Xpf and Not the Fanconi Anaemia Proteins or Rev3 Accounts for the Extreme Resistance to Cisplatin in Dictyostelium discoideum

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Abstract

Organisms like Dictyostelium discoideum, often referred to as DNA damage “extremophiles”, can survive exposure to extremely high doses of radiation and DNA crosslinking agents. These agents form highly toxic DNA crosslinks that cause extensive DNA damage. However, little is known about how Dictyostelium and the other “extremophiles” can tolerate and repair such large numbers of DNA crosslinks. Here we describe a comprehensive genetic analysis of crosslink repair in Dictyostelium discoideum. We analyse three gene groups that are crucial for a replication-coupled repair process that removes DNA crosslinks in higher eukarya: The Fanconi anaemia pathway (FA), translesion synthesis (TLS), and nucleotide excision repair. Gene disruption studies unexpectedly reveal that the FA genes and the TLS enzyme Rev3 play minor roles in tolerance to crosslinks in Dictyostelium. However, disruption of the Xpf nuclease subcomponent results in striking hypersensitivity to crosslinks. Genetic interaction studies reveal that although Xpf functions with FA and TLS gene products, most Xpf mediated repair is independent of these two gene groups. These results suggest that Dictyostelium utilises a distinct Xpf nuclease-mediated repair process to remove crosslinked DNA. Other DNA damage-resistant organisms and chemoresistant cancer cells might adopt a similar strategy to develop resistance to DNA crosslinking agents.

Introduction

DNA interstrand crosslinks are complex lesions that covalently link the two complementary strands of DNA. Agents that cause this type of lesion can originate from an endogenous source such as reactive species generated by lipid peroxidation, or as a consequence of exposure to exogenous mutagens [1–5]. For this reason the cytotoxicity of DNA crosslinks is exploited in cancer chemotherapy, where drugs such as cisplatin, mitomycin C and melphalan are administered as potent DNA crosslinking agents. DNA crosslinks are extremely cytotoxic because they form an absolute barrier to replication [6]. In addition, a crosslink present in a gene coding sequence, will also block transcription. Apart from cell death, DNA crosslinks can also lead to cell senescence and dysfunction [7,8]. These features are observed in humans born with defective crosslink repair as such individuals exhibit growth retardation, stem cell attrition and symptoms consistent with premature aging [9]. These phenotypic features may be due to the accumulation of unrepaired crosslinks in genomic DNA.

Crosslinks can also form between adjacent bases on the same DNA strand, which are referred to as intrastrand crosslinks. Of the two classes of crosslinks, interstrand crosslink is believed to be the more cytotoxic. Crystal structures of lesions formed by reacting cisplatin with DNA have now been solved showing that these lesions cause substantial helix distortion. In terms of DNA repair, genetic and biochemical studies have shown that intrastrand crosslinks are largely repaired by nucleotide excision repair [10]. Repair of interstrand crosslinks is much more complex and poorly understood. Much of the work here is underpinned by genetic studies of classes of mutants that in certain organisms render cells selectively or generally sensitive to chemical crosslinking agents. Four clear repair gene groups in vertebrates stand out in this manner: the Fanconi anaemia (FA) genes, the translesion DNA polymerases Rev1 and Rev3, homologous recombination (HR) repair genes and finally the structure-specific nucleases subcomponents XPF and Mus81 [11–13]. Taking this knowledge into account a replication-coupled model for interstrand crosslink repair has been proposed. This model suggests that replication pausing at or near a crosslink initiates a cleavage (a step commonly referred to as unhooking), which is followed by lesion bypass over the crosslinked base by translesion DNA synthesis (TLS). An intact chromatid is therefore created and can now be used as a template to complete repair by HR [11,13].

Not all the gene groups that function in vertebrate crosslink repair are conserved in yeast. Apart from FANCN of the other 12 Fanconi anaemia genes appears to have orthologues in this organism [14,15]. This limits the use of yeast in understanding crosslink repair in higher eukaryotes. Crosslink repair has therefore been largely studied in immortalised vertebrate cell lines (such as chicken DT40 cells or Chinese hamster ovary cells). A drawback of some of these systems is however that they contain mutations in other genes such as p53 that may influence repair. For these reasons some workers have turned to worms and flies [16,17], as both organisms are genetically tractable and have some of the vertebrate crosslink repair groups conserved. A potential limitation of these
DNA Crosslink Repair in Dictyostelium discoideum

Author Summary

Organisms are constantly exposed to environmental and endogenous molecules that chemically modify the DNA in their genomes. A particularly pernicious chemical modification is when the two strands of DNA are crosslinked. These crosslinks must be removed so that genomes can be copied, and the damage caused by their persistence is often exploited in cancer chemotherapy. It is also no surprise that all organisms have developed effective means to remove these lesions, and work in prokaryotes and eukaryotes has shown that crosslinks are removed by the concerted action of certain DNA repair pathways. Whilst the obvious route of accumulating crosslinks is by exposure to anti-cancer drugs, these lesions may also arise spontaneously in DNA. This could be why inherited inactivation of one of the crosslink repair pathways results in the catastrophic human illness Fanconi anaemia. Here we determine how the social amoeba Dictyostelium discoideum, an organism that is unusually resistant to DNA-damaging agents, removes crosslinks. Our results indicate that this organism has evolved a distinct strategy to remove these lesions. More specifically, we discover that a particular nuclease subcomponent removes the crosslinks by a distinct repair process. We postulate that this strategy to remove crosslinks could be used by other DNA damage-resistant organisms and also by cancer cells that have developed resistance to chemotherapy.

Inactivation in humans leads to Fanconi anaemia—an illness that leads to developmental defects, stem cell attrition and cancer predisposition [5, 29, 30]. There are 13 known FA genes in humans. Most of them (FANCA, B, C, E, F, G, L, M, FAAP100 and FAAP24) assemble into a nuclear complex—hitherto referred to as the FA core complex. This complex interacts with the E2 enzyme Ube2t [31, 32], and monoubiquitinates two other FA proteins FANCd2 and FANCi. Both proteins form a complex and co-localise at sites of DNA damage with FANCd1 (BRCA2), FANCN (PALB2) and the FANCj helicase [30]. All the FA proteins are highly conserved in vertebrates. As a first step to dissect crosslink repair in Dictyostelium we delineated the pattern and depth of their conservation in all eukaryotes. A clear picture emerges from this analysis (Figure 1A): a minimal FA pathway consists of FANCd2 (FncD2), FANCj (FncJ), FANCn (FncN), FANCm (FncM), FANCf (FncF), Ube2t (Ube2t) and FancD1 (BRCA2); the later appears to have evolved in the ancestral eukaryote. Additional components, including most of the FA core complex proteins, evolved later in the ancestral metazoan. With respect to Dictyostelium, this analysis suggests a simplified FA pathway may operate in this organism (Figure 1B).

Next, we proceeded to establish a functional role for the minimal FA pathway in Dictyostelium. We bioinformatically identified the genomic loci of the Dictyostelium FA genes and using these information generated knockouts of orthologues of FANCd2, FANCg, FANCj and Ube2t (Figures S1, S2, and S3, and Table S1). To study the response to DNA crosslinks, the various Dictyostelium strains were exposed to cisplatin. After one hour exposure to a range of doses, the amoebae were diluted, plated out onto bacterial lawns and allowed to grow for 4 days. Surviving amoebae form distinct plaques on the bacterially coated agar plates, each of which represents a colony arisen from a single cell. The number of plaques was counted and survival was expressed as a percentage of plaques formed by mock-treated cells. This assay is very much like the standard colony survival assay used in toxicity studies with vertebrate cell lines. The data in Figure 2 shows that most of the FA knockout strains show a moderate sensitivity to cisplatin. A notable exception is the fncl knockout, which does not seem to be sensitive. Also of note is the dose of mutagen required to compromise wild type cells, which is in the range of 500 μM. This is a very large dose considering that human and chicken cells show sensitivities in the 1–40 nM range. This difference becomes even more striking when considering the chicken fancl knockout, which has a D50 of 5 nM (8 fold more sensitive compared to wild type), to its Dictyostelium counterpart, which has a D50 value of 165 μM (2 fold more sensitive than wild type). We can conclude that, firstly, Dictyostelium is much more resistant to cisplatin than vertebrate cells. Secondly, the identifiable FA genes are functionally required for this resistance, though unlike in vertebrates their overall contribution is much less marked.

An FncL protein complex monoubiquitinates Fncd2 in Dictyostelium

The monoubiquitination of FANCd2 is a key biochemical step in the FA pathway. In vertebrates this step requires the complete FA core complex, with FANCN and Ube2t forming the catalytic core of this reaction [33]. Studies in at least two non-vertebrate model organisms (flies and worms) confirm that FANCd2 monoubiquitination is conserved [16, 17]. Both these organisms appear to have lost many core complex genes, once again raising the possibility of a minimal FA pathway operating in simpler organisms. Dictyostelium provides a unique opportunity to test if this is true since it lacks obvious orthologues of so many FA genes. Our first step was to establish whether FncD2 is monoubiquitinated and then to
To determine the genetic requirements for this, we developed a FncD2 reporter strain where a YFP-tag was knocked in frame after the penultimate codon in the last exon of this gene (Figure 3A). Western blot analysis (Figure 3B) and cisplatin survival data (Figure S5) confirm that this strain expresses functional FANCD2-YFP and is not sensitive to cisplatin. In order to detect monoubiquitinated FncD2 we expressed HA-tagged ubiquitin in the FncD2-YFP strain.

Cell extracts prepared from cisplatin or mock-treated cells were immunoprecipitated with an anti-YFP antibody and Western blotted for the HA-tag. A single DNA damage-inducible band, which corresponded to the size of FncD2-YFP was detected (Figure 3C). We then knocked out fncL in this strain and found that monoubiquitinated FncD2-YFP was no longer detectable (Figure 3D). Our next step was to determine if FncL acted alone or as part of a complex. Our bioinformatics analysis presented in Figure 1 revealed a possible DNA Crosslink Repair in Dictyostelium discoideum.
FANCE orthologue - FncE (Figure S4). FANCE is an essential component of the vertebrate FA nuclear complex. We deleted this gene and found that the resultant \(\Delta fncE\) strain was moderately sensitive to cisplatin (Figure 2) and that monoubiquitinated FncD2-YFP was no longer detectable (Figure 3D). Finally, we needed to determine if any one of these FA core complex proteins exists in a complex. To assay for this, we generated a strain that expresses N-terminal TAP-tagged FncL (Figure S5). Whole cell extract from this strain was subjected to size exclusion chromatography and fractions were blotted for TAP-FncL. The data in Figure 3E clearly show that TAP-FncL is present in two large molecular size peaks of approximately 800 kDa and 140 kDa respectively (Figure 3E). In summary, this data shows that FncL and FncE are required for FncD2 monoubiquitination (Figure 3F). Since FncL appears to reside in a protein complex it is unclear whether a truly ‘minimal’ FA pathway operates in this simple organism.

Rev3 functions with FncD2 to repair crosslinks

A recent study surveyed the relative sensitivity of a large number of DNA repair mutants generated in the isogenic chicken cell line

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**Figure 2. Disruption of the FA genes in *Dictyostelium* leads to sensitivity to the crosslinking agent cisplatin.** (A) The genetic loci for fncD2, fncL, fncI, fncJ, fncM, and ube2T were disrupted and deletion strains were confirmed by Southern analysis (Figure S1, S2, and S3). Viability was estimated by colony survival after exposure to a dose titration of cisplatin for 1 hour prior to plating on bacterial agar lawns. Each survival curve consists of a triplicate experimental data set run in parallel with the wild type (Ax2) control. (B) Table of \(D_{50}\) toxicity values for all deletion strains. X-fold represents the difference in the \(D_{50}\) value of the relevant strain relative to the wild type control. KO: deletion strain, SE: standard error, n: number of independent experiments carried out.

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Figure 3. An FncL complex is required for FncD2 monoubiquitination in Dictyostelium. (A) Map of the fncD2 locus (DDB_G0268216) and design of 3’ YFP-knockin strategy. The fncD2 coding region is shown in dark. (B) An anti-YFP Western blot performed on extracts obtained from Ax2 cells, two FncD2-YFP knockin clones, and cells expressing a YFP-tagged PHD finger protein (positive control). FncD2-YFP is expressed from its endogenous locus and migrates only as a single band. (C) To detect monoubiquitinated FncD2-YFP, HA-ubiquitin was constitutively expressed in FncD2-YFP knockin cells. Whole cell lysates prepared from mock treated (-) cells or cells exposed to 400 mM cisplatin (+) for 1 hour were immunoprecipitated with an anti-YFP antibody and then Western blot with anti-HA (top) or anti-YFP (bottom) antibodies respectively. DNA damage induces FncD2 expression and this correlates with increased monoubiquitination. (D) Disruption of fncL and fncE in the fncD2-YFP+HA–ubiquitin strain results in undetectable FncD2 monoubiquitination. (E) Whole cell extract from in situ tagged TAP-FncL cells was subjected to size exclusion chromatography. Fractions were collected and Western blotted with an anti-TAP antibody to detect the tagged FncL protein. TAP-FncL fractionates in large molecular mass peaks. (F) A schematic diagram summarising the FA crosslink repair pathway in Dictyostelium.

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DT40 [34]. This comparison revealed that the most sensitive mutants are those that lack the translesion polymerases Rev1 and Rev3, followed closely by mutants that lack the FA genes. Analysis of double mutants within these two groups of genes in DT40 shows that they participate in a common process to repair crosslinks [11]. These observations prompted us to establish the role of TLS in Dictyostelium crosslink repair. The Dictyostelium genome appears to contain a smaller complement of TLS enzymes than vertebrates. However a Rev3 orthologue (rev3) was easy to identify. We disrupted the rev3 locus, the ensuing Δrev3 strain (Figure 4A and 4B) was viable, grew normally in culture and showed normal development (Figure 5B). We then tested the Δrev3 for sensitivity to cisplatin and were surprised to see that it was only moderately sensitive to this agent (3 fold over WT) (Figure 4C). We then disrupted fncD2 in this strain to test the genetic interaction between these two crosslink repair genes. fncD2 deficiency makes no additional or synergistic impact in the Δrev3 strain (Figure 4C and 4D), indicating that both genes function in a common process to repair crosslinks. However, notably the Δrev3 strain (like the FA mutants) was not strongly sensitive to crosslinks, once again

Figure 4. The TLS polymerase Rev3 and FncD2 function in the same process to repair crosslinks in Dictyostelium. (A) Map of the intact and disrupted rev3 locus (DDB_G0271608). rev3 exons (dark) are largely deleted by the used knockout strategy. (B) Southern blot of Ax2 cells and two Δrev3 cell lines. Genomic DNA was digested with EcoRV and probed with the DNA region marked in (A). The wildtype band is 8.6 kb, which is converted into 2.8 kb band if gene targeting was successful. (C) Colony survival of wild type, ΔfncD2, Δrev3, and Δrev3ΔfncD2 double knockout cells following exposure to cisplatin. Knocking out rev3 in a ΔfncD2 knockout background has no synergistic impact on sensitivity to crosslinks. The survival curve consists of a triplicate experimental data set run in parallel with the wild type (Ax2) control. (D) Graphical representation of the D50 values of the strains in (C), calculated from five independent colony survival experiments.

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Figure 5. Xpf functions largely independently of either Rev3 or FncD2 in crosslink repair. (A) The ΔfncD2Δxpf and the Δrev3Δxpf double knockout strains are as sensitive to cisplatin as the single Δxpf knockout (assayed by colony survival). Results shown are from a single experiment, which is representative of at least three independent experiments. Error bars indicate variation between triplicate plating. The kill curves are shown with a concentration range from 0–100 μM (top of the panel) and also 0–10 μM (bottom). (B) Development is not impaired in Δxpf and Δrev3 strains. doi:10.1371/journal.pgen.1000645.g005
The nucleotide excision repair nuclease subcomponent XPF is essential for crosslink repair

The fact that Dictyostelium mutants of FA and TLS genes are only moderately sensitive to cisplatin surprised us. This organism may be resistant to cisplatin because of reduced bioavailability of the drug (reduced uptake/enhanced breakdown). In addition, it is noteworthy that the Dictyostelium genome is very AT-biased [20]. Since cisplatin crosslinks form at mainly GC sequences, this could mean that very few lesions are produced. Alternatively, crosslink repair may be carried out by another process that does not use the FA and TLS genes studied here. One obvious pathway would be HR. However, to date we and others have not been able to knockout core genes in this pathway [35,36]. Perhaps this could be due to an essential role for HR in cell viability. Another candidate group of genes are those involved in nucleotide excision repair. Vertebrate cell lines lacking NER show differential requirements for crosslink repair. Certain genes like xpa and xpc play at best only a minor role [9,37], whilst the nucleosome subcomponent XPF appears to be very important. Indeed, all models of crosslink repair invoke the action of a nuclease in cutting on either side of the crosslink, a step referred to as unhooking. In addition to XPF/ERCC1, the Mus81/EME1 nuclease complex is also believed to be important in vertebrate crosslink repair [12,38,39]. We therefore set out to disrupt XPF (xpf), XPG (xpc) and Mus81 (mus81). The respective orthologues were identified, their loci mapped and disrupted (Figure 6A–6D and Figure S6A and S6B). All three mutant cell lines were then tested for sensitivity to cisplatin. The Δxpc and Δmus81 strains were not sensitive (Figure 6E and Figure S6C), in contrast to the Δxpf/ mutant, which was extremely sensitive to cisplatin. The Δxpc values reflect this with Δxpf giving a value of 4 μM, in contrast with Δxpc (342 μM) and wild type (290 μM) (Figure 6F). In summary, the excision repair nuclease subcomponent XPF is essential for repairing crosslinks in Dictyostelium. This activity is not due to the role of this gene in global NER since the xpc mutant is not at all sensitised to cisplatin.

xpf functions largely independently of rev3 and fncD2

The current models for crosslink repair all involve a nuclease(s) carrying out the excision step. Our discovery of an essential role played by xpf and not mus81 in crosslink repair makes xpf a very good candidate component for the nuclease implicated in such a step. In a Xenopus cell free system, a crosslinked plasmid was repaired in a replication process that involves excision and TLS [40]. This important study therefore raises the question regarding the identity of the nuclease(s) involved in this excision step. A genetic test to determine if XPF might be involved here is to generate a ΔxpfΔrev3 strain and to establish genetic epistasis between these genes. If the double mutant is as sensitive as the single Δxpf strain then this indicates that xpf functions with rev3 in crosslink repair. This is indeed what we see since disruption of rev3 does not impact further on the sensitivity to cisplatin in the Δxpf strain (Figure 5A). In addition, we also demonstrated that xpf is epistatic with respect to fncD2 (Figure 5A). The single Δxpf mutant is 20–30 fold more sensitive than either Δrev3 or ΔfncD2 strains, respectively, which indicates that most crosslink repair requires xpf but not rev3 or fncD2. Finally there is considerable evidence for the role of XPF in HR repair [41–43]. To test whether HR repair is compromised in Δxpf we analysed gene targeting efficiency into two independent loci. The data in Table 1 clearly shows that homologous gene targeting is compromised to varying degrees in both loci analysed in the Δxpf strain compared to wild type AX2.

Discussion

The studies presented in this paper establish the genetic requirements for crosslink repair in Dictyostelium. This simple unicellular genotoxin-resistant eukaryote shares the important groups of crosslink repair genes that function in humans. In contrast to vertebrates, the FA proteins and TLS enzymes appear to play only a minor role in repairing crosslinks. However, the most striking discovery reported here is that the nucleotide excision repair gene xpf is essential for crosslink repair in Dictyostelium.

So far our analysis has confirmed that at least two known proteins that are crucial for the function of the FA core complex, FANCE and FANCL, are conserved in this organism. FANCE links the FA complex to its main substrate FANCID2 [44] and FANCL is the E3 subunit in the complex [45]. An unresolved question is whether Dictyostelium has a truly simplified FA pathway, Certainly the FncL protein exists in a high molecular mass protein complex. Such a complex may consist only of FncL and FncE. Another possibility is that there are other proteins in this complex. Such proteins could be other FA core complex orthologues that have simply evaded detection by bioinformatics as they may have diverged at the amino acid sequence level but not at a structural level. Alternatively, both FncL and FncE could be embedded in a complex consisting of new proteins or into a known surrogate multiprotein E3 ligase complex. Purification and identification of components of the FncL complex should address these possibilities.

It has long been appreciated that Dictyostelium is an unusually DNA damage-resistant organism. The work presented here further illustrates this. Conceivably factors such as reduced bioavailability of cisplatin and the number of crosslinks introduced into the genome may contribute towards this resistance. However, the profound cisplatin sensitivity as a result of xpf inactivation clearly showed that a sufficient number of crosslinks are formed to cause lethal damage. We were surprised that the FA and the TLS proteins seem to only contribute a minor activity towards this crosslink resistance. This finding contrasts with what has been observed with vertebrate cells, where both groups of genes are crucial for tolerance to DNA crosslinking agents. The profound sensitivity of the xpf-deficient strain raises the question how xpf contributes to tolerance to DNA crosslinking agents repair. The genetic interactions between xpf, fncD2 and rev3 show that there is a minor repair process involving all three genes, but the dominant mechanism of xpf-dependent crosslink repair remains to be determined.

During vegetative growth, Dictyostelium displays a skewed cell cycle distribution. The vast majority of cells are found in the G2 phase of the cell cycle [46,47]. The G1 phase appears to be very short. This, in terms of DNA content, means that most of the vegetative cells possess a duplicated genome. Upon exposure to cisplatin, it is very likely that lesions form at only one copy per site. Under such conditions, the most straightforward means of repair would be to excise the crosslink creating a double strand break. The undamaged copy could then be used as the template for HR-mediated double strand break repair (Figure 7). Such a model predicts the requirement for HR genes in crosslink repair, a proposition that is currently difficult to address, since we and others thus far have been unable to disrupt HR genes in Dictyostelium [35,36]. In addition, this model of crosslink repair may require additional nucleases to not only to create but also to process DNA double strand breaks, remove flaps or resolve secondary structures. All these activities would be quite distinct from the unhooking step itself.

Considerable work in both yeast and vertebrates point to a critical role for XPF and its orthologues in homologous recombination repair [42,48]. ES cells and yeast knockouts appear to be defective at homologous gene targeting though it is
Figure 6. Disruption of the NER nuclease subcomponent xpf but not xpc results in profound sensitivity to cisplatin in *Dictyostelium*.

(A) Map of the intact and disrupted xpf locus (DDB_G0284419). xpf exons (dark) are largely deleted by this strategy. (B) Southern blot of Ax2 cells and two xpf knockout clones. Genomic DNA was digested with BamHI and probed with the DNA region marked on the map in (A). The WT band is 6.6 kb, which is converted into a 2.5 kb band with successful gene targeting. (C) Map of the intact and disrupted xpc locus (DDB_G0292296). xpc exons (dark) are largely deleted by this strategy. (D) Southern blot of Ax2 cells and two xpc knockout clones. Genomic DNA was digested with XbaI and probed with the DNA region marked on the map in (C). The WT band is 10.6 kb. With successful gene targeting the band is converted into a 3.3 kb band. (E) Colony survival of wild type, Δxpc, and Δxpf cells after exposure to cisplatin. Knocking out xpf results in a profound sensitivity to cisplatin. (F) A table showing the doubling time (T2) and D50 values for Δxpf and Δxpc strains. SE: standard error.

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important to appreciate that this is not always a consistent feature [41,43]. Recombinant XPF/ERCC1 also function in processing recombination intermediates as well as synthetic replication forks [49]. Indeed we also find a defect in gene targeting in the Δxpff strain indicating that like in other organisms, Xpf does play a role in HR in Dictyostelium. It is therefore possible that it is the HR functions of Xpf that determines why it is so crucial for the tolerance of crosslinks in Dictyostelium. Finally, it is noteworthy that there are many organisms that share resistance to DNA damaging agents with Dictyostelium. The dependence of an excision nuclease-based repair mechanism may be responsible for such resistance. Such a mechanism may not just be limited to such organisms but also to human cancers which develop resistance to cisplatin [50]. Induction of such an excision repair pathway may account for the acquired resistance to chemical crosslinking agents. Future work will aim to address the genetic requirements and elucidate the mechanism of the Xpf-dependent crosslink repair pathway.

### Materials and Methods

#### Strain generation

All targeting constructs were generated using pLPBLP as the backbone [51]. 5' and 3' homology arms were generated by PCR amplification from Ax2 genomic DNA using Pwo polymerase and inserted into the plasmid on either side of the blasticidin-resistance cassette (bsr). The HA-ubiquitin overexpression construct was generated using pDXA-3C as backbone [52]. The wild-type strain

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**Table 1.** Measured gene-targeting frequencies into two loci in WT and Δxpff cells.

| Strain | Gene Targeted | DDB_G0293840 | DDB_G0267916 |
|--------|---------------|---------------|---------------|
| WT     | 24% (12/50)   | 80% (131/170) |               |
| Δxpff  | 0% (0/25)     | 2% (12/518)   |               |

The table depicts the gene-targeting efficiencies into two independent genetic loci of WT and Δxpff cells. For each locus multiple transfections were performed and drug-resistant clones were analysed by PCR for proper targeting. The number of gene-targeting events is shown alongside the total number of drug resistant clones. Gene-targeting efficiency is expressed as a percentage value of the total number of clones. doi:10.1371/journal.pgen.1000645.t001

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**Figure 7. Model for an Xpf-dependent crosslink repair process in Dictyostelium.** Model outlining a potential mechanism of crosslink repair that may operate in Dictyostelium. A small proportion of repair is channelled through a combined FA, TLS, and HR route stimulated by replication fork stalling. However, as Dictyostelium spends most of its time in the G2 phase of the cell cycle it is likely that a crosslink may form at only one site on a particular chromosome. A simple mechanism may operate whereby Xpf/Ercc1 together with other nucleases may cut out the crosslinked section creating a double strand break. HR using the sister chromatid as a template might then complete this repair process. It is also possible that Xpf plays an additional role in the HR step. doi:10.1371/journal.pgen.1000645.g007
and the parent of all strains generated in this study was the Kay laboratory version of Ax2. Transformants were created by electrotransfection (GenePulser Xcell Bio-Rad) of 17.5 μg of the targeting cassette or 25 μg of the overexpression plasmid. Potential homologous recombinants were selected for blasticidin resistance (10 μg/ml) at limiting dilution in 96-well plates, whereas overexpression lines were selected as a pool of transformants in the presence of 10 μg/ml G418. After approximately 10 days, the content of positive wells were cloned out onto SM agar plates in association with K. aerogenes. Colonies were picked and analysed by PCR. Genomic DNA was prepared from approximately 3 x 10⁸ cells using the GenElute Mammalian Genomic DNA Miniprep Kit (Sigma) according to manufacturer’s protocol. Two screening primers were designed per strain, one placed just upstream of the 5’ homology arm (primer X) and another just downstream of the 3’ homology arm (primer Y) in the genomic sequence. Each of the two primers was paired with a primer of the appropriate sense that bound within the bar cassette (BSR1B and BSR2B). The generation of a product by primer X and BSR1B, and primer Y and BSR2B indicated that the bar cassette integrated into the correct genomic loci.

BSR1B - 5’ – CATTGTAATCTTCTCTGTGCAGTACTTCCGAC – 3’
BSR2B - 5’ - GTGTAGGGAGTTGATTTCAGACTATGGTACCA - 3’

All disrupted strains were confirmed by Southern analyses according to standard protocol. Genomic DNA was extracted using a method adapted from a universal, rapid high-salt extraction protocol [53]. When further genetic manipulation (either gene disruption or in situ tagging) of a knockout strain was required, the bar cassette was removed from the parental strain by transfection with pDEX-NLS-Cre [51] and selecting for G418 resistance (10 μg/ml). After approximately 10 days of selection, resistant cells were cloned out onto SMagar plates in the presence of K. aerogenes and tested for blasticidin (10 μg/ml) and sensitivity in axenic media.

Cell culture

All strains were routinely grown at 22°C in axenic medium [19] supplemented with vitamins (0.1 mg/l B12, 0.02 mg/l Biotin, 0.2 mg/l Riboflavin) in the presence of tetracycline (10 μg/ml) and streptomycin (200 μg/ml), either in tissue culture plates or in conical flasks shaken at 180 rpm (shaken suspension). Strains can also be cultured in association with Rhizobium meliloti on SM agar plates. Strains carrying pDXA-3C-based neoR-expressing plasmids were grown in axenic medium supplemented with G418 (10 μg/ml).

Dictyostelium development

Axenically grown cells in log phase (2.5 x 10⁵ cells/ml) were harvested by centrifugation (200 g, 2 minutes) and washed twice with KK2 buffer (16.5 mM KH₂PO₄, 3.9 mM K₂HPO₄, pH 6.1). Cells were resuspended in KK2 plus 0.1 mM CaCl₂ to 2.5 x 10⁵ cells/ml and 4 ml (10⁶ cells) were plated per agar plate (1.5 x 10⁶ cells/cm²) in duplicate. Cells were allowed to settle on the agar for 15 minutes before the buffer was aspirated. Plates were then incubated in a moist box at 22°C with light. Photographs were taken with a Nikon Coolpix 4500 camera mounted on a Wild M10 microscope at the indicated time points in development.

Colony survival assay

Cells in logarithmic growth phase (2–6 x 10⁶ cells/ml) were harvested, resuspended at 1 x 10⁹ cells/ml in Pt buffer (3 mM NaCl, 1 mM NaPO₄, pH 6.5) and treated with cisplatin (Sigma) or mock-treated for 1 hour at 22°C in shaken suspension in the dark. The cisplatin solution was prepared in the dark immediately prior to use by dissolving in Pt buffer to a concentration of 1 mg/ml (3.5 mM). After treatment, cells were serially diluted in KK2 buffer and 50 μl of two dilutions shown to contain approximately 50 viable cells in preliminary experiments were plated in triplicate on SM agar plates with 100 μg/ml of two-day old K. aerogenes culture. The plates were incubated at room temperature and the number of complete colonies per plate was scored 4 days after plating. An average was taken between the triplicate plates. Viability was calculated as a percentage of the estimated number of cells plated, which was then normalised against that of the mock-treated culture.

Immunoprecipitation

Typically, 10⁷ cells were harvested by centrifugation and washed twice with 1 ml KK2 buffer. The cell pellet was resuspended in 500 μl NETN lysis buffer (20 mM Tris-HCl pH 8.0, 150 mM NaCl, 1 mM EDTA, 1 mM DTT, 0.5% NP40, 10% glycerol, 1 x Protease Inhibitor Cocktail [Roch], 5 mM NEM [Sigma]). The lysate was drawn through a 25G needle four times to ensure complete lysis of the cells and to shear the genomic DNA. The resulting whole cell extract was cleared by centrifugation and the protein concentration was determined by Bradford assay. Protein concentrations across all samples were equalised and the total extract volume was made up to 540 μl with NETN lysis buffer, 1 μl of rabbit polyclonal antibody to GFP (Abcam ab6556) was added and mixed by rotation during an 1 hour incubation at 4°C. 200 μl of 50 mg/ml freshly prepared protein A-agarose beads (GE Healthcare) were then added and the samples mixed and incubated as the previous step. The beads were then pelleted by centrifugation, washed four times with 1 ml NETN lysis buffer and finally resuspended in 100 μl 2 x SDS loading buffer.

Western blotting

Protein samples were run on 10% NuPAGE Bis-Tris pre-cast gels (Invitrogen) in 1 x MOPS buffer (Invitrogen). The separated proteins were transferred onto nitrocellulose membrane (Millipore). After blocking with 5% milk/PBST, the blot was incubated with the appropriate primary and secondary antibody diluted in PBST (PBS with 0.03% v/v Tween-20) for 1 hour each at room temperature. The following antibodies and dilutions were used: rabbit anti-GFP antibody ab6556 (Abcam; 1:2000), goat anti-rabbit IgG HRP-conjugated antibody (Southern Biotech; 1:1000–1:2000), mouse monoclonal anti-HA (clone 12CA5) HRP-conjugated antibody (Roche; 1:1000), rabbit anti-TAP antibody (Sigma-Aldrich; 1:600).

Size exclusion chromatography

2 x 10⁸ exponentially growing cells were harvested and washed three times with KK2 buffer before flash-freezing in liquid nitrogen and storing at −80°C until use. The cell pellet was resuspended in 10 ml high salt buffer (50 mM HEPES pH 7.9, 5 mM MgCl₂, 420 mM NaCl, 0.2 mM EDTA, 25% glycerol, 2 mM DTT, 1 x Protease Inhibitor Cocktail [Roch] on ice. The suspension was taken up in a syringe and forced through a 3 μm Nucleofector filter (Whatmann) and absorbent pad (Millipore) to complete cell lysis, and was subsequently passed through a 26G needle to lyse the nuclei. The resulting lysate was mixed gently at 4°C for 30 minutes to extract nuclear protein and cleared by centrifugation (16,000 g, 10 minutes at 4°C). 2 ml whole cell extract was filtered through a 0.2 μm filter and applied to a Superose 6 XK 16/70 column (GE Healthcare) equilibrated with high salt buffer. 4 ml fractions were collected and 25 μl of each fraction was resolved on 10% Bis-Tris polyacrylamide gels and analysed by Western blotting.
Computational methods

Orthologue searches were done using two publicly available databases – NCBI BLAST Link (BLink) and Kyoto Encyclopaedia of Genes and Genomes (KEGG) Orthology. NCBI BLink – http://www.ncbi.nlm.nih.gov/sites/entrez KEGG Orthology – www.genome.jp/kegg/

PSI-BLAST searches were carried out using the NCBI Blastp suite.

http://blast.ncbi.nlm.nih.gov/Blast.cgi

Structure of the Dictyostelium FncE orthologue was predicted using Phyre.

http://www.sbg.bio.ic.ac.uk/–phyre/

Sequence alignments were carried out using ClustalW [54] and displayed using JalView [http://www.jalview.org/].

Supporting Information

Figure S1 Generation and verification of the ΔfncD2 and ΔfncJ null strains. (A) Generation and verification of the ΔfncD2 strain. Schematic representation of the targeting construct used for knocking out fncD2 (DDB_G0268216) and location of the probes and restriction sites used for Southern blot analysis. This analysis resulted in a 10.7 kb band for WT cells and a 7.1 kb band for ΔfncD2 strains. (B) Generation and verification of the ΔfncJ strain. Schematic representation of the targeting construct used for knocking out fncJ (DDB_G0293476) and location of the probes and restriction sites used for Southern blot analysis. This analysis resulted in a 10.7 kb band for WT cells and a 7.1 kb band for ΔfncJ strains.

Found at: doi:10.1371/journal.pgen.1000645.s001 (2.75 MB TIF)

Figure S2 Generation and verification of the ΔfncM and ΔfncL null strains. (A) Generation and verification of the ΔfncM strain. Schematic representation of the targeting construct used for knocking out fncM (DDB_G0274841) and location of the probes and restriction sites used for Southern blot analysis. This analysis resulted in an 11.2 kb band for WT cells and a 4.6 kb band for ΔfncM strains. (B) Generation and verification of the ΔfncL strain. Schematic representation of the targeting construct used for knocking out fncL (DDB_G0292744) and location of probes and restriction sites used for Southern blot analysis. This analysis resulted in a 13.9 kb band for WT cells and a 4.4 kb band for ΔfncL strains.

Found at: doi:10.1371/journal.pgen.1000645.s002 (5.55 MB TIF)

Figure S3 Generation and verification of the ΔfncJ and Δube2T null strains. (A) Generation and verification of the ΔfncJ strain. Schematic representation of the targeting construct used for knocking out fncJ (DDB_G0286621) and location of the probes and restriction sites used for Southern blot analysis. This analysis resulted in a 9.1 kb band for WT cells and a 4.7 kb band for ΔfncJ strains. (B) Generation and verification of the Δube2T strain. Schematic representation of the targeting construct used for knocking out ube2T (DDB_G0291199) and location of the probes and restriction sites used for Southern blot analysis. This analysis resulted in a 6.3 kb band for WT cells and a 7.3 kb band for Δube2T strains.

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Figure S4 Generation and verification of the ΔfncE strain. (A) ClustalW alignment of the FncE sequences of Homo sapiens, Mus musculus, Gallus gallus, Danio rerio, Arabidopsis thaliana, and Dictyostelium discoideum. The Dictyostelium FncE sequence is highlighted by dashed red lines. (B) Generation and verification of the ΔfncE strain. Schematic representation of the targeting construct used for knocking out fncE (DDB_G0279669) and location of the probes and restriction sites used for Southern blot analysis. (C) Verification of the ΔfncE strain by Southern blot. This analysis resulted in a 4.0 kb band for WT cells and a 5.1 kb band for ΔfncE strains.

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Figure S5 Generation and verification of the TAP-FncL strain. (A) Schematic representation of the targeting construct used for N-terminal in situ tagging of FncL with TAP. (B) Western blot showing TAP-FncL expression and specific detection of FANCL by the anti TAP antibody. Ax2 lysate was included as a negative control. The lysate of 7.5×10^5 cells was loaded per lane.

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Figure S6 Generation and verification of the Δmus81 strain. (A) Schematic representation of the targeting construct used for knocking out mus81 (DDB_G0276519). (B) Verification of Δmus81 clones by Southern blot analysis. This analysis resulted in a 7.1 kb band for WT cells and a 6.0 kb band for Δmus81 strains. (C) The Δmus81 strain is not sensitive to cisplatin as assayed by colony survival. Results shown are from a single experiment. Error bars indicate variation between triplicate plating.

Found at: doi:10.1371/journal.pgen.1000645.s006 (1.42 MB TIF)

Figure S7 The FncD2-GFP strain is not sensitive to cisplatin. The FncD2 C-terminal GFP tagged strain does not show sensitivity to cisplatin.

Found at: doi:10.1371/journal.pgen.1000645.s007 (0.29 MB TIF)

Table S1 Table of all the strains generated and used in this study. The systematic strain name (HMxxxx) is based on the nomenclature used in R. R. Kay’s lab. Parental strain, genotype (∆ = deletion), overexpression plasmid present, and drug resistance of each strain are presented.

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Author Contributions

Conceived and designed the experiments: XYZ JL KJP. Performed the experiments: XYZ JL. Analyzed the data: JL DT KJP. Contributed reagents/materials/analysis tools: DT MMB RRK. Wrote the paper: KJP.

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