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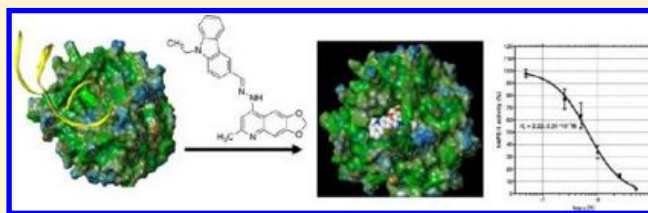
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Identification and Characterization of Human Apurinic/Apyrimidinic Endonuclease-1 Inhibitors

Ajay Srinivasan,[†] Lirong Wang,[†] Cari J. Cline,[†] Zhaojun Xie,[†] Robert W. Sobol,^{§,⊥} Xiang-Qun Xie,^{†,‡,||} and Barry Gold^{*,†,||}

[†]Department of Pharmaceutical Sciences, [‡]Computational Chemical Genomics Screening Center, ^{||}University of Pittsburgh Drug Discovery Institute, [§]University of Pittsburgh Cancer Institute, Hillman Cancer Center, and [⊥]Department of Pharmacology & Chemical Biology, University of Pittsburgh, Pittsburgh, Pennsylvania 15261, United States

ABSTRACT: The repair of abasic sites that arise in DNA from hydrolytic depurination/depyrimidination of the nitrogenous bases from the sugar–phosphate backbone and the action of DNA glycosylases on deaminated, oxidized, and alkylated bases are critical to cell survival. Apurinic/aprimidinic endonuclease-1/redox effector factor-1 (APE-1; aka APE1/ref-1) is responsible for the initial removal of abasic lesions as part of the base excision repair pathway. Deletion of APE-1 activity is embryonic lethal in animals and is lethal in cells. Potential inhibitors of the repair function of APE-1 were identified based upon molecular modeling of the crystal structure of the APE-1 protein. We describe the characterization of several unique nanomolar inhibitors using two complementary biochemical screens. The most active molecules all contain a 2-methyl-4-amino-6,7-dioxolo-quinoline structure that is predicted from the modeling to anchor the compounds in the endonuclease site of the protein. The mechanism of action of the selected compounds was probed by fluorescence and competition studies, which indicate, in a specific case, direct interaction between the inhibitor and the active site of the protein. It is demonstrated that the inhibitors induce time-dependent increases in the accumulation of abasic sites in cells at levels that correlate with their potency to inhibit APE-1 endonuclease excision. The inhibitor molecules also potentiate by 5-fold the toxicity of a DNA methylating agent that creates abasic sites. The molecules represent a new class of APE-1 inhibitors that can be used to probe the biology of this critical enzyme and to sensitize resistant tumor cells to the cytotoxicity of clinically used DNA damaging anticancer drugs.



Abasic sites created by hydrolytic depurination/depyrimidination and excision of lesions by base excision repair (BER*) DNA glycosylases are both cytotoxic and mutagenic.^{1,2} It is estimated that more than 10^4 abasic sites are formed per mammalian cell per day.^{3,4} The repair of abasic lesions in mammalian cells is predominantly mediated by the initial action of apurinic/aprimidinic endonuclease-1/redox effector factor-1 (APE-1),⁵ which cleaves the phosphodiester linkage that is 5' to the abasic site, leaving a single strand break (SSB) with 3'-hydroxyl and 5'-deoxyribose phosphate (dRP) termini.⁶ This repair intermediate is then processed by Pol β , which removes the 5'-DRP structure to afford a 5'-phosphate and then adds the appropriate complementary base to the 3'-terminus.⁷ In the final step, DNA ligase seals the nick. While cells and animals can survive without the different DNA glycosylases, albeit with increased sensitivity to DNA damaging agents,^{8–11} the genetic deletion of APE-1, which is expressed ubiquitously, is lethal in cells.¹² In mice, embryos terminate at postimplantation following blastocyst formation, and without developmental defects.^{13,14} Heterozygous mice are viable but become sensitized to DNA damaging agents that induce the formation of abasic sites.^{15–17} Deletion of Pol β , which is also critical in BER,¹⁸ causes neonatal lethality due to defective neurogenesis characterized by apoptotic cell death in the developing central and peripheral nervous systems,¹⁹ indicating the critical need

for cells to maintain functional BER during embryogenesis. Zebrafish knockdown of AP endonuclease (Apex) using siRNA also terminates during development.²⁰ Of interest is the observation that Pol β appears to be translationally coupled to Apex since the mRNA for the polymerase is present in the null fish but the protein is absent.²¹ Whether this is also the case in mammalian cells is not known.

The endonuclease function of APE-1 is located toward the C-terminus of the protein. The N-terminal domain is associated with the redox center (a.k.a., ref-1) that regulates the activity of specific transcriptional factors by maintaining them in a reduced state.^{22–26} In addition, APE-1 has been linked to several other functions, including RNA processing²⁷ and in Ca²⁺-dependent gene expression and regulation.²⁸ The lethality of APE-1 knockouts has been attributed to loss of the repair activity, and the mechanism of cell death involves apoptosis.²⁹ Overexpression of APE-1 makes cells resistant to alkylating agents.¹² There is also evidence that APE-1 expression can be induced by genotoxic agents, including cancer drugs.³⁰ These data raise the question of whether APE-1 expression is associated with tumor

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resistance to DNA damaging agents. In this regard, the lethality of clinically used anticancer treatments can be enhanced by a temporal decrease in APE-1 using antisense technology.^{31–34} Therefore, molecules that modulate APE-1 activity could be important adjuvants to clinically used DNA damaging antineoplastic agents. Recently, it has been reported that inhibitors of APE-1 endonuclease activity can create a synthetic lethality in cells defective in double-strand break repair, that is, BRCA1, BRCA2, and ATM.³⁵ This result is not unexpected since homologous recombination (HR) mutants are particularly sensitive to methylation damage repaired by BER.^{36,37} In fact, yeast cells that lack HR tolerate DNA alkylation damage better if there is no BER, indicating the biological consequences of BER in the absence of HR.³⁸ This result with APE-1 induced synthetic lethality is similar to the interaction between BRCA defective cells and PARP inhibitors.^{39,40}

A number of small molecule inhibitors of APE-1 that have been identified and characterized.^{41–47} In many cases the inhibitors identified in screens are dicarboxylic acids or related analogues (Figure 1). These molecules potentially mimic the

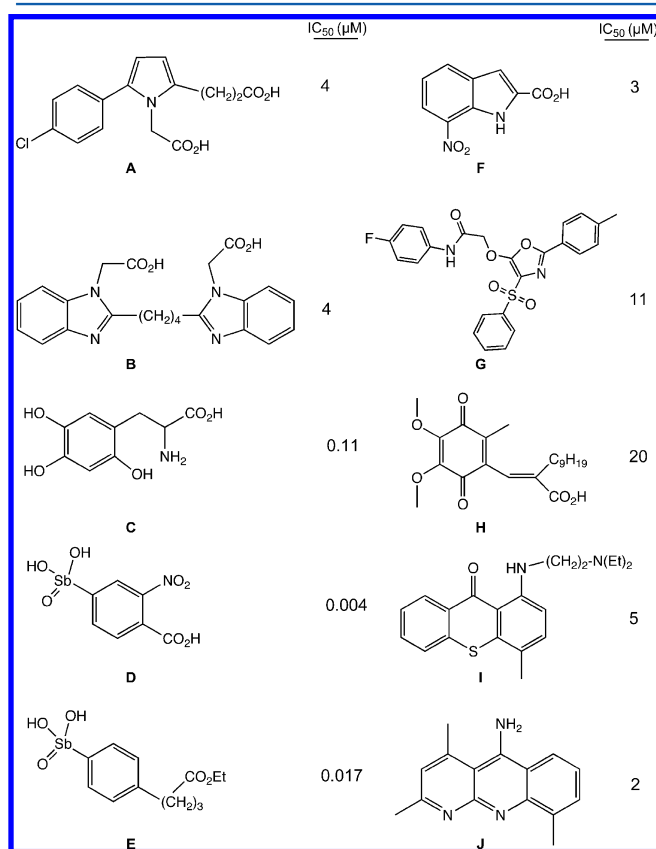


Figure 1. Structures of some previously reported inhibitors of APE-1 (and their calculated IC₅₀ values): A (ref 46), B (ref 46), C (ref 45), D (ref 44), E (ref 44), F (ref 85), G (ref 85), H (ref 79), I (ref 48), J (ref 86).

phosphate linkages flanking the abasic lesion on the DNA (Figure 2), which participate in salt bridges with the cationic face of the enzyme. Included in these inhibitors are a series of arylstibonic acids, which though extremely potent in biochemical experiments, lacked activity in cells.⁴⁴ Lucanthone inhibits APE-1 activity⁴¹ and binds to the protein⁴⁸ but also interacts with a number of other cellular targets, including DNA, so the mechanism of action remains uncertain.⁴⁹ The

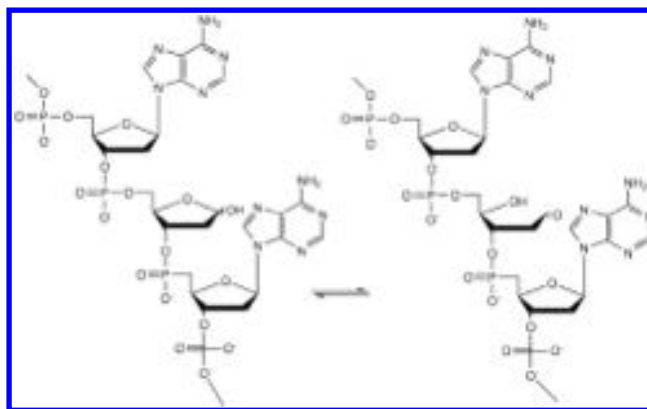


Figure 2. Structure of closed and ring-opened abasic site in DNA.

dimethoxyquinone, E3330⁵⁰ (Figure 1), which also has a number of cellular activities,⁵¹ has been shown to inhibit both the endonuclease and redox activity of APE-1.^{52–54} Of note is that NMR data indicate that the quinone molecule binds near the active site associated with the endonuclease activity.⁵⁴ How the redox and endonuclease activities, which are in physically distinct regions of the protein, are inhibited by a single molecule remains unclear. Another approach to block APE-1 activity is to chemically modify the abasic lesion using aldehyde reactive compounds, such as methoxylamine. Methoxylamine reacts to form an imine with the aldehyde group in the ring-open form of the abasic lesion (Figure 2), which prevents the protein mediated cleavage of the abasic site.^{55,56} Though methoxylamine will react with any cellular aldehyde or ketone, it is being clinically evaluated in a phase-1 trial as an adjunct therapy in combination with DNA alkylating agents.⁵⁷

Using the crystal structure of APE-1,⁵⁸ we have computationally constructed molecules that would sterically block its endonuclease site. Upon the basis of the modeling, potential inhibitor molecules were identified and sequentially assayed in two complementary biochemical screens. Fluorescence studies, including displacement assays, indicate that one of the inhibitors directly interacts with the APE-1 endonuclease site. The toxicity of a range of inhibitors was evaluated in a resistant T98G glioma cell line. To complement the inhibition and toxicity studies, we measured the levels of abasic lesions in the untreated T98G cells and in cells treated with the APE-1 inhibitors alone or with a methylating agent that selectively generates N3-methyladenine lesions⁵⁹ that are processed via abasic intermediates.^{60,61}

EXPERIMENTAL PROCEDURES

Materials. Potential APE-1 inhibitor compounds were obtained via the Developmental Therapeutics Program of the National Cancer Institute, National Institute of Health, Bethesda, MD. Chemicals, solvents, and CellLytic NuCLEAR Extraction kit were purchased from Sigma Aldrich Chemicals (St. Louis, MO). Oligonucleotide sequences were custom-synthesized by Integrated DNA Technologies (Coralville, IA) or Sigma Aldrich (St. Louis, MO). Nickel nitrilotriacetate (Ni-NTA) sepharose resin was procured from Qiagen Inc. (Valencia, CA). pENTR, pDEST17, *Escherichia coli* DH5α, precast polyacrylamide gels, DNazol reagent, cell culture components, and CyQUANT Cell Proliferation Assay kit were obtained from Invitrogen (Carlsbad, CA). Human glioblastoma (T98G) cells were obtained from American

Type Culture Collection (ATCC) (Manassas, VA). CellTiter 96 AQueous One Solution Reagent was obtained from Promega Corp, Madison, WI. DNA Damage Quantification kit (abasic site counting) was obtained from Dojindo Molecular Technologies (Rockville, MD). *Escherichia coli* C41(DE3) cells were obtained from Lucigen Corp. (Middleton, WI). His-Pur Cobalt resin, Trypan Blue stain, BCA protein assay kit, dialysis cassettes, and molecular biology grade buffers, as well as plastic and glassware were obtained from Fisher Scientific (Pittsburgh, PA). The sequence-specific alkylating agent, 1-methyl-4-[1-methyl-4-(3-methoxysulfonylpropanamido)pyrrole-2-carboxamido]pyrrole-2-carboxamido}propane (MeLex), was synthesized as previously described.⁵⁹

Virtual Screening Strategy for Small-Molecule Inhibitors of APE-1. The 3D structural model of APE-1 was constructed based on the X-ray crystallographic data for IDEW in the PDB⁵⁸ and was further modeled by refining the low-resolution region using our previously reported protocols.⁶² All computations were performed using Sybyl 8.2 (Tripos Associates) on a Dell 32 CPU Linux Cluster. After the generation of Surflex-Dock protomol using the reported binding residues and default parameters, a virtual docking screen⁶³ was carried out on the reported 3D virtual chemical compound library.⁶⁴ For each compound, 20 optimal conformers were generated by docking calculations. The conformer with the best total Hammerhead score (8.21)^{65,66} was chosen for further biochemical and cellular studies.

Cloning, Expression, and Purification of APE-1. The coding sequence for the human apurinic endonuclease (*ape-1*) was previously amplified by reverse transcriptase-polymerase chain reaction (RT-PCR) and cloned into pENTR, a directional TOPO-cloning Gateway entry vector (R. W. Sobol et al., unpublished). The *ape-1* coding sequence was subcloned into the destination vector pDEST17 via the LR reaction, which introduced an N-terminal polyhistidine (6x) coding sequence. The pDEST17-*ape-1* construct, amplified and purified from *Escherichia coli* DH5 α cells, was used to transform "OverExpress" *E. coli* C41(DE3) cells for expression and purification of recombinant (His)₆-APE-1 protein.

Transformed *E. coli* C41(DE3) was grown overnight at 37 °C and with constant shaking (100 rpm) in 5 mL of Luria-Bertani (LB) medium containing 100 μ g mL⁻¹ ampicillin. The overnight culture was used to seed 500 mL of LB medium and the cells were allowed to grow at 37 °C with constant shaking (100 rpm, 2 h) until the culture reached midlog phase ($\lambda_{600\text{nm}} \sim 0.5$). Isopropyl β -D-1-thiogalactopyranoside (IPTG) at 0.8 mM was used to induce APE-1 expression, at 25 °C, under constant stirring (100 rpm, ~ 2 h). At 16 h postinduction, the cells were harvested by centrifugation (5000 rpm, 10 min), washed free of media with 1 \times phosphate buffered saline (PBS) and resuspended in 5.77 mM sodium phosphates (4.3 mM Na₂HPO₄ + 1.47 mM NaH₂PO₄, pH 7.4), 2.7 mM KCl, 330 mM NaCl, 10% (v/v) glycerol, 0.5% (v/v) Tween-20, 0.1% (w/v) lysozyme, 10 units mL⁻¹ benzonase (nuclease), and incubated for 30 min at 25 °C. Intact cells were lysed by ultrasonication (5 min, 30% amplitude), centrifuged at 10 000 rpm, for 30 min, and clear supernatant (cell lysates) collected. APE-1 protein in cell lysates was purified by immobilized metal affinity chromatography using Ni- or Co-chelated resins to bind to the N-terminus His₆ tag. Immobilized APE-1 protein was eluted using a 10–250 mM gradient of imidazole in 10 mM HEPES (pH 7.4), 50 mM KCl. Pooled fractions containing APE-1 protein were dialyzed against 10 mM HEPES (pH 7.4),

250 mM KCl, 10% (v/v) glycerol, and protein concentration was estimated colorimetrically (BCA protein assay kit) as well as by UV absorbance ($\lambda_{280\text{nm}}$). Homogeneity of purified proteins was checked by sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE), followed by staining with 0.05% Coomassie Brilliant Blue R-250, 50% (v/v) methanol, 10% (v/v) acetic acid. Purified proteins were stored at –80 °C (long-term) or kept at –20 °C during regular use.

Molecular Beacon Assay for Screening of APE-1 Inhibitors. A fluorescence-based molecular beacon assay to measure APE-1 protein activity and inhibition was performed as described previously^{44,67} with some modifications.⁶⁸ APE-1 (2 nM) in 10 mM HEPES (pH 7.8) 0.5 mM MgCl₂, 100 mM NaCl, 100 mM KCl, 0.0005% (w/v) BSA, was incubated with 20 nM 5'-(F)-CGACTXTTGAATTGACACGCCATGTCGATCAATTCAATAGTCG-(D)-3' [X = tetrahydrofuran (THF) abasic site modification; F, 6-carboxyfluorescein (6-FAM); and D, DABCYL] that forms a hairpin with a 15 base pair stem and a 13 base loop. The THF modification is a known substrate for APE-1 excision. APE-1 mediated cleavage of the phosphodiester backbone at the THF-abasic site led to release of the 5' 6-FAM (fluorophore) containing pentanucleotide, disrupting the Förster resonance energy transfer (FRET) mediated quenching of the 6-FAM by DABCYL in the native hairpin oligomer. Since the signal was generated by continuous liberation of the fluorophore-labeled excision fragment, progress of this reaction could be monitored in real-time using a Cary Eclipse fluorescence spectrometer (Varian, Palo Alto, CA) or a Synergy H1 multiwell plate reader (BioTek, Winooski, VT). Reactions were optimized to yield $\sim 90\%$ substrate turnover in 10 min, and fluorescence intensities were measured every 12 s at 495 nm excitation and 520 nm emission. Data from the first 1–3 min of the assays (linear phase) were used to determine rate of substrate turnover (V_0). The Michaelis constant (K_M) was deduced experimentally, by incubating 2 nM APE-1 protein with 2–500 nM hairpin DNA substrate and measuring V_0 — the experimentally observed V_0 were analyzed as a function of substrate concentration [S] via nonlinear regression using Prism (v.4; GraphPad Software, Inc., La Jolla, CA) and the K_M was calculated as per the algorithm in the software package. To assay inhibition, 2 nM APE-1 protein was preincubated with 0.05–10.00 μ M of the candidate inhibitors at 25 °C for 60 min, before initiating the excision assay by addition of hairpin DNA substrate. V_0 was calculated at various concentrations of inhibitors and represented as inverse function of increasing inhibitor concentration [I]. Data were plotted on Prism and analyzed by nonlinear regression to obtain values for inhibitor concentration at 50% inhibition (IC_{50}), and the inhibitory constant (K_i) was approximated by the formula, $K_i = [IC_{50}]/(1 + [S]/K_M)$, where [S] was the substrate concentration.⁶⁹ All assays were carried out in three independent sets to obtain mean data and standard error of mean.

Gel-Based Assay for APE-1 Inhibitors. In addition to the real-time fluorescence assay for studying inhibition of APE-1 activity, we also employed a gel-based assay to visualize and quantify repair activity of APE-1 and its inhibition. Briefly, 20 nM APE-1, in 10 mM HEPES (pH 7.8) 0.5 mM MgCl₂, 100 mM NaCl, 100 mM KCl, 0.0005% (w/v) BSA, was incubated with 100 nM of a duplex DNA that was formed by annealing 5'-(F)-CGATCATCACTXTTGAAGACTGACACTGACC-3', which contained the THF abasic site modification (X), a 5' 6-FAM fluorophore (F), and the complementary sequence, 5'-

GGTCAGTGTCTCAGTCTCAATAGTGATGATCG-3'. Cleavage by APE-1 at the THF abasic site led to release of the 5'-fluorophore tagged 11-nucleotide fragment, which could be resolved from the uncleaved 30-nucleotide fragment (intact substrate) by gel electrophoresis under denaturing conditions. To assay inhibition of APE-1 activity, the APE-1 protein was preincubated with 5–100 μ M of the candidate inhibitors for 60 min, before initiation of the excision assay by addition of duplex DNA substrate. Reaction products were separated on a 20% denaturing (urea) polyacrylamide gel, and the bands were visualized by fluorescence emission of the 6-FAM under ultraviolet illumination using a GelDoc EZ Imager (BioRad, Hercules, CA) and quantified using the Image Lab Software (BioRad, Hercules, CA). Inhibitory activity was calculated by correlating fluorescence intensity of the 11-nucleotide fragment product as the (inverse) function of inhibitor concentration. All assays included a positive control (complete digestion of duplex DNA in absence of any inhibitor) as well as a negative control (uncleaved duplex DNA), which corresponded to 100% and 0% substrate turnover, respectively. Substrate turnover in the presence of candidate inhibitors were calculated as a percentage of positive control. Data were plotted on Prism, analyzed by nonlinear regression, and the IC_{50} values were obtained from the dose response curves. All assays were carried out in three independent measurements.

Fluorescence and UV Spectrometry Binding Studies.

The fluorescence spectra were measured at room temperature on a Cary Eclipse fluorescence spectrometer (Varian, Palo Alto, CA). The fluorescence properties of 1 and 4 were measured in 10 mM HEPES, 0.5 mM $MgCl_2$, 100 mM KCl, 100 mM NaCl, 2% glycerol, in the absence or presence of wild type (unmodified) hairpin DNA (5'-CGAATTCGCAGGACCGAATTCG-3') or THF-modified hairpin DNA (5'-CGA χ TTTCGCAGGACCGAATTCG-3') or APE-1 protein. A solution of 0.5 μ M of either DNA sequence was titrated against 1–10 μ M ethidium bromide (EtBr) to determine saturation of EtBr by fluorescence (530 nm excitation with 610 nm emission) and the subsequent displacement of EtBr (4 μ M) by 1–20 μ M 1 or 4 was also assayed. The APE protein–inhibitor binding fluorescence was assayed by titrating 0.1–20 μ M of 4 against 1 μ M of APE-1 protein. Displacement of 4 (6 μ M) from 1 μ M APE-1 protein by increasing amounts (0.75–30 μ M) of E3330 was also measured.

The effect of the small molecules on DNA stabilization was determined by measuring UV absorption of DNA at 260 nm as a function of temperature in a Cary 300 UV–visible spectrophotometer (Varian, Palo Alto, CA) under similar conditions as previously described.⁷⁰ The transition temperatures (T_M) was calculated from the first derivative analysis.

Isolation and Assay of APE-1 Protein from Nuclear Extracts of Glioma Cells. Human glioblastoma cells (T98G) were maintained in growth medium, that is, Eagle's Minimum Essential Medium (EMEM) with 10% fetal bovine serum (FBS), 50 μ g μ L⁻¹ gentamycin, 1 \times MEM nonessential amino acids, and 1 mM sodium pyruvate. Cells were harvested by trypsin-EDTA treatment, washed with 1 \times PBS, and nuclear proteins isolated using the CellLytic NuCLEAR Extraction kit (Sigma-Aldrich) as per manufacturer's instructions. Isolated nuclear proteins were estimated colorimetrically (BCA protein assay) and stored at –80 °C (long-term) or at –20 °C during regular use. To assay APE-1 activity in T98G cells, nuclear extracts were serially diluted in 10 mM HEPES, 50 mM KCl, 1 mM DTT, 20% (v/v) glycerol, and the activities were measured

via the molecular beacon assay (as described above). The final dilution used for all assays (EC_{2nM} , effective concentration of 2 nM) gave ~90% turnover of 25 nM hairpin substrate DNA in 10 min, which was equivalent to the activity of 2 nM recombinant APE-1 protein in the molecular beacon assay. EC_{2nM} of nuclear extracts was incubated with 2–500 nM hairpin DNA substrate and the V_0 was measured for each value of [S]. Finally, the apparent K_M^* was deduced using the software algorithm in Prism (as described previously), and the K_M^* was used to determine the apparent K_i^* . Inhibition of EC_{2nM} APE-1 activity by the inhibitors was assayed via the molecular beacon assay (described above).

Cytotoxic Effects of APE-1 Inhibitors on Cultured Human Glioblastoma (T98G) Cells. Human glioblastoma cells (T98G) were maintained in growth medium as described above. To assay short-term cytotoxicity, T98G cells maintained in growth medium, were harvested by trypsin-EDTA treatment, resuspended in fresh growth medium at a density of 10⁴ cells mL⁻¹, and 200 μ L (2000 cells) of this suspension was seeded into the wells of a sterile 96-well tissue culture plate. The cells were allowed to attach and grow for 24 h, and then were treated with serial dilutions of the drug for 72 h. After treatment, viable cells were measured by an MTS assay using the CellTiter 96 AQueous kit (Promega) as per manufacturer's instructions. To assay long-term cytotoxicity, T98G cells maintained in growth medium were harvested by Trypsin-EDTA treatment, resuspended in fresh growth medium at a density of 500 cells mL⁻¹, and 200 μ L (100 cells) of this suspension was seeded into the wells of a sterile 96-well tissue culture plate. The cells were allowed to attach and grow for 24 h and treated with serial dilutions of the drug for 240 h. After treatment, viable cells were measured using the CyQUANT cell proliferation assay kit (Invitrogen) as per manufacturer's instructions.

Cytotoxic Potentiation of Various DNA Damaging Agents. T98G cells maintained in growth medium were resuspended in fresh growth medium and seeded into 96-well plates at 2000 cells/well (short-term cytotoxicity) or 100 cells/well (long-term cytotoxicity) as described in the previous section. After 24 h, the cells were treated with serial dilutions of various DNA damaging drugs in the absence or presence of APE-1 inhibitors. At the end of the treatment period, cell survival was assayed using CellTiter 96 AQueous or CyQUANT kits for the short- and long-term studies respectively.

Quantification of Abasic Sites. T98G cells maintained in growth medium were harvested by Trypsin-EDTA treatment, resuspended in fresh growth medium at a density of 2.5 \times 10⁵ cells mL⁻¹, and 2 mL (5 \times 10⁵ cells) of this suspension was seeded into the wells of a sterile 6-well tissue culture plate. Cells were allowed to grow for 24 h, treated with the various compounds for different time points and/or at different concentrations. At the end of each treatment, adherent cells were harvested by trypsin-EDTA treatment and pooled with live and dead cells in culture supernatant. Pooled cells in suspension were pelleted by centrifugation, washed with 1 \times PBS, and genomic DNA was isolated using the DNAzol reagent (Invitrogen). DNA in solution was quantified by UV absorbance (λ_{260nm}) using a Synergy H1 multiwell plate reader (BioTek, Winooski, VT). Abasic sites in purified DNA were then reacted with the biotin-conjugated aldehyde reactive probe (ARP), immobilized onto 96-well ELISA plates, and detected using horseradish peroxidase (HRP)-conjugated streptavidin

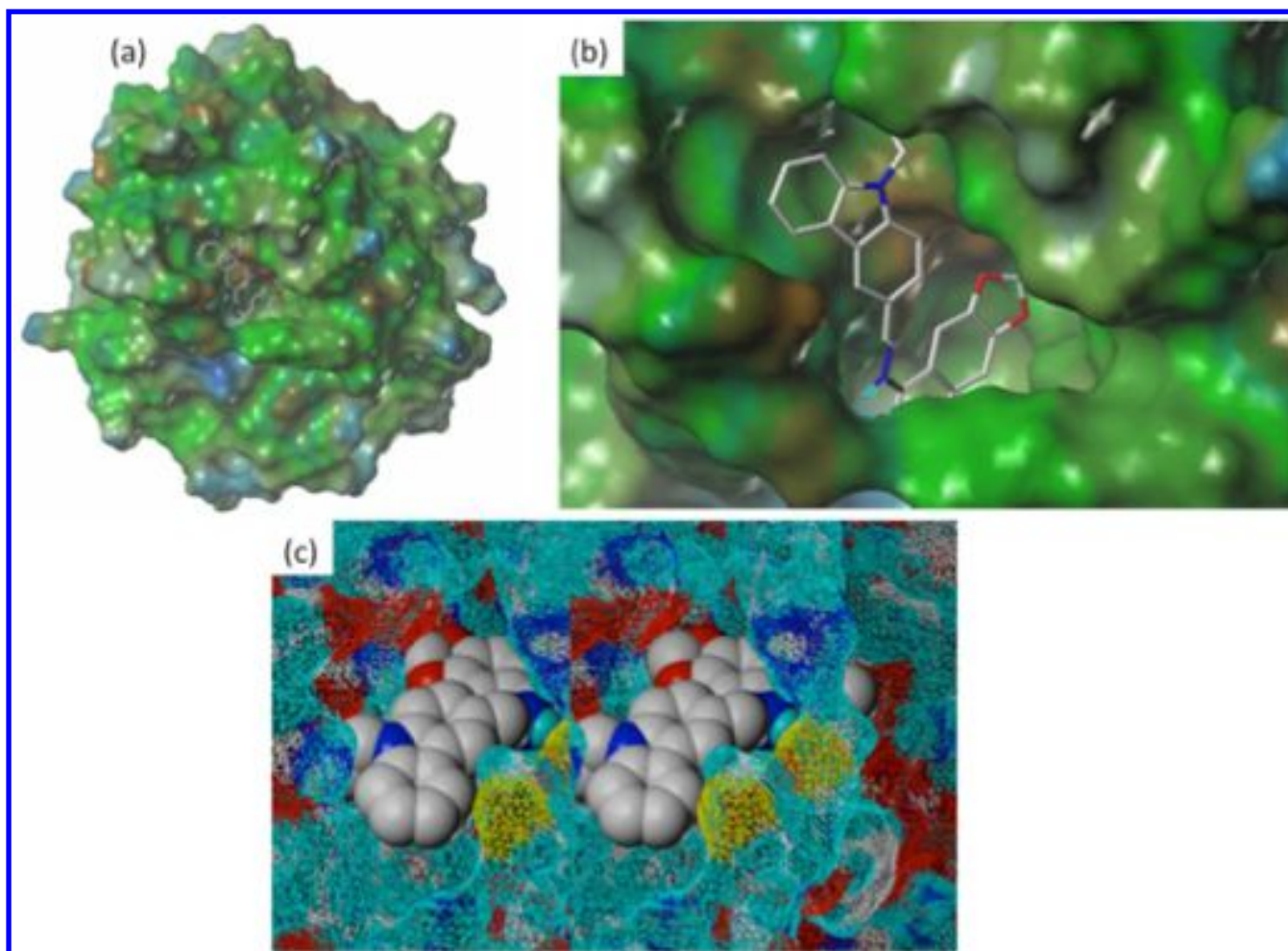


Figure 3. Docking results of 4 with APE-1: (a) View of compound 4 docked into the X-ray crystallographic structure of APE-1 (1DEW.pdb);⁵⁸ (b) enlarged docking pose of 4 showing methylenedioxy ring inserted inside the binding pocket and carbazole ring pointing outward, forming strong H-bonding and hydrophobic interactions; (c) relaxed stereoview of the N-ethylcarbazole ring of compound 4 pointing out with the benzodiazole ring system buried into the active site of APE-1.

using a colorimetric substrate as per manufacturer's instructions (Dojindo Molecular Technologies, Rockville, Maryland).

RESULTS AND DISCUSSION

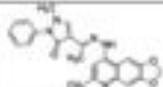
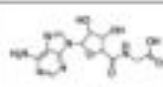
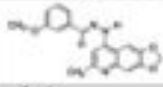
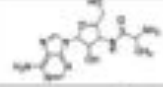
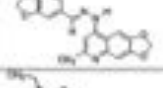
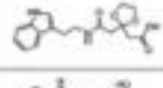
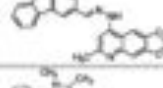

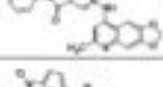
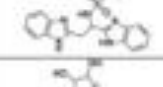
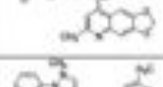
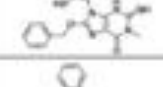
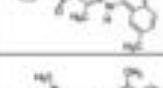
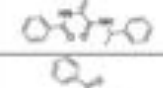
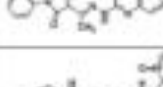
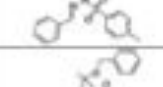
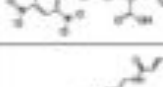
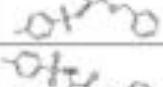
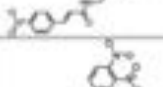
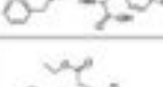
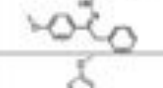
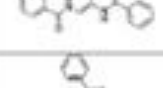
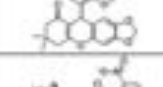
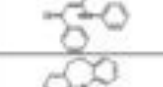

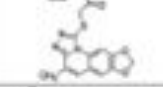
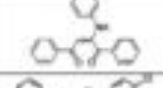
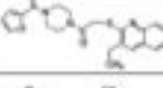

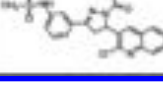
The development of small molecule inhibitors of APE-1 has been challenging because of the structure around the protein's active site that is rich in basic amino acid residues, which make salt bridge contacts with the DNA phosphate backbone. An Arg residue on a flexible loop region moves into the DNA stack and occupies the vacancy created when the abasic lesion is extruded into the protein's active site.⁵⁸ The hydrophobic face of the synthetic THF abasic site analogue in the crystal structure of APE-1 interacts with amino acid residues that create a hydrophobic pocket; however, the physiologically relevant 2-hydroxy-THF abasic site could form an H-bond via the 1'- α -hydroxyl group that is missing in the THF lesion.

Virtual Screening for Small-Molecule Inhibitors of APE-1. In silico screening was performed using the Surflex-dock program^{65,66} and established docking protocols⁶³ based on the X-ray crystallographic data of APE-1 (1DEW.PDB).⁵⁸ The predicted binding pocket was validated through 3D database docking searches and confirmed through experiments using the in silico screened hits (discussed below). Figure 3 shows a docking pose of inhibitor 4 (Table 1) in the predicted

binding pocket with the methylenedioxy ring mimicking the furan ring of the abasic site substrate in the X-ray cocrystallographic structure of APE-1.⁵⁸ Analysis of the docking results reveal H-bonding interactions of compound 4 with Asn174 (2.33 Å), Thr268 (2.67 Å), and hydrophobic interactions (2.66–3.0 Å) of the dioxoloquinoline moiety with Phe266 and Trp280 as well as the carbazole ring with a hydrophobic domain (Trp280, Met271, and Val278). In the X-ray structure, the hydrophobic face of the extra-helical abasic lesion packs within a complementary APE-1 pocket formed by the side chains of Phe266, Trp280, and Leu282. In addition, Asn174, Asn212, and His309 also interact through H-bonds with the phosphate that is 5' of the abasic site target. Our docking also predicted that the basic Arg177 side chain could interact with the aromatic quinoline ring of compound 4 via a Π -cation interaction (3.42 Å).

Screening for APE-1 Inhibitors. The initial assessment of the molecules suggested by the computational study employed a molecular beacon approach similar to that previously described.^{47,67,68} In this assay, the DNA hairpin substrate has a 6-FAM fluorophore on the 5'-terminus, a DABACYL fluorescence quencher on the 3'-terminus, and a THF abasic site located six base pairs from the 5'-terminus in the stem region (Figure 4). APE-1 activity is monitored as an increase in

Table 1. Structures of Compounds Identified from in Silico Screen and APE-1 Endonuclease Inhibition (K_i)

Name	NCI#	Structure	K_i (μ M) ^a	Name	NCI#	Structure	K_i (μ M)
1	NSC332398		0.18	16	NSC163444		ND
2	NSC332384		0.22	17	NSC115605		ND
3	NSC332389		0.19	18	NSC128335		ND
4	NSC332395		0.12	19	NSC372329		ND
5	NSC332396		0.68	20	NSC283787		ND ^d
6	NSC332397		0.64	21	NSC11847		ND
7	NSC300598		10.89	22	NSC117589		ND
8	NSC332410		ND ^b	23	NSC126939		ND ^d
9	NSC89640		13.00	24	NSC296950		ND ^d
10	NSC107215		17.00	25	NSC343032		ND
11	NSC131534		6.70	26	NSC374123		ND
12	NSC375491		32.00	27	NSC63636		ND ^d
13	NSC614430		4.90	28	CS-965570 ^c		ND
14	NSC402686		37.00	29	CS-365759 ^c		ND
15	NSC658900		ND	30	CS-2504743 ^c		ND

^aEndonuclease activity. ^bNo detectable activity ($K_i > 100 \mu$ M). ^cChempidder accession number. ^dInsufficient solubility in buffer containing 0.25% DMSO to determine if $K_i > 100 \mu$ M.

fluorescence due to the release of the 5'-(6-FAM)-GCACT fragment as a function of time. Compounds that had good inhibitory activity ($K_i \leq 10 \mu$ M) in the molecular beacon screen were then retested using a gel-based assay to directly measure DNA excision as a function of inhibitor concentration (Figure 5). This second assay, which directly measures the amount and

site of DNA nicking, confirms that the endonuclease processing of the abasic lesion is inhibited and eliminates the possibility of artifacts that can arise in the fluorescence measurements or from random nuclease activity. In this assay, substrate turnover was calculated from relative intensities of cleaved vs intact oligomers and correlated with inhibitor concentration. Using

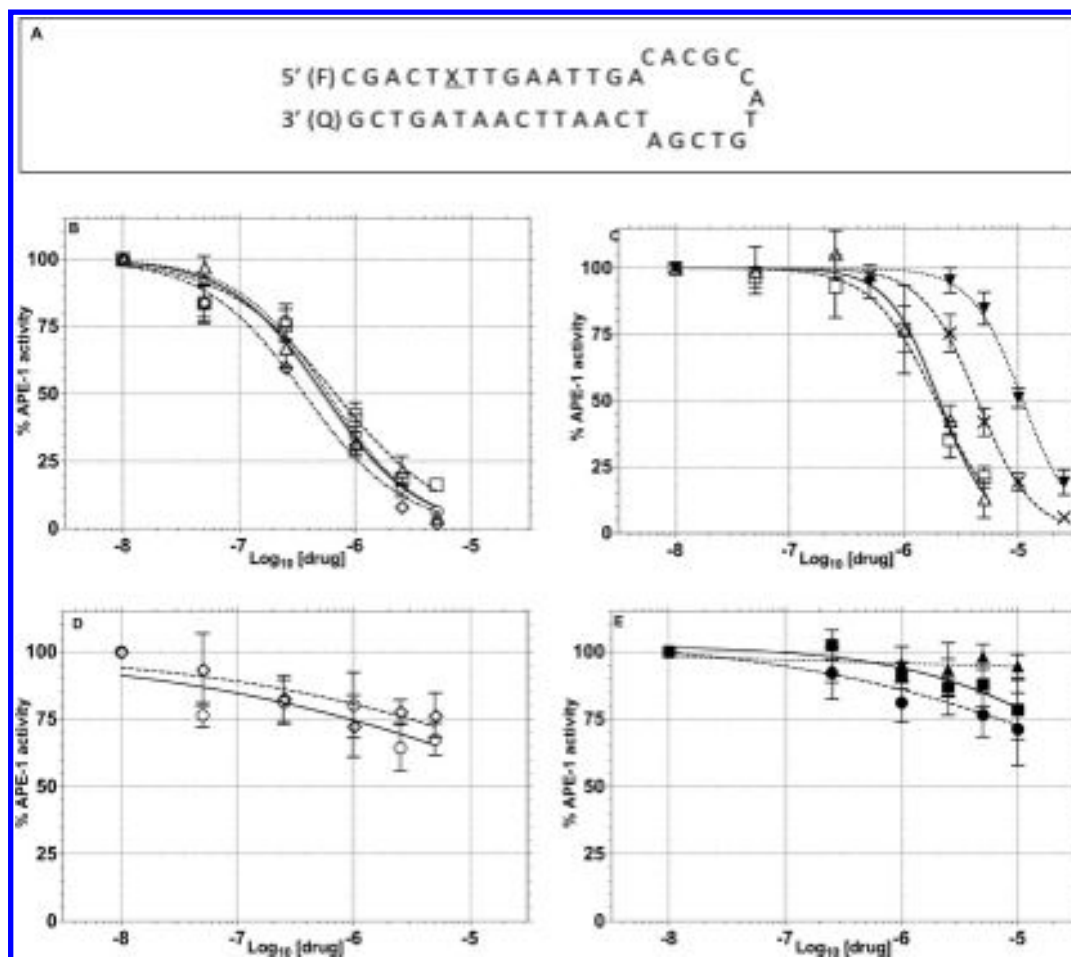


Figure 4. Inhibition of APE-1 endonuclease activity by 1–5, 8, 28–30, E3330, and EtBr in the molecular beacon assay (K_i values determined from this assay were calculated as described in Experimental Procedures and are shown in Table 1). (A) Sequence of THF modified hairpin DNA substrate, where, F, fluorophore and Q, quencher. (B–D) Changes in APE-1 endonuclease activity as a function of varying concentrations of potential APE-1 inhibitor molecules: (B) 1 (△), 2 (□), 3 (○), 4 (◇); (C) 5 (△), 6 (□), EtBr (▼), E3330 (X); (D) 7 (○), 8 (◇); and (E) 28 (■), 29 (●), and 30 (▲).

this approach, the IC_{50} values (Table 2) were estimated for APE-1 inhibition by 1 (4.4 μ M), 2 (10.1 μ M), 3 (2.9 μ M), 4 (4.1 μ M), 5 (34.4 μ M), 8 (>100 μ M), and E3330 (14.7 μ M). The inhibitory activity (K_i) observed in the molecular beacon assay (Table 1, Figure 5) correlated in all cases with the IC_{50} activity in the gel assay (Table 2). Of the 30 compounds (Table 1) evaluated based on the modeling, six showed a K_i of <1.0 μ M in the molecular beacon assay (Table 1, Figure 4). It was not possible to fully characterize some of the compounds because of their limited solubility. The relatively high hit rate in the molecular beacon assay is attributed to the success of the molecular modeling to predict structures in contrast to a random screen of a large chemical library. The importance of the benzo[d][1,3]dioxole structure and H-bond donor at position-N⁴ on the quinoline ring, which were predicted from the modeling, was evident from the assays. For example, compounds 1 and 8 are identical except for the methylenedioxy substitution on the quinoline ring, yet their effect on APE-1 activity differs by more than 50-fold. The dramatic difference between 1 and 7 also illustrates the important role of the dioxole ring. The appendage on the hydrazone had less impact on the *in vitro* inhibition. Compounds 1–5 and 8 were selected for additional evaluation based on the range of their *in vitro* activity.

In addition to the compounds identified from the modeling studies, E3330, a previously reported inhibitor that has been shown to interact with the endonuclease active site of APE-1,⁵⁴ was tested for comparative purposes. The K_i for E3330 in our system was 2.2 μ M in the molecular beacon assay (Figure 4C) and the IC_{50} in the gel assay was 14.7 μ M (Figure 5). In order to probe the potential role of DNA intercalation as a mechanism for APE-1 inhibition (see below), ethidium bromide (EtBr) was evaluated. It showed a K_i of 1.8 μ M in the molecular beacon assay (Figure 4C), but its intercalating and fluorescence properties hindered analysis in the gel assay.

Inhibition of APE-1 Activity in Nuclear Lysates. The ability of 1–5 and 8 to block APE-1 endonuclease activity in a complex mixture of proteins from T98G cell nuclear lysates was tested using the molecular beacon assay. The APE-1 activity in the lysates was standardized to give ~90% turnover of 25 nM substrate in 10 min. This activity is similar to that of 2 nM recombinant APE-1 used in the molecular beacon assay (Figure 4). The K_M and V_{max} of EC_{2nM} APE-1 activity in nuclear lysates were determined to be 24.4 nM and 8.5 fluorescence units (arbitrary) min⁻¹, respectively. Using these values, the K_i for 1–5 was calculated to be 0.45, 0.24, 0.28, 0.21, and 1.41 μ M, respectively. For comparison E3330 had a K_i of 2.93 μ M. Compounds 1–4 clearly blocked excision of the target DNA

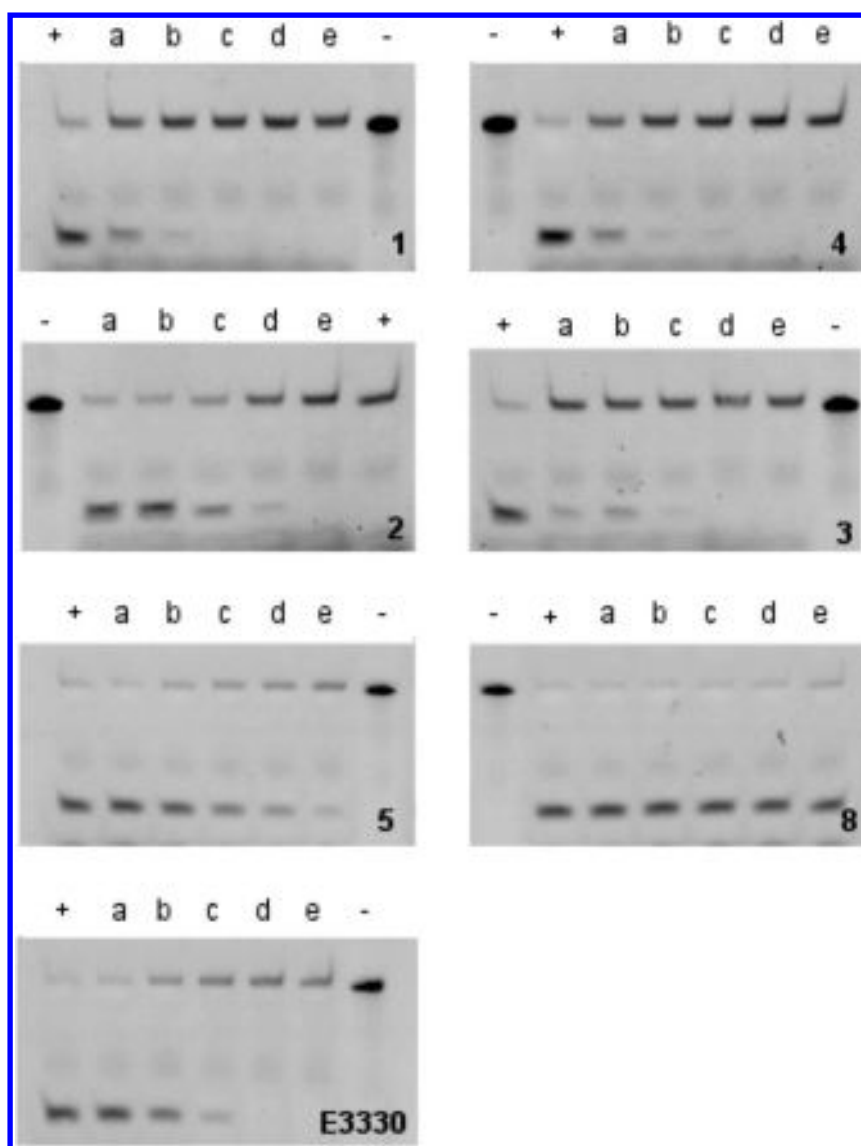


Figure 5. Gel based assay measuring the inhibition of APE-1 endonuclease activity by compounds 1–5, and 8 and E3330 (upper band, uncleaved 30 nucleotide THF-modified DNA; lower band, endonuclease cleaved 11 nucleotide fragment): +, control (no inhibitor); –, uncleaved target; a–e, 5, 10, 25, 50, and 100 μ M compound, respectively.

Table 2. IC_{50} Values for Compounds 1, 2, 3, 4, 5, 8, and E3330 Were Calculated from the in Gel Excision Assay by Quantifying the Relative Intensities of the APE-1 Cleaved Fragment versus Full-Length DNA Substrate in the Absence or Presence of Potential APE-1 Inhibitors

compound	IC_{50} (μ M)
1	4.4
2	10.1
3	2.9
4	4.1
5	34.4
8	>100
E3330	14.7

with the abasic site analogue in the presence of the nuclear lysate that contain APE-1 activity (Figure 6). These results indicate that compounds 1–5 have reasonable selectivity even in the complex mixture of proteins in the nuclear lysates.

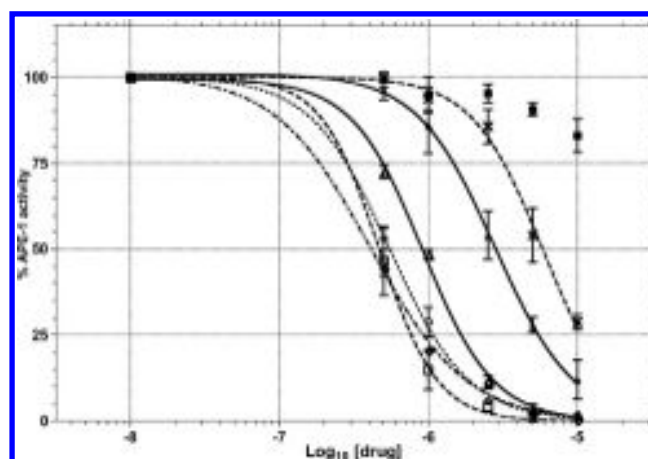


Figure 6. Inhibition of APE-1 endonuclease activity by 1 (Δ), 2 (\square), 3 (\circ), 4 (\diamond), 5 (\blacktriangle), 8 (\blacksquare), and E3330 (\times) in T98G cell nuclear extracts in molecular beacon assay (see Figure 4 for details).

Interaction of Inhibitors with APE-1 Protein. To establish whether there is a direct interaction between an inhibitor and APE-1, the fluorescence spectra of **4** was determined in the absence and presence of increasing amounts of the protein. Compound **4** is the only potent inhibitor that is fluorescent in buffer (containing 0.5% DMSO) with an excitation at 395 nm and a broad emission maximum at 485–520 nm. In the presence of APE-1, there is a concentration-dependent enhanced fluorescence of **4** (Figure 7), which suggests a physical interaction with the protein that

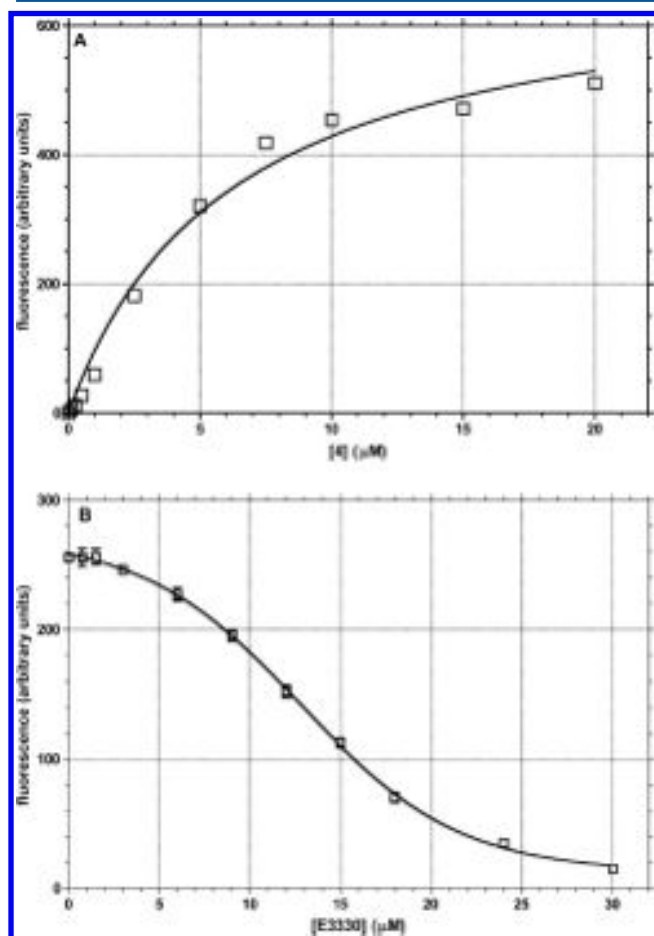


Figure 7. Concentration-dependent change in the fluorescence of **4** in the presence of 1 μM APE-1: (A) binding of **4** to APE-1 as measured by the increase in fluorescence (excitation at 395 nm, emission at 510 nm); K_D of 6.15 μM ; (B) displacement of **4** from APE-1 by E3330 measured by a decrease in the fluorescence of **4**.

reduces fluorescence quenching due to interaction with solvent. The emission maximum also shifts with the 485 nm band increasing to a smaller extent than the long wavelength band, which is displaced by 10–510 nm. These data suggest direct binding between the protein and the inhibitor. E3330 is an APE-1 inhibitor for which there are NMR data showing an interaction with the protein's active site.⁵⁴ Therefore, we incubated **4** (6.0 μM) with APE-1 (1 μM) and measured the change in fluorescence upon the addition of 0.5–10.0 μM E3330 (Figure 7). The results show that there is an E3330 concentration-dependent displacement of **4** from the protein as indicated by the decrease in **4**'s fluorescence intensity. To obtain approximately 50% decrease in the fluorescence of 6 μM **4** requires approximately 12 μM E3330. E3330 binds to APE-1

with a K_D of 55 μM from the NMR study and causes a 3-fold inhibition of APE-1 endonuclease activity at 100 μM concentration in a gel based excision assay.⁵⁴ The IC_{50} for E3330 redox inhibition is 10 μM .⁴⁷ The displacement of **4** from APE-1 by E3330 indicates that **4** binds to, or near, the endonuclease site of the protein.

Interaction of Inhibitors with DNA. A potential mechanism by which compounds can block APE-1 activity involves the interaction with the DNA substrate by intercalation. Such binding to DNA could sterically reduce the accessibility of the protein to the abasic site substrate. The intercalation of compounds that have planar aromatic systems is common and relevant because the abasic lesion potentially provides a high affinity site for the planar compounds to enter into the DNA base stack. The interaction of **1**, which exhibits no fluorescence in solution, with an unmodified DNA oligomer and one with an abasic site causes a concentration-dependent increase in fluorescence intensity (Figure 8). Compound **4**

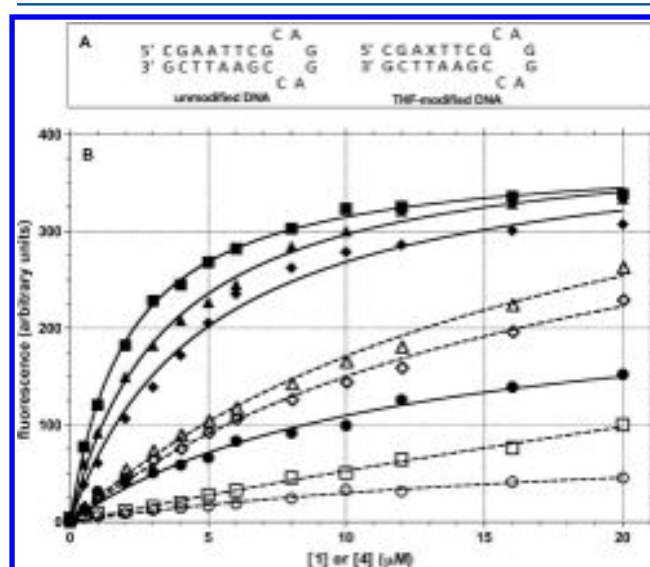


Figure 8. The interaction of compounds **1** and **4** with DNA was evaluated by changes in their fluorescence in the absence and presence of EtBr: (A) sequence of DNA hairpins used in the fluorescence measurements (X = THF abasic site); (B) fluorescence of compounds **1** (395 nm excitation and 460 nm emission) and **4** (400 nm excitation and 500 nm emission) with 0.5 μM DNA was evaluated by fluorescence increases in the absence (●, **1** with unmodified DNA; ■, **1** with THF modified DNA; ◆, **4** with unmodified DNA; ▲, **4** with THF modified DNA) and presence of 4 μM EtBr (○, **1** with unmodified DNA and EtBr; □, **1** with THF modified DNA and EtBr; ◇, **4** with unmodified DNA and EtBr; △, **4** with THF modified DNA and EtBr).

shows a similar fluorescence response. To further probe the potential role of intercalation, the effect of EtBr on the fluorescence of **1** and **4** was determined. Ethidium is a well-characterized intercalator that has a K_D near 1 μM with double-stranded DNA,⁷¹ although it binds almost 10-fold more strongly to DNA with a bulge.⁷² However, we observed similar K_D 's for EtBr with the unmodified and THF modified hairpin DNA of 1.6 and 1.8 μM , respectively (data not shown). The fluorescence enhancement of **1** and **4** in the presence of DNA was significantly muted by the presence of EtBr suggesting competitive binding to the DNA. Of note is that **1** appears to exhibit a strong preference for DNA containing the THF abasic

site based on the difference in the fluorescence intensities (Figure 8). The small changes in the fluorescence of **4** with both DNAs indicate little discrimination for the THF lesion. A similar pattern was observed in the effect of **1** and **4** on the thermal stability of DNA as determined by UV melting experiments using several DNA substrates (data not shown). With the self-complementary sequence 5'-GA-GAGCGCTCTC, **1** increased the T_M from 36.4 to 40.8 °C while the T_M for this sequence with **4** was unchanged (37.3 °C). Using the unmodified hairpin DNA as shown in Figure 8B, we observed a similar effect: ΔT_M was 4 °C with **1** and 0.5 °C with **4**. The stability of the hairpin with the THF abasic site increased by >9 °C in the presence of **1** but not appreciably with **4**. These data suggest weak binding between DNA and **4**, while **1** clearly has a stronger interaction with both unmodified DNA and DNA with a THF abasic site.

In summary, both **1** and **4** can bind to DNA, presumably by intercalation, but the binding is relatively weak, as compared to EtBr, and nonspecific and cannot explain their ability to inhibit APE-1 endonuclease activity at nanomolar concentrations in the different assays. Consistent with this conclusion is the weaker inhibition of APE-1 endonuclease activity by EtBr, which has a higher affinity for DNA than **1** or **4**. The overall evidence is consistent with inhibitor **4** acting by binding directly to the protein. The mechanism of action for some of the other inhibitors remains to be experimentally determined.

Toxicity. It was assumed that inhibition of APE-1 activity would lead to toxicity since genetic deletion of the protein is lethal to cells.⁸ Therefore, the toxicities of compounds **1**–**5** and E3330 were determined in human T98G glioma cells using both short-term (3-day) MTS (Figure 9a) and long-term (10-day) CyQUANT (Figure 9b) assays. The toxicities were also determined in combination with the DNA minor groove methylating agent, MeLex. MeLex efficiently and selectively generates N3-methyladenine, which is converted into an abasic site by a BER glycosylase^{73–75} and/or hydrolytic depurination.⁷⁶ The T98G cell line is relatively resistant to DNA damaging anticancer agents due to expression of multiple drug resistance proteins and O⁶-alkylguanine-DNA alkyltransferase.^{77,78} Compound **4** was again the most active with an LD₅₀ of 960 nM in the MTS assay and 190 nM in the clonogenic assay. The toxicities of compounds **1**–**3** were all approximately 500 nM in the long term assay and <5 μ M in the short term assay, while **5** was almost 20-fold less toxic than **4**. Of note, E3330 was approximately 250- and 750-fold less toxic to the cells than **4** in the short- and long-term assays, respectively. The previously reported E3330 IC₅₀ for cell growth in HEYC-2 and SKOV-3X ovarian tumor cells is ~35 μ M.⁷⁹ As a reference, the methylating agent, MeLex showed an LD₅₀ of 250 and 12 μ M in the short and long-term cytotoxicity assays, respectively. The mechanism responsible for the toxicity of the inhibitor compounds remains to be determined, but their activities generally correlate with the K_i for in vitro APE-1 inhibition (Table 1) suggesting that the decrease in APE-1 activity is involved.

A goal of the development of APE-1 inhibitors is to determine if they can be used to enhance the activity of DNA alkylating agents. Therefore, we tested how the LD₁₀ concentrations of compounds **1** (2.20 μ M), **4** (0.25 μ M), and **5** (6.50 μ M) affected the toxicity of MeLex (Figure 10) in the 72 h MTS assay. The LD₅₀ of MeLex dropped from 267 μ M by itself to 54, 49, and 80 μ M in the presence of **1**, **4** and **5**, respectively (Figure 10A). When an LD₁₀ concentration of

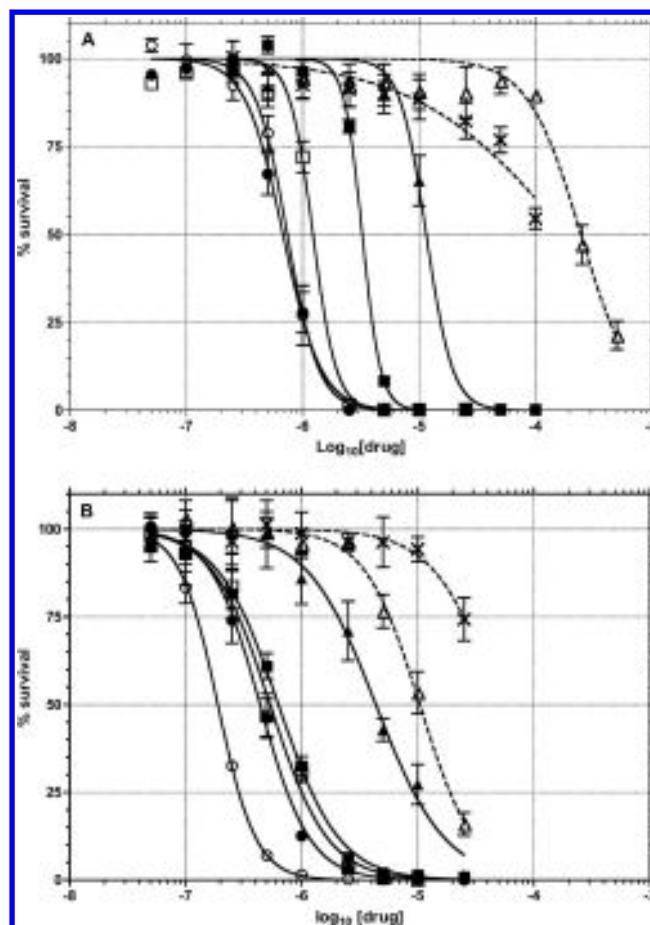


Figure 9. The toxicity of compounds **1** (■), **2** (□), **3** (●), **4** (○), **5** (▲), MeLex (△), and E3330 (×) were determined in T98G cells: (A) short-term (72 h) MTS assay; (B) 10-day Cyquant assay.

E3330 (6.9 μ M) was combined with MeLex, the cells were only 2-fold more sensitive to MeLex (128 μ M). In a long-term assay the corresponding LD₁₀ concentrations of **1** (0.17 μ M), **4** (0.08 μ M), and **5** (0.90 μ M) caused a decrease in the LD₅₀ of MeLex (9.9 μ M) to 2.0, 1.6, and 3.6 μ M respectively. E3330 (14.80 μ M) had little or no contribution toward potentiating the toxicity of MeLex (Figure 10B). Overall the maximum potentiation (i.e., with **4**) of MeLex was approximately 5-fold for both the short and long-term assays.

Some of the compounds that we tested are closely related to a series of molecules evaluated for antimalarial⁸⁰ and antitubercular activity.⁸¹ The IC₅₀ for the toxicity of compound **3** in a VERA kidney cell line was determined to be 630 nM, which compares to the 540 nM that we observed in the glioma T98G line (Table 1). The origin of the toxicity in the VERA cells was not determined but the inhibition of APE-1 would be predicted to be toxic in all cells.

Formation of Abasic Sites. To confirm that the inhibitors were actually affecting the cellular processing of abasic sites, a biotinylated aldehyde reactive probe (Biotin-ARP)⁸² was used to measure the level of aldehyde reactive sites (ARS) in genomic DNA isolated from T98G cells that were incubated with the different inhibitors. The biotin tagged ARS are quantified using an avidin–biotin assay, followed by colorimetric detection with horseradish peroxidase conjugated to avidin. While the method does not specifically measure abasic DNA sites, it is assumed that most of the increase in signal

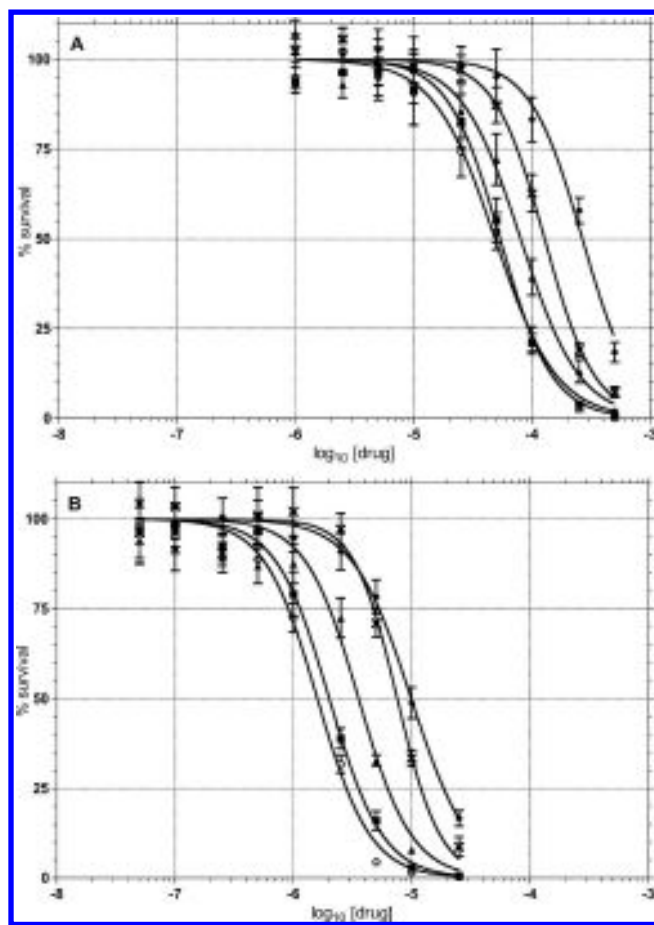


Figure 10. The potentiation of MeLex cytotoxicity was determined in T98G cells treated with different concentrations of MeLex alone (◆) or in combination with LD₁₀ concentrations of 1 (■), 4 (○), 5 (▲), or E3330 (×). (A) short-term (72 h) MTS assay; (B) long-term (10 days) CyQUANT assay.

results from an increase in the ring-opened form of abasic sites (Figure 2). The background level of ARS in untreated or 0.25% DMSO treated T98G cells was $1/10^5$ nucleotides, which is similar to previous reports (Figure 11a).^{82,83} In the presence of 0.25 μ M 4 (Figure 11), which corresponds to less than the LD₁₀ concentration due to the 25-fold increase in the number of cells used in the ARS vs the toxicity assay (Figure 9), there was a continuous increase over the initial 6 h. The level of approximately 17 ARS/ 10^5 nucleotides was maintained until the final 18 h time point. Because the concentration of the inhibitors is below the LD₁₀, it is unlikely that a significant percentage of the ARS measured are in dead cells. However, it cannot be ruled out that ARS levels may rise in cells entering or in apoptosis. Therefore, it is not possible to unequivocally say that the accumulation of ARS leads to cell death or vice versa. A more detailed time course for markers of cell death and the build up of ARS may resolve this issue.

At 0.25 μ M, 1 showed a similar effect, but 5, which is a weaker inhibitor of APE-1 endonuclease activity (Table 1), had a less dramatic effect with an increase in ARS of approximately 4-fold above background. The number of ARS with 0.25 μ M 8 was the same as background. For comparison, 0.25 μ M E3330 caused an increase in ARS similar to 1 and 4 at 2 h but then the number of lesions began to drop and returned to control levels by 18 h. EtBr was also evaluated at similar concentrations to determine if a strong intercalator could affect ARS levels. It had

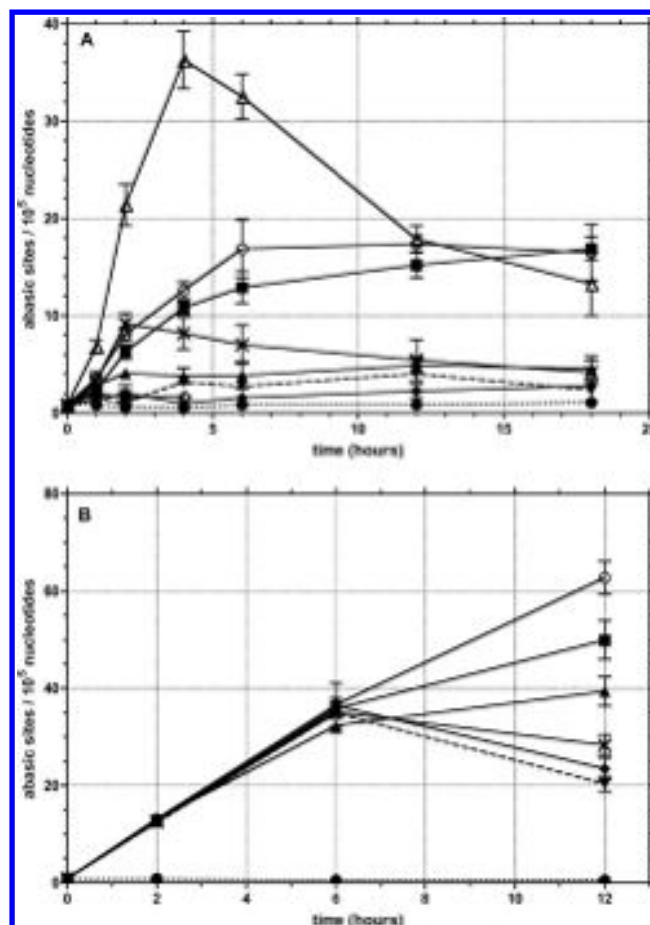


Figure 11. Formation of aldehyde reactive sites (ARS, abasic sites) was quantified in (A) cells treated with 0.25 μ M 1 (■), 4 (○), 5 (▲), 8 (◇), E3330 (×), EtBr (▼) or 80 μ M MeLex (△); (B) cells treated with combinations of 80 μ M MeLex (LD₁₀) and 0.25 μ M of 1 (■), 4 (○), 5 (▲), E3330 (×), EtBr (▼). Controls employed in both sets were untreated cells (●) or cells treated with 0.2% DMSO (◆).

no affect on the formation of ARS. As a positive control for the formation of ARS, the cells were treated with 80 μ M of MeLex. This MeLex treatment afforded a rapid induction of ARS (36/ 10^5 nucleotides) through the first 4 h that then decreased through 18 h to a value similar to that observed with 1 and 4. The rapid increase in ARS from MeLex through the first 4 h is consistent with the DNA methylation time course,⁸⁴ while the gradual decrease of ARS levels is attributed to their removal by functional BER.

The effect of APE-1 inhibitors in combination with 80 μ M MeLex was also determined (Figure 11b). At the time points employed in our assays, the level of ARS reflects the kinetics of 3-mA lesion formation due to MeLex, and BER processing of the lesion by MPG excision, APE-1 cleavage at the abasic site and dRP lyase or flap endonuclease removal of the aldehyde functionality from the DNA. The level of ARS induced by MeLex by itself or with the different inhibitor compounds was not significantly different at the 2 or 6 h time points. However, the number of lesions dropped off with MeLex alone or with either 0.25 μ M E3330 or EtBr. In contrast, the levels of ARS continued to increase through at least 12 h when MeLex was combined with 0.25 μ M of the more potent inhibitors 1 (50 sites/ 10^5 nucleotides) or 4 (63 sites/ 10^5 nucleotides). Therefore, the combination of DNA methylation and APE-1

inhibition has the effect of sustaining an elevated level of ARS, which correlates with the enhancement of MeLex toxicity.

APE-1 Redox Activity. APE-1 has a redox activity that is associated with the regulation of specific transcriptional factors. To probe whether any of the molecules identified as APE-1 endonuclease inhibitors affected the redox activity, the ability of APE-1 to affect Jun/Fos binding to an SP-1 cognate DNA sequence was determined using a gel shift assay as described previously.⁷⁹ E3330 was used as a positive control. We observed no effect with 100 μ M concentration of 1–5 and 8 (data not shown), which is far above the concentration required to inhibit endonuclease activity. In summary, the compounds that are potent inhibitors of APE-1 endonuclease activity increase the persistence of ARS and synergistically affect the toxicity of a DNA methylating agent that generates lesions that are repaired by BER. Structural studies are underway to establish the binding mode of the different APE-1 inhibitors that will be used to synthesize more potent and selective APE-1 inhibitors.

AUTHOR INFORMATION

Corresponding Author

*E-mail: goldbi@pitt.edu. Telephone: 1-412-383-9593.

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ABBREVIATIONS USED

APE-1, human apurinic endonuclease-1/redox effector factor-1; ARS, aldehyde reactive sites; BER, base excision repair; DMSO, dimethyl sulfoxide; E3330, (2E)-2-[(4,5-dimethoxy-2-methyl-3,6-dioxo-1,4-cyclohexadien-1-yl)methylene]undecanoic acid; EtBr, ethidium bromide; HEPES, 4-(2-hydroxy-ethyl)-1-piperazineethanesulfonic acid; HR, homologous recombination; MeLex, methyl 3-(1-methyl-5-(1-methyl-5-(propylcarbamoyl)-1H-pyrrol-3-ylcarbamoyl)-1H-pyrrol-3-ylamino)-3-oxopropane-1-sulfonate; MMS, methyl methane-sulfonate; PBS, phosphate buffered saline; THF, tetrahydrofuran

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