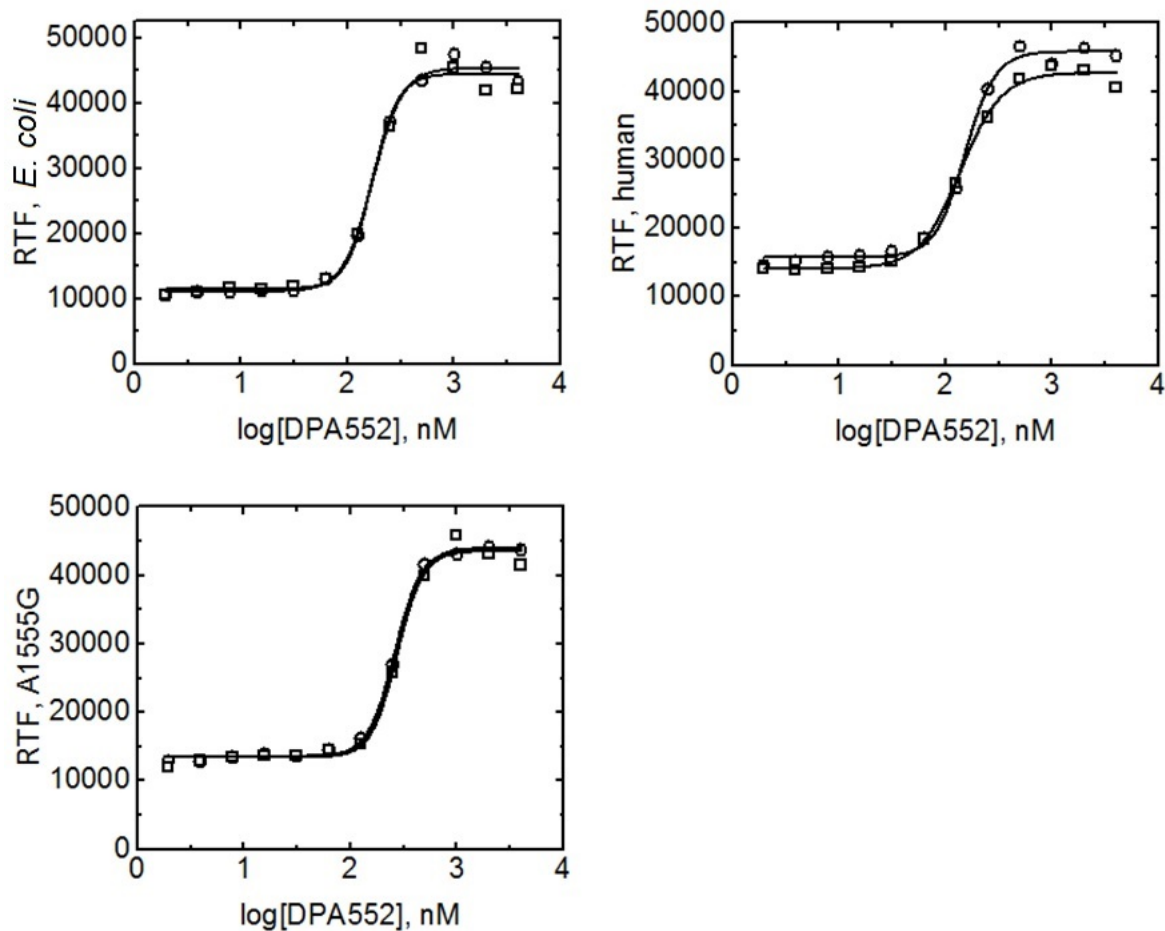


## SUPPORTING INFORMATION

### Antimicrobial Activity, AME Resistance and A-site Binding Studies of Anthraquinone-Neomycin Conjugates

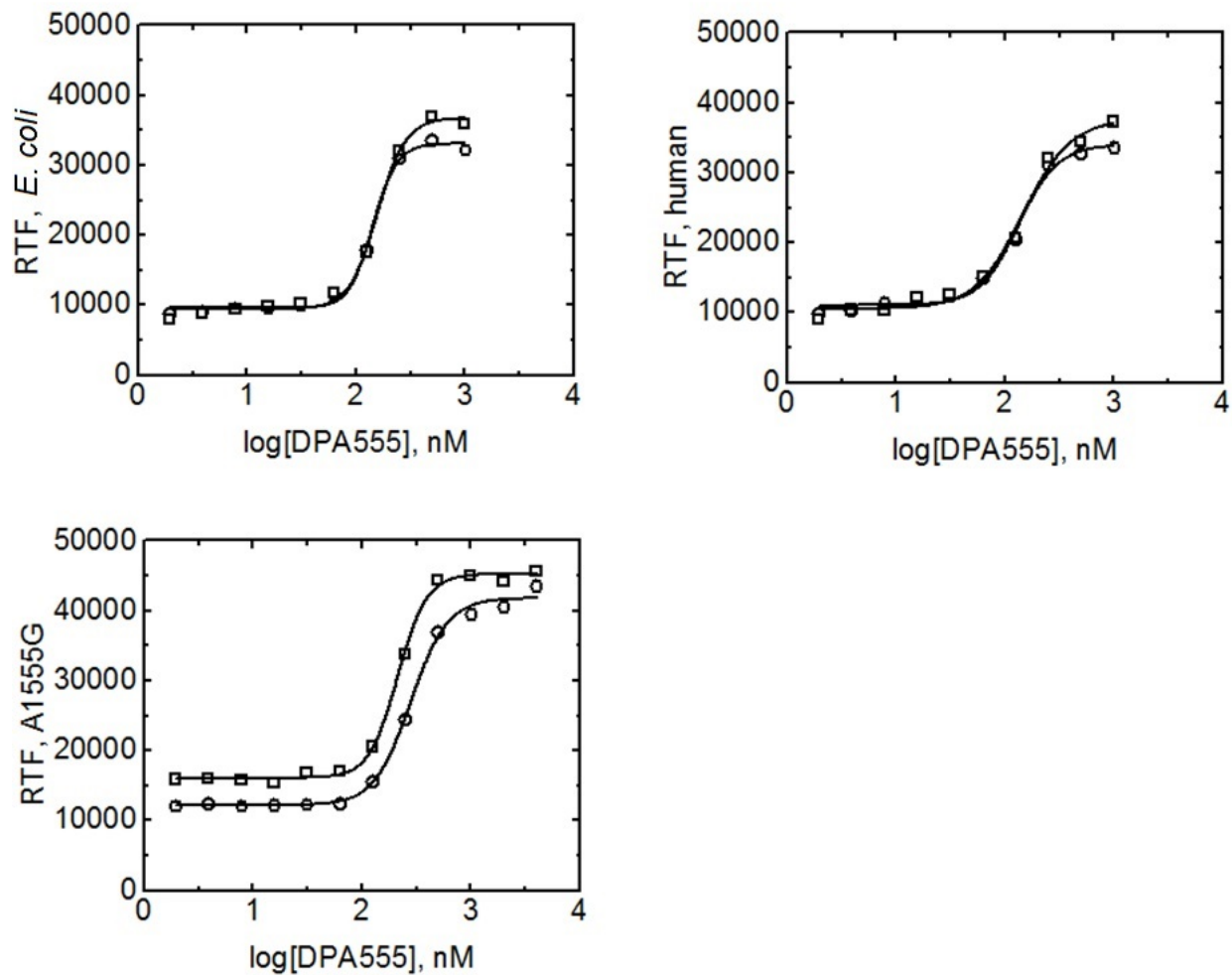
Natalya N. Degtyareva,<sup>a</sup> Changjun Gong,<sup>b</sup> Sandra Story,<sup>a</sup> Nathanael S Levinson<sup>d</sup>,  
Adegboyega K Oyelere<sup>d</sup>, Keith D. Green,<sup>c</sup> Sylvie Garneau-Tsodikova,<sup>c</sup> and Dev P. Arya,<sup>a,b</sup>

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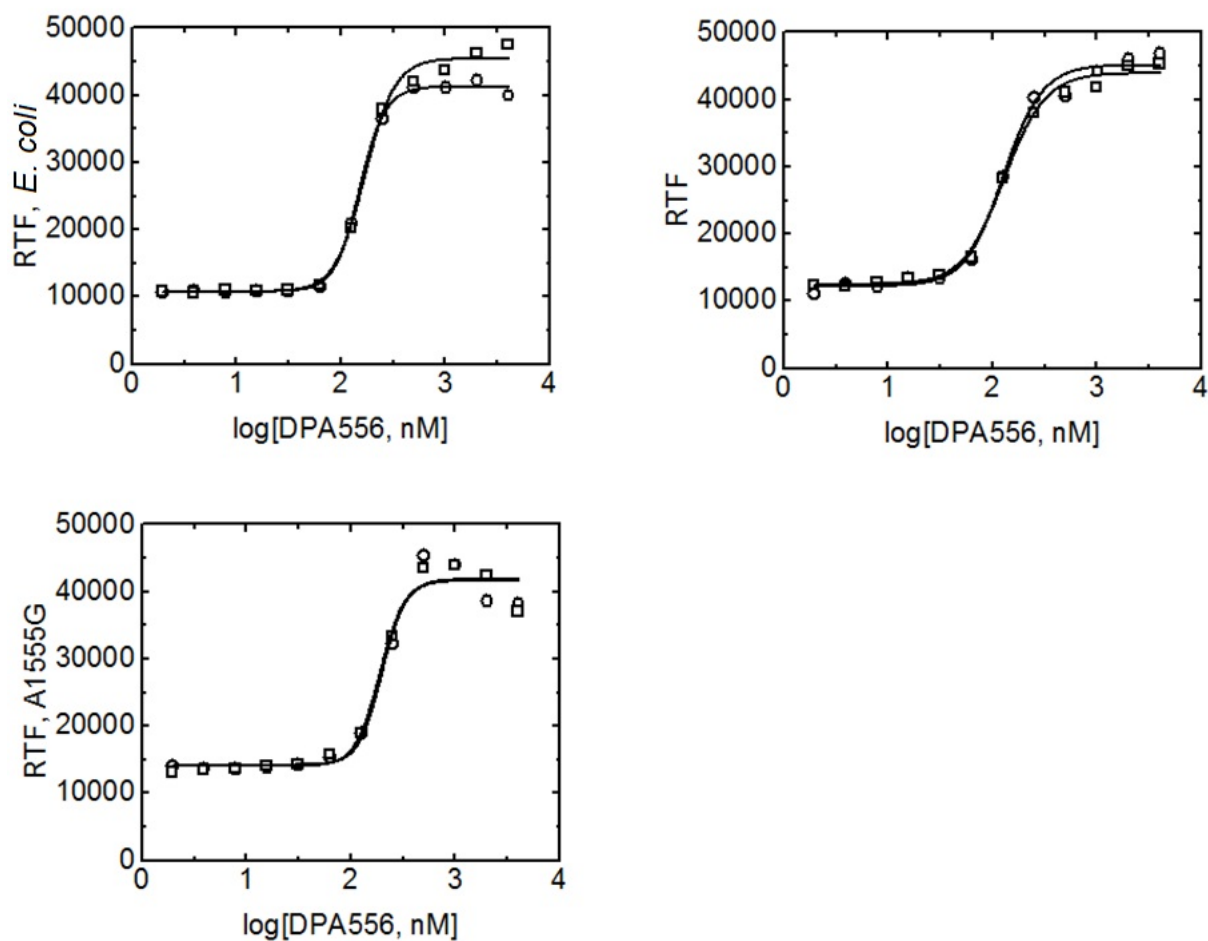


**Figure S1.** Titration plots of 100 nM A-site:F-NEO complex with compound **2**. IC<sub>50</sub> values for *E. coli*, human cytosolic, and A1555G A-sites were 173 ± 1 nM, 146 ± 10 nM, and 270 ± 10 nM, respectively.



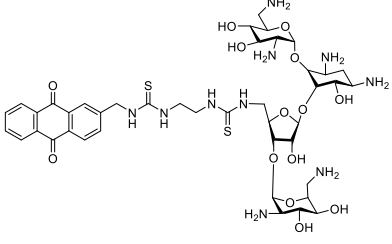
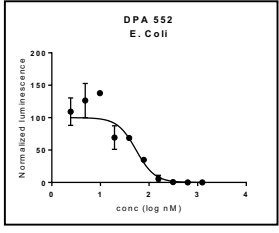
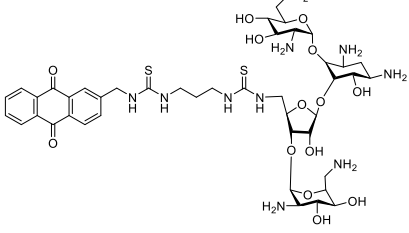
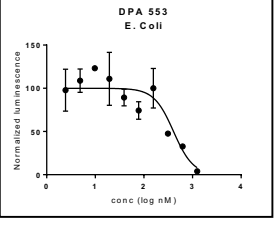
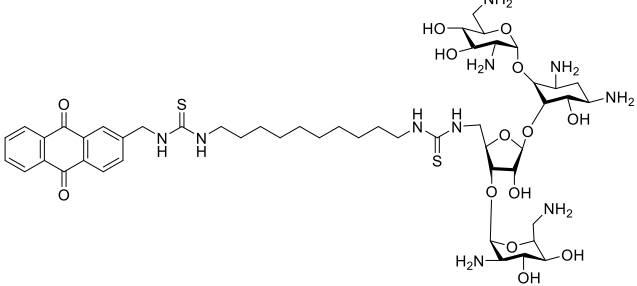
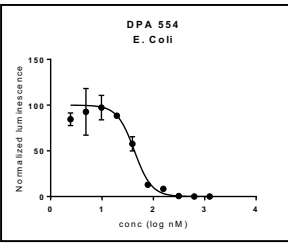
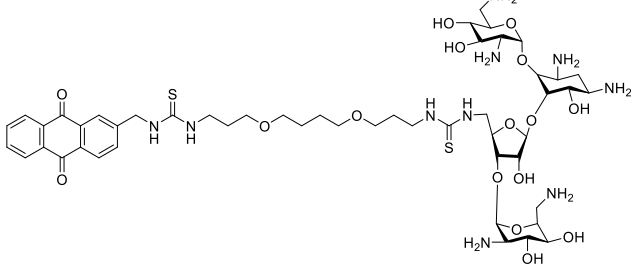
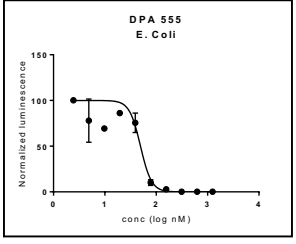
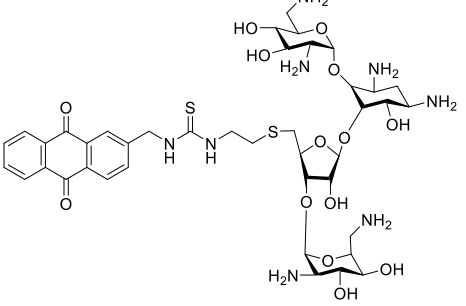
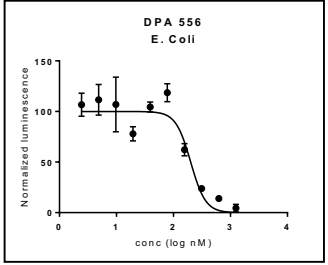


**Figure S2.** Titration plots of 100 nM A-site:F-NEO complex with compound **5**. IC<sub>50</sub> values for *E. coli*, human cytosolic, and A1555G A-sites were 151 ± 10 nM, 140 ± 10 nM, and 249 ± 47 nM, respectively.

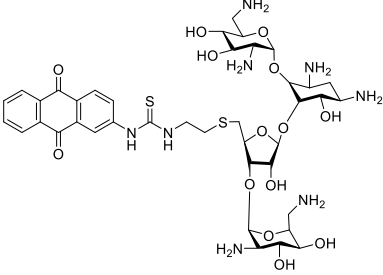
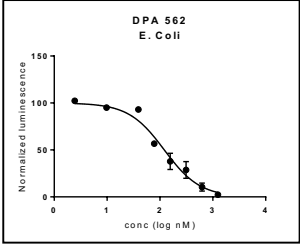
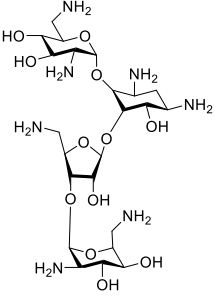
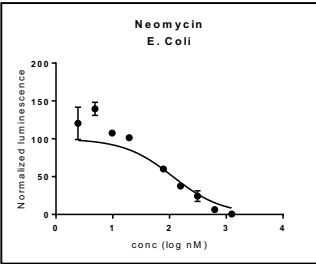
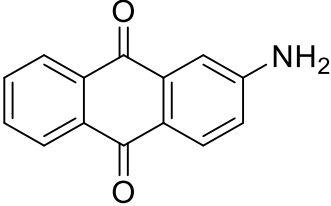
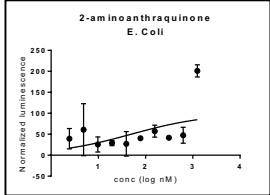
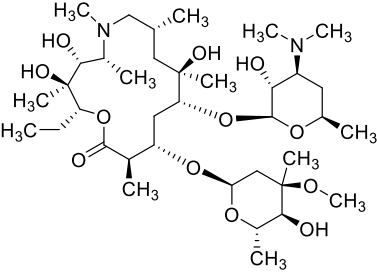
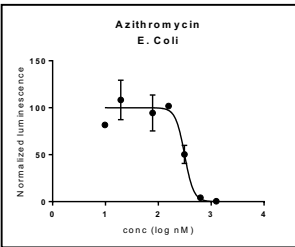


**Figure S3.** Titration plots of 100 nM A-site:F-NEO complex with compound **6**. IC<sub>50</sub> values for *E. coli*, human cytosolic, and A1555G A-sites were 162 ± 13 nM, 129 ± 1 nM, and 196 ± 4 nM, respectively.

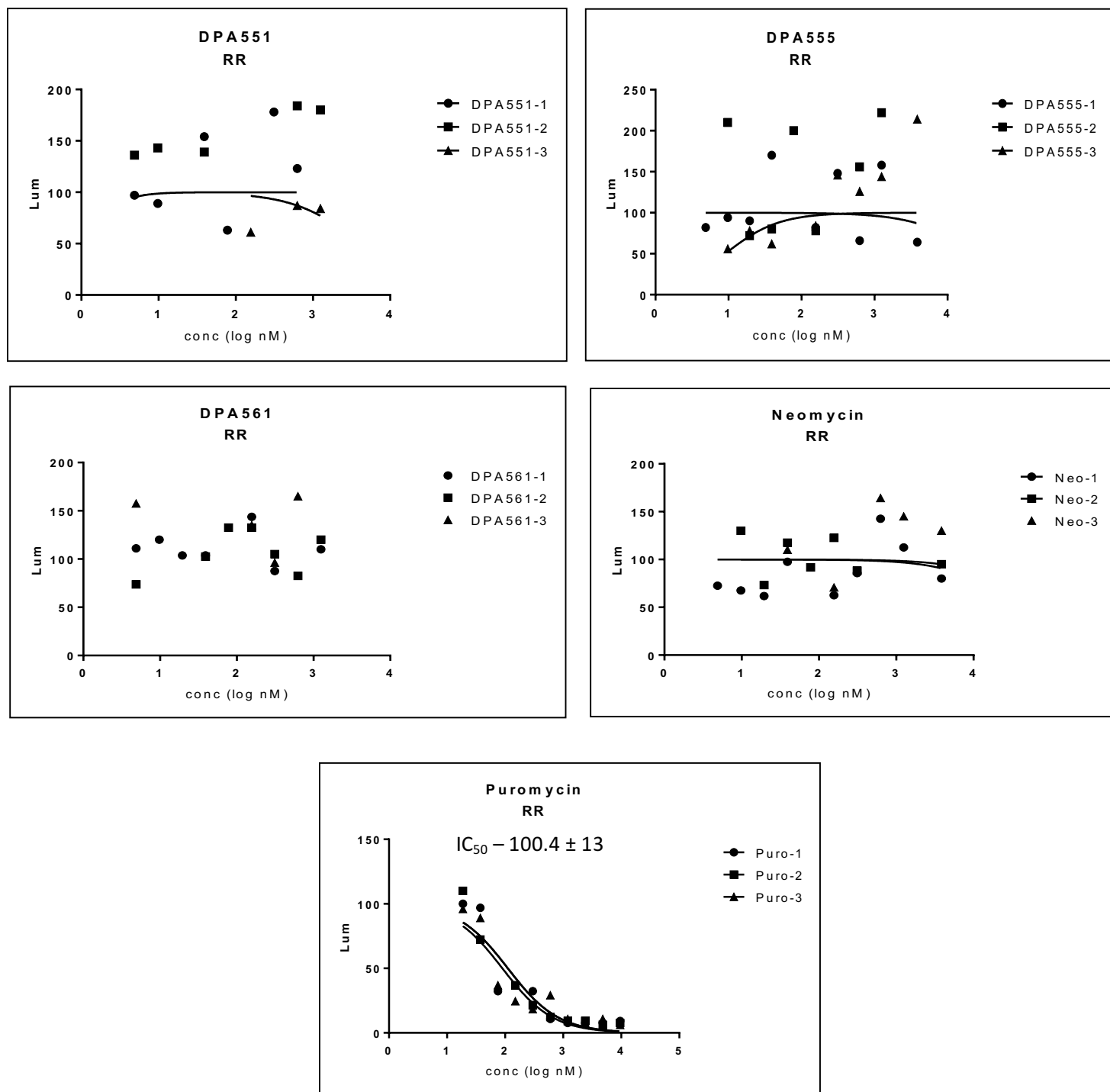
Compound	Structure	Cell Free Data	IC <sub>50</sub> (nM)
DPA551 1			NI

<p>DPA552</p> <p>2</p>			<p>50±8.8</p>
<p>DPA553</p> <p>3</p>			<p>303±28</p>
<p>DPA554</p> <p>4</p>			<p>40.3±6.7</p>
<p>DPA555</p> <p>5</p>			<p>35.4±3.5</p>
<p>DPA556</p> <p>6</p>			<p>178±18</p>

<p>DPA557</p> <p>7</p>			<p>91±1.7</p>
<p>DPA558</p> <p>8</p>			<p>101±16</p>
<p>DPA559</p> <p>9</p>			<p>250±35</p>
<p>DPA560</p> <p>10</p>			<p>71.4±2</p>
<p>DPA561</p> <p>11</p>			<p>63.7±8.5</p>

<p>DPA562 12</p>			<p>164±8</p>
<p>Neomycin NEO</p>			<p>139.7±10</p>
<p>2-amino anthraquinone</p>			<p>NI</p>
<p>Azithromycin</p>			<p>337.1±37</p>

**Figure S4.**  $IC_{50}$  of anthraquinone-neomycin conjugates in cell-free luciferase inhibition assays for prokaryotic systems.  $IC_{50}$  is the concentration of the compound that inhibits translation of luciferase as determined in the cell-free translation assay described in the Material and Methods.



**Figure S5.** IC<sub>50</sub> of anthraquinone-neomycin conjugates in cell-free luciferase inhibition assays for eukaryotic systems. IC<sub>50</sub> is the concentration of the compound that inhibits translation of luciferase as determined in the cell-free translation assay described in the Material and Methods.