Synergistic Anticancer Activity of Arsenic Trioxide with Erlotinib Is Based on Inhibition of EGFR-Mediated DNA Double-Strand Break Repair

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Abstract
Arsenic trioxide (ATO), one of the oldest remedies used in traditional medicine, was recently rediscovered as an anticancer drug and approved for treatment of relapsed acute promyelocytic leukemia. However, its activity against nonhematologic cancers is rather limited so far. Here, we show that inhibition of ATO-mediated EGF receptor (EGFR) activation can be used to potently sensitize diverse solid cancer types against ATO. Thus, combination of ATO and the EGFR inhibitor erlotinib exerted synergistic activity against multiple cancer cell lines. Subsequent analyses revealed that this effect was based on the blockade of ATO-induced EGFR phosphorylation leading to more pronounced G2–M arrest as well as enhanced and more rapid induction of apoptosis. Comparable ATO-sensitizing effects were also found with PI3K/AKT and mitogen-activated protein/extracellular signal-regulated kinase (MEK) inhibitors, suggesting an essential role of the EGFR-mediated downstream signaling pathway in cancer cell protection against ATO. H2AX staining and comet assay revealed that erlotinib significantly increases ATO-induced DNA double-strand breaks (DSB) well in accordance with a role of the EGFR signaling axis in DNA damage repair. Indeed, EGFR inhibition led to downregulation of several DNA DSB repair proteins such as Rad51 and Rad50 as well as reduced phosphorylation of BRCA1. Finally, the combination treatment of ATO and erlotinib was also distinctly superior to both monotreatments against the notoriously therapy-resistant human A549 lung cancer and the orthotopic p31 mesothelioma xenograft model in vivo. In conclusion, this study suggests that combination of ATO and EGFR inhibitors is a promising therapeutic strategy against various solid tumors harboring wild-type EGFR. Mol Cancer Ther; 12(6); 1073–84. ©2013 AACR.

Introduction
In 2000, arsenic trioxide (ATO; Fig. 1A) was approved by the U.S. Food and Drug Administration for the treatment of relapsed and refractory acute promyelocytic leukemia (APL; ref. 1) and, thus, represents besides platinum compounds the only metal-based anticancer drug in clinical use. Induction of intracellular reactive oxygen species (ROS) and reactive nitrogen species (RNS) based on interference with the intracellular redox balance is one of the main mechanisms of action underlying the anti-cancer activity of ATO (1, 2). In addition, the selective efficacy of ATO against APL was shown to be due to degradation of the APL-specific promyelocytic leukemia and retinoic acid receptor-α (PML-RARA) fusion oncoprotein (3, 4). On the basis of its successful clinical application, the anticancer activity of ATO either as monotherapy or in combination with other agents was also intensively studied in various other hematologic malignancies (5, 6) and several solid cancer types (7). However, reports from phase II clinical trials on patients with hepatocellular carcinoma (HCC), metastatic renal cell carcinoma, and metastatic melanoma suggested so far that ATO has only limited efficacy against solid tumors (8–12). Although multiple resistance mechanisms for arsenic have been described on the cellular level (1), the reasons underlying the lack of efficacy of ATO in solid tumor types are still widely unclear.

In addition to its use as anticancer agent, arsenic came into the focus of interest as unintentional exposure to inorganic arsenic salts (mainly via drinking water) was found to exert potent carcinogenic activity (13). Recently, arsenic was reported to induce activation of the EGFR receptor (EGFR) pathway in nonmalignant tissues of exposed humans (such as lung, bladder, and prostate;
Consequently, this EGFR response was suggested to be one of the main drivers of the carcinogenic effect of inorganic arsenic salts (16). In addition, stimulation of EGFR was also observed in several malignant cancer types after treatment with arsenic (17, 18). The mechanisms underlying the recently described stimulation of EGFR by arsenic are still a matter of discussion. On the one hand, treatment with arsenic might stimulate the release of the EGFR ligand, heparin-binding EGF (19). On the other hand, the metal was shown to induce c-Src, which subsequently phosphorylates the EGFR (18, 20).

With regard to cancer, the EGFR pathway is a major player in cell survival, cell-cycle progression, tumor invasion, and angiogenesis (21). Moreover, there are several reports that the stimulation of the EGFR pathway is associated with enhanced DNA damage repair (22). Thus, based on the importance of the EGFR signaling for cell survival in malignant tissues and the discovery of several mutations leading to constitutive EGFR activation (23), this pathway was identified as an ideal target for cancer treatment. Consequently, several EGFR inhibitors have already been successfully developed and approved for the treatment of solid human malignancies especially...
non-small-cell lung carcinoma (NSCLC) at the disseminated stage (e.g., erlotinib, gefitinib; ref. 23). On the basis of the known EGFR stimulation by environmental arsenic exposure and the role of EGFR-mediated signals in multiple survival pathways, we hypothesized that EGFR might be involved in the insensitivity of solid tumors to ATO treatment. Consequently, the aim of the present study was to investigate the effects of erlotinib (Fig. 1B) on the anticancer activity of ATO against diverse solid cancer types and to gain more insights into the underlying molecular mechanisms.

Materials and Methods

Chemicals
ATO was purchased from Sigma-Aldrich, gefitinib from AstraZeneca, MK-2206 from Selleck Chemicals LLC, U0126 from Cell Signaling, and all other kinase inhibitors were from LC laboratories. For in vitro studies, ATO was dissolved in 1 mol/L NaOH, whereas for all other substances dimethyl sulfoxide (DMSO) stocks were prepared. The stock solutions were further diluted into culture media at the indicated concentrations. The final DMSO and NaOH concentrations were always less than 1%.

Cell culture
The following human cancer cell lines were used in this study: the NSCLC cell lines A549 and A427 [from American Type Culture Collection (ATCC)]; the mesothelioma cell line P31 (donated by Prof. K. Grankvist, Umea University, Umea, Sweden); the mesothelioma cell lines VMC6, VMC12, VMC23, and VMC31 (established at the Institute of Cancer Research, Vienna, Austria); the HCC HepG2 (from ATCC); the colorectal cancer line cells HCT116 (donated by Dr. B. Vogelstein, John Hopkins University, Baltimore, MD), SW620 and SW480 (from ATCC); osteosarcoma cell lines IOS (donated by Dr. Manara, Istituti Ortopedici Rizzoli, Bologna, Italy) and MG63 (from ATCC); the thyroid carcinoma cell lines SW1736 and SW579 (from ATCC); and the cervix carcinoma KB-3-1 (donated by Dr. Shen, Laboratory of Cell Biology, National Cancer Institute, National Institutes of Health, Bethesda, MD). HCT116 cells were grown in McCoy’s culture medium. A427, HepG2, SW480, and P31 were grown in minimum essential medium. All other cell lines were cultivated in RPMI-1640. Culture media were supplemented with 10% fetal calf serum (PAA). Cell cultures were periodically checked for Mycoplasma contamination. Cell line authentication has been done either by array-comparative genomic hybridization and/or short tandem repeat fingerprint.

Cytotoxicity assays
Cells were plated (2 × 10³ cells/well) in 96-well plates and allowed to recover for 24 hours. Subsequently, the dissolved drugs were added. After 72-hour exposure, the proportion of viable cells was determined by MTT assay following the manufacturer’s recommendations (EZ4U; Biomedica). Cytotoxicity was expressed as IC₅₀ values calculated from full dose–response curves. Synergism was determined by calculation of the combination index (CI) according to Chou and Talalay (24) using the CalcuSyn software (Biosoft). CI < 0.8, CI = 0.8–1.2, or CI > 1.2 represent synergism, additive effects, and antagonism of the 2 investigated substances, respectively.

Cell-cycle analysis
Cells (2 × 10⁶) were seeded into 6-well plates and allowed to recover for 24 hours. After drug exposure, cells were collected, washed with PBS, fixed in 70% ice-cold ethanol, and stored at −20°C. To analyze cell-cycle distribution, cells were transferred into PBS, incubated with RNase A (10 µg/mL) for 30 minutes at 37°C, followed by 30-minute treatment with 5 µg/mL propidium iodide (PI). Fluorescence levels were analyzed by flow cytometry using FACS Calibur (Becton Dickinson). The resulting DNA histograms were quantified using Cell Quest Pro software (Becton Dickinson and Company).

Mitochondrial membrane potential detection
Breakdown of the mitochondrial membrane potential (ΔΨm) was determined by fluorescence-activated cell sorting (FACS) analysis using JC-1 (5,5′,6,6′-tetramethyl-1,3,3′,3′-tetraethylbenzimidazol-carbocyanine iodide). For this purpose, the Mitochondrial Membrane Potential Detection Kit (Stratagene) was used according to the manufacturer’s instructions. A549 cells (5 × 10⁵) were seeded in T25 cm² culture flasks and treated with the tested drugs after 24-hour recovery. After trypsinization and PBS washing, cells were incubated for 10 minutes in freshly prepared JC-1 solution (10 µg/mL in cell culture media) at 37°C. Spare dye was removed by washing in PBS and cell-associated fluorescence measured via FACS.

Hoechst 33258–PI staining
A549 cells (2 × 10⁴ cells/well) were seeded in 24-well plates and allowed to recover for 24 hours. Cells were treated with ATO and erlotinib for the indicated exposure times. Then, the cells were stained with 2 µL/mL HOEPI mix [ratio 1:1—Hoechst 33258 (1 mg/mL) and PI (2.5 mg/mL)] for 1 hour at 37°C. Live photomicrographs of treated cells were made using fluorescent equipment on the Nikon Eclipse Ti inverted microscope system. Triplicate photomicrographs for each treatment were captured with 4',6-diamidino-2-phenylindole (DAPI)-, CY3-filters, and phase contrast. Morphologic features of more than 300 cell nuclei for each treatment were counted.

³H-thymidine incorporation assay
A549 cells (5 × 10⁵ cells/well) were seeded in a 96-well plate and treated 24 hours later with drugs for another 24 hours. Medium was replaced by a 2 nmol/L ³H-thymidine solution (diluted in full culture medium; radioactivity: 25 Ci/mmol/L). After 1-hour incubation at 37°C, cells were washed 3 times with PBS. Cell lysates were prepared and the radioactivity determined as described previously (25).
Western blot analysis

Proteins were isolated, resolved by SDS/PAGE, and transferred onto a polyvinylidene difluoride membrane for Western blotting as previously described (26). The following antibodies were used: EGFR, pEGFR (Tyr992 and Tyr1068), AKT, pAKT (Ser437), extracellular signal-regulated kinase (ERK)1/2 [p44/42 mitogen-activated protein kinase (MAPK)], pERK (Thr202/Tyr204), pH2AX (Ser139), cyclin B1, Rad50, Rad51, pBRCA1 (Ser1524), PARP, cl. PARP, and p21 Waf/Cip1 (all polyclonal rabbit; Cell Signaling Technology), p53 (DO-1; monoclonal mouse; Thermo Scientific), β-actin and α-tubulin monoclonal mouse (Sigma) were all used in 1:1,000 dilutions. In addition, horseradish peroxidase–labeled secondary antibodies from Santa Cruz Biotechnology were used at working dilutions of 1:10,000.

Immunofluorescence

A total of 2 × 10^5 A549 cells were seeded in 8-well chamber slides (BD Falcon). After 24 hours recovery cells were treated for another 24 hours with ATO, erlotinib, and their combination. Treated cells were washed with PBS and fixed in methanol/acetone at 4°C for 10 minutes before washing 3 times for 10 minutes in PBS with 1% bovine serum albumin (BSA). Fixed cells were incubated with phospho-H2AX antibody [diluted 1:100 in PBS with 1% BSA (Cell signaling)] for 1 hour at room temperature. After 3 further washing steps with PBS (1% BSA), cells were incubated (1 hour) with the second antibody (anti-rabbit immunoglobulin G (IgG) labeled with fluorescein isothiocyanate (FITC); Sigma-Aldrich), diluted 1:1,000 in PBS with 1% BSA. Cells were counterstained with DAPI before mounting. Images were obtained using a Leica DMRXA fluorescence microscope (Leica Mikroskopie und System) equipped with appropriate epifluorescence filters and a COHU charge-coupled device camera. Subsequent image handling was carried out in Adobe Photoshop CS4.

Comet assay

DNA double-strand breaks (DNA DSB) were analyzed using alkaline comet assay according to the guidelines of Tice and colleagues (27). A549 cells were seeded in 6-well plates (2.5 × 10^5 cells/well) and allowed to recover for 24 hours. After treatment for 6 hours, cells were trypsinized and washed with PBS. Cells treated with 100 μmol/L H2O2 were trypsinized after 30 minutes. Comet assay was then conducted with some modifications as described by Heffeter (28).

Animals

Six- to 8-week-old female CB-17 scid/scid [severe combined immunodeficient mouse (SCID)] mice were purchased from Harlan Laboratories. The animals were kept in a pathogen-free environment and every procedure was done in a laminar airflow cabinet. The experiments were carried out according to the regulations of the Ethics Committee for the Care and Use of Laboratory Animals at the Medical University Vienna (Vienna, Austria), the U.S. Public Health Service Policy on Human Care and Use of Laboratory Animals as well as the United Kingdom Coordinating Committee on Cancer Prevention Research’s Guidelines for the Welfare of Animals in Experimental Neoplasia.

Xenograft experiments

For the local tumor growth experiments, A549 cells (1 × 10^6) were injected subcutaneously into the right flank. For orthotopic models, P31 cells (6 × 10^6) were injected intraperitoneally. Animals were randomly assigned to treatment groups and therapy was started when tumor nodules reached a mean size of 25 mm^3 (A549) or 1 week after injection (P31). Animals were treated with erlotinib (orally 25 mg/kg dissolved in Cremophor EL diluted 1:1 in 96% ethanol) and then diluted 1:10 with deionized water before administration; 5 times a week for 2 weeks), ATO (intraperitoneally i.p.) 5 mg/kg dissolved in 1 mol/L NaOH and further diluted in PBS 1:10. pH value was adjusted at 7.0–7.5 with HCl before administration; 5 times a week for 2 weeks), or with a combination of both drugs. Animals in the control group received the Cremophor EL solvent orally and PBS intraperitoneally. Animals were controlled for distress development every day and tumor size was assessed regularly by caliper measurement (A549). Tumor volume was calculated using the formula: (length × width^2)/2. P31 tumors were weighted after mice were sacrificed. The P31 experiment was terminated after 62 days from transplantation when the first animal had to be sacrificed because of weight loss as previously shown in Hoda and colleagues (29).

Statistical analysis

All data are expressed as mean ± SD. Results were analyzed and illustrated with GraphPad Prism (version 5; GraphPad Software). Statistical analyses were performed using one- and two-way ANOVA with drug treatment and time as independent variables and conducted with Bonferroni posttests to examine the differences between the different drug treatment regimens and the diverse responses. The statistical significance is either described in the respective figure legends, or indicated with asterisks (*, P < 0.05; **, P < 0.01; ***, P < 0.001).

Results

ATO and erlotinib synergistically reduce cancer cell viability

The cell lines used in this study and their respective IC_{50} values after 72-hour treatment with ATO and erlotinib (as single agents or combined) are listed in Table 1. In addition, genetic characteristics are shown in Supplementary Table S1. Selected full dose–response curves are shown in Fig. 1C and in Supplementary Fig. S1A–S1C. ATO monotherapy exerted distinct anticancer activity against the tested cells in the low μmol/L range. The most resistant cell lines were IOS (osteosarcoma; IC_{50} > 25 μmol/L), VMCl31 (mesothelioma; IC_{50} = 18.87 μmol/L),

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and A549 (NSCLC; IC₅₀ = 17.58 μmol/L), whereas SW579 (thyroid carcinoma; IC₅₀ = 1.95 μmol/L), MG63 (osteosarcoma; IC₅₀ = 2.87 μmol/L), and P31 (mesothelioma; IC₅₀ = 2.52 μmol/L) exhibited the strongest ATO sensitivity. As all cell lines harbor wild-type EGFR, the anticancer effect of erlotinib was minor and resulted in general in IC₅₀ values above 20 μmol/L (Table 1).

Combination of ATO and erlotinib had synergistic activities as compared with monotherapy with ATO (up to 9.4-fold decrease in IC₅₀ of ATO alone by IC₅₀ of ATO with 2.5 μmol/L erlo).

Table 1. IC₅₀ values of different cells treated with ATO, erlotinib, or their combination

<table>
<thead>
<tr>
<th>Cell line</th>
<th>Histology</th>
<th>ATO (μmol/L) IC₅₀</th>
<th>Erlotinib (μmol/L) IC₅₀</th>
<th>ATO with 2.5 μmol/L</th>
<th>Fold decrease</th>
</tr>
</thead>
<tbody>
<tr>
<td>A427</td>
<td>NSCLC</td>
<td>5.2 ± 0.4</td>
<td>&gt;20</td>
<td>4.5 ± 0.3</td>
<td>1.2</td>
</tr>
<tr>
<td>A549</td>
<td>NSCLC</td>
<td>17.6 ± 0.9</td>
<td>&gt;20</td>
<td>5.7 ± 0.8</td>
<td>3.1</td>
</tr>
<tr>
<td>HepG2</td>
<td>HCC</td>
<td>5.9 ± 0.7</td>
<td>&gt;20</td>
<td>0.8 ± 0.1</td>
<td>7.4</td>
</tr>
<tr>
<td>HCT116</td>
<td>CC</td>
<td>7.5 ± 0.3</td>
<td>&gt;20</td>
<td>0.8 ± 0.1</td>
<td>9.4</td>
</tr>
<tr>
<td>SW480</td>
<td>CC</td>
<td>8.6 ± 0.1</td>
<td>&gt;20</td>
<td>2.8 ± 0.2</td>
<td>3.1</td>
</tr>
<tr>
<td>SW620</td>
<td>CC</td>
<td>8.4 ± 0.3</td>
<td>&gt;20</td>
<td>4.2 ± 0.1</td>
<td>2.0</td>
</tr>
<tr>
<td>IOS</td>
<td>OS</td>
<td>&gt;25</td>
<td>&gt;20</td>
<td>9.3 ± 0.1</td>
<td>&gt;2.7</td>
</tr>
<tr>
<td>MG63</td>
<td>OS</td>
<td>2.8 ± 0.1</td>
<td>&gt;20</td>
<td>1.2 ± 0.1</td>
<td>2.3</td>
</tr>
<tr>
<td>KB-3-1</td>
<td>Cervical carcinoma</td>
<td>6.3 ± 0.3</td>
<td>&gt;20</td>
<td>2.0 ± 0.1</td>
<td>3.2</td>
</tr>
<tr>
<td>SW579</td>
<td>TC</td>
<td>1.9 ± 0.1</td>
<td>&gt;20</td>
<td>0.8 ± 0.1</td>
<td>2.4</td>
</tr>
<tr>
<td>SW1736</td>
<td>TC</td>
<td>8.5 ± 0.1</td>
<td>&gt;20</td>
<td>4.7 ± 0.1</td>
<td>1.8</td>
</tr>
<tr>
<td>P31</td>
<td>Mesothelioma</td>
<td>2.5 ± 0.1</td>
<td>&gt;20</td>
<td>0.5 ± 0.1</td>
<td>5.0</td>
</tr>
<tr>
<td>VMC6</td>
<td>Mesothelioma</td>
<td>11.8 ± 0.1</td>
<td>&gt;20</td>
<td>4.4 ± 0.3</td>
<td>2.7</td>
</tr>
<tr>
<td>VMC12</td>
<td>Mesothelioma</td>
<td>11.3 ± 0.3</td>
<td>&gt;20</td>
<td>3.4 ± 0.1</td>
<td>3.3</td>
</tr>
<tr>
<td>VMC23</td>
<td>Mesothelioma</td>
<td>12.3 ± 0.2</td>
<td>16.9 ± 0.2</td>
<td>4.7 ± 0.3</td>
<td>2.6</td>
</tr>
<tr>
<td>VMC31</td>
<td>Mesothelioma</td>
<td>18.9 ± 0.2</td>
<td>&gt;20</td>
<td>4.8 ± 0.2</td>
<td>3.9</td>
</tr>
</tbody>
</table>

Abbreviations: CC, colon carcinoma; OS, osteosarcoma; TC, thyroid carcinoma.

*Fold decrease of ATO IC₅₀ calculated by dividing IC₅₀ of ATO alone by IC₅₀ of ATO with 2.5 μmol/L erlo.

Erlotinib inhibits ATO-induced EGFR stimulation

To investigate whether, in accordance to other studies (15–19), treatment with ATO leads to induction of EGFR signaling, the EGFR phosphorylation status of ATO-treated A549 cells was analyzed by Western blotting. This cell line was selected, as it is known for its notorious drug resistance against multiple chemotherapeutics. As presented in Fig. 2C and Supplementary Fig. S2A, respectively, ATO time- and dose-dependently increased EGFR phosphorylation at Y1068 and Y992 sites with the most pronounced phosphorylation detected after 16 hours. Notably, also in the case of erlotinib, a slight increase of EGFR phosphorylation was detected at the Y992 but not the Y1068 site. Treatment with erlotinib completely inhibited ATO-induced EGFR phosphorylation at both analyzed tyrosine sites (Y1068 and Y992).

AT0 synergizes with inhibitors of the EGFR downstream signaling

EGFR signaling is known to result in the activation of the MAPK and the PI3K/AKT kinase pathways (23). To gain more insights into the role of these EGFR downstream signaling pathways in the synergistic effects of ATO with erlotinib, the phosphoinositide 3-kinase (PI3K)-inhibitor Ly294002 and the AKT-inhibitor MK-2206 were used to block the PI3K/AKT pathway, whereas for the MAPK pathway the mitogen-activated protein/extracellular signal-regulated kinase (MEK)-inhibitor U0126 was applied (Fig. 2D–F and Supplementary Fig. S2B–S2D). Notably, strong synergism was observed by combination of ATO with all 3 kinase inhibitors with CI values between 0.73 and 0.02 for MK-2206, 0.93 and 0.3 for Ly294002, and 0.9 and 0.1 for U0126. With regard to pathway stimulation, ATO monotherapy induced phosphorylation of AKT (in accordance to Liu and colleagues, ref. 30), which was inhibited by cotreatment with erlotinib as well as both inhibitors of the PI3K/AKT pathway (Fig. 2D and E, bottom). Notably, no stimulation of ERK phosphorylation by ATO monotherapy was observed (Fig. 2F, bottom). However, although ATO monotherapy had no visible effect on the ERK phosphorylation status, the combination...
with erlotinib further enhanced the erlotinib-induced ERK inhibition indicating that ATO also modulates the anticancer activity of erlotinib.

**Effects of erlotinib on ATO-induced apoptosis and cell-cycle arrest**

In a next step, it was investigated whether the observed synergism was due to enhanced apoptosis induction or the inhibition of cell proliferation (Fig. 3A). Notably, no increase of apoptosis rates was observed after 24-hour drug exposure in any of the experimental groups (Supplementary Fig. S3A and S3B). Also, after 48 hours, no induction of programmed cell death was observed in A549 cells treated with either ATO or erlotinib monotreatment. In contrast, combination of the 2 drugs resulted in up to 10-fold raise of the apoptosis levels (Supplementary Fig. S3C). This effect was further increased after 72-hour drug exposure leading to up to 90% PI-positive cells in the highest combination setting (10 μmol/L of each drug; Fig. 3B). This time-dependent effect was confirmed by detection of caspase-dependent PARP cleavage (Fig. 3C).

To investigate whether ATO in combination with erlotinib influences DNA synthesis, ³H-thymidine incorporation assays were conducted (Fig. 3F). After 24-hour incubation, only a slight decrease of DNA synthesis was detected after ATO monotreatment, whereas 5 and 10 μmol/L erlotinib blocked replication by 22% and 43%, respectively. In contrast, ATO cotreatment with erlotinib induced a strong synergistic decrease of ³H-thymidine incorporation at all concentrations resulting in CI values
less than 0.5. This block of DNA synthesis was accompanied by distinct changes in the cell-cycle distribution. Thus, cells treated with 10 μmol/L ATO and 5 or 10 μmol/L erlotinib for 24 hours revealed a distinct increase in the amount of G2–M phase cells (from 12% to 22% and 26%, respectively; Supplementary Fig. S3D). This G2–M phase arrest further increased to 42% and 43% of G2–M cells after 48-hour combination treatment with the indicated drugs. D, the influence of ATO/erlotinib on cell-cycle distribution of A549 cells was determined by FACS analyses after 48-hour drug exposure. E, the impact of 24-hour drug exposure on the expression levels of cyclin B in A549 cells was determined by Western blot analysis. F, the impact of ATO and erlotinib on DNA synthesis was measured by 3H-thymidine incorporation after 24-hour incubation with the indicated drugs. Data are compared with those for cell viability determined by MTT assay. The statistical significance was calculated using two-way ANOVA (*, P < 0.05; **, P < 0.001; ***, P < 0.001; ns, not significant). If not otherwise indicated, significance is given in comparison with control group. C, Western blot analysis of PARP cleavage in A549 cells was determined after 16-, 24-, and 48-hour treatment with the indicated drugs.

Effects of the drug combination on DNA damage and repair

Besides its oncogenic activity, EGFR has recently been shown to promote DNA DSB repair by regulation of nonhomologous end-joining (34–36) as well as homologous end-joining (22, 37). To detect the amount of DNA DSBs after drug treatment, phosphorylation of H2AX histones was analyzed by immunostaining. Notably, at the concentrations used the amount of DSBs induced by the single agents was very minor reflecting the distinct resensitization of A549 cells to ATO by erlotinib.

Synergistic Anticancer Activity of ATO with Erlotinib

Synergistic Anticancer Activity of ATO with Erlotinib

Figure 3. Impact of the ATO and erlotinib (erlo) drug combination on apoptosis induction, cell-cycle distribution, and DNA synthesis. A, phase-contrast microscopy pictures depict A549 cells after 72-hour treatment with ATO, erlotinib, or their combination at the indicated concentrations. B, apoptotic A549 cells were counted after ATO/erlotinib exposure for 72 hours on live cells stained with Hoechst 33258 and PI. The statistical significance was calculated using two-way ANOVA (*, P < 0.05; **, P < 0.001; ***, P < 0.001; ns, not significant). If not otherwise indicated, significance is given in comparison with control group. C, Western blot analysis of PARP cleavage in A549 cells was determined after 16-, 24-, and 48-hour treatment with the indicated drugs. D, the in...
therapy resistance of A549 cells against both drugs. In contrast, the drug combination considerably induced H2AX nuclear foci formation (Fig. 4A). Phosphorylation of H2AX was also confirmed by Western blotting after 16-, 24-, and 48-hour drug treatment (Fig. 4B). Also, the comet assays confirmed that the amount of DNA DSBs was significantly higher after 6-hour treatment with the drug combination setting than in the monotherapies (Fig. 4C). In accordance with these data, the combination of ATO with erlotinib led to an earlier induction of DNA damage response signals such as p53 and p21 (Fig. 4B). Notewor-thy, the enhancement of DNA damage was not based on generation of hydroxyl or superoxide radicals upon combi-nation of ATO with erlotinib (data not shown). In contrast, the drug combination led to distinct downregu-lation of the DNA DSB repair proteins Rad50 and Rad51 as well as a concentration-dependent decrease of BRCA1 phosphorylation (Fig. 4D).

Synergistic activity of ATO and erlotinib in vivo

On the basis of the promising cell culture data, the efficacy of the ATO/erlotinib combination was tested against A549 xenografts in SCID mice. All treatment settings were well tolerated with no signs of toxicity, distress, or impact on activity/behavior parameters (data not shown). This is also reflected by the unaltered body...
weight in all experimental groups (Fig. 5A). With regard to the anticancer activity, due to the high resistance of A549 cells to the single agents, neither ATO nor erlotinib alone had any anticancer activity (Fig. 5B). In contrast, treatment with the drug combination resulted in significant reduction of tumor growth. In addition, the therapeutic efficacy of the new combination strategy was also tested in the orthotopic mesothelioma model P31 as this cell line was among the most responsive ones to ATO treatment in the MTT assays. Figure 5C depicts mean intraperitoneal tumor mass after 2 weeks of drug treatment (example pictures are shown in Supplementary Fig. S4). Also, in this xenograft, erlotinib monotherapy was found to be widely ineffective. In contrast, ATO monotherapy already had some growth inhibitory potential resulting in a significant reduction from a mean of 2.40 g to a mean of 0.96 g in the ATO-treated mice. Combination of ATO with erlotinib further enhanced this activity (to a mean of 0.52 g) emphasizing the potential of the treatment regimen.

Discussion

Intrinsic and acquired resistance is still one of the major obstacles for successful anticancer therapy (38). Besides the development of new therapeutic approaches, one of the most promising options to improve treatment efficacy is the use of combination strategies (39). The anticancer activity of arsenic is well known and resulted in the recent approval of ATO against APL (1). However, the activity of ATO against solid cancer types is so far rather limited (8–11, 40). Some recent reports indicate that exposure to inorganic arsenic salts leads to activation of EGFR in nonmalignant as well as in malignant cells, for example, via c-Src kinase-mediated signals (14–19). In many solid tumor types, EGFR is one of the most important oncogenic receptor tyrosine kinases significantly supporting cell proliferation and survival via the MAPK and the PI3K/AKT axes, respectively (23, 41, 42). Consequently, we hypothesized that EGFR stimulation might play an important role in the resistance of solid cancer types against ATO. Indeed, our data show that inhibition of EGFR by erlotinib and gefitinib sensitizes diverse and even highly ATO-resistant solid tumor types of epithelial and mesenchymal origin against ATO, independent of their TP53 or RAS mutation status (Supplementary Table S1). Moreover, our results suggest that these effects are mainly based on inhibition of EGFR-mediated DNA DSB repair associated with higher levels of DNA damage, subsequent cell-cycle arrest predominantly in G2–M phase, and apoptosis induction. The synergistic activity of this drug combination was finally also confirmed using 2 human solid tumor xenograft models in vivo. A schematic
The synergism between either EGFR or MAPK/P38K inhibition and ATO has never been attributed to enhanced DNA damage based on reduced DSB repair so far. This is surprising, as both ATO and inhibition of EGFR and its downstream signals have been shown to result in compromised DNA repair processes (22, 34–37). Interestingly, we could not detect an enhanced generation of ROS in the drug combination despite synergistic induction of DNA DSBs and subsequent phosphorylation of H2AX. These data again suggest that reduced repair rather than enhanced damage caused a synergistic proliferation arrest and apoptosis induction by the ATO/erlotinib combination.

In general, the role of DNA damage in the anticancer activity of arsenic is controversially discussed (47, 48). Although it is widely assumed that the carcinogenic effects of chronic arsenic exposure are associated with arsenic-induced reactive species, which subsequently result in single-strand as well as DNA DSBs (47, 48), the underlying mechanisms seem to be complex. In addition, ROS-mediated DNA damage might be enhanced on the basis of downregulation of several ssDNA damage repair mechanisms by arsenic (e.g., decreased expression of the nucleotide excision repair (NER) proteins ERCC1, XBP, and XPF in lymphocytes from exposed individuals (49)). Accordingly, a high number of indirect indications for enhanced DNA damage after arsenic exposure have been reported. For example, ATO induced base excision repair (BER)-specific DNA polymerase β (Polβ) activity indicative for ROS-mediated single-strand breaks (44). In addition, there is increasing evidence that especially deficiency in DNA DSB repair is associated with increased sensitivity to arsenic (50). Notably, besides multiple other functions, the EGFR pathway has been recently shown to be involved in the regulation of DNA DSB repair by positive regulation of both the homologous recombination as well as the nonhomologous end-joining (22, 37). Thus, activation of EGFR resulted in a decreased number of residual DNA DSBs, whereas the number of H2AX-positive DSB foci was clearly increased when EGFR was blocked by erlotinib in A549 and other lung cancer cell lines (22, 34). Moreover, erlotinib also attenuated DNA damage-induced Rad51 foci and resulted in cytoplasmic retention of BRCA1 (37), both essential components of the DSB repair machinery. With regard to our data, these findings strongly support the hypothesis that the synergism between ATO and EGFR inhibition observed in diverse solid tumor models is mainly caused by a synthetic lethal interaction between ATO and EGFR-induced DNA damage and loss of EGFR-mediated repair capacity. Accordingly, the combination of ATO with erlotinib resulted in our hands in downregulation of the DNA DSB repair proteins Rad51 and Rad50 as well as reduced phosphorylation of BRCA1 associated with a significant increase of DNA DSBs, profound activation of pH2AX, and p53/p21–mediated cell-cycle arrest. The fact that this combination is strongly active even against the notoriously drug-resistant NSCLC model A549 indicates that such cross-talk might also exist for other DNA-damaging anticancer drugs. Moreover, we show for the first time that a synergistic anticancer activity of ATO is also achievable by coadministration of the EGFR inhibitor erlotinib in vivo when treating subcutaneous and orthotopic solid tumor xenografts. On the one hand, this implicates that ATO as anticancer agent might expand toward novel indications against solid tumors when combined with EGFR inhibitors. On the other hand, it has to be considered that EGFR tyrosine kinase inhibitors such as erlotinib have been successfully used predominantly in tumors harboring constitutively activating EGFR-mutations (23). However, the widespread sensitization toward erlotinib by coadministration of ATO reported in this study both in vitro and in vivo was observed in cell models without EGFR-activating mutations. Thus, the combination with ATO might expand the beneficial application of EGFR inhibitors toward patient populations harboring cancers with wild-type EGFR background. Moreover, it has been hypothesized that EGFR activation might play a role in the carcinogenic effects of chronic exposure to arsenic (16). Thus, one might hypothesize that combination with EGFR inhibitors might not only sensitize cancer cells toward ATO but at the same time reduce the potential procarcinogenic effects of ATO cancer therapy.

Summarizing, our data suggest that the combination of ATO with erlotinib or other EGFR inhibitors might be a promising strategy to enhance the susceptibility of solid cancer cells to ATO treatment and, thus, overcome drug resistance. This is of special interest as both, ATO as well as erlotinib, are already in clinical use, which facilitates the performance of clinical combination trials.
Disclosure of Potential Conflicts of Interest

No potential conflicts of interest were disclosed.

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References


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