO final concentration) were added to the reactions and incubated for 20 min before the addition of duplex DNA and spermidine. An additional lane of human RAD51 was tested in the absence of DMSO under the same reaction conditions. The reactions incubated for 60 min before deproteinization. The reaction products were resolved and analyzed as described above.

2.12 mHop2-Mnd1 pull-downs

To determine the interaction of *ehRad51* with mHop2-Mnd1, an affinity pull-down was performed as described [37]. Briefly, mHop2-Mnd1 (3.5 mg) and BSA (5 mg) were immobilized on 0.5 ml of Affi-gel 15 matrix (BioRad), separately, according to the manufacturer's instructions. *ehRad51* (7 μ g) was incubated with 17.5 μ l of Affi-mHop2-Mnd1 or Affi-BSA in Buffer B containing 100 mM KCl at 4°C with agitation. After a 60 min incubation at 4°C, the supernatant was removed, and the beads were washed three times with Buffer B containing 100 mM KCl. An

equal volume of SDS loading dye (160 mM Tris-HCl pH 6.8, 60% glycerol, 4% SDS (w/v)) was added to each sample, and an aliquot of the supernatant, wash, and eluate were separated using 12% SDS-PAGE. The gel was stained with Coomassie blue.

2.13 D-loop assay

ehRad51 (2 μ M) was incubated with the 32 P-radiolabeled OL90 (4.5 μ M) in Buffer D containing 16 mM creatine phosphate and 36 μ g/ml creatine kinase for 5 min at 37°C. CaCl₂ (0.5 mM) was added to the reaction (where indicated) followed by a 5 min incubation at 37°C. This was followed by the addition of mHop2-Mnd1 (0.16 μ M; H2M1), where indicated, and another 5 min incubation at 37°C. The reactions were then initiated by the addition of pBluescript (35 μ M base pairs). At 0.5, 1, 2, 6, and 14 min the reactions were deproteinized with the addition of SDS (0.5%) and Proteinase K (0.5 μ g/ml). Reaction products were resolved using 0.9% agarose gel electrophoresis, dried on DE81 anion exchange paper (GE Healthcare), and analyzed using a phosphorimager (GE Healthcare Life Sciences Typhoon FLA 7000).

2.14 Statistical analysis

The *in vitro* assays each had a minimum of three separate experiments, and error bars for each graph represents SEM. The effect of DIDS on viability and encystation of *E. invadens* was carried out as three biological replicates. All values are given as means $\pm$ SD. Statistical analyses were performed using GraphPad Prism Version 6.05. Comparisons of two groups were carried out using an unpaired two-tailed T-test. *P* values less than 0.05 were considered statistically significant.

3. Results

3.1 Isolation and purification of *ehRad51*

E. histolytica was reported to overexpress *ehRad51* in response to DNA damage [29] suggesting that *ehRad51* was a functional recombinase. Lopez-Casamichana *et al.* (2008) provided evidence that partially purified recombinant *ehRad51* bound DNA and was capable of homologous DNA pairing to form a D-loop [29]. While these studies were important to show that *ehRad51* was indeed a functional recombinase, the lack of biochemical characterization is an impediment to understanding its role in HR in *E. histolytica*. Here, we sought to provide a thorough biochemical examination of *ehRad51* and its activities. We first cloned the *RAD51* gene from the *E. histolytica* genome. The sequence encoding the 6 histidine epitope tag was fused to the 5' end of the *ehRAD51* gene by PCR and was inserted into a bacterial expression plasmid to be used for protein production [37]. Using a combination of affinity and ion exchange chromatography, *ehRad51* was purified to near homogeneity (Figure 1A).

3.2 ATP hydrolysis by *ehRad51* is stimulated by the presence of DNA

Analysis of the primary sequence of the *RAD51* gene from *E. histolytica* revealed the presence of the conserved Walker A and B motifs (Figure 1 in [42]) known to be important for binding and hydrolysis of ATP. To determine if *ehRad51* possessed the ability to hydrolyze ATP, purified *ehRad51* was incubated with saturating amounts of [³²P-γ]-ATP. The reaction products were resolved using thin-layer chromatography, and the results from three independent experiments indicated relatively weak hydrolysis (7%; Figure 1B). Previous reports indicated the presence of DNA stimulated the ATP hydrolysis activity of human and yeast Rad51 [43, 44]. To determine if DNA stimulated the ATPase activity of *ehRad51*, we incubated *ehRad51* with saturating amounts of φX174 (+) virion ssDNA or linearized φX174 replicative form I dsDNA and [³²P-γ]-ATP. Our results show that DNA stimulated the ATP hydrolysis activity of *ehRad51* (Figure 1B). The greatest stimulation occurred in the presence of ssDNA (32%) while a modest

stimulation of ATP hydrolysis (17%) occurred in the presence of dsDNA (Figure 1B). These results are in agreement with those reported for Rad51 proteins from other organisms [43-47].

3.3 *ehRad51* binds DNA

The stimulation of the ATP hydrolysis activity of *ehRad51* by the presence of ssDNA or dsDNA suggested *ehRad51* might bind DNA. To address this notion, an electrophoretic mobility shift assay (EMSA) was performed. In this assay, increasing concentrations of *ehRad51* were incubated with ³²P-labeled ssDNA or dsDNA. The reaction products were resolved using non-denaturing polyacrylamide gel electrophoresis. Our results show that *ehRad51* binds both ssDNA and dsDNA in concentration-dependent manner (Figure 1C and 1D, respectively). The *ehRad51*-DNA complexes migrated with reduced mobility through the acrylamide gel. At sub-saturating concentrations, *ehRad51* appeared to display cooperative binding. *EhRad51* showed a preference for ssDNA over dsDNA as 5-fold more *ehRad51* was required to reach saturation than for ssDNA (Figure 1C and 1D, respectively). Our results show that *ehRad51* bound both ssDNA and dsDNA with a strong preference for ssDNA.

3.4 *ehRad51* forms a presynaptic filament

RecA and its eukaryotic orthologs, Rad51 and Dmc1, form right-handed helical filaments on ssDNA in the presence of ATP [13, 47-51]. To test the formation of a presynaptic filament, we utilized a nuclease protection assay [15, 37, 45, 52]. In this assay, *ehRad51* is incubated with ³²P-labeled ssDNA in the presence of ATP to allow for filament formation, followed by the addition of DNase I to degrade any exposed DNA. If *ehRad51* forms a filament, there will be protection of the ³²P-labeled ssDNA from DNase I. As shown in Figure 2A, *ehRad51* forms a stable nucleoprotein filament within one minute and maintains protection of the ³²P-labeled ssDNA during the course of the experiment (Figure 2A, lanes 3-7) in the presence of ATP. To determine

what nucleotides facilitated presynaptic filament formation by *ehRad51*, we performed the nuclease protection assay as described above with the exception that ATP was substituted with the slowly hydrolyzable ATP analog, ATP- γ -S; the non-hydrolyzable ATP analog, AMP-PNP; or ADP. Our results show that *ehRad51* formed a stable presynaptic filament in the presence of ATP and AMP-PNP (Figure 2B, lanes 3 and 6). The integrity of the filament was slightly compromised in the presence of ATP- γ -S (Figure 2B, lane 5). Surprisingly, in the presence of ADP, *ehRad51* was quite adept at forming a presynaptic filament that protected the 32 P-labeled ssDNA (Figure 2B, lane 7). The absence of nucleotide did not support filament formation by *ehRad51* (Figure 2B, lane 4). Taken together, these data suggest that formation of a stable *ehRad51* presynaptic filament is dependent upon binding a nucleotide but not its hydrolysis.

3.5 *ehRad51* mediates plasmid length DNA strand exchange

The capability of *ehRad51* to form a presynaptic filament led us to ask if *ehRad51* possessed the ability to promote DNA strand exchange. To address this question, we used a three-strand assay that utilizes ϕ X174 DNA viral (+) ssDNA and linearized ϕ X174 dsDNA that is 5.4 kilobase pairs in length to monitor DNA strand exchange [37, 44, 48]. In this assay, *ehRad51* was incubated with circular ϕ X174 ssDNA to allow for filament formation followed by the addition of RPA. RPA is a ssDNA binding protein that prevents the formation of secondary structure in ssDNA. The DNA strand exchange reaction is initiated by the addition of linearized ϕ X174 dsDNA. Early in strand exchange, a joint-molecule forms. The joint-molecule is extended by *ehRad51*, displacing the homologous linear ssDNA to produce a nicked-circular molecule (Figure 3A). In a concentration dependent manner, *ehRad51* catalyzed the formation of joint-molecules (Figure 3B, lane 2) that were extended by DNA strand exchange to yield a nicked-circular dsDNA product (Figure 3B, lanes 3-8). The optimal DNA strand exchange activity occurred at 10 μ M

ehRad51 (Figure 3B, lane 5 and Figure 3C). This corresponds to the expected molar ratio of three nucleotides per monomer of *ehRad51* protein that was reported for human and *Saccharomyces cerevisiae* Rad51 [44, 51, 53]. These findings demonstrate that *ehRad51* catalyzed robust DNA strand exchange over thousands of base pairs.

3.6 *ehRad51* catalyzes nucleotide-dependent DNA strand exchange

The ability of Rad51 to catalyze DNA strand exchange is dependent upon the ability to bind, but not hydrolyze ATP in many species [43, 45, 54-56]. Since *ehRad51* formed a presynaptic filament in the presence of ATP, AMP-PNP, ATP- γ -S, and ADP, we wished to determine if *ehRad51* required ATP binding or hydrolysis to catalyze DNA strand exchange using an oligonucleotide DNA strand exchange system. In this assay, *ehRad51* was incubated with an unlabeled ssDNA oligonucleotide to allow filament formation. The reaction was initiated by the addition of radiolabeled duplex DNA, where the ^{32}P -radiolabeled strand of the duplex DNA is homologous to the unlabeled ssDNA located within the *ehRad51*-ssDNA filament. If *ehRad51* promotes DNA strand exchange, the ^{32}P -oligonucleotide will be displaced from the dsDNA complex and exchanged for the unlabeled strand within the *ehRad51* filament (Figure 4A). We found that *ehRad51* catalyzed DNA strand exchange in the presence of ATP in a concentration dependent manner (Figure 4B, lanes 2-5). In the absence of a nucleotide, DNA strand exchange was not observed (Figure 4B, lane 6). When AMP-PNP was substituted for ATP, there was only a slight reduction (4%) in the amount of product formed (Figure 4B, compare lane 8 to 5). Substitution of ATP with ATP- γ -S or ADP failed to support *ehRad51*-mediated DNA strand exchange (Figure 4B, lanes 7 and 9). These results indicate that *ehRad51* facilitates DNA strand exchange that is dependent on nucleotide binding and not its hydrolysis.

3.7 *ehRad51* promotes weak D-loop formation

The fact that *ehRad51* can form a stable presynaptic filament that invades duplex DNA to promote DNA strand exchange suggests that it should be able to catalyze the formation of D-loops through homologous DNA pairing. To confirm this idea, a D-loop formation assay was performed [15, 37, 48]. In this assay, *ehRad51* was incubated with a ³²P-radiolabeled ssDNA oligonucleotide to form a presynaptic filament. The ssDNA oligonucleotide is homologous to a region within the supercoiled pBluescript dsDNA plasmid. Upon addition of the dsDNA plasmid, the *ehRad51* presynaptic filament searches for homology between the radiolabeled ssDNA and the dsDNA. Once homology is located, the presynaptic filament invades the duplex DNA, displacing the homologous strand to produce a D-loop (Figure 5A). Using this assay, a time course analysis was performed which revealed *ehRad51* weakly forms D-loop (0.5%, Figure 5B, lanes 2-6). The weak D-loop forming activity of *ehRad51* seen using our D-loop formation assay resembles the relatively low level of D-loop formation activity reported for human RAD51 [13]. In the same report, the presence of calcium enhanced the D-loop formation of human RAD51 [13]. Guided by this study, we examined the effect of calcium on *ehRad51* D-loop formation. Our results show that calcium increased the D-loop forming activity of *ehRad51* 4-fold (2%; Figure 5B, lanes 8-12).

3.8 *ehRad51* interacts with mHop2-Mnd1

HR accessory factors are known to modulate the activity of the Rad51 and Dmc1 recombinases [6]. One of these accessory factors, HOP2-MND1, interacts with and greatly enhances the D-loop formation activity of the human RAD51 and DMC1 recombinases [57, 58]. Petukhova *et al.* (2005) [40] showed that murine Hop2-Mnd1 could physically interact with both human RAD51 and DMC1 and stimulate their D-loop formation activity. This is likely due to the high degree of homology between the human and murine RAD51 (99% identity) and DMC1 (97% identity) recombinases (Figure 2 in [42]). *E. histolytica* has the *HOP2* and *MND1* genes and their

expression is upregulated during encystation in *E. histolytica* at the same time as *ehRad51* and *ehDmc1* [33]. Based on these studies, we wanted to determine if *ehHop2-Mnd1* would interact with *ehRad51* to promote the D-loop formation activity of *ehRad51*. Unfortunately, purified *ehHop2-Mnd1* protein complex is not currently available. Notably, most of what is known about Hop2-Mnd1 is derived from studies using murine Hop2-Mnd1 (mHop2-Mnd1) with human RAD51 and DMC1 [17, 20, 40, 59]. In addition to these studies, Ploquin *et al.* (2007) showed mHop2-Mnd1 interacted with *Schizosaccharomyces pombe* Rad51 (*SpRad51*) [19] despite lower sequence identity (74%) between *SpRad51* and murine Rad51 (Figure 2 in [42]). The ability of mHop2-Mnd1 to interact with human RAD51, human DMC1, *SpRad51*, and *ehDmc1* [17, 19, 37, 40, 60] combined with the sequence identity of *ehRad51* with murine RAD51 (65% identity) that is close to that of *SpRad51* and murine RAD51 (74% identity) (Figure 2 in [42]), led us to reason that mHop2-Mnd1 may interact with *ehRad51* thereby potentially enhancing its recombination activities. We first determined if mHop2-Mnd1 interacted with *ehRad51* by performing an affinity pull-down assay using either bovine serum albumin (BSA) or purified mHop2-Mnd1 immobilized on Affi-gel matrix [37]. In this reaction, *ehRad51* was incubated with Affi-mHop2-Mnd1 beads. The beads were extensively washed to remove non-specifically bound proteins. The remaining proteins bound to the Affi-mHop2-Mnd1 beads were eluted. As shown in Figure 6A, *ehRad51* was located in the elution fraction, indicating a physical interaction with mHop2-Mnd1 (Figure 6A, lanes 2-4). When *ehRad51* was incubated with Affi-BSA, *ehRad51* was located in the supernatant, indicating there was no physical interaction with BSA or non-specific interaction of *ehRad51* with the Affi-gel matrix (Figure 6A, lanes 5-7). Our findings indicate mHop2-Mnd1 interacts specifically with *ehRad51*. The sequence homology between the human and *E. histolytica* recombinases is 66% for Rad51 and 61% for Dmc1, and the conservation

between human and *E. histolytica* Hop2 and Mnd1 is 31% and 43%, respectively. The interaction between mHop2-Mnd1 and both *E. histolytica* recombinases [37] illustrates the evolutionary importance of the interaction between Hop2-Mnd1 and Rad51.

3.9 mHop2-Mnd1 stimulates *ehRad51* homologous DNA pairing

The interaction of mHop2-Mnd1 with *ehRad51* and the fact that mHop2-Mnd1 is reported to greatly enhance the D-loop formation catalyzed by human RAD51 [40, 57-59], human DMC1 [18, 40, 60], and *ehDmc1* [37] led us to ask if mHop2-Mnd1 would enhance *ehRad51*-mediated D-loop formation. Using our D-loop formation assay described above (Figure 5A), we show that inclusion of the mHop2-Mnd1 complex in the reaction greatly stimulated the homologous pairing activity of *ehRad51* (33%; Figure 6B, lanes 2-6). Addition of calcium to the reaction containing mHop2-Mnd1 only provided a slight increase (2-3%) in the amount of D-loop formed by *ehRad51* (Figure 6B, lanes 8-12). These data suggest *ehRad51* D-loop formation is strongly enhanced by mHop2-Mnd1 regardless of the presence of calcium. The ability of murine Hop2-Mnd1 to strongly enhance D-loop formation by the Rad51 recombinase from *E. histolytica* demonstrates that the role of Hop2-Mnd1 in HR is evolutionarily conserved and would suggest that upon availability, *ehHop2-Mnd1* would likely enhance the activity of *ehRad51*.

3.10 mHop2-Mnd1 and calcium stimulate *ehRad51* oligonucleotide strand exchange

To shed light on the effect calcium had on *ehRad51*-mediated recombination, we performed a time course analysis using our oligonucleotide DNA strand exchange assay described above (Figure 4A). Our results show that DNA strand exchange by *ehRad51* is weakly stimulated by the presence of calcium with DNA strand exchange product detected as early as 5 minutes (Figure 7A and 7B). The presence of mHop2-Mnd1 resulted in more than a 2-fold stimulation of *ehRad51*-mediated DNA strand exchange with DNA strand exchange product observed at 5

minutes (Figure 7A and 7B). The presence of both calcium and mHop2-Mnd1 provided the greatest enhancement (3-fold) of *ehRad51* activity with DNA strand exchange product detectable as early as 2.5 min (Figure 7A and 7B). These data, along with the D-loop results above, suggest that calcium and mHop2-Mnd1 function in separate but additive ways to promote DNA strand exchange mediated by *ehRad51*.

3.11 DIDS inhibits *ehRad51* strand exchange

Previously, we showed that the commercially available human RAD51 inhibitor DIDS (4,4'-diisothiocyanostilbene-2,2'-disulfonic acid) inhibited *ehDmc1* recombinase activities *in vitro* [37]. To determine if DIDS was able to inhibit *ehRad51*, we performed the oligonucleotide strand exchange assay (Figure 4A) with increasing concentrations of DIDS. DIDS inhibited *ehRad51* DNA strand exchange in a concentration dependent manner (Figure 8A) with an $IC_{50}=18.5 \pm 2.7$ μ M. Recently, a small molecule, B02, was reported to inhibit human RAD51 [36]. We wanted to determine if B02 could inhibit *ehRad51*-mediated DNA strand exchange. Although the strand exchange of human RAD51 is slightly attenuated in the presence of 6% DMSO (Figure 8B and 8C, compare lane 2 in both), B02 inhibited the strand exchange activity of human RAD51 (Figure 8B, lanes 3-5). In contrast, we show that the strand exchange activity of *ehRad51* is not affected by the presence of B02 at concentrations that inhibit human RAD51 (Figure 8B, lanes 7-10). Since we previously reported *ehDmc1* was sensitive to inhibition of D-loop formation by DIDS [37], we wanted to investigate whether B02 had an effect on *ehDmc1*. Similar to *ehRad51*, B02 had no effect on *ehDmc1*-mediated DNA strand exchange with concentrations that were able to inhibit human RAD51 (Figure 8B, lanes 12-15). Importantly, the ability of B02 to inhibit human RAD51 and not *ehRad51* or *ehDmc1* suggests there may be selectivity among inhibitors of Rad51 between species.

3.12 DIDS disrupts *ehRad51* activities through interference with DNA binding

To characterize the effect of DIDS on *ehRad51* functions, we first investigated the effect of DIDS on *ehRad51* ATP hydrolysis activity. Time course analysis revealed that DIDS stimulated the ATP hydrolysis activity of *ehRad51* from 7% to 16% (Figure 9A), similar to that of human RAD51 [61]. In the presence of dsDNA, there was minimal inhibition (1%) of *ehRad51* in the presence of DIDS. However, DIDS diminished the amount of ATP hydrolyzed by *ehRad51* in the presence of ssDNA by 35% (Figure 9A). This suggested DIDS may be binding to one of the DNA binding sites in *ehRad51*, which would interfere with the ability of *ehRad51* to bind ssDNA or dsDNA. To test this idea, we performed an EMSA to monitor the effect of DIDS on the DNA binding activity of *ehRad51*. Here, *ehRad51* was incubated with ³²P-labeled ssDNA or dsDNA in the presence of increasing amounts of DIDS. Our results show that DIDS similarly inhibited ssDNA and dsDNA binding by *ehRad51*. Specifically, DIDS started to inhibit ssDNA binding at 20 μ M (3:1 ratio of DIDS to *ehRad51*; Figure 9B, lane 6) and was completely abolished at 40 μ M DIDS (6:1 ratio of DIDS to *ehRad51*; Figure 9B, lane 8). By comparison, the ability of *ehRad51* to bind dsDNA began to be inhibited at 100 μ M DIDS (3:1 ratio of DIDS to *ehRad51*; Figure 9C, lane 7) and was complete at 150 μ M DIDS (4:1 ratio of DIDS to *ehRad51*; Figure 9C, lane 7). Interference of *ehRad51* ssDNA binding would likely interfere with *ehRad51* presynaptic filament dynamics. To test this idea, we used our nuclease protection assay described previously to test the stability of the *ehRad51* presynaptic filament in the presence of increasing concentrations of DIDS. Our results show that addition of 30 μ M DIDS resulted in complete loss of protection of the ssDNA from DNase I, indicating disruption of *ehRad51* presynaptic filament formation (4:1 ratio of DIDS to *ehRad51*; Figure 9D, lane 8). This concentration of DIDS is in agreement with the concentration required to disrupt ssDNA binding (Figure 9B). Together, these data indicate that DIDS interferes

with the ability of *ehRad51* to bind ssDNA. Our results are in agreement with the mechanism of DIDS inhibition reported for human RAD51 [61] and *ehDmc1* [37].

3.13 DIDS inhibits encystation but not proliferation in *E. invadens*

Encystation is thought to involve HR to reduce ploidy [33], and the key enzymes of HR, *ehRad51* and *ehDmc1* are both expressed during encystation [29]. Because of sensitivity to DIDS *in vitro*, we reasoned that inhibition of the *ehRad51* and *ehDmc1* recombinases with DIDS may disrupt encystation. Unfortunately, induction of encystation in *E. histolytica* has not been reported. Instead, we used the reptilian parasite *Entamoeba invadens* that is often used as a model for studying encystation in *Entamoeba* due to its ability to encyst *in vitro*. To determine the effect of DIDS on encystation of *E. invadens*, cells were induced to encyst in the absence or presence of DIDS (30 μ M or 150 μ M). In control cells, spherical refractile cysts were readily visualized by light microscopy and, on average, we observed 27.5 $\pm$ 9.7% encystation efficiency in control cells (data not shown) after 48 hr in encystation medium. In contrast, very few cysts were observed in DIDS-treated cultures; DIDS significantly inhibited encystation in a dose-dependent fashion (Figure 10A). To determine the effect of DIDS on the viability of *E. invadens*, cells were induced to encyst in the absence or presence of DIDS (30 or 150 μ M) for 48 hr, and the viability was measured as a percent of control cells. The final concentrations of control cells (5.6 $\pm$ 3.4 $\times 10^6$ cells/ml) or treated cells (30 μ M: 5.3 $\pm$ 0.98 $\times 10^6$ cells/ml; 150 μ M: 6.7 $\pm$ 3.6 $\times 10^6$ cells/ml) were not significantly different suggesting that the proliferation rate and viability were not hampered by DIDS (Figure 10B). Therefore, inhibition of encystation was not simply due to cell death.

4. Discussion

The results presented in this report significantly expand on the preliminary findings using partially purified *ehRad51* reported by Lopez-Casamichana *et al.* (2008). Intriguingly we found that many of the biochemical properties of *ehRad51* are similar to human RAD51, which is in agreement with the high degree of sequence conservation between the *ehRad51* and human RAD51 (66%; Figure 2 in [42]). For example, in our study, we confirmed that *ehRad51* binds ssDNA and dsDNA [29], and our results demonstrate that *ehRad51* preferentially bound ssDNA over dsDNA similar to human RAD51 [43]. We show the ATP hydrolysis activity of *ehRad51* was enhanced by the presence of dsDNA and ssDNA, with the greatest stimulation occurring in the presence of ssDNA, comparable to human and other Rad51 orthologs [43, 44]. Interestingly, the nucleoprotein complexes formed between *ehRad51* and ssDNA entered the gel, failing to form large aggregates, which was reported to occur for human RAD51 [43, 45]. Nonetheless, these nucleoprotein complexes form stable presynaptic filaments in the presence of ATP and AMP-PNP, providing nearly complete protection of the ssDNA in a nuclease protection assay. Although ADP provided some protection of the ssDNA, the presynaptic filament formed with this nucleotide was not as stable and failed to support DNA strand exchange, like human RAD51 [45]. While it is not clear why some recombinases fail to catalyze strand exchange in the presence of ATP- γ -S, it is possible that the *ehRad51* filament that formed in the presence of ATP- γ -S or ADP formed partially stable yet nonfunctional or inactive presynaptic filaments. In support of this idea, human RAD51 presynaptic filaments formed in the presence of ATP- γ -S and visualized with electron microscopy, are less extended than an *E. coli* RecA presynaptic filament [53] and resemble the inactive form of *E. coli* RecA filaments found in the absence of a nucleotide cofactor [62]. Studies with human RAD51 and *E. coli* RecA show that the ADP-bound RAD51 or RecA presynaptic filaments are in the inactive conformation, which results in inability to perform DNA strand

exchange [13, 63]. Human Dmc1 [48] and *ehDmc1* [37] also fail to form D-loop in the presence of ATP- γ -S, whereas human RAD51 was shown to have weak strand exchange activity in the presence of ATP- γ -S [43]. However, in an ATP-dependent manner, *ehRad51* effectively facilitated DNA strand exchange over several thousand bases with optimal activity occurring around three nucleotides per *ehRad51* monomer as seen with human and yeast Rad51 [44, 51, 53]. The amount of nicked circular dsDNA formed by *ehRad51* (~30%) is similar to that reported for *S. cerevisiae* Rad51 (36%, [44]).

The initial report on *ehRad51* by Lopez-Casamichana *et al.* (2008) revealed that partially purified *ehRad51* catalyzed D-loop formation [29]. We confirmed these results and show that *ehRad51* struggled to catalyze the formation of D-loop even in the presence of calcium, which is known to activate human RAD51 [13]. However, when the HR accessory factor Hop2-Mnd1 was present, *ehRad51* was able to efficiently form D-loops. The ability of Hop2-Mnd1 to enhance the homologous DNA pairing and DNA strand exchange activities of human RAD51 [40], *S. pombe* Rad51 [19], and *ehRad51* suggests the function of Hop2-Mnd1 is highly conserved. Because calcium adds to the stimulatory effect of Hop2-Mnd1 we suggest that calcium and Hop2-Mnd1 do not overlap in their mechanism of stimulation. Hop2-Mnd1 is reported to promote a conformational change in human RAD51 that promotes nucleotide and DNA binding [20], and calcium is also proposed to promote a conformational change in the human RAD51 filament [13]. It is possible that mHop2-Mnd1 induces a conformational change in *ehRad51* that allows more efficient binding of calcium. Our results demonstrating that mHop2-Mnd1 stimulates the activity of *ehRad51*, suggest the interaction of Hop2-Mnd1 with Rad51 is evolutionarily conserved. It is reasonable to propose that *ehHop2-Mnd1* will likely have a similar effect with *ehRad51*. Upon

availability of purified *ehHop2-Mnd1* protein complex, it will be of particular importance to test this hypothesis.

The fact that *ehRad51* and *ehDmc1* are simultaneously present in *E. histolytica* [29, 33, 37] suggested the pair of recombinases have complementary activities. Not surprisingly, the DNA binding and ATP hydrolysis activities between *ehRad51* and *ehDmc1* [37] were similar. Both *ehRad51* and *ehDmc1* formed stable presynaptic filaments in the presence of ATP and AMP-PNP. Though, the ability to form a presynaptic filament in the presence of ATP- γ -S and ADP was unique to *ehRad51*. D-loop formation in the absence or presence of calcium was much stronger with *ehDmc1* than *ehRad51*. The plasmid length DNA strand exchange catalyzed by *ehRad51* was significantly stronger than reported for *ehDmc1* [37]. *mHop2-Mnd1* robustly enhanced both *ehRad51* and *ehDmc1* mediated D-loop formation. Both *ehRad51* and *ehDmc1* catalyzed oligonucleotide DNA strand exchange, yet we found that *ehDmc1* was much more responsive to the presence of calcium, with or without *mHop2-Mnd1* present [37]. While these observations are solely from *in vitro* experiments, it is interesting to speculate that *ehDmc1* may be involved in early steps of HR that include invasion of the homologous partner and D-loop formation, while *ehRad51* may be more adept at later steps such as DNA strand exchange. Taken together, *ehRad51* and *ehDmc1* may complement each other during HR in *E. histolytica*. Future work will be necessary to test these ideas.

Previously, we demonstrated that DIDS, a small molecule inhibitor of human RAD51 [61], inhibited *ehDmc1*-mediated activities by disrupting DNA binding [37]. During the course of these studies, the small molecule B02 was reported to be a potent inhibitor of human RAD51 both *in vitro* [36] and *in vivo* [64, 65]. We examined the effect of the DIDS and B02 inhibitors on *ehRad51* activities. While DIDS inhibited *ehRad51* similar to the inhibition we observed with *ehDmc1*

[37], B02 failed to inhibit either *ehRad51* or *ehDmc1*. The failure of B02 to inhibit either *ehRad51* or *ehDmc1* is significant because it shows that small molecules such as B02 can discriminate between different Rad51 orthologs. Since our studies showed that DIDS inhibited *ehRad51* and *ehDmc1*-mediated activities [37] *in vitro*, we asked if DIDS would have an effect *in vivo*. While we did not see a decrease in cell viability under the conditions tested, there was a significant decrease in the level of encystation that occurred in the related *E. invadens*. This observation is in agreement with the current idea that HR is important for encystation to occur in *Entamoeba* [33]. Although, DIDS inhibits the activity of *ehDmc1* and *ehRad51* *in vitro* and decreases the extent of encystation, we are aware that DIDS may be inhibiting encystation through an unknown mechanism that does not depend on the recombinases [66]. In either case, our results demonstrate that small molecules may be tailored to target a specific recombinase, which could be beneficial in identifying potential therapeutics and for studying the role HR plays in the *E. histolytica* life cycle.

The majority of our biochemical understanding of the Rad51 recombinase comes from studies on human and yeast Rad51 proteins. A few biochemical studies on Rad51 from the eukaryotic pathogens *Plasmodium falciparum* [67], *Leishmania infantum* [68], *Ustilago maydis* [69], *Toxoplasma gondii* [70] reveal that Rad51 from these pathogens possess ATP hydrolysis activity, the ability to bind DNA and/or catalyze D-loop formation. Although the Rad51 recombinases from pathogens have relatively high identity and are closely related to other Rad51 recombinases (Figure 2 and 3 in [42]), there are differences in their activity. For example, *L. infantum* Rad51 was reported to preferentially bind ssDNA over dsDNA in a manner similar to *ehRad51* DNA binding [35]. Contrastingly, the D-loop formation activity of *L. infantum* Rad51 is significantly greater than observed for human RAD51 [68], *S. cerevisiae* Rad51 [14], or

ehRad51 in the absence of calcium. While human RAD51 and *ehRad51* are both stimulated by the presence of calcium [13], to our knowledge, it is not known whether this is the case for *L. infantum* RAD51 or any other Rad51 from a pathogen. Interestingly, *L. infantum* Rad51, *ehRad51* and human RAD51 possess ATP hydrolysis activity in the absence of DNA [35, 53]. This is distinct from *S. cerevisiae* and *T. gondii* Rad51 recombinases that possess DNA-dependent ATP hydrolysis activity [44, 45, 70]. It is possible that these differences observed in the activities of RAD51 from pathogens and human RAD51 may be exploited to develop a potential therapeutic.

HR is important to generate genetic diversity and promote proper chromosome segregation during meiosis. HR also plays a critical role in the repair of DSBs, interstrand cross-links, or DNA adducts in close proximity on opposite strands. DSBs can arise from exogenous sources such as exposure to ionizing radiation and endogenous sources as in the case of a stalled replication fork that occurs due to an attempt to replicate over single-strand break. These stalled replication forks associated with DSBs are reported to be repaired by HR [71-73]. If DSBs are left unrepaired, cell cycle checkpoints are activated potentially leading to cell death [74]. Therefore, it is critical that a DSB is recognized and efficiently repaired. During the life cycle of *Entamoeba*, a uninucleated trophozoite will undergo several rounds of replication to form a tetra-nucleated cyst. The additional rounds of replication increase the probability a stalled or damaged replication fork could occur. Since Rad51 is the key enzyme involved in the search for homology in the sister chromatid, which serves as a template for the repair of the DSB, it is likely to be important to reactivate stalled replication forks [75, 76] as *Entamoeba* goes through its life cycle. The relative importance of four nuclei in the cysts of *Entamoeba* species is unknown. Furthermore, genetic exchange among these nuclei has not been reported. However, the presence of transcriptionally active genes that encode the major meiosis-specific genes, including recombinases in *E. histolytica*, [29] supports

the idea that such genetic exchange could occur in this stage of the life cycle. Extensive genetic exchange among the nuclei could promote genetic diversity by creating polymorphisms. In support of this possibility, several studies demonstrate that the *E. histolytica* genome possesses extensive polymorphic sequences [77-79]. Indeed, some of these polymorphisms could positively impact fitness in the diverse environments both inside and outside the host. Examples of fitness gains that would benefit the parasite in the host may include acquisition of drug resistance or loss of immune-modulatory antigens. Genetic exchange mediated by *ehDmc1* [37] and *ehRad51* (current study) is likely to contribute significantly to the process of generating diverse genomes in multiple life stages of this pathogen.

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Figure Legends

Figure 1. *ehRad51* hydrolyzes ATP and binds DNA. (A) Purified recombinant *ehRad51* (0.5 μ g) on a 12% SDS-polyacrylamide gel stained with Coomassie blue. (B) Time course analysis of *ehRad51* ATPase activity in the absence and presence of ϕ X174 ssDNA or linearized ϕ X174 dsDNA. (C) Increasing concentrations of *ehRad51* (lanes 2-6) were incubated with 32 P-labeled ssDNA, and were resolved on a 12% polyacrylamide gel. (D) Increasing concentrations of *ehRad51* (lanes 2-6) were incubated with 32 P-labeled dsDNA. The samples were resolved on a 12% polyacrylamide gel. The results for C and D were quantified using a phosphorimager and graphed. Lane 1 for C and D contained no protein, and lane 7 for C and D was treated with SDS/PK (S/P). Error bars represent SEM (n = 3).

Figure 2. *ehRad51* forms a nucleoprotein filament. (A) Time course analysis for *ehRad51* presynaptic filament formation on 32 P-labeled ssDNA (ss) in the presence of DNase I (lanes 3-7). (B) Presynaptic filament formation in the absence and presence of ATP, AMP-PNP, ATP- γ -S, or ADP, as indicated. Samples were resolved on native polyacrylamide gels and analyzed using a phosphorimager. For both (A) and (B), lane 1 contained only 32 P-labeled ssDNA, and lane 2 contained 32 P-labeled ssDNA and DNase I. The results are from three independent experiments and are presented as percent protection. Error bars represent SEM. Deg, degraded 32 P-labeled ssDNA.

Figure 3. *ehRad51* facilitates plasmid length homologous DNA pairing and DNA strand exchange. (A) Schematic of plasmid length strand exchange assay. C_{ss}, circular ssDNA; l_{ds}, linearized dsDNA; j_m, joint-molecule; n_c, nicked-circular; l_{ss}, linearized ssDNA. (B) Increasing concentrations of *ehRad51* (lanes 2-8) were incubated with circular ϕ X174 ssDNA (c_{ss}). The reaction was initiated with the introduction of linearized ϕ X174 dsDNA (l_{ds}) and was deproteinized after 90 min. Reaction products were resolved on an agarose gel and stained with ethidium bromide. Lane 1 contained no protein. (C) The percent product of nicked-circular and total product (j_m + n_c) were graphed.

Figure 4. *ehRad51* mediates oligonucleotide DNA strand exchange. (A) Schematic of oligonucleotide DNA strand exchange assay; * represents ³²P-radiolabeled oligonucleotide. (B) *ehRad51* (0.25 μ M, lane 2; 0.5 μ M, lane 3; 0.75 μ M, lane 4; 1 μ M, lanes 5-9) was incubated with unlabeled ssDNA in the presence (lanes 2-5) or absence (lane 6) of ATP, or in the presence of ATP- γ -S (lane 7), AMP-PNP (lane 8), or ADP (lane 9) followed by the addition of radiolabeled dsDNA complex. Lane 1 contained no protein. Reaction products were deproteinized and separated on a 12% non-denaturing polyacrylamide gel followed by analysis of the gels using a phosphorimager. Results from three separate experiments were graphed with DNA strand exchange activity presented as percent product of the displaced ³²P-ssDNA. Error bars represent SEM.

Figure 5. Calcium stimulates the homologous DNA pairing activity of *ehRad51*. (A) Schematic of D-loop formation assay. * represents ³²P-radiolabeled oligonucleotide. (B) *ehRad51* was incubated with ³²P-ssDNA (ss) in the absence (lanes 2-6) or presence of calcium (lanes 8-12). Reactions were initiated by the addition of supercoiled duplex DNA (sc) and were deproteinized at the indicated times. Lane 1 contained no protein, and lane 7 contained calcium and no protein. Reaction products from three independent experiments were analyzed by resolving with agarose gel electrophoresis and quantified using a phosphorimager. Error bars represent SEM.

Figure 6. mHop2-Mnd1 physically interacts with *ehRad51* and enhances *ehRad51* homologous DNA pairing activity. (A) *ehRad51* was incubated with mHop2-Mnd1 (H2M1) immobilized on Affi-gel or bovine serum albumin (BSA) immobilized on Affi-gel. The supernatant was removed, the beads were washed, and interacting proteins eluted with SDS. Aliquots of the supernatant (S), wash (W), and eluate (E) for Affi-mHop2-Mnd1 (lanes 2-4) and Affi-BSA (lanes 5-7) were separated using 12% SDS-PAGE and stained with Coomassie blue. Lane 1 contained molecular weight markers (sizes indicated). (B) Time course analysis of *ehRad51* incubated with ³²P-radiolabeled ssDNA for filament formation in the absence (lanes 2-6) or presence (lanes 8-12) of calcium, followed by the addition of mHop2-Mnd1 and supercoiled duplex DNA. Reactions were deproteinized at the indicated times, subjected to agarose gel electrophoresis, and analyzed using a phosphorimager. Lane 1 contained mHop2-Mnd1 and no *ehRad51*, and lane 7 contained mHop2-Mnd1 and calcium but no *ehRad51*. Results are represented as percent D-loop formed from three independent experiments. Error bars represent SEM.

Figure 7. *ehRad51* oligonucleotide DNA strand exchange activity is stimulated by calcium and mHop2-Mnd1. (A) Time course analysis of *ehRad51* in the absence (top row) or presence of calcium (Ca²⁺, second row), mHop2-Mnd1 (H2M1; third row), or both calcium and mHop2-Mnd1 (Ca²⁺ + H2M1, bottom row). Lane 1 for each condition contained no *ehRad51*. Reactions were deproteinized at the indicated time points, resolved using native polyacrylamide gel electrophoresis, and analyzed using a phosphorimager. (B) Results from three independent experiments of each condition were quantified and graphed as percent product. Error bars represent SEM.

Figure 8. *ehRad51* DNA strand exchange activity is inhibited by DIDS but not B02. (A) DNA strand exchange analysis of *ehRad51* with increasing concentrations of DIDS (2.5, 5, 10, 15, 20, 30, and 40 μ M; lanes 3-9, respectively). (B) DNA strand exchange analysis in the presence and absence of increasing concentrations of B02 (100, 200, 300 μ M) with human RAD51 (hRAD51; lanes 3-5), *ehRad51* (lanes 8-10), and *ehDmc1* (lanes 13-15). Each B02 reaction contained 6% DMSO. (C) A control reaction for human RAD51 was performed in the absence of DMSO. The reactions were initiated by the addition 32 P-radiolabeled duplex DNA. After deproteinization, the reaction products were resolved using non-denaturing PAGE. Lane 1 for A and C and lanes 1, 6, and 11 for B contained no protein. Lane 2 for A and C and lanes 2, 7, and 12 for B were tested in the absence of DIDS or B02 for the indicated recombinase. Percent product formation was graphed, and error bars represent SEM (n = 3).

Figure 9. DIDS disrupts the DNA binding activity of *ehRad51*. (A) Time course analysis of *ehRad51* ATPase activity in the presence or absence of DIDS (67 μ M), with and without ϕ X174 ssDNA or linearized ϕ X174 dsDNA. Reactions were stopped with the addition of EDTA at the indicated times prior to separation with thin-layer chromatography and phosphorimager analysis. (B) *ehRad51* (7 μ M) incubated with 32 P-radiolabeled ssDNA and increasing amounts of DIDS (5 μ M, 10 μ M, 15 μ M, 20 μ M, 30 μ M, 40 μ M; lanes 3-8 respectively). Lane 1 contained no protein or DIDS. Lane 2 contained no DIDS. Lane 9 contained 40 μ M DIDS without *ehRad51*. (C) *ehRad51* (35 μ M) incubated with the 32 P-radiolabeled dsDNA and increasing concentrations of DIDS (20 μ M, 40 μ M, 80 μ M, 100 μ M, 150 μ M, 200 μ M; lanes 3-8 respectively). Lane 1 lacked protein and DIDS, lane 2 contained no DIDS and lane 9 contained 200 μ M DIDS and no protein. (D) *ehRad51* (7 μ M) was incubated with radiolabeled ssDNA in the absence (lane 3) and presence of increasing concentrations of DIDS (5 μ M, 10 μ M, 15 μ M, 20 μ M, 30 μ M, and 40 μ M; lanes 4-9) prior to the addition of DNase I. The reaction were deproteinized, and the products were separated using non-denaturing PAGE. Lane 1 contained radiolabeled ssDNA alone, lane 2 contained DNase with radiolabeled ssDNA, and lane 10 contained radiolabeled ssDNA in the presence of 40 μ M DIDS and DNase I. Error bars represent SEM (n = 3).

Figure 10. DIDS treatment attenuates *E. invadens* encystation but not viability. (A) Cyst formation was scored after 48 hr in encystation medium with or without DIDS (* P <0.05). (B) Cells were incubated with DIDS for 48 hr, after which viability was determined. The data from both experiments represent the means $\pm$ SD (n = 3).

Table 1. List of oligonucleotide names, sequences, and the assays in which they were used.

Name	Assay	Sequence (5' – 3')
H3	DNA binding	TTGATAAGAGGTCATTTGAATTCATGGCTTAGAGCTTAA TTGCTGAATCTGGTGCTGGGATCCAACATGTTTTAAATAT G
H3c	DNA binding	CATATTTAAACATGTTGGATCCCAGCACCAGATTCAGC AATTAAGCTCTAAGCCATGAATTCAAATGACCTCTTATC AA
OL83-1	Strand exchange	AAATGAACATAAAGTAAATAAGTATAAGGATAATACAA AATAAGTAAATGAAAAACATAGAAAATAAAGTAAAGGA TATAAA
OL83-2	Strand exchange	TTTATATCCTTTACTTTATTTTCTATGTTTATTCATTTACT TATTTTGTATTATCCTTATACTTATTTACTTTATGTTTCATT T

OL90	D-loop/Nuclease protection	AAATCAATCTAAAGTATATATGAGTAAACTTGGTCTGAC AGTTACCAATGCTTAATCAGTGAGGCACCTATCTCAGCG ATCTGTCTATTT
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