Introduction

The retinoblastoma protein (commonly called Rb) is widely recognized as one of the most important tumor suppressors in humans. Along with the similar “pocket” proteins p107 and p130, it is responsible for regulating cell cycle progression [1]. The Rb family regulates cell cycle through binding and inhibiting the transcriptional activity of early 2 factors (E2Fs) and its tumor suppressor activity is tightly related to this function [2]. When phosphorylated (typically by CDKs 2, 4, and 6 in G1), Rb becomes inactivated, thus freeing E2Fs and allowing cell cycle progression [3]. In order to avoid aberrant cell cycle entry, CDK inhibitors such as CDKN2A (commonly known as p16) prevent CDKs from phosphorylating Rb and force cells to remain in G1 [4].

Depending upon context, E2F family members can serve as transcriptional activators that drive cell cycle progression or transcriptional repressors which restrain cell cycle progression [5]. As activators, E2Fs are recognized as being important in proliferation through their transcriptional activation of S phase genes. E2Fs activate transcription via association with histone acetyltransferase (HAT) activity [6,7]. As repressors, E2Fs inhibit transcription of genes utilized in S phase entry by binding to their promoters and repressing transcription through their ability to bind to Rb and other pocket proteins, which in turn recruit chromatin modifiers such as histone deacetylases (HDACs) [6–8]. Of all the E2F family members, E2F3 is the only one individually required for cellular proliferation to occur [9–13]. E2F3 is important for transcription of various genes for S phase entry and has been shown to have roles in transcribing genes needed for G2/M phases (such as Aurora kinase A [14], CDC2 [15], and cyclin B1 [15,16]). There are two E2F isoforms, E2F3a and E2F3b, each resulting from transcription at two different promoters. E2F3b levels remain constant throughout the cell cycle, whereas E2F3a levels fluctuate and reach peak expression levels around the G1/S transition [17–19]. Mouse knockout studies reveal that E2F3a and E2F3b are generally compensatory for one another [20,21], but deletion of both isoforms is lethal [9,20]. Finally, E2F3 is more highly expressed in multiple cancers (see [5] for a review), including lung [22] and its activity has been correlated with increased sensitivity to taxane treatment in ovarian cancers [23] and ER-negative breast cancer [24].

The CDK/Rb/E2F pathway is disrupted in virtually every instance of human lung cancer, the leading cause of cancer-related death worldwide [25]. In small cell lung cancer (SCLC), which accounts for approximately 15% of lung cancers, the most common mechanism of Rb pathway disruption is mutation of the Rb protein itself. In fact, approximately 90% of small cell lung cancers lack a functional Rb protein [26,27]. In contrast, in non-small cell lung cancer (NSCLC), Rb mutation accounts for 15–30% of these cancers [26,28], and deregulation of the CDK/Rb/E2F pathway more commonly occurs via silencing of the CDK inhibitor p16 [29–32]. In most cases of NSCLC where the RB1 gene is intact, inhibitors of CDK4 and 6 would represent a
potential way to target this pathway. This hypothesis has been examined in several clinical trials and preliminary results in breast cancer are very promising [33–35], suggesting that CDK/Rb/E2F pathway inhibitors may have an important role to play in the treatment of various cancers.

The benefit of CDK inhibitors may be limited to tumors in which the Rb protein remains WT and functional, and thus, reagents that could target this pathway downstream of Rb might be useful in cancers where Rb is commonly mutated (such as lung cancer). To test this hypothesis, we have examined the activity of HLM006474 against a panel of lung cancer cell lines. HLM006474 (also discussed here as 6474) is a small molecule pan-inhibitor of E2F-DNA binding [36]. Although the IC50 of HLM006474 is relatively high (30 μM), it has found use as a tool compound [37–40]. In vivo studies indicate that the effects of 6474 treatment on different cell lines included a reduction in cell proliferation and an increase in apoptosis and reduced invasion in a three-dimensional tissue culture model system [36]. HLM006474 may be useful in cancer prevention by leading to an increase in apoptosis and decrease of proliferation in tumorigenic human embryonic stem cells [39] as well as leading to a decrease in tumor formation in mouse embryos prone to retinoblastoma [40]. Together, these results suggest that interference with E2F activity using small molecules may have clinical application in cancer therapy. In the current work, we provide a more thorough characterization of 6474 in the context of lung cancer. HLM006474 reduces the viability of a broad array of cell lines with a biological IC50 that varies between 15 and 75 μM. Combination of HLM006474 with common chemotherapeutic agents cisplatin and gemcitabine demonstrates little synergy; however, HLM006474 synergizes with paclitaxel. Taken together, these results suggest that HLM006474 may have efficacy against cancers in which E2F is deregulated and may be useful in combination with other drugs that target cell cycle components.

Materials and Methods

Cell lines and chemical compounds

Lung cancer cell lines were obtained from ATCC or originators and were provided by the Moffitt SPORE in Lung Cancer Cell Core facility. All lines are authenticated by genotyping and maintained free of *Mycoplasma*. SCLC cell lines were grown in RPMI 1640 with 10% FBS and 1% pen/strep. NSCLC cell lines were cultured in RPMI 1640 with 10% FBS without antibiotics. HLM006474 was synthesized and validated by the Moffitt Chemistry Core as previously described [36]. Gemcitabine (Gemzar, Eli Lilly) was purchased through the Moffitt Pharmacy and dissolved in water. Cisplatin and paclitaxel were purchased from Sigma and concentrated stock solutions were prepared in dimethyl sulfoxide.

siRNA knockdown

Cells plated at ~50% confluency in 6-well plates were transfected using Lipofectamine 2000 (Life Technologies) according to vendor instructions with 200 pmol of either siGENOME Non-Targeting siRNA #2 (Dharmacon) or siRNA targeting E2F3a or E2F3b as described in a paper by Hurst et al [41]. Cells were harvested four hours after transfection for cell viability assay (as described below). The remaining cells were re-plated and harvested for Western blot analysis 24 hours after transfection.

Viability assays

The antiproliferative activity of compounds and their combinations was evaluated using a high-throughput CellTiter-Blue cell viability assay. The IC50 was calculated using Clonetics Procell1 software. The average IC50 of the NSCLC and SCLC groups are calculated in bold along with STDEVs. The differences are not statistically significant.

Table 1. HLM006474 IC50 values across various cell types.

| Cell Line | Tumor Type | IC50 | STDEV |
|-----------|------------|------|-------|
| A549      | NSCLC      | 31.80| 12.90 |
| NCI-H1299 | NSCLC      | 27.30| 16.50 |
| NCI-H1650 | NSCLC      | 34.00| 3.60  |
| NCI-H1975 | NSCLC      | 44.30| 12.10 |
| NCI-H292  | NSCLC      | 28.90| 3.10  |
| NCI-H358  | NSCLC      | 19.10| 4.60  |
| NCI-H641  | NSCLC      | 15.50| 3.40  |
| NCI-H661  | NSCLC      | 23.00| 3.20  |
| DMS-79    | SCLC       | 22.30| 3.10  |
| SCLC-16HC | SCLC       | 24.90| 4.00  |
| SCLC-16HV | SCLC       | 51.40| 10.90 |
| SCLC-86M  | SCLC       | 15.70| 2.40  |
| DMS114    | SCLC       | 23.80| 1.50  |
| NCI-H209  | SCLC       | 21.90| 7.19  |
| NCI-H69   | SCLC       | 53.70| 5.44  |
| NCI-H82   | SCLC       | 21.30| 3.02  |
| NCI-N417  | SCLC       | 75.10| 6.96  |

Average IC50:
- NSCLC: 27.99 ± 7.43
- SCLC: 34.46 ± 4.95
- Overall: 31.41 ± 6.11

The IC50 of various cell lines was determined as described in the methods. STDEV refers to the standard deviation of indicated values calculated using Excel STDEV function. The average IC50 of the NSCLC and SCLC groups are calculated in bold along with STDEVs. The differences are not statistically significant.

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An E2F Inhibitor Synergizes with Paclitaxel

A. 

|     | 0 | 3 | 6 | 9 | 12 | 24 |
|-----|---|---|---|---|----|----|
| H1299 |   |   |   |   |    |    |
| H292  |   |   |   |   |    |    |

B. 

|     | Total-E2F3 |
|-----|------------|
| H1299 |   |   |   |   |    |    |
| H292  |   |   |   |   |    |    |

C. 

|     | Total-E2F1 |
|-----|------------|
| H1299 |   |   |   |   |    |    |
| H292  |   |   |   |   |    |    |

D. 

|     | Total-E2F4 |
|-----|------------|
| H1299 |   |   |   |   |    |    |
| H292  |   |   |   |   |    |    |

E. 

|     | Total-MCM10 |
|-----|-------------|
| H1299 |   |   |   |   |    |    |
| H292  |   |   |   |   |    |    |

F. 

|     | Total-MCM2 |
|-----|------------|
| H1299 |   |   |   |   |    |    |
| H292  |   |   |   |   |    |    |

G. 

|     | Total-CCNE2 |
|-----|-------------|
| H1299 |   |   |   |   |    |    |
| H292  |   |   |   |   |    |    |
viability assay (Promega), as previously described [42]. For the assays, 1000 cells in 24 μL were plated in 384-well plates and incubated overnight at 37°C, 5% CO2. The next day, the drugs were diluted in media and 6 μL of these dilutions added to appropriate wells using an automated pipetting station. Four replicate wells were used for each drug concentration. The cells were incubated with the drug for 120 hours, at which time, 5 μL CellTiter-Blue reagent (Promega Corp., Madison, WI) was added. Fluorescence was read with a Synergy HT microplate reader (BioTek Instruments, Inc., Winooski, VT). IC50s were determined using a sigmoidal equilibrium model regression using XLfit version 4.3.2 (ID Business Solutions Ltd.) and is defined as the concentration of drug required for a 50% reduction in growth/viability. Combination indices CI were performed in triplicate and repeated at least three times.

Combined indices

The IC50 values obtained from single drug cell viability assays were used to design combination experiments. For 6474 combinations with cisplatin, gemcitabine and paclitaxel these ratios were 1:1, 500:1 and 4000:1, respectively. For drug combination experiments, the cell viability assays were performed and the results analyzed for synergistic, additive, or antagonistic effects using the combination index (CI) method developed by Chou and Talalay [43]. Combination indices CI<1, CI = 1, and CI>1 indicate synergistic, additive effects, and antagonism, respectively.

Antibodies and western blotting

Western blots were performed as previously described [36,44,45]. Briefly, whole cell lysates were subjected to SDS-PAGE and transferred to PVDF membrane. Detection of proteins was accomplished using horseradish-peroxidase-conjugated secondary antibodies and enhanced chemiluminescence (ECL) purchased from Amersham or Thermo Scientific. Various antibodies used included E2F1 (C-20, sc-193, Santa Cruz), E2F3 (C-18, sc-678, Santa Cruz), PARP (#9542L, Cell Signaling Technology), and monoclonal β-actin (clone AC-15, cat no: A5441, Sigma). Adobe Photoshop CS was used to quantify Western blot band intensity readings directly from exposed films using the rectangular marquee tool/histogram and the inverted scanned film image. This same square was used for all further band readings in order to ensure that the same area was analyzed for each band. The readings were then adjusted to account for actin and background and arbitrarily normalized to the cell line H239 (assigned a value of 1).

Real-time polymerase chain reaction

RNA was harvested from cells using the RNeasy mini kit from Qiagen. Reverse transcription polymerase chain reaction (PCR) was then performed using the iScript cDNA synthesis kit (Bio-Rad). This cDNA was then utilized in real-time PCR using either iQ SYBR Green Supermix (Bio-Rad) or PerfeCTa SYBR Green SuperMix (Quanta Biosciences, VWR). E2F1, E2F3, E2F4, tubulin, MCM2, MCM10, CCNE2, and GAPDH primers were ordered from Integrated DNA Technologies. Primer sequences are as follows: E2F1 F (5'-GCTGGAACCATCAGTGAATATC-3'), E2F1 R (5'-TCTGCAATGCTAGGAAGGTCTTG-3'), E2F3a/b F (5'-CAGGTGAGTGGTCAGAGGT-3'), E2F3a/b R (5'-GGGCTGGGTTGGAATAGGGAAAA-3'), tubulin R (5'-TGGCACTGGCTCTGGGTTCG-3'), MCM2 F (5'-CTGTGTGTTGAGGAGCACG-3'), MCM2 R (5'-CTTTGTTCCGTGACCTTGTG-3'), MCM10 F (5'-CTGTCAGTGGACAGATTG-3'), MCM10 R (5'-TCCGGTGCCATTTTAGAG-3'), CCNE2 F (5'-CAGGTTTGGAGGAGGAGG-3'), CCNE2 R (5'-ACTCTCCTCCAGTACGGGAAAA-3'), GAPDH F (5'-GAGTGCAACGGATTTGGTCTG-3'), and GAPDH R (5'-TTGATTTTGAGGGATCTCG-3').

Statistical analysis

For the real-time PCR analysis for the time point experiment, the difference between expression of each experimental gene (E2F1, E2F3, E2F4, CCNE2, MCM2, and MCM10) and expression of the control gene (tubulin) was calculated for each cell line at each time point. The difference between each of the non-0 hour time points and the 0 hour time point readings for each gene in each cell line was calculated using T-Tests with Welch’s correction. The paclitaxel IC50s were log-transformed to improve normality. The correlation of E2F3 mRNA and protein expression with log paclitaxel IC50s was calculated using Pearson correlation coefficient. Wilcoxon rank-sum tests were used to explore the difference of cell viability in control siRNA treatment with either E2F3a or E2F3b siRNA treatment.

Results

HLM006474 has broad antiproliferative activity

To examine the potential of 6474 in the treatment of lung cancer, we determined the 6474 IC50 in seventeen lung cancer cell lines (Table 1). This analysis included eight non-small cell lung cancer lines (NSCLC) and nine small cell lung cancer lines (SCLC). In this viability assay, cells were plated at low density on day zero and were grown in the presence of various 6474 concentrations for five days. After five days, the relative viability of cells was determined by staining with CT-Blue (Promega). The results reveal that the 6474 IC50 ranges from ~15 to ~75 μM across the seventeen cell lines. The average biological IC50 of all the cell lines was 31.4 (±6.1) μM, which is essentially identical to the biochemical IC50 of 29.8 (±7.6) μM, previously reported [36]. There was no statistically significant difference between SCLC and NSCLC.
Figure 2. HLM006474 synergizes with paclitaxel, but not with cisplatin or gemcitabine. A–C. H1299 cells were subjected to viability assays in the presence of 6474 (HLM) combined with cisplatin (A, CisPt), gemcitabine (B, Gem) and paclitaxel (C, Pac), as indicated (see methods). Results reveal synergy with paclitaxel (CI average 0.98) and antagonism with cisplatin and gemcitabine (CI average 1.40 and 1.39, respectively). D. Western blotting reveals that in H1299 cells treated for 72 hours with 20 μM 6474 alone, 5 nM paclitaxel alone, or the combination of the two, 6474 and paclitaxel synergize in the induction of PARP cleavage. E–G. H292 cells were tested in similar viability assays to determine the efficacy of 6474 (HLM) combined with cisplatin (E, CisPt), gemcitabine (F, Gem) and paclitaxel (G, Pac). As seen in H1299 cells, 6474 synergizes with paclitaxel (CI average 0.96), but is antagonistic with cisplatin (CI average 1.51) and gemcitabine (CI average 1.46). doi:10.1371/journal.pone.0096357.g002
Figure 3. Sensitivity to paclitaxel correlates with E2F3 levels in NSCLC. A. cDNA from ten NSCLC cell lines were utilized in real-time PCR to detect endogenous E2F3 expression levels (as compared to GAPDH as a control). The expression levels were then compared to the paclitaxel logIC₅₀ of each line and graphed as shown. B. Lysates from ten NSCLC cell lines were prepared and ran in a Western blot to detect endogenous E2F3 levels. C. Densitometrically analyzed values of the protein expression levels (as compared to β-actin as a control) were then graphed against the paclitaxel logIC₅₀ for each line as shown.

Figure 4. Depletion of E2F3 alters sensitivity to paclitaxel. A. H1299 cells were transiently transfected with 200 pmol control, E2F3a, or E2F3b siRNA and harvested after 24 hours. Western blots of these lysates demonstrate the extent of E2F knockdown. B. MTS assays were used to determine the sensitivity of each cell line to 50 nM paclitaxel. Cells with diminished E2F3 levels were more viable in the presence of paclitaxel than control cells. Notes: * represents p<0.05, ** represents p<0.01.

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Short treatments with HLM006474 lead to increased expression of several known E2F-regulated genes

As a component of our analysis of HLM006474, we examined the expression of E2F family members following treatment by Western blotting. NSCLC cell lines H292 and H1299 were treated with 60 µM HLM006474 for 0, 3, 6, 9, 12, and 24 hours and analyzed via Western blot (Figure 1 A). Surprisingly, protein levels of both E2F3a and b isoforms increased in early time points (typically around 6–9 hours). Levels of the E2F1 protein increased more modestly following treatment, peaking between 6 and 12 hours and returning to baseline levels after 24 hours. In real-time PCR analysis with tubulin as a control, E2F3 mRNA levels increased significantly after 3 hours of treatment and then decreased in each subsequent time point (Figure 1 B), while E2F1 mRNA expression levels were significantly increased after short treatment times in H292 alone (Figure 1 C) and E2F4 levels remained constant or decreased at each time point (Figure 1 D). It was also noted that some genes commonly regulated by E2Fs; MCM10 (Figure 1 E), MCM2 (Figure 1 F), and CCNE2 (Figure 1 G), were more highly expressed in early time points in a manner comparable to the changes seen in E2F3 mRNA expression. The results shown in Figure 1 suggest that treatment with an inhibitor of E2F-DNA binding can result in the activation of E2F-regulated targets, including auto-regulated E2F family members. The E2F family is known to actively repress transcription [46–49], and thus, we propose that treatment with HLM006474 may displace E2F-repressive complexes and thereby activate transcription of E2F-regulate genes that are predominantly repressed by E2F complexes. To explore this possibility, we used siRNA specifically against E2F1, E2F3a, E2F3b, E2F4 and Rb to deplete two NSCLC cell lines of various E2F complexes. Microarray was performed to examine the effect of these siRNAs on the expression of an E2F signature previously defined based on E2F3 overexpression [50]. The results, which will be published elsewhere, demonstrate that depletion of individual E2Fs results in many E2F signature genes being activated, as would be expected from a de-repression model.

HLM006474 synergizes with paclitaxel

Having established that 6474 has anti-proliferative effects on lung cancer cell lines, but may influence E2F-regulated genes in a complex manner, we sought to determine if 6474 would synergize with chemotherapeutic drugs commonly used in lung cancer treatment. H1299 cells were treated with 6474 alone and in combination with cisplatin, gemcitabine and paclitaxel. Combinations were chosen based on the IC50 of the cells to the individual compounds and combination indices were calculated [43]. Figure 2 reveals that there is antagonism between 6474 and cisplatin (panel 2A, CI average 1.40) and gemcitabine (panel 2B, CI average 1.39). In contrast, 6474 weakly synergizes with paclitaxel (panel 2C, CI average 0.98). To further explore the nature of the synergy between 6474 and paclitaxel, H1299 cells were treated with modest doses of each drug alone or in combination and utilized in Western blot analysis. The appearance of cleaved PARP in cells treated with the drug combination confirms that the combination of 6474 and paclitaxel induces more apoptosis than either drug alone (Figure 2D). To explore whether this synergy would be observed in additional cell lines, we also examined the NSCLC cell line H292. As in the case of H1299 cells, 6474 synergized with paclitaxel (panel 2G, CI average 0.96), but showed no synergy with cisplatin (panel 2E, CI average 1.51) or gemcitabine (panel 2F, CI average 1.46).

E2F3 levels impact sensitivity to paclitaxel

The observations that an E2F inhibitor could synergize with paclitaxel and the previously discussed increase in E2F3 levels following early time point treatments with HLM006474 suggested that E2F3 activity might play a role in paclitaxel sensitivity. Real-time PCR was used to analyze the endogenous expression of E2F3 (with GAPDH serving as an internal control), and then these values were correlated to the paclitaxel logIC50 of each cell line (Figure 3A). Lysates from ten NSCLC cell lines were run in Western blots (Figure 3B) and densitometrically analyzed using β-actin as a control. These E2F levels were then plotted against the corresponding paclitaxel logIC50 values (Figure 3C). In both the real-time PCR and Western blot analyses, a strong inverse correlation between E2F3 and paclitaxel IC50 was noted. To more formally test this hypothesis, we used siRNA to deplete H1299 cells of E2F3a and E2F3b and then determined their sensitivity to paclitaxel as measured in MTS assays. Western blotting reveals an almost complete knockdown of the targeted E2Fs in the H1299 cells (Fig 4A). Control siRNAs did not affect E2F expression. As expected, Fig 4B reveals that the cells with decreased E2F3 levels were less sensitive to paclitaxel.

Conclusions

The CDK/Rb/E2F pathway represents a good target for the treatment of various solid tumors. Although development has been slow due to toxicity of early compounds, CDK inhibitors are starting to gain traction in clinical trials [33,51,52]. We propose that targeting the CDK/Rb/E2F pathway even further downstream, at the E2F level, may also be of value. Thus, we have examined the potential of a pan-E2F inhibitor, HLM006474, in the treatment of lung cancer.

We propose the model in Figure 5 to explain our results. First, we observe that treatment with 6474 leads to a transient increase in not only E2F3 protein and mRNA expression levels, but also an increase in many other E2F-regulated transcripts. These counter-
intuitive observations are reasonable based on the long-known observation that E2Fs are active repressors of transcription [46–49]; however, they do raise concerns that pan-E2F complex inhibition may have unwanted consequences. Thus, future E2F-targeted compounds should selectively block transcription activating E2F complexes and spare transcriptionally repressing E2F complexes. Second, we believe the increased levels of E2F3 (and likely other E2F-regulated genes) increase the sensitivity of the cells to paclitaxel. We base this conclusion on the correlation observed between normal E2F3 levels and paclitaxel sensitivity, as well as the results of E2F3 siRNA experiments. This documents the first link between levels of E2F3 and paclitaxel in NSCLC, though a relationship between high levels of E2F3 activity and increased sensitivity to paclitaxel has been previously observed in ovarian [23] and ER-negative breast cancer [24]. The mechanism by which E2F3 generates sensitivity to paclitaxel is unknown. One possibility is that it relates directly to E2F3’s role in cell cycle. For example, in cells with high E2F3 levels, it would be expected that the cells would proliferate more, thus giving cells a greater opportunity to enter M phase (where paclitaxel would be most effective). However, this explanation alone would suggest that these cells should be more sensitive to gemcitabine as well due to entering S phase more often. It might be more likely that greater sensitivity to paclitaxel is due to apoptosis-regulating genes becoming highly expressed due to the increase in E2F3. Also, it has been previously noted that overexpression of E2F3 leads to an enrichment of genes that are microtubule-related [23], so this could perhaps explain the correlations we see between E2F3 levels and paclitaxel sensitivity. Likewise, as mentioned previously, E2F3 has been noted to have a role at the G2/M checkpoint through its regulation of expression of Aurora kinase A [14], CDC2 [15], and cyclin B1 [15,16], which may also point to higher levels of E2F3 leading to an increase in cells entering that phase of the cell cycle and perhaps then increasing paclitaxel sensitivity.

We have demonstrated that HLM006474 is effective in lung cancer cell lines. Furthermore, we have shown that 6474 synergizes well with paclitaxel, potentially due to 6474’s effects on E2F3 levels. Taken together, these results suggest that potent, specific activator E2F inhibition may be useful in the treatment of NSCLC in the future (especially in combination with other agents), and that E2F3 levels may be a good predictor of paclitaxel sensitivity in NSCLC.

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Author Contributions

Conceived and designed the experiments: CAK WDC. Performed the experiments: CAK. Analyzed the data: LC. Wrote the paper: CAK WDC.
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