In-Depth Analysis Shows Synergy between Erlotinib and miR-34a

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Abstract

Tyrosine kinase inhibitors directed against epidermal growth factor receptor (EGFR-TKI), such as erlotinib, are effective in a limited fraction of non-small cell lung cancer (NSCLC). However, the majority of NSCLC and other cancer types remain resistant. Therapeutic miRNA mimics modeled after endogenous tumor suppressor miRNAs inhibit tumor growth by repressing multiple oncogenes at once and, therefore, may be used to augment drug sensitivity. Here, we investigated the relationship of miR-34a and erlotinib and determined the therapeutic activity of the combination in NSCLC cells with primary and acquired erlotinib resistance. The drug combination was also tested in a panel of hepatocellular carcinoma cells (HCC), a cancer type known to be refractory to erlotinib. Using multiple analytical approaches, drug-induced inhibition of cancer cell proliferation was determined to reveal additive, antagonistic or synergistic effects. Our data show a strong synergistic interaction between erlotinib and miR-34a mimics in all cancer cells tested. Synergy was observed across a range of different dose levels and drug ratios, reducing IC50 dose requirements for erlotinib and miR-34a by up to 46-fold and 13-fold, respectively. Maximal synergy was detected at dosages that provide a high level of cancer cell inhibition beyond the one that is induced by the single agents alone and, thus, is of clinical relevance. The data suggest that a majority of NSCLC and other cancers previously not suited for erlotinib may prove sensitive to the drug when used in combination with a miR-34a-based therapy.

Introduction

Lung cancer accounts for the most cancer-related deaths in both men and women [1]. Targeted therapies are used depending on the cancer genotype or stage of disease and includes erlotinib, a small molecule inhibitor directed against epidermal growth factor receptor (EGFR). Erlotinib functions as competitive inhibitor of ATP-binding at the active site of the EGFR kinase [2]. Clinical trials investigating EGFR inhibitors revealed that responses occurred in a selective fraction of lung cancer patients, preferentially in never-smokers diagnosed with activating mutations in the EGFR gene and adenocarcinoma or bronchioalveolar histotype [3,4]. However, a majority of non-small cell lung cancer (NSCLC) patients remained resistant. Primary and secondary resistance has been associated with activating KRAS mutations that may co-exist with EGFR mutations despite the fact that KRAS and EGFR mutations appeared to be predominantly mutually exclusive [5,6], an acquisition of a second mutation in the catalytic domain of EGFR (usually T790M) [7], an amplification and overexpression of receptor kinase MET and its ligand HGF, providing signals into the PI3K pathway and substituting for an inactivation of EGFR [8], increased expression of the receptor kinase AXL and its ligand GAS6 [9,10], and several others [11–15].

The therapeutic use of tumor suppressor miRNAs continues to raise much attention because they can interfere with many different oncogenic pathways at once by broadly targeting multiple oncogenes [16,17]. Certain miRNAs can also suppress the properties of cancer stem cells, a cell species particularly resistant to many several cancer therapies [18,19]. Because of these features, synthetic mimics of tumor suppressor miRNAs have become attractive tools to battle cancer as mono-therapies but also to break resistance mechanisms, making current treatment regimens more effective. An example is provided by miR-34a that inhibits a broad range of cancer cell types in culture and in preclinical animal studies [20]. A liposomal formulation, MRX34, loaded with miR-34a mimics has recently entered clinical testing for patients with hepatocellular carcinoma (HCC) and other cancers with metastatic lesions in the liver, including cancers of the lung [21]. There are several studies demonstrating how miR-34a can sensitize cancer cells to conventional cytotoxic chemotherapies [18,22–26]. However, the combination of tumor suppressor miRNAs with targeted therapies is a less well studied approach. Due to the broad anti-oncogenic activity of miR-34a, and the fact that both MET and AXL are directly repressed by miR-34a [27–29], we rationalized that miR-34a can sensitize cancer cells to erlotinib.

Here, we employed multiple analytical approaches to distinguish between additive, antagonistic and synergistic drug interactions and studied the effects of the erlotinib-miR-34a combination in a panel of NSCLC and HCC cell lines with primary or acquired...
erlotinib resistance. Our data indicate strong synergy in all cell lines tested for various drug-drug ratios. Importantly, miR-34a and erlotinib cooperated synergistically at dose levels that induce maximal cancer cell inhibition, one that is greater than the inhibition achieved by either agent alone. Thus, our results demonstrate how the therapeutic application of erlotinib can be expanded to other cancers and point to a novel combination therapy that could quickly be implemented in the clinic.

Materials and Methods

Cell Lines

Human non-small cell lung cancer (NSCLC) cell lines A549, H460, H1299, H226, HCC827 parental and HCC827res were used to assess the combinatorial effects of miR-34a and EGFR-TKIs. The particular cell lines were selected based on the high IC50 values of EGFR-TKIs in these cells, their oncogenic properties and susceptibility to miRNAs. These cell lines are either erlotinib resistant (A549, H460, H1299, H226) or sensitive (HCC827). In addition, cell lines with acquired resistance were created by applying increased selective pressure of erlotinib over ten weeks, starting at an equivalent of IC10 and ending at an IC50 equivalent. As cellular proliferation exhibited normal doubling rates under IC50 selection, the resistant cells were plated at a low dilution (HCC827res) or high dilution to create near-pure, erlotinib resistant clones (HCC827res-3, 5, 6 and 7). To study effects in hepatocellular carcinoma (HCC) cells, Hep3B, Huh7, C3A and HepG2 were used. Huh7 cells were acquired from the Japanese Collection of Research Bioresources Cell Bank. All other parental and resistant clones (HCC827res) were purchased from the American Type Culture Collection (ATCC, Manassas, VA) and cultured according to the supplier’s instructions.

RNA Isolation and qRT-PCR

Total RNA from cell pellets was isolated using the mirVANA PARIS RNA isolation kit (Ambion, Austin, TX) following the manufacturer’s instructions. RNA concentration was determined by absorbance measurement (A260) on a Nanodrop ND-1000 (Thermo Scientific, Wilmington, DE). For the quantification of miRNA and mRNA by quantitative reverse-transcription polymerase chain reaction (qRT-PCR), we used commercially available reagents. The RNA was converted to cDNA using MMLV-RT (Invitrogen, Carlsbad, CA) under the following conditions: 4°C for 15 min; 16°C for 30 min; 42°C for 30 min; 85°C for 5 min. Following cDNA synthesis, qPCR was performed on 2 μL of cDNA on the ABI Prism 7900HT SDS (Applied Biosystems, Life Technologies, Foster City, CA) using Platinum Taq Polymerase (Invitrogen) under the following cycling conditions: 95°C for 1 min (initial denature); then 50 cycles of 95°C for 5 sec, 60°C for 30 sec. TaqMan Gene Expression Assays and TaqMan MicroRNA Assays were used for expression analysis of miRNA and mRNA in all lung and liver cell lines. For miRNA expression, additions to the manufacturers’ reagents include DMSO (final concentration of 6%) and tetramethylammoniumchloride (TMAC; final concentration of 50 mM in both RT and PCR) to improve the slope, linearity and sensitivity of the miRNA assays. Expression levels of both miRNA and mRNA were determined by relative quantitation to the HCC827 parental cell line. The raw Ct values of the miRNA and mRNA targets were normalized to selected housekeeping genes to create delta-Ct values, converted to linear space and then expressed as percentage expression.

miRNA and Erlotinib Treatment

Erlotinib hydrochloride was purchased from LC Laboratories (Woburn, MA). Synthetic miR-34a and miR-NC mimics were manufactured by Life Technologies (Ambion, Austin, TX). To determine the IC50 value of each drug alone, 2,000–3,000 cells per well were seeded in a 96-well plate format and treated with either erlotinib or miR-34a as follows. (i) miR-34a mimics were reverse-transfected in triplicates in a serial dilution (0.03–30 nM) using RNAiMax lipofectamine from Invitrogen according to a published protocol [30]. As controls, cells were also transfected with RNAiMax alone (mock) or in complex with a negative control miRNA mimic (miR-NC). Cells were incubated with AlamarBlue (Invitrogen) 4 days or 6 days post transfection to determine cellular proliferation of lung or liver cancer cells, respectively. Proliferation data were normalized to mock-transfected cells. (ii) Erlotinib, prepared as a 10 and 20 mM stock solution in dimethyloxide (DMSO), was added to cells one day after seeding at a final concentration ranging from 0.1 and 100 μM. Solvent alone (0.5% final DMSO in H226 and HCC827, 1% final DMSO in all other cell lines) was added to cells in separate wells as a control. Three days thereafter, cellular proliferation was measured by AlamarBlue and normalized to the solvent control.

Regression Trendlines & IC50 Values

Linear and non-linear regression trendlines were generated using the CompuSyn (ComBioSyn, Inc, Paramus, NJ) and Graphpad (Prism) softwares, respectively. The non-linear trendlines provided a better fit for the actual data and were used to calculate IC50, IC25 and other drug concentrations (ICn), although both softwares generated similar values.

Combination Effects Determined by the “Fixed Concentration” Method

The “Fixed Concentration” method was used for cell lines with acquired resistance (HCC827res). Cells were reverse-transfected with miR-34a using the miRNA at a fixed, weak concentration (~IC25) as described above. The following day, cells were treated with erlotinib in a serial dilution (0.01–100 μM). Cell proliferation inhibition was analyzed 3 days later by AlamarBlue. To measure the effects of the single agents and to correct for effects potentially contributed by lipid carrier or vehicle, cells were also treated with miR-34a in combination with solvent (0.5% DMSO in HCC827res, 1% DMSO in all other cell lines) or erlotinib in combination with mock-transfection. All proliferation data was normalized to mock-transfected cells treated with solvent (DMSO). The combinatorial effect was evaluated by a visual inspection of the erlotinib dose-response curve and a shift of the IC50 value in the presence or absence of miR-34a (graphed and calculated using Graphpad).

Combination Effects Determined by the “Fixed Ratio” Method

Cells were treated with 7 concentrations of erlotinib each in combination with 7 concentrations of miR-34a. Each drug was used at a concentration approximately equal to its IC50 and at concentrations within 2.5-fold (NSCLC) or 2-fold (HCC) increments above or below. This matrix yielded a total of 49 different combinations representing 13 different ratios. Each drug was also used alone at these concentrations. miR-34a and erlotinib were added as described above, and cellular proliferation was determined by AlamarBlue. Each data point was performed in triplicates.
Calculation of Combination Index (CI) Values

Combination index (CI) values based on Loewe’s additivity model were determined to assess the nature of drug-drug interactions that can be additive (CI = 1), antagonistic (CI>1), or synergistic (CI<1) for various drug-drug concentrations and effect levels (Fa, fraction affected; inhibition of cancer cell proliferation) [31–33]. Both linear regression and non-linear regression trendlines were used to calculate and compare CI values. CI values based on linear regression analysis was done using the CompuSyn software (ComboSyn Inc., Paramus, NJ), following the method by Chou et al., whereby the hyperbolic and sigmoidal dose-effect curves are transformed into a linear form [31,34]. CI values derived from non-linear regression trendlines were calculated using Equation 1 in which CAx and CBx are the concentrations of drug A and drug B in the combination to produce effect X (Fa). ICxA and ICxB are the concentrations of drug A and drug B used as a single agent to produce that same effect.

\[
CI = \frac{C_{Ax}}{IC_{50,A}} + \frac{C_{Bx}}{IC_{50,B}}
\] (1)

Drug concentrations required in Equation 1 to determine CI values (CAx, CBx, ICxA and ICxB) were calculated using the Hill equation [Equation 2], IC50 and Hill slope value (n) derived from non-linear regression trendlines (Graphpad).

\[
E = E_{max} \times \frac{C^n}{IC50^n + C^n}
\] (2)

Isobolograms

To describe the dose-dependent interaction of erlotinib and miR-34a, isobolograms at effect levels of 50% and 80% inhibition of cancer cell proliferation were created. Since the single agents – alone or in combination – usually reached 50% cancer cell inhibition, the 50% isobologram provided an actual comparison of the single use vs. the combination. The 80% isobologram was used to illustrate the utility of the combination at a high effect level that have practical implications in oncology. In each of these, additivity was determined by extrapolating the dose requirements for each drug in combination from its single use (IC50, IC90). Data points above or below the line of additivity indicate antagonism or synergy, respectively. For all 49 combinations, drug concentrations required in the combination were compared to those of the single agents alone to reach the same effect and expressed as a fold change (dose reduction index, DRI).

Curve Shift Analysis

To allow a direct comparison of the dose-response curves and to identify synergistic drug-drug interaction, non-linear regression trendlines of each drug alone or of the combination (IC50/IC90 ratio or other ratios where indicated) were normalized to its own IC50 value and referred to as IC50 equivalents (IC50 eq). IC50 equivalents of the combination were calculated using Equation 3 and described in [35]. Data of the single agents and in combination were graphed in the same diagram to illustrate lower drug concentrations required to achieve any given effect relative to the single agents. This is represented in a left-shift of the dose-response curve and indicates synergy [35].

\[
IC_{50eq} = \frac{C_{Ax}}{IC_{50,A}} + \frac{C_{Bx}}{IC_{50,B}}
\] (3)

Statistical Analysis

Statistical analysis was done using the Excel (Microsoft), CompuSyn and Graphpad softwares. Averages and standard deviations were calculated from triplicate experiments. Goodness of fit of linear and non-linear regression trendlines was described by R (CompuSyn) and R² (Graphpad) values, respectively, and were >0.9 for most cell lines except H226 and HepG2 cells due to limiting drug insensitivity.

Results

miR-34a Restores Sensitivity to Erlotinib in Non-small Cell Lung Cancer Cells

To study drug resistance in cells with acquired resistance, we used HCC827 cells that express an activating EGFR mutation (deletion of exon 19 resulting in deletion of amino acids 746–750). HCC827 are highly sensitive to erlotinib with an IC50 value of 0.022 µM (Fig. 1A). Erlotinib-resistant cell lines were developed by exposing the parental HCC827 cells to increasing erlotinib concentrations over the course of 10 weeks until the culture showed no signs of growth inhibition at a concentration that is equivalent to IC50 in the parental cell line (Fig. 1B). During this process, individual cell clones (HCC827™#2, #6, #7) as well as a pool of resistant cells (HCC827™®) were propagated. Total RNA was isolated and probed by quantitative PCR for levels of miR-34 family members and genes known to induce resistance. In agreement with previously published data, HCC827 cells resistant to erlotinib showed increased mRNA levels of MET and its ligand HGF that presumably function to bypass EGFR signaling (Fig. S1) [8]. In contrast, expression levels of other genes also associated with resistance, such as AXL, GAS6, KRAS, FGFR1, ERBB3, PIK3CA, and EGRF itself, were not elevated. Levels of miR-34b/c family members were reduced in several of the resistant HCC827 cells (Fig. S1). Interestingly, miR-34a was not reduced in erlotinib-resistant HCC827 cells suggesting that miR-34a does not play a causal role in the onset of resistance in these cells which can occur independently of miR-34 by amplification of the MET gene [8].

Since both MET and AXL are directly repressed by miR-34, and because inhibition of AXL can antagonize erlotinib resistance [10,36], the introduction of synthetic miR-34 mimics may restore erlotinib sensitivity. To explore this possibility, HCC827™ cells were exposed to increasing erlotinib concentrations, ranging from 0.03–100 µM, either in the absence or presence of miR-34a used at a fixed, weak concentration (0.3 nM). The effects of erlotinib were expected to be concentration-dependent, such that erlotinib in combination with miR-34a produced lower IC50 values relative to erlotinib alone. As shown in Figure 1C, erlotinib was not very potent in HCC827™ cells (IC50 = 25.2 µM). However, when used in combination with miR-34a, the erlotinib IC50 value decreased to 0.094 µM. This result suggests that adding a small amount of miR-34a is capable of restoring erlotinib sensitivity that is similar to the one of parental HCC827 cells. The effects were specific to the miR-34a sequence as the addition of a negative control miRNA (mir-NC) did not improve the potency of erlotinib (Fig. 1C). Thus, the data generated in HCC827™ cells indicate that miR-34a can sensitize cancer cells with acquired erlotinib resistance.
To determine whether the miRNA can also counteract primary resistance mechanisms, we used H1299 cells that have mutations in the NRAS and TP53 genes [37]. In these cells, erlotinib produced an IC\textsubscript{50} value of 11.0 \textmu M (Fig. 1D) in accord with previous results showing EGFR-TKI IC\textsubscript{50} values between 8.6 and 38 \textmu M [38,39]. In combination with 0.3 nM miR-34a, the erlotinib dose-response curve shifted along the x-axis, indicating an approximately 4-fold lower IC\textsubscript{50} value (3.0 \textmu M). This result is in contrast to miR-NC that did not alter the potency of erlotinib, and suggests that miR-34a sensitizes non-small lung cancer cells with both acquired as well as primary resistance.

miR-34a and Erlotinib Synergize in Non-small Cell Lung Cancer Cells

The shift of the erlotinib IC\textsubscript{50} value demonstrated how a fixed miR-34a concentration can improve the potency of erlotinib. However, this model, also known as “Fixed-Concentration-Model”, does not allow the assessment of synergy. To investigate whether both drugs can enhance each other, we employed the “Fixed-Ratio-Model” that is based on Loewe’s concept of additivity [31–33]. In this model, combination index (CI) values are calculated based on the slope and IC\textsubscript{50} value of each dose-response curve (drug alone or in combination) and define whether the drug-drug interactions are synergistic (CI<1), additive (CI = 1), or antagonistic (CI>1). Since the accuracy of the CI values depends on the fit of the dose-response curve trendline, CI values were calculated by two methods using either linear or non-linear regression trendlines (see Materials and Methods). Four erlotinib-resistant cell lines were used, all of which differ in their genetic make-up: A549 (mutations in KRAS, STK11, CDKN2A), H460 (mutations in KRAS, STK11, CDKN2A, PIK3CA), H1299 (mutations in NRAS, TP53), and H226 (mutations in CDKN2A) [37]. A qRT-PCR analysis showed a marked increase of AXL, GAS6 and FGFR1 mRNA levels in these cells relative to erlotinib-sensitive HCC827 cells, further providing an explanation for erlotinib resistance (Fig. S1). Levels of miR-34 were significantly reduced in H1299 and H460 cells. In a first step, erlotinib or miR-34a were added to cells in a serial dilution to determine IC\textsubscript{50} values of each drug alone. For erlotinib, these ranged between 4.2 and >50 \textmu M and are within the range of published data [38–42] (Fig. S2). The IC\textsubscript{50} values of miR-34a ranged from 0.4 to 15.6 nM. Neither drug was capable of 100% cancer cell inhibition, nor did the maximal activity of either drug exceed 75%. Erlotinib and miR-34a were least effective in H226 cells,
yielding theoretical IC$_{50}$ values as a result of an extrapolation of the dose-response curve. In a second step, each drug was used at a concentration equal to its own approximate IC$_{50}$ value, as well as at multiples thereof above and below (fixed ratio). As controls, each drug was used at these concentrations alone. Both linear and non-linear regression models produced CI values that were well below 1.0 in all cell lines tested indicating strong synergy (Fig. 2A). CI values we considered relevant are those below 0.6. In most cell lines, synergy was observed at higher dose levels and at higher magnitude of cancer cell inhibition. This is critical because a practical application of the drug combination calls for synergy at maximal cancer cell inhibition (75% inhibition or greater). In general, the non-linear regression trendline provided a better fit for the actual data, although both models generated similar results.

Next, we generated isobolograms and determined the dose requirements for each drug at 50% and 80% cancer cell inhibition as a read-out for synergy. The 50% effect level was chosen because the potency of a drug is frequently assessed at its IC$_{50}$ and because in our studies each drug alone was capable of inhibiting most cancer cells by 50%, allowing a comparison of each drug alone with the combination within the range of actual data. The 80% effect level was chosen because it is important to demonstrate synergy at high inhibitory activity for oncology applications. Although the concentrations of each drug alone to achieve 80% inhibition are based on an extrapolation of the dose-response curves and are theoretical in nature, the miR-34a-erlotinib combination readily achieved 80% inhibition or greater and is within the range of actual data. Since the two drugs by themselves were not very effective in H226 cells, isobolograms at 30% and 50% inhibition were created for H226 data. As shown in Figure 2B, the isobole of the combination was well below the additive isobole for every cell line and effect level indicating strong synergy. The dose requirement for erlotinib decreased to 2 µM or less in most cell lines to achieve 50% inhibition, reducing the dose by 4- to 46-fold. Likewise, the required concentration of miR-34a was also substantially less in the combination relative to miR-34a alone, reducing its dose by 7- to 13-fold. This reduction in dose level, along referred to as dose reduction index (DRI), was markedly evident at 80% inhibition at which the dose requirements were reduced by up to 26-fold (erlotinib) and 39-fold (miR-34a).

Third, we performed curve-shift analyses whereby the concentration of each drug has been normalized to its own IC$_{50}$ value [35]. This conversion of drug concentrations into IC$_{50}$ equivalents (IC$_{50}$ eq) allows a direct comparison of each dose-response curve from the single agents and the combination. Trendlines were generated and span effect levels from 0–100% inhibition. The slope of the trendline indicates drug potency, and the maximal activity can be gaged from actual data points. Synergy is identified when IC$_{50}$ equivalents of the combination are lower to achieve an equal effect relative to the single agents [35]. This is visually indicated by a left-shift of the combination trendline. As seen in Figure 2C, the combination is well separated from the single agents indicating synergy. In H460 and H226 cells, the IC$_{50}$ equivalents of the combination are greater at low effect levels (0–25%) and lower at effect levels above 30% compared to those of the single agents. This observation agrees with data from CI plots showing antagonism below 25% inhibition and synergy above 25% inhibition in these cells (Fig. 1A). Thus, the analysis reveals synergistic effects for drug concentrations that induce a high level of cancer cell inhibition. A benefit for the combination is further demonstrated by the fact that the actual level of inhibition is greater for the combination relative to the single agents – the maximal activity of the single drugs is no greater than 75% and can be extended beyond 90% when used in combination.

**Multiple Ratios of Erlotinib and miR-34a Cooperate Synergistically**

Our analysis suggests that erlotinib and miR-34a synergize when the two drugs are combined at a ratio derived from their IC$_{50}$ values. Because drug-drug interactions can change depending on the relative amounts, we explored the effects of multiple erlotinib-miR-34a ratios by combining erlotinib at concentrations from 0.41–100 µM with miR-34a at concentrations from 0.12–30 nM. Drug doses were increased in 2.5-fold increments, and each drug was also used alone as controls. This matrix yielded 49 drug combinations representing 13 different drug ratios (Fig. 3A). Levels of cancer cell inhibition, CI and DRI values were determined for each combination and graphed in CI plots, isobolograms and curve-shift diagrams. For the sake of this exercise, we focused on combinations in which miR-34a and erlotinib were added in an IC$_{50a}$/IC$_{50b}$ ratio (molar ratio 1:333) and the following molar-based ratios: 1:533, 1:1333, 1:8333 and 1:20833.

Calculated CI values predict that erlotinib and miR-34a combined at all of these ratios provide strong synergy (Fig. 3B). At effect levels greater than 75% inhibition, CI values were below 0.2. The ratios that contained higher amounts of erlotinib provided lower synergy at effect levels below ~75% and were slightly superior at effect levels above 75% inhibition. Similarly, the isobologram indicates strong synergy for various erlotinib-miR-34a ratios (Fig. 3C). Actual data points demonstrate that 30 nM miR-34a or 100 µM erlotinib are required to induce ~80% cancer cell inhibition when used as single agents. In contrast, the required dose levels of erlotinib in the combination were substantially decreased as miR-34a amounts were increased. For instance, merely 2.56 µM erlotinib was needed to induce ~80% inhibition when used with 12 nM miR-34a, thereby reducing the dose requirement of erlotinib by ~40-fold. Further evidence for the synergistic action of these ratios comes from curve-shift analyses that reveal much lower IC$_{50}$ equivalents of the combination compared with IC$_{50}$ values of the single agents alone (Fig. 3D). The IC$_{50}$ eq data correlate with CI data showing dose-dependent degrees of synergy among various ratios: low ratios show lower synergy at low effect levels which is reversed at high levels of cancer cell inhibition.

The full range of 49 combinations was also tested in H1299, H460 and H226 cells and confirmed the results obtained with A549 cells (Fig. S3). Multiple ratios provided good synergy, and the ones with higher potency clustered to the ones with higher drug concentrations. Among these were many that met our cutoffs and produced >75% cancer cell inhibition, CI <0.6, and DRI >2 for each drug.

**Erlotinib and miR-34a Cooperate Synergistically in Hepatocellular Carcinoma Cells**

To investigate whether the cooperative activity of erlotinib and miR-34a has utility in other cancer indications, we probed this combination in cell models of hepatocellular carcinoma. Liver cancer was chosen as test platform because erlotinib is moderately effective in patients with advanced liver as a single agent and failed to prolong overall survival and time-to-progression in combination with sorafenib [43–45]. In addition, MRX34, a miR-34a liposome currently in clinical testing, effectively eliminated liver tumors in preclinical animal studies and therefore may be an attractive agent in combination with erlotinib. Cell models used included Hep3B,
C3A, HepG2 and Huh7, several of which showed an upregulation of erlotinib-resistance genes, AXL, HGF, FGFR1 and ERBB3 in comparison to an erlotinib-sensitive lung cancer line (Fig. S4). Collectively, levels of miR-34 family members were low or undetectable in liver cancer cells. In agreement with our expectation, IC50 values of erlotinib were 25 μM or greater in these four cell lines (Fig. S5). The IC50 values of miR-34a ranged between 0.3 and 2.3 nM and, thus, were similar to those in lung cancer cells. These values were used as a guide to combine erlotinib and miR-34a at a fixed ratio of IC50:IC50 and to produce CI, isoboles and IC50 eq values (Fig. 4). In addition, each combination was also tested in a matrix of different concentrations to assess the combinatorial effects across multiple ratios (Fig. S6). Our data predict strong synergy between erlotinib and miR-34a in all cell lines tested. Synergy was observed at high levels of cancer cell inhibition and, hence, occurs within the desirable range of activity (Fig. 4A). This result is confirmed by the IC50 eq curve shift analyses indicating synergy at higher dose and effect levels. The analysis also shows that the maximal inhibitory activity of the combination is substantially expanded compared to those of the single agents (Fig. 4C). Isobolograms demonstrate a stark reduction of the erlotinib dose when used with miR-34a to induce

**Figure 2. Synergistic effects between miR-34a and erlotinib in NSCLC cells.** (A) Combination index (CI) analysis. CI values were generated by linear regression (red) and non-linear regression methods (blue). Trendlines indicate CI values at any given effect (Fa, fraction affected, % inhibition), and symbols represent CI values derived from actual data points. CI = 1, additivity; CI > 1, antagonism; CI < 1, synergy. (B) Isobologram analysis. The diagonal, dotted line indicates additivity, and the red symbol shows dose requirements to achieve 50% and 80% (A549, H1299, H460) or 30% and 50% (H226) cancer cell inhibition, respectively. Data points below the line of additivity indicate synergy, data points above denote antagonism. (C) Curve shift analysis. Data derived from non-linear regression trendlines were normalized to IC50 values of the single agents (IC50 eq) and plotted in the same graph. Left and right shifts of the dose-response curves of the combination (red dotted line) relative to the dose-response curves of the single agents (grey, black) indicate synergy or antagonism, respectively. Actual experimental data points are shown (symbols). doi:10.1371/journal.pone.0089105.g002
50% inhibition or greater, such as 80% (Fig. 4B). In combination, erlotinib can be used at concentrations as low as 2 \( \mu M \) to inhibit cancer cells by 50%, thereby lowering its dose by 75-fold compared to its single use (see HepG2). Synergy is not limited to a specific ratio but is apparent across most ratios tested (Fig. S6). Thus, the data are similar to those generated in lung cancer cells and predict enhanced efficacy for the erlotinib-miR-34a combination in cancers where erlotinib alone is insufficient.

**Discussion**

An accurate evaluation of drug-drug interactions is complex because outcomes depend on drug ratios, drug concentrations and desired potency [31]. To investigate the pharmacological relationship between miR-34a mimics and erlotinib, we used multiple analytical approaches to reveal drug enhancements (‘‘Fixed Concentration’’ model) and to distinguish between additivity, antagonism and synergy (‘‘Fixed Ratio’’ model). We examined CI values, isobolograms and IC\(_{50}\) equivalents derived from linear or non-linear data regression. Our data show that miR-34a augments the sensitivity to erlotinib in all cancer cells tested – whether they were associated with primary or secondary/acquired resistance. A plausible explanation is provided by the fact that tumor suppressor miRNAs inhibit numerous cancer pathways. In support of this hypothesis, AXL and MET, gene products specifically linked to erlotinib resistance, are directly repressed by miR-34a [27–29]. Unexpectedly, erlotinib also enhanced the therapeutic effects of the miR-34a mimic, despite existing evidence implicating miR-34a in the control of multiple oncogenic signaling pathways, including the EGFR pathway [46]. Thus, this result demonstrates that a miRNA mimic can synergize with a single gene-directed therapy and invites the search for other combinations. Based on our data, we speculate that miR-34a may also synergize with other EGFR inhibitors, such as gefitinib, afatinib, panitumumab and cetuximab, as well as HER2 inhibitors such as lapatinib, pertuzumab and trastuzumab.

In lung cancer cells with acquired resistance (HCC827 res), adding a small amount of miR-34a was capable of reducing erlotinib IC\(_{50}\) values below 0.1 \( \mu M \). This is a remarkable result and suggests that miR-34a can render this cell line equally erlotinib-sensitive compared to parental HCC827 cells. In lung cancer cells with primary resistance, the IC\(_{50}\) dose requirement for erlotinib decreased by 4- to 46-fold and was approximately 2 \( \mu M \). This may be within the range of concentrations that have clinical utility [2]. Erlotinib is given as a daily, oral dose of up to 150 mg. Although the clinical dose level of MRX34 has yet to be established, the molar ratios between miR-34a and erlotinib used in the clinic are likely within the range of ratios that have shown synergy in our cell studies. Further studies are warranted to confirm the superior activity of the combination in vivo which will consider administration routes, absorption and pharmacokinetic properties specific to each drug.

Erlotinib is currently used as a first-line therapy for NSCLC patients with activating EGFR mutations. It is also used as a maintenance therapy after chemotherapy and second- and third-line therapy for locally advanced or metastatic NSCLC that has failed at least one prior chemotherapy regimen. Clinical trials failed to demonstrate a survival benefit of erlotinib in combination with cisplatin/gemcitabine or carboplatin/paclitaxel compared to conventional chemotherapy alone [2,47]. A recent Phase III trial, investigating erlotinib plus sorafenib in HCC, also did not meet its endpoint [43]. Thus, other approaches for combination therapies are desired. Our data show that the erlotinib plus miR-34a combination is particularly effective and may substantially...
broaden the NSCLC patient population that can be treated with erlotinib. The combination was similarly synergistic in HCC cells, suggesting that the synergistic interaction is presumably a result of their molecular mechanisms of action and can also be applied to cancers outside of NSCLC. Since MRX34, a liposomal nanoparticle loaded with synthetic miR-34a mimics, has recently a phase 1 clinical trial [21], clinical testing of the erlotinib-miR-34a combination could be quickly initiated.

Supporting Information

Figure S1 Endogenous miR-34 and mRNA levels of genes controlling erlotinib resistance in NSCLC cells. Total RNA was used in triplicate qRT-PCR to measure miR-34a/b/c and mRNA levels of genes implicated in erlotinib resistance. Data were normalized to house-keeping miRNAs and mRNAs, respectively, and expressed as percent change compared to levels in HCC827 cells. u, undetected.

Figure S2 Dose-response curves of the single agents in NSCLC cells resistant to erlotinib. Cells were treated in triplicates with erlotinib or miR-34a alone at indicated concentrations. Cellular proliferation was measured 3 days or 4 days after erlotinib treatment or miR-34a reverse-transfection, respectively. Non-linear regression trendlines were generated using Graphpad, and IC50 and IC25 values were calculated. Goodness of fit of non-linear regression trendlines is indicated by R² values. The asterisk denotes theoretical IC50 values derived from an extrapolation of the dose-response curve (H226).

Figure S3 Summary table showing potency, CI and DRI values of erlotinib and miR-34a combined at various
concentrations and ratios in NSCLC cells. Combinations that yield \( Fa > 65\% \), \( CI < 0.6 \), \( DRI > 2 \) are highlighted in grey and are considered relevant. \( Fa \), fraction affected (% inhibition of cellular proliferation); \( CI \), combination index; 
\( DRI \), dose reduction index.

(TIF)

Figure S4 Endogenous expression of miR-34 and mRNAs of genes controlling erlotinib resistance in HCC cells. Total RNA was used in triplicate qRT-PCR to measure miR-34a/b/e and mRNA levels of genes implicated in erlotinib resistance. Data were normalized to house-keeping miRNAs and mRNAs, respectively, and expressed as percent change compared to levels in HCC327 cells. u, undetected.

Figure S5 Dose-response curves of the single agents in HCC cells resistant to erlotinib. Cells were treated in triplicates with erlotinib or miR-34a alone at indicated concentrations. Cellular proliferation was measured 3 days or 6 days after erlotinib treatment or miR-34a reverse-transfection, respectively.

Author Contributions
Conceived and designed the experiments: JZ KK AGB. Performed the experiments: JZ KK. Analyzed the data: JZ KK AGB. Wrote the paper: JZ KK AGB.

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