An Evolutionarily Conserved Synthetic Lethal Interaction Network Identifies FEN1 as a Broad-Spectrum Target for Anticancer Therapeutic Development

Derek M. van Pel1,2, Irene J. Barrett1, Yoko Shimizu3, Babu V. Sajesh3, Brent J. Guppy3, Tom Pfeifer3, Kirk J. McManus4, Philip Hieter1*

1 Michael Smith Laboratories, University of British Columbia, Vancouver, Canada, 2 Department of Biochemistry and Molecular Biology, University of British Columbia, Vancouver, Canada, 3 Department of Screening, Centre for Drug Research and Development, Vancouver, Canada, 4 Manitoba Institute of Cell Biology, Department of Biochemistry and Medical Genetics, University of Manitoba, Winnipeg, Canada

Abstract

Harnessing genetic differences between cancerous and noncancerous cells offers a strategy for the development of new therapies. Extrapolating from yeast genetic interaction data, we used cultured human cells and siRNA to construct and evaluate a synthetic lethal interaction network comprised of chromosome instability (CIN) genes that are frequently mutated in colorectal cancer. A small number of genes in this network were found to have synthetic lethal interactions with a large number of cancer CIN genes; these genes are thus attractive targets for anticancer therapeutic development. The protein product of one highly connected gene, the flap endonuclease FEN1, was used as a target for small-molecule inhibitor screening using a newly developed fluorescence-based assay for enzyme activity. Thirteen initial hits identified through in vitro biochemical screening were tested in cells, and it was found that two compounds could selectively inhibit the proliferation of cultured cancer cells carrying inactivating mutations in CDC4, a gene frequently mutated in a variety of cancers. Inhibition of flap endonuclease activity was also found to recapitulate a genetic interaction between FEN1 and MRE11A, another gene frequently mutated in colorectal cancers, and to lead to increased endogenous DNA damage. These chemical-genetic interactions in mammalian cells validate evolutionarily conserved synthetic lethal interactions and demonstrate that a cross-species candidate gene approach is successful in identifying small-molecule inhibitors that prove effective in a cell-based cancer model.

Introduction

Cancerous cells carry somatic mutations that genotypically distinguish them from surrounding noncancerous cells, and this provides an opportunity that can be exploited for therapeutic development. One strategy for the specific targeting of cancer genotypes relative to nonmutated somatic cells is to exploit synthetic lethal interactions [1]. For example, breast cancer cells with mutations in BRCA1 or BRCA2 are extremely susceptible to knockdown or chemical inhibition of PARP1, which encodes poly(ADP)ribose polymerase [PARP] [2,3].

While exploiting synthetic lethality has the potential to be an effective approach to treating tumors, a major challenge is the identification of clinically relevant small-molecule inhibitors. One approach, pioneered by the National Cancer Institute, is to screen many thousands of unknown potential therapeutics on cancer cell lines [4]. Compounds generate a “fingerprint” of activity against certain cell lines, which can then be deconvoluted, usually by mutation sequencing, to yield novel gene-drug interactions, in a so-called “bottom-up” approach. Alternatively, a “top-down” approach applies compounds of known target or mode of action to known genotypes, again to identify new gene-drug interactions. Recently, two groups used such an approach to screen more than 100 compounds against hundreds of cancer cell lines whose mutational status was known [5,6], observing that gene-drug interactions tended to be more significant for targeted therapies, such as compounds targeting the BCR-ABL fusion protein, than for generally cytotoxic drugs, such as DNA damaging agents or antimicrobials [6]. Thus, screening for compounds targeting a specific genetic lesion is preferable to developing new cytotoxic agents. Such targeted compounds can then be deployed as first-line anticancer therapeutics either singly or in a combination regimen that would lessen the likelihood of drug-resistant clones developing within the tumor cell population [7,8].
Identification of Therapeutic Genetic Interactions

Author Summary

Anticancer therapeutic discovery is a major challenge in cancer research. Because cancer is a disease caused by somatic genetic mutations, the search for anticancer therapeutics is often driven by the ability to exploit genetic differences specific to tumor cells. Recently, cancer therapeutic development has sought to exploit synthetic lethality, a situation in which the combination of two independently viable mutations results in lethality. If a compound can be found to selectively kill a specific genotype via inhibition of a specific gene product, this is known as a chemical-genetic interaction, and it mimics a synthetic lethal genetic interaction. The ideal therapeutic would be broad spectrum, that is, active against multiple cancer genotypes within a tumor type and/or across a variety of cancers. We have developed an approach, taking advantage of the evolutionary conservation of synthetic lethal interactions, to identify “second-site” targets in cancer: genes whose chemical inhibition leads to selective killing of tumor cells across a broad spectrum of cancer genotypes. We identified small-molecule inhibitors of one such target, FEN1, and showed that these compounds were able to selectively kill human cells carrying cancer-relevant mutations. This approach will facilitate the development of anticancer therapeutics active against a variety of cancer genotypes.

Many different cancer mutations lead to a limited repertoire of cancer phenotypes, such as chromosome instability, checkpoint dysfunction, and hyperplasia [9]. It is possible to identify a gene target that results in synthetic lethality with a large number of unlinked gene mutations by screening for targets that result in synthetic lethality with a common tumor phenotype. For example, chromosome instability (CIN), an increase in the rate of gain or loss of whole or parts of chromosomes, is observed in the form of aneuploidy in more than 90% of solid tumors and over 75% of blood cancers [10]. As the maintenance of genomic stability is an essential cellular process, CIN represents a phenotype that could potentially be leveraged towards selective killing of cancerous cells relative to normal cells. A gene that is synthetic lethal with a large number of cancer-related CIN genes would be an attractive therapeutic target in a large fraction of tumors.

Genetically tractable model organisms, such as the budding yeast *Saccharomyces cerevisiae*, facilitate the identification of human CIN genes, via identification and sequencing of their human orthologs. For example, identification of yeast CIN genes [11] led to the sequencing of the human homologs of 200 yeast CIN genes in human colorectal cancers, and it was discovered that human homologs of the yeast CIN genes SMC1, SMC3, NIPBL, STAG3, RFC20, FBX17/CDC4, MRE11A, RAD54B and BLM have been found to be mutated in colorectal cancer, and together account for approximately 25% of the CIN mutational spectrum of this disease [13–15,20–22]. Protein BLAST was used to identify the budding yeast orthologs of these human genes (Table 1) and we constructed a synthetic lethal interaction network (Figure 1A), using literature and publicly available genetic interaction data (BioGrid and the Saccharomyces Genome Database) [18,23]. To investigate the conservation of this network between yeast and a human cell line, we used siRNA-mediated knockdown of potential synthetic lethal gene pairs in the cell line HCT116. Knockdown efficiencies were evaluated by Western blots (Figure S1A). All pair-wise combinations between the three “central” synthetic lethal partner genes, WDHD1, FEN1, and CHTF8, and the ten outer cancer-mutated CIN genes were evaluated for synthetic lethality (Figure 1B, 1C, 1D). (CHTF8 was selected as a representative of the alternative RFC6CHTF8, comprised of Dcc1, Ctf8, and Ctf18 in *S. cerevisiae*).

Of the 30 possible synthetic lethal interactions among the genes tested, 22 have been reported in yeast [18,23]. We found 16 of the predicted interactions (73%) were conserved between yeast and human cells, and 6 predicted interactions did not appear conserved in our assay (27%). Furthermore, one interaction, between FEN1 and STAG1, was not predicted based on yeast data; however, we detected a genetic interaction between these genes (Figure 1F and Table S1). No interactions were observed with STAG3, which functions primarily in human meiosis [24]. As in yeast, all three central genes – WDHD1, FEN1, and CHTF8 – were highly connected to sister chromatid cohesion genes (e.g. cohesin and/or cohesin loaders) (Figure 1F).

As FEN1 encodes an enzyme, whereas WDHD1 and CHTF8 do not; it may be amenable to biochemical inhibitor screening. Thus, we sought to further validate genetic interactions between FEN1 and other genes in the network. To ensure that these observed interactions were not cell line-dependent, we attempted to recapitulate interactions between FEN1 and each of CDC4, RAD54B, and RFC20 in the karyotypically stable, immortalized fibroblast cell line hTERT. As in HCT116 cells, genetic interactions were observed following knockdown of all three gene pairs (Figure 1E, Table S2, Figure S1C). We found that individual siRNAs could recapitulate the genetic interactions observed with the pooled siRNAs (Table S3). These data validate a subset of genetic interactions identified in the HCT116 cells and thus confirm FEN1 as a strong candidate therapeutic target.

Results

A cross-species approach reveals conserved genetic interaction partners of cancer genes

The human genes *SMC1, SMC3, NIPBL, STAG3, RFC20, FBX17/CDC4, MRE11A, RAD54B* and *BLM* have been found to be mutated in colorectal cancer, and together account for approximately 25% of the *S. cerevisiae* mutational spectrum of this disease [13–15,20–22]. Protein BLAST was used to identify the budding yeast orthologs of these human genes (Table 1) and we constructed a synthetic lethal interaction network (Figure 1A), using literature and publicly available genetic interaction data (BioGrid and the Saccharomyces Genome Database) [18,23].

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Table 1. Yeast and human gene orthologs.\textsuperscript{A}

| Yeast gene | Human ortholog(s) | % identity | % similarity | BLASTP expectation |
|------------|-------------------|------------|--------------|--------------------|
| BBE1       | RNF20             | 21%        | 42%          | 4 × 10\textsuperscript{-26} |
| CDC4       | CDC4 (FBXW7)      | 29%        | 51%          | 4 × 10\textsuperscript{-61} |
| CTF18      | CHTF18            | 24%        | 42%          | 8 × 10\textsuperscript{-36} |
| CTF4       | WDHD1 (AND1)      | 21%        | 36%          | 9 × 10\textsuperscript{-18} |
| CTF8       | CHTF8             | 21%        | 37%          | 6 × 10\textsuperscript{-18} |
| DCC1       | DCC1              | 24%        | 39%          | 10\textsuperscript{-10}   |
| MRE11      | MRE11A            | 41%        | 59%          | 5 × 10\textsuperscript{-12} |
| RAD27      | FEN1              | 60%        | 76%          | 2 × 10\textsuperscript{-137} |
| RDHS4      | RAD54B            | 37%        | 52%          | 3 × 10\textsuperscript{-114} |
| SCC1 (MCD1)| RAD21             | 35%        | 54%          | 2 × 10\textsuperscript{-7} |
| SCC2       | NIPBL             | 20%        | 39%          | 10\textsuperscript{-18}   |
| SCC3 (IRR1)| STAG1, STAG2, STAG3| 26%      | 46%          | 10\textsuperscript{-20}   |
| SG1        | BLM, WRN          | 39%        | 60%          | 4 × 10\textsuperscript{-12} |
| SMC1       | SMC1A             | 30%        | 54%          | 10\textsuperscript{-163}  |
| SMC3       | SMC3              | 32%        | 55%          | 0 \textsuperscript{B}      |

\textsuperscript{A}Names indicated are the names used in this work. Names in parentheses indicate common alternative gene names. Members of the cohesin complex (and SMC2/NIPBL, a cohesin loader) are indicated in boldface type.

\textsuperscript{B}Identified with DELTA BLAST algorithm.

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Developing a high-throughput in vitro assay for FEN1 activity

FEN1 (Flap ENdouclease 1) encodes an enzyme previously shown to be amenable to biochemical assay development in vitro [25] that has been implicated in almost all DNA transactions, including DNA repair and replication [26]. Adapting a previous radiolabel-based in vitro assay, we developed an in vitro assay for FEN1 activity based on fluorescence quenching [25]. In this assay, three oligonucleotides are annealed to generate the synthetic substrate, positioning a fluorophore and fluorescent quencher in close proximity. The flap endonuclease activity of FEN1 cleaves the 5'- flap to which the fluorophore is attached, allowing it to diffuse away from the quencher and fluoresce (Figure 2). Using a potent, previously described in vitro FEN1 inhibitor, compound 16 from Tumey, et al. [27], we observed significant inhibition of flap endonuclease activity (Figure 3, upper left panel). A screen of 30,000 compounds, from libraries containing known and FDA-approved drugs, and the Canadian Chemical Biology Network library, yielded approximately 90 hits, following a counterscreen using a quencherless substrate to eliminate false positives caused by fluorescent compounds and fluorescent quenchers. Ultimately, 13 compounds were selected for further investigation based on structural diversity and having drug-like properties (as described by Lipinski’s “Rule of Five”; [28]). These compounds were found to have mid-nanomolar to low micromolar IC\textsubscript{50}s in vitro (Figure 3, remaining panels).

Flap endonuclease inhibitors recapitulate genetic interactions in cells

We next sought to determine whether the flap endonuclease inhibitors we identified could recapitulate any of the genetic interactions found previously (Figure 1F). We first targeted the interaction between FEN1 and CDC4, owing to the fact that CDC4 has been shown to be a CIN gene mutated in many tumor types [11,29–32]. We took advantage of a matched pair of cell lines in which both copies of CDC4 had been inactivated in HCT116 cells [13]. siRNA-mediated knockdown of FEN1 in this cell pair resulted in selective proliferation inhibition (Figure 4A). We applied the small-molecule hits from the screen to this matched pair of cell lines and found six compounds that selectively inhibited the proliferation of CDC4-knockout HCT116 cells relative to wild type cells (Figure 4B and Figure S2B). To ensure that these results were not cell line-specific, we utilized another matched pair of cell lines with inactivated CDC4, this time in DLD-1 cells. The six compounds showing selective proliferation inhibition of CDC4-knockout HCT116 cells were applied to CDC4-knockout and wild type DLD-1 cells [13], and RF00974 and NSC645851 were found to selectively inhibit the proliferation of CDC4-knockout DLD-1 cells relative to wild type (Figure 4C).

To further test the idea that CDC4 activity is responsible for the observed effect, cells in which CDC4 had been inactivated in a heterozygous state were also treated with RF00974 and NSC645851. As with homozygous CDC4\textsuperscript{-/-} cells, heterozygous CDC4\textsuperscript{+/--} cells displayed a statistically significant decrease in proliferation relative to wild type CDC4\textsuperscript{+/+} cells, albeit lesser in magnitude (Figure 5).

We next attempted to recapitulate the interaction between FEN1 and MRE11A, as MRE11A has been shown to be mutated at a frequency of 4% in colorectal cancers [15]. We treated cells in which MRE11A had been depleted via siRNA with the more potent of the two flap endonuclease inhibitors described above, RF00974, and found that MRE11A depletion sensitized cells to flap endonuclease inhibitor treatment (Figure 6A). We also found that treatment with a previously-described small-molecule inhibitor of MRE11A, mirin [33], was able to sensitize cells to treatment with RF00974 (Figure 6B). Taken together, these data suggest that inhibition of flap endonuclease activity is sufficient to recapitulate evolutionarily conserved, colorectal cancer-relevant synthetic lethal genetic interactions.

Finally, we wished to characterize the mechanism by which inhibition of flap endonuclease activity may lead to cell death. Given the role of FEN1 in DNA replication and repair, we asked whether endogenous DNA damage increases as a result of FEN1
inhibition. We used HCT116 cells in which 53BP1 had been stably tagged with mCherry to ask whether 53BP1 focus formation, indicative of DNA repair centers [34,35], increased. We found a statistically significant (p < 0.05) increase in the frequency of cells with many 53BP1 foci following siRNA-mediated knockdown of FEN1. Furthermore, we observed a similar increase (p < 0.05) following treatment with the flap endonuclease inhibitor RF00974 (Figure 6C). We next measured the level of H2AX phosphorylation (γ-H2AX), an independent indicator of DNA damage [36], in HCT116 CDC4+/− and CDC4−/− cells in response to RF00974. We found that, similar to increasing 33BP1 focus formation, RF00974 treatment increased H2AX phosphorylation (Figure S3). H2AX phosphorylation was increased even in untreated HCT116 CDC4−/− cells, so no increase in phosphorylation was observed. In order to determine whether RF00974 leads to an increase in apoptosis in CDC4-deficient cells, we asked whether PARP cleavage, a marker of apoptosis [37], is increased following RF00974 treatment. We found that RF00974 treatment did not increase PARP cleavage in either wild type or CDC4-deficient cells. Taken together, these results suggest that loss of FEN1, or inhibition of flap endonuclease activity, lead to an increase in endogenous DNA damage that inhibits the proliferation of CDC4-deficient cells by non-apoptotic means.

Discussion

In this study, we used a cross-species candidate approach to identify new anticancer therapeutic targets for small-molecule inhibition having a potentially broad spectrum of applicability. We
found that a yeast CIN synthetic lethal interaction network is largely conserved between \emph{S. cerevisiae} and a human tumor cell line. Based on this network, we screened for \emph{in vitro} inhibitors of the highly connected enzyme FEN1. Flap endonuclease inhibitors discovered in this screen recapitulated synthetic lethal interactions between \textit{FEN1} and each of \textit{CDC4} and \textit{MRE11A}, demonstrating that evolutionarily conserved genetic interactions in a core cellular process, such as the maintenance of genomic stability, can be exploited as a means to inhibit the proliferation of tumor cells carrying specific and cancer-relevant mutations.

The idea of using the unique genetic profile of tumor cells relative to somatic cells to selectively kill cancer has been applied by various groups, such as in the case of the chemical-genetic interaction between \textit{BRCA1/2} and PARP inhibitors [2,3]. Several studies have focused on DNA damage, usually by identifying inhibitors of DNA damage response proteins that either directly kill tumor cells, or that potentiate the effects of DNA damaging agents [38–42]. Recently, two large-scale studies examining chemical-genetic interactions between new or established anti-cancer treatments and cancer cell lines of known genotype demonstrated the promise of such top-down approaches by identifying previously unknown sensitivities of many cancer genotypes, such as between Ewing’s sarcomas and PARP inhibitors [5,6]. An alternative means to construct genetic interaction networks for the discovery of therapeutic targets is to take a cross-species candidate approach in a genetically tractable model organism. In \textit{S. cerevisiae}, defined genetic changes can be introduced and subsequently screened in a high-throughput manner [16,43] (though mammalian genome editing technologies are advancing rapidly [44,45]). The nearly 75% [16,22] conservation of synthetic lethal interactions we found between yeast and human cells is similar to the degree of conservation of genetic interactions between \textit{S. cerevisiae} and the model metazoan \textit{Caenorhabditis elegans} in a related network, identified by our group and others [18,19,46], and expands upon previous proof-of-principle work by our group [47]. Although we ultimately targeted the highly conserved flap endonuclease \textit{FEN1} in the current study, yeast genetic data has the potential to implicate biological processes, as opposed to specific proteins, as therapeutic targets; in this way, targets can be identified that are not conserved in \textit{S. cerevisiae}. For example, we recently demonstrated that mutation of cohesin genes in yeast was synthetic lethal with mutation of proteins playing a role in replication fork stability. siRNA-mediated knockdown of cohesin genes was found to sensitize human cells to inhibition of PARP, a protein involved in replication fork progression, but without a known ortholog in yeast [19]. Thus, the versatility of yeast synthetic lethal networks to predict therapeutic targets makes our approach complementary to large-scale screening for gene-drug interactions [4–6].

Therapeutics that target a specific genotype, such as EGFR family inhibitors in the case of \textit{ERBB2} (also known as \textit{HER2}) amplification, produce more significant gene-drug interactions than more general cytotoxic agents [6]; however, the indications for such agents are limited to a handful of genotypes. \textit{FEN1} plays a critical role in nearly all DNA transactions, including DNA replication via Okazaki fragment maturation [48,49], long-patch base excision repair [50,51], the prevention of trinucleotide repeat expansions [25,52], and restart of stalled replication forks [53]. Yeast RAD27 is one of the most highly genetically connected genes in the yeast genome (Tables S5 and S6); many of these interactors are CIN genes [11], and many of the corresponding human orthologs may prove to be mutated and cause CIN in tumours. Given that the majority of the genetic interactions were conserved in the CIN synthetic lethal interaction network interrogated here, \textit{FEN1} may be a widely applicable target in cancers harboring mutations in a variety of CIN genes. More generally, DNA repair and replication protein inhibitors are being actively developed as anticancer therapeutics [2,3,41,54] and the process of DNA replication forms a genetic hub in \textit{S. cerevisiae} [16,23,43,55]. The critical role of \textit{FEN1} in DNA transactions is analogous to that of PARP, a protein playing a role in DNA repair and the protection of stalled DNA replication forks [56,57]. PARP is synthetic lethal with mutations in \textit{BRCA1/2} [2,3], and its therapeutic range has been extended more recently to include cells with mutations in \textit{PTEN} [38] and cohesins [19]. Thus, like

Figure 2. Screening for FEN1 inhibitors \emph{in vitro}. Schematic representation of the fluorescence-based assay employed to identify FEN1 inhibitors. In the absence of inhibitor, FEN1 cleaves the 5’ flap to which the 6-FAM fluorophore is attached, allowing it to diffuse away from the BHQ-1 quencher and fluoresce. Activity is read as increasing fluorescence over time. doi:10.1371/journal.pgen.1003254.g002
PARP, FEN1 potentially represents a potent, broadly-applicable target for anticancer therapeutic development.

In turn, the ideal anticancer therapeutic would have a broad spectrum, suggesting it would be more advantageous to target a phenotype common in cancer. CIN in the form of aneuploidy is seen in >90% of solid tumors [10] and represents a sub-lethal mutation in an otherwise essential process. Of relevance to the current work, moderate aneuploidy and CIN correlate with poor prognosis in cancer, but extreme aneuploidy correlates with improved patient outcomes [58,59]. Yeast RAD27 is a CIN gene [17], and FEN1 mutation in various systems leads to CIN and has been associated with cancer [17,60]; thus, inhibition of FEN1 in cancers that already exhibit CIN could lead to a level of CIN incompatible with viability. In the present study, flap endonuclease inhibitors were found to recapitulate the synthetic lethal interactions between FEN1 and each of CDC4 and MRE11A [18,23]. We observed that both depletion and inhibition of flap endonuclease activity led to an increase in endogenous DNA damage. Recent reports have shown that γ-H2AX levels are not increased following FEN1 depletion [61]; however, we observed increases in DNA damage using two independent assays following two means of FEN1 inhibition, and attribute these results to cell background differences, such as the mismatch repair deficiency present in HCT116 cells. Furthermore, this increase in DNA damage led to a non-apoptotic inhibition of proliferation. Thus, one explanation for the lethality in combination with inactivation of CDC4 is that the cell is inappropriately driven through the cell cycle, owing to elevated levels of cyclin E [13], when otherwise it would arrest to try to repair DNA damage. Likewise, increased endogenous DNA damage combined with loss of MRE11A, a protein playing a critical role in the first steps of the DNA damage response [62], could lead to a level of DNA damage or mutation that is incompatible with proliferation. CDC4 has been reported to be mutated in a wide variety of tumor types, at frequencies ranging from 6% to >30%, depending on the tumor type [13,21,29,63,64]. Recently, it has been suggested that reduction of CDC4 activity to some level below that of wild type, but above complete abrogation of function, is optimal for tumor progression [63]. Thus, the fact that two flap endonuclease inhibitors described here were able to selectively inhibit the proliferation of both heterozygous and homozygous CDC4-knockout cell lines suggests that CDC4 loss, whether complete

![Figure 3. IC50 curves of flap endonuclease inhibitors. FEN1 assays were carried out as described in Materials and Methods. Compound names are indicated above each graph, and structures are given to the right of each graph. Tumey 16 (top-left panel) was included as a positive control for flap endonuclease inhibition. doi:10.1371/journal.pgen.1003254.g003](https://www.plosgenetics.org/psp/article/10.1371/journal.pgen.1003254.g003)
or partial, sensitizes cells to inhibition of flap endonuclease activity. As well, the fact that both genotypes were sensitive to inhibition of flap endonuclease activity adds weight to the suggestion that this response is specific to CDC4 activity, in the same way that changing response following alteration in dosage in biochemical screening is suggestive of target identity [65].

In summary, here we have presented a rational, cross-species approach to the identification of anticancer therapeutic targets by targeting CIN, a common cancer phenotype. The use of conserved synthetic lethal interaction networks to identify highly-connected second-site targets is an accessible alternative to large scale screens: it narrows down the number of synthetic lethal gene pairs to be directly retested from tens of thousands to dozens, and is based on strong synthetic lethal interactions discovered in yeast networks. We have demonstrated the potential of this approach to identify targets and therapeutics, such as FEN1 and the flap endonuclease inhibitors described here, having potentially broad applicability in the treatment of cancer.

Materials and Methods

Cell culture

HCT116 cells were purchased from ATCC. HCT116 derivatives, DLD-1 and DLD-1 derivatives were gifts of Dr. Bert Vogelstein (Johns Hopkins University). (Importantly, we observed that the deleted exon in CDC4 in these cell lines is not exon 5, as previously reported [13], but exon 8. We attribute the difference to changing annotations in public sequence databases between 2004 and the present.) 53BP1-mCherry HCT116 cells were a gift of Dr. Sam Aparicio (UBC). These cells were grown in McCoy’s 5A medium with 10% FBS.
**Figure 5.** NSC645851 and RF00974 selectively inhibit the proliferation of HCT116 and DLD-1 cells with both homozygous and heterozygous inactivating mutations of **CDC4**. Experiments were carried out as described in Figure 4B. Data were analyzed by one-way ANOVA followed by a Tukey test. Shown is mean ± SEM. *p < 0.05; **p < 0.01; ***p < 0.001.

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**Figure 6.** RF00974 recapitulates the interaction between **FEN1** and **MRE11A**, and leads to increased endogenous DNA damage. (A) siRNA-mediated knockdown of MRE11A sensitizes HCT116 cells to treatment with RF00974. siRNA transfection experiments were carried out as described in Materials and Methods. Cells were fixed and imaged four days following siRNA transfection, after cells had been incubated in compound for 48 hours. Data were analyzed by one-way ANOVA followed by a Tukey test. Shown is mean ± SEM. *p < 0.05; **p < 0.01; ***p < 0.001. (B) Chemical inhibition of MRE11A sensitizes cells to RF00974. Cells were incubated in the indicated compounds at the indicated concentrations for three days prior to fixation and imaging as described in Materials and Methods. Data were analyzed by one-way ANOVA followed by a Tukey test. Shown is mean ± SEM. *p < 0.05; **p < 0.01; ***p < 0.001. (C) Inhibition of flap endonuclease activity with RF00974 mimics siRNA-mediated knockdown of FEN1 by increasing endogenous DNA damage. Data were analyzed by Student's t test. Shown is mean ± SEM.

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Immortalized (telomerase) BJ normal human skin fibroblasts, hTERT [66], were generously provided by Dr. C.P. Case (University of Bristol) and were grown in DMEM containing 10% FBS. Mirin was purchased from Sigma-Aldrich. RF00974 was purchased from Maybridge, Ltd.

Western blotting

Western blots were performed as detailed elsewhere [47]. Antibodies used for Western blots are described in Table S4.

RNA interference

Subconfluent and asynchronous cells were transiently transfected with siRNAs. HCT116 cells were transfected with ON-TARGETplus siRNA pools at a total siRNA concentration of 25 nM using DharmaFECT 1 (Dharmacon). In dual siRNA experiments, the total siRNA concentration was 50 nM. Cultures were replenished with fresh medium 11 hours after transfection. hTERT cells were transfected with ON-TARGETplus siRNA pools, or independent duplexes, at a total siRNA concentration of 100 nM using RNAiMax (Invitrogen). Cultures were replenished with fresh medium 24 hours after transfection.

Synthetic lethal assays, cell imaging, and compound incubation

HCT116 cells were harvested 24 hours after siRNA transfection and re-plated in 96-well optical bottom plates. hTERT cells were transfected directly in 96-well plates. HCT116 cells were fixed four days after transfection, and hTERT cells were fixed seven days after transfection, in 4% paraformaldehyde/PBS. Nuclei were labelled with Hoechst 33342. Stained nuclei were counted using a Cellomics Arrayscan VTI fluorescence imager as described previously [47] or a Zeiss AxioObserver Z1 equipped with an LED Colibri light source, a 20 x apochromat dry lens (numerical aperture = 0.8) and AxioVision v4.8 software. Images were analyzed using the Physiology Analyzer (Assaybuilder) option within the AxioVision software. Data were normalized to GAPDH-silenced controls and conventional statistics (e.g. column statistics and Student’s t-tests) were performed. Experiments were performed twice; indicated numbers are averaged from at least 6 wells.

To determine the presence of a synthetic lethal interaction, the proliferative defect was calculated, and is defined as

\[
\frac{1 - \text{Proliferation predicted by a multiplicative model}}{\text{Observed proliferation}}
\]

where the predicted proliferation was the product of the proliferation of the two individual gene knockdowns, following a multiplicative model of genetic interactions [67]. Synthetic lethal interactions were scored as a proliferative defect of three times the average SEM of the experiment or greater.

During compound incubation experiments, cells were incubated in compound of interest in 96-well optical bottom plates for approximately three days prior to fixation and analysis. Data (from six independent wells) were analyzed using a one-way ANOVA followed by a Tukey test.

FEN1 purification

FEN1 was expressed in BL21 E. coli from pET28b(+) (a generous gift from R. Bambara, University of Rochester) using 1 mM IPTG. Bacteria were lysed in lysis buffer (50 mM NaH₂PO₄, 300 mM NaCl, 10 mM imidazole, pH 8.0 containing 2× protease inhibitor) via a French press at 10 000 psi. The lysate was clarified and passed through a 0.22 μm filter before being loaded onto a HiTrap FF column (1 mL, GE Healthcare) in an AKTAFLC P-920 system (GE Healthcare). The column was washed in 10 volumes of wash buffer (lysis buffer+20 mM imidazole), and FEN1 was eluted with 5 volumes of elution buffer (lysis buffer+125 mM imidazole). The lysate was diluted with 9 volumes HI buffer (30 mM HEPES-KOH, 0.5% sxy-inositol, pH 7.8) with 30 mM NaH₂PO₄ and concentrated in a protein concentrator (Amicon). It was then loaded onto a hydroxyapatite resin (HA Ultrogel, Pall Life Sciences). The hydroxyapatite resin was washed with 10 volumes of HI-30 mM KCl and FEN1 was eluted with 5 volumes of HI-200 mM KCl. The eluate was diluted with 5 volumes HI-30 mM KCl prior to concentration, and then loaded onto a strong cation exchange column (1 mL HiTRAP SP FF FPLC, GE Healthcare Life Sciences). The column was washed with 10 volumes of HI-30 mM KCl, then 10 volumes of HI-200 mM KCl, and FEN1 was eluted with a gradient from HI-200 mM KCl to HI-500 mM KCl over 10 column volumes. Purified FEN1 was concentrated in FEN1 dilution buffer (30 mM HEPES-KOH, 5% glycerol, 0.1 mg/mL BSA, 0.01% NP-40), and aliquots of known concentration were frozen at −80°C.

In vitro FEN1 inhibition assay

Oligonucleotides used were as follows: “template”, 5'-GGTGACGGTGGATTGAAATTTAGGCTGGCACGGT CG-3', “upstream”, 5'-CGACCGTGCCAGCCTAAATTTCA ATC-3', “downstream”, 5'-6-FAM-CCAGGGCCACCCGTC AC-BHQ-1-3'. (6-FAM is 6-carboxyfluorescein; BHQ-1 is black hole quencher 1.) The three oligonucleotides were annealed at equimolar amounts in annealing buffer (50 mM Tris, 50 mM NaCl, 1 mM DTT, pH 8.0) by heating to 94°C, cooling to 70°C, and gradually cooling to room temperature. FEN1 assays were carried out with 6 pmol FEN1 and 20 nM annealed substrate in FEN1 buffer (50 mM Tris pH 8.0, 50 mM NaCl, 8 mM MgCl₂, 0.1 mg/mL BSA, 2 mM DTT). Assays were carried out at room temperature and kinetic reads were taken over approximately ten minutes in a Varioskan plate reader (Thermo Fisher Scientific), using excitation and emission wavelengths of 492 nm and 517 nm, respectively.

Fluorescent imaging

53BP1-mCherry cells were grown on cover slips. Following desired treatment (either two hours of bleomycin treatment at 5 μg/mL, four days following siRNA transfection, or after 24 hours of RF00974 treatment at 10 μM), cells were fixed for five minutes in 4% paraformaldehyde/PBS, mounted in Vectashield mounting medium containing DAPI (500 ng/mL), and imaged on a Zeiss Axioplan microscope with a Coolscan HQ camera, using appropriate filters and controlled by Metamorph software.

Apoptosis analysis

Cells were treated with RF00974 for 48 hours prior to harvesting of medium and cells in lysis buffer (50 mM Tris, 150 mM NaCl, 1% Triton-X-100, pH 7.5). Lysates were sonicated and clarified by centrifugation at 13 000 rpm for 15 minutes at 4°C. As a positive control, HCT116 cells were treated with 1 μM staurosporine prior to harvesting. Lysates were subjected to Western blotting as described above.
Synthetic genetic array

Synthetic genetic array analysis of rad27A against a collection of yeast essential DAnP alleles [68] and temperature sensitive alleles [69] was carried out as described previously [11,19].

Supporting Information

Figure S1 Western blots demonstrating knockdown of gene products targeted in this study. Cells were transfected with siRNA SMARTpools targeting the genes of interest. Proteins were harvested 3 days after transfection and Western blots were performed as detailed in Materials and Methods. Anti-β-tubulin was used as a loading control. (A) Knockdown of siRNA pools in HCT116 cells. (B) Knockdown of individual siRNA duplexes in hTERT cells. (DOC)

Figure S2 (A) The effect of non-silencing siRNA versus GAPDH siRNA on HCT116 cells. Cells were transfected with the indicated siRNAs, transferred to 96-well plates, fixed, and imaged as in Materials and Methods. (B) Response of HCT116 cells to selected siRNA on HCT116 cells. Cells were transfected with the indicated compounds. Cells were incubated with compound at the indicated concentrations for 72 hours in optically clear 96-well plates prior to fixation and imaging as described in Materials and Methods. Data were analyzed by one-way ANOVA followed by a Tukey test. Shown is mean ± SEM. (TIF)

Figure S3 Treatment of cells with RF00974 leads to an increase in phosphorylated H2AX, but not to an increase in apoptosis. Cells were treated with the indicated concentrations of RF00974 for 48 hours before protein was harvested and subjected to Western blot. Staurosporine (Stau) was used as a control to initiate apoptosis. (TIF)

Table S1 siRNA pool silencing in HCT116 cells. Horizontal lines indicate experiments carried out on different days. (DOC)

Table S2 siRNA pool silencing in hTERT cells. (DOC)

Table S3 Synthetic Lethality between FEN1 and cancer genes in hTERT cells. Horizontal lines indicate experiments carried out on different days. (DOC)

Table S4 Antibodies employed in Western blots in this study. (DOC)

Table S5 Genetic interactors of rad27A and cancer mutations. (XLS)

Table S6 Raw data from rad27A SGA against a collection temperature-sensitive and DAnP alleles of essential genes. #Spots, number of times allele was represented on array. E-C, experimental value minus control value (negative values indicate double mutant grows more slowly than control). Pval, p value of E-C. (XLS)

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Author Contributions

Conceived and designed the experiments: DMvP IJB YS BVS BJG TP KJM PH. Performed the experiments: DMvP IJB YS BVS BJG TP KJM PH. Analyzed the data: DMvP IJB YS BVS BJG TP KJM PH. Contributed reagents/materials/analysis tools: DMvP TP KJM PH. Wrote the paper: DMvP IJB YS BVS BJG TP KJM PH.

References

1. Hartwell LH, Szankasi P, Roberts CJ, Murray AW, Friend SH (1997) Integrating genetic approaches into the discovery of anticancer drug. Science 278: 1064–1068. 10.1126/science.278.5340.1064.
2. Farmer H, McCabe N, Lord CJ, Tutt AN, Johnson DA, et al. (2005) Targeting the DNA repair defect in BRCA mutant cells as a therapeutic strategy. Nature 434: 917–921.
3. Bryant HE, Schultz N, Thomas HD, Parker KM, Flower D, et al. (2005) Specific killing of BRCA2-deficient tumours with inhibitors of poly(ADP-ribose) polymerase. Nature 434: 913–917.
4. Weinstein JN, Myers TG, O’Connor PM, Friend SH, Fornace AJ, Jr., et al. (1997) An information-intensive approach to the molecular pharmacology of cancer. Science 275: 349–354.
5. Barretina J, Caponigro G, Stranges N, Venkatesan K, Margolin AA, et al. (2012) The Cancer Cell Line Encyclopedia enables predictive modelling of antitumour drug sensitivity. Nature 483: 605–607.
6. Garnett MJ, Edelman EJ, Heidorn SJ, Greenman CD, Dastur A, et al. (2012) Systematic identification of genomic markers of drug sensitivity in cancer cells. Nature 483: 570–575.
7. Grebien F, Hantschel O, Wojcik J, Kaupe I, Kovacic B, et al. (2011) Targeting the SH2-kinase interface in Bcr-Abl inhibits leukemogenesis. Cell 147: 506–519.
8. Zhao G, Chen A, Jamieson CH, Fershter T, Abrahamsson A, et al. (2009) Hedgehog signalling is essential for maintenance of cancer stem cells in myeloid leukaemia. Nature 458: 776–779.
9. Hanahan D, Weinberg RA (2011) Hallmarks of cancer: the next generation. Cell 144: 646–674.
10. Weaver BA, Cleveland DW (2006) Does aneuploidy cause cancer? Curr Opin Cell Biol 18: 658–667.
11. Sterling PC, Bloom MS, Solanki-Patil T, Smith S, Sipahimalani P, et al. (2011) The complete spectrum of yeast chromosome instability genes identifies candidate CIN cancer genes and functional roles for ASTRA complex components. PLoS Genet 7: e1002057. doi:10.1371/journal.pgen.1002057.
12. Cahill DP, Lengauer C, Vu J, Riggins GJ, Wilson JK, et al. (1998) Mutations of mitotic checkpoint genes in human cancers. Nature 392: 300–303. 10.1038/32088.
13. Rajagopalan H, Jallerpalvi PV, Rago C, Velezseucune KE, Vankler KW, et al. (2004) Inactivation of KCDC1 can cause chromosomal instability. Nature 428: 77–81. 10.1038/nature02313.
14. Barber TD, McManus K, Yuen KW, Reis M, Parmigiani G, et al. (2008) Chromatid cohesion defects may underlie chromosome instability in human colorectal cancers. Proc Natl Acad Sci U S A 105: 3443–3448.
15. Wang Z, Cummings JM, Shen D, Cahill DP, Jallerpalvi PV, et al. (2004) Three classes of genes mutated in colorectal cancers with chromosomal instability. Cancer Res 64: 2998–3001.
16. Tong AH, Evangelista M, Parsons AB, Xu H, Bader GD, et al. (2001) Systematic genetic analysis with ordered arrays of yeast deletion mutants. Science 294: 2364–2368.
17. Yuen KW, Warren CD, Chen O, Kenvit T, Hieter P, et al. (2007) Systematic genome instability screens in yeast and their potential relevance to cancer. Proc Natl Acad Sci U S A 104: 3925–3930.
18. McLellan J, O’Neil N, Tarailo S, Stoepel J, Bryan J, et al. (2009) Synthetic lethal genetic interactions that decrease somatic cell proliferation in Caenorhabditis elegans identify the alternative RFC CTF18 as a candidate cancer drug target. Mol Biol Cell 20: 5306–5313.
19. McLellan J, O’Neil N, Barrett I, Ferree E, van Pel DM, et al. (2012) Synthetic lethality of cohesins with PARPs and replication fork mediators. PLoS Genet 8: e1002574. doi:10.1371/journal.pgen.1002574.
20. Hiramoto T, Nakamichi T, Sumiyoshi T, Fukuda T, Matsuura S, et al. (1999) Mutations of a novel human RAD54 homologue, RAD54B, in primary cancer. Oncogene 18: 3422–3426.
21. Kemp Z, Rowan A, Chambers W, Wortham N, Halford S, et al. (2005) CDC4 mutations occur in a subset of colorectal cancers but are not predicted to cause loss of function and are not associated with chromosomal instability. Cancer Res 65: 11361–11366.
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22. Cahill DP, Kindler KW, Vogelstein B, Lengauer C (1999) Genetic instability and darwinian selection in tumours. Trends Cell Biol 9: M57–M60. 10.1016/S0962-8924(99)01661-X.

23. Costanzo M, Baryshnikova A, Bellay J, Kim Y, Spear ED, et al. (2010) The genetic landscape of a cell. Science 327: 425–431.

24. Prieto I, Suja JA, Perzzi N, Kremer L, Martinez N, et al. (2001) Mammalian STAG3 is a cohesin specific to sister chromatid arms in meiosis I. Nat Cell Biol 3: 761–766.

25. Liu Y, Bamba RA (2003) Analysis of human flap endonuclease 1 mutants reveals a mechanism to prevent triplet repeat expansion. J Biol Chem 278: 17279–17379.

26. Zheng L, Jin J, Finger LD, Guo Z, Zer C, et al. (2011) Functional regulation of FEN1 nucleolytic activity and its link to cancer. Nucleic Acids Res 39: 781–789.

27. Tumey LN, Bom D, Huck B, Gleason E, Wang J, et al. (2005) The identification and optimization of a N-hydroxy urea series of flap endonuclease 1 inhibitors. Bioorg Med Chem Lett 15: 277–281.

28. Lipinski CA, Lombardo F, Dominy BW, Ferren PJ (2001) Experimental and computational approaches to estimate solubility and permeability in drug discovery and development settings. Adv Drug Deliv Rev 46: 3–26.

29. Akhooei S, Sun D, von der LN, Apostolidou S, Klotz K, et al. (2007) FBXW7/hCDC4 is a general tumor suppressor in human cancer. Cancer Res 67: 9006–9012.

30. Koh JL, Ding H, Costanzo M, Baryshnikova A, Toufighi K, et al. (2010) DRUGVISION: a database of quantitative genetic interaction networks in yeast. Nucleic Acids Res 38: D502–D507.

31. Miyaki M, Yamaguchi T, Iijima T, Takahashi K, Matsumoto H, et al. (2009) Somatic mutations of the CDC4 (FBXW7) gene in hereditary colorectal tumors. Oncology 76: 430–434.

32. Milne AN, Leguit R, Corver WE, Morsink FH, Polak M, et al. (2010) Loss of CDC4/FBXW7 in gastric carcinoma. Cell Oncol 32: 347–359.

33. Dupre A, Boyer-Chatenet L, Sattler RM, Modi AP, Lee JH, et al. (2008) A forward chemical genetic screen reveals an inhibitor of the Mre11-Rad50-Nbs1 complex. Nat Cell Biol 4: 119–125.

34. Anderson L, Henderson C, Adachi Y (2001) Phosphorylation and rapid relocation of 53BP1 to nuclear foci upon DNA damage. Mol Cell Biol 21: 1719–1729.

35. Rappold I, Isobeuchi K, Date T, Chen J (2001) Tumor suppressor p53 binding protein 1 (53BP1) is involved in DNA damage-signaling pathways. J Cell Biol 153: 615–628.

36. Markova E, Schultz N, Belyaev IY (2007) Kinetics and dose-response of residual DNA mismatch repair proteins MSH2 or MLH1. Cancer Cell 17: 235–248.

37. Lazebnik YA, Kaufmann SH, Desnoyers S, Poirier GG, Earnshaw WC (1994) DNase IV (FEN1) structure-specific endonuclease. EMBO J 13: 1235–1246.

38. Mendes-Pereira AM, Martin SA, Brough R, McCarthy A, Taylor JR, et al. (2007) FBXW7/hCDC4 is a general tumor suppressor in human cancer. Cancer Res 67: 9006–9012.

39. Reamer PM, Griffiths MR, Long JM, Charlier JD, Mascornick S, et al. (2011) Selective killing of ATM- or p53-deficient cancer cells through inhibition of ATR. Nat Chem Biol 7: 498–506.

40. Zheng L, Dai H, Hegde ML, Zhou M, Guo Z, et al. (2011) Fen1 mutations that specifically disrupt its interaction with PCNA cause aneuploidy-associated cancer. Cell Res 21: 1858–1870.

41. Lee AJ, Endesfelder D, Rowan AJ, Walther A, Birkbak NJ, et al. (2011) Chromosomal instability confers intrinsic multidrug resistance. Cancer Res 71: 621–628.

42. Duxin JP, Moore HR, Sidorova J, Karanja K, Honaker Y, et al. (2012) An integrated platform of genomic assays reveals small-molecule bioactivities. Nat Chem 4: 498–506.

43. Strom CE, Johansson F, Uhlén M, Szigyarto CA, Erixon K, et al. (2011) Poly (ADP-ribose) polymerase (PARP) is not involved in base excision repair but PARP inhibition traps a single-strand intermediate. Nucleic Acids Res 39: 3166–3175.

44. Tang S, Hanby FC, Helleday T (2012) Mre11-dependent degradation of stalled DNA replication forks is prevented by BRCA2 and PARP1. Cancer Res 72: 2014–2021.

45. Bouchal NJ, Ekland AC, Li Q, McClelland SE, Endesfelder D, et al. (2011) Paradoxical relationship between chromosomal instability and survival outcome in cancer. Cancer Res 71: 3447–3452.

46. Lee AJ, Endesfelder D, Rowan AJ, Walther A, Birkbak NJ, et al. (2011) Chromosomal instability confers intrinsic multidrug resistance. Cancer Res 71: 1858–1870.

47. Harrington JJ, Lieber MR (1994) The characterization of a mammalian DNA structure-specific endonuclease. EMBO J 13: 1235–1246.

48. Murante RS, Rumbaugh JA, Barnes CJ, Norton JR, Bamba RA (1996) Calf RTH1 nuclease can remove the initiator RNAs of Okazaki fragments by endonuclease function. J Biol Chem 271: 23898–23897.

49. Harrington JJ, Lieber MR (1994) The characterization of a mammalian DNA structure-specific endonuclease. EMBO J 13: 1235–1246.

50. Kim K, Biale S, Matsunoto Y (1986) Involvement of flap endonuclease 1 in base excision DNA repair. J Biol Chem 273: 8942–8948.

51. Klangland A, Lindahl T (1997) Second pathway for completion of human DNA base excision-repair: reconstitution with purified proteins and requirement for DNAase IV (FEN1). EMBO J 16: 3341–3348.

52. Singh P, Zheng L, Chavez Y, Qiu J, Shen B (2007) Concerted action of exonuclease and Gap-dependent endonuclease activities of FEN-1 contributes to the resolution of triplet repeat sequences (CTGs)n- and (GA)n-derived secondary structures formed during maturation of Okazaki fragments. J Biol Chem 282: 3465–3477.

53. Zheng L, Zhou M, Chai Q, Parrish J, Xue D, et al. (2005) Novel function of the flap endonuclease 1 complex in processing stalled DNA replication forks. EMBO Rep 6: 83–89.

54. Jaiswal AS, Banerjee S, Aneja R, Sarkar FH, Ostrov DA, et al. (2011) DNA Polymerase beta as a Novel Target for Chemotherapy Intervention of Colorectal Cancer. PLoS ONE 6: e16691. doi:10.1371/journal.pone.0016691

55. Tong AH, Lesage G, Bader GD, Ding H, Xu H, et al. (2004) Global mapping of the yeast genetic interaction network. Science 303: 808–813. 10.1126/science.1091317.

56. Strom CE, Johansson F, Uhlén M, Szigyarto CA, Erixon K, et al. (2011) Poly (ADP-ribose) polymerase (PARP) is not involved in base excision repair but PARP inhibition traps a single-strand intermediate. Nucleic Acids Res 39: 3166–3175.

57. Zhang L, Zhou M, Chai Q, Parrish J, Xue D, et al. (2005) Novel function of the flap endonuclease 1 complex in processing stalled DNA replication forks. EMBO Rep 6: 83–89.

58. Tumey LN, Bom D, Huck B, Gleason E, Wang J, et al. (2005) The identification and optimization of a N-hydroxy urea series of flap endonuclease 1 inhibitors. Bioorg Med Chem Lett 15: 277–281.