Activity of Megazol, a Trypanocidal Nitroimidazole, Is Associated with DNA Damage

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New drugs are urgently required for treatment of human African trypanosomiasis (22). One compound with promise is megazol (5, 12, 13, 18), a nitroheterocyclic compound that forms a nitro radical anion upon reduction (31). Megazol appears to enter Trypanosoma brucei via passive diffusion (3), but little is known about its mode of action. We determined the effect of megazol in assays that measured oxidative and reductive stress. We also selected parasites resistant to megazol in order to assess cellular changes associated with resistance.

Bloodstream forms (17) and procyclic forms (6) of T. brucei brucei (strain 427) were cultivated by using standard techniques. Assays measuring drug sensitivity were carried out to determine the drug concentration producing a 50% decrease in cell proliferation (IC50) and the resistance factor (average concentration of 107 cells ml−1 in culture medium. Megazol (0.15 to 30 μM) or buthionine sulfoximine (BSO; 10 to 20 μM) was added to the parasite suspensions, and the suspensions were incubated for 2 h. Reduced thiol levels in control and drug-treated parasites were then determined by derivatization with monobromobimane (thiolyte) and separation by high-pressure liquid chromatography as previously described (30).

BSO (16) reduces glutathione (2) and trypanothione (15) levels in trypanosomatids. When added to cell cultures 24 h prior to evaluation of IC50s, BSO enhanced the susceptibilities of both wild-type and megazol-resistant trypanosomes to megazol. At 10 and 20 μM, respectively, BSO caused the IC50 for wild-type procyclics to drop from 0.28 ± 0.01 μM and for bloodstream forms the IC50 was 0.15 ± 0.02 μM.

To induce megazol resistance in vitro, trypanosomes were exposed to drug concentrations that doubled every 2 weeks, starting at 0.008 and 0.01 μM for bloodstream forms and procyclies, respectively. After 6 months of cultivation, procyclic and bloodstream forms that tolerated 10 and 1 μM megazol, respectively, were cloned by limiting dilution.

The IC50 for the cloned, resistant, procyclic line was 29.24 ± 3.2 μM (105-fold-reduced sensitivity), while the IC50 for the bloodstream form line was 3.2 ± 0.48 μM (21-fold-reduced sensitivity).

Cross-resistance to megazol and other trypanocides was also studied (Table 1). Moderate levels of cross-resistance to two nitroheterocyclic trypanocides, the nitrofuran nifurtimox and the nitroimidazole benznidazole, were detected. Significant, albeit low-level, cross-resistance to the melaminophenyl arsenic compounds cymelarsan, melarsen oxide, and melarsoprol, the diamidine compounds pentamidine and beraenil, and suramin was also observed.

Verapamil (20), PAK-104P (7), and two phenothiazine derivatives (prochlorperazine and trifluparazine) (14) all inhibit cellular extrusion pumps but, when used at 5 μM, failed to reverse megazol resistance in T. brucei. Efflux pumps are thus unlikely to play a significant role in the megazol resistance characterized here.

It was recently shown that megazol exposure reduces trypanothione levels in Trypanosoma cruzi (23). Bloodstream form T. brucei obtained from infected Wistar rats (21) and procyclic forms grown in vitro (6) were resuspended at a concentration of 107 cells ml−1 in culture medium. Megazol (0.15 to 30 μM) or buthionine sulfoximine (BSO; 10 to 20 μM) was added to the parasite suspensions, and the suspensions were incubated for 2 h. Reduced thiol levels in control and drug-treated parasites were then determined by derivatization with monobromobimane (thiolyte) and separation by high-pressure liquid chromatography as previously described (30).

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TABLE 1. Cross-resistance of bloodstream and procyclic forms of T. brucei brucei strain 427 to megazol and other trypanocidesa

<table>
<thead>
<tr>
<th>Drug</th>
<th>IC50 (μM)</th>
<th>Bloodstream forms</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Wild type</td>
<td>Resistant (factor of resistance)</td>
</tr>
<tr>
<td>Megazol</td>
<td>0.28 ± 0.01</td>
<td>29.24 ± 3.18 (104)*</td>
</tr>
<tr>
<td>Nifurtimox</td>
<td>4.89 ± 0.12</td>
<td>37.12 ± 3.84 (7.6)*</td>
</tr>
<tr>
<td>Benznidazole</td>
<td>2.89 ± 0.20</td>
<td>12.87 ± 1.29 (4.5)**</td>
</tr>
<tr>
<td>Suramin</td>
<td>15.59 ± 2.61</td>
<td>66.06 ± 4.15 (4.2)**</td>
</tr>
<tr>
<td>Berenil</td>
<td>10.99 ± 1.50</td>
<td>41.14 ± 1.31 (3.7)*</td>
</tr>
<tr>
<td>Pentamidine</td>
<td>2.14 ± 0.03</td>
<td>5.06 ± 0.23 (2.4) (NS)</td>
</tr>
<tr>
<td>Melarsen oxide</td>
<td>0.02 ± 0.00</td>
<td>0.07 ± 0.08 (3.5)*</td>
</tr>
<tr>
<td>Melarsoprol</td>
<td>2.38 ± 0.08</td>
<td>2.60 ± 0.02 (1.1) (NS)</td>
</tr>
<tr>
<td>Cyemelarsan</td>
<td>0.06 ± 0.00</td>
<td>0.13 ± 0.00 (2.2)**</td>
</tr>
<tr>
<td>Ethidium bromide</td>
<td>12.15 ± 1.52</td>
<td>16.89 ± 1.73 (1.4) (NS)</td>
</tr>
<tr>
<td>Isomotamidium</td>
<td>9.80 ± 0.79</td>
<td>11.44 ± 0.79 (1.2) (NS)</td>
</tr>
<tr>
<td>Brilliant green</td>
<td>0.016 ± 0.00</td>
<td>0.053 ± 0.00 (3.3)</td>
</tr>
</tbody>
</table>

a Sensitivities of wild-type and selected megazol-resistant lines to a number of drugs were determined. The factors of resistance indicated in parentheses were calculated as ratios of IC50 for resistant parasites to those for wild-type parasites. Values that were signiﬁcantly different from that for the control, by Student’s t test, are designated by an asterisk(s) (*, P < 0.001; **, P < 0.01; ***, P < 0.05; NS, not significant at P = 0.05). Values are means ± standard deviations (n = 3 independent experiments).

TABLE 2. Activity of megazol against wild-type and RAD51−/− T. brucei bloodstream forms

<table>
<thead>
<tr>
<th>Drug</th>
<th>IC50 (μM)</th>
<th>RAD51−/− mutants</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Wild type</td>
<td></td>
</tr>
<tr>
<td>Megazol</td>
<td>0.13 ± 0.0025</td>
<td>0.029 ± 0.002</td>
</tr>
<tr>
<td>Megazol with NAC</td>
<td>0.13 ± 0.0200</td>
<td>ND</td>
</tr>
<tr>
<td>Nifurtimox</td>
<td>3.37 ± 0.19</td>
<td>3.40 ± 0.15</td>
</tr>
<tr>
<td>Nifurtimox with NAC</td>
<td>12.72 ± 1.1</td>
<td>ND</td>
</tr>
</tbody>
</table>

a Values are means ± standard deviations (n = 3 independent experiments). The effect of 0.5 mM NAC was assessed by adding this concentration to wild-type cells only prior to addition of drug. ND, not determined.

The nitroheterocyclic compounds are generally believed to exert their cytotoxic effects only after activation by single electron reduction of their corresponding nitro-anion radicals (27). In bacteria, the toxic effects of metronidazole are accompanied by damage to DNA (25, 29) and cells defective in DNA repair become hypersensitive to the action of metronidazole (11). A principal enzyme involved in eukaryotic DNA repair is RAD51. T. brucei mutants lacking this enzyme (24) were hypersensitive to metronidazole, indicating that the toxic effects of megazol are accompanied by damage to DNA (Table 2). In contrast, the susceptibility of RAD51 deletion mutants to the trypanocidal nitrofuran nifurtimox was similar to that of wild-type cells. N-acetylcysteine (NAC) antagonizes oxidative stress by interacting with a number of the reduced oxygen species (4, 8). When coadministered at 0.5 mM (the highest concentration at which no detrimental effect on trypanosomes was induced by this compound), NAC had no antagonistic effect on the action of megazol while it did have a modest protective effect against the action of nifurtimox (Table 2).

The activity of nifurtimox in T. brucei therefore appears to be mediated through reduced oxygen species, produced during futile redox cycling with reduced drug. Megazol’s activity, however, derives from direct damage exerted by the reduced megazol nitro anion radical derivatives. In T. cruzi, nifurtimox is also known to exert its activity through redox cycling (9, 10). Recently it was shown that nifurtimox stimulated an increase in intracellular trypanothione, which is the main antioxidant in trypanosomatids. Mol. Biochem. Parasitol. 62:15–20.

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REFERENCES