

Overexpression of BLM promotes DNA damage and increased sensitivity to platinum salts in triple negative breast and serous ovarian cancers

NJ Birnbak^{1,2,*}, Y Li^{2,*}, S Pathania^{2*}, A Greene-Colozzi², M Dreze², C Bowman-Colin², Z Sztupinski¹, M Krzystanek¹, M Diossy¹, N Tung³, PD Ryan⁴, JE Garber², DP Silver^{2,3}, JD Iglehart^{2,5}, ZC Wang^{2,5}, D Szuts⁶, Z Szallasi^{1,7#}, AL Richardson^{2,5#}

¹ Department of Bio and Health Informatics, Technical University of Denmark, Lyngby, Denmark

² Dana-Farber Cancer Institute, Harvard Medical School, Boston Massachusetts, USA

³Beth Israel Deaconess Medical Center, Harvard Medical School, Boston Massachusetts, USA

⁴Texas Oncology, The Woodlands, Texas, USA

⁵Brigham and Women's Hospital, Harvard Medical School, Boston Massachusetts, USA

⁶Institute of Enzymology, Research Center for Natural Sciences, Hungarian Academy of Sciences, Budapest, Hungary.

⁷ Computational Health Informatics Program (CHIP) Boston Children's Hospital Harvard Medical School, Boston Massachusetts, USA

*These primary authors contributed equally to this work.

#These senior authors contributed equally to this work.

Correspondence should be sent to:

Zoltan Szallasi, Computational Health Informatics Program (CHIP) Boston Children's Hospital, Harvard Medical School, 300 Longwood Ave., Boston Massachusetts, USA, 02215, e-mail: Zoltan.szallasi@childrens.harvard.edu, +1-617-355-2179

or Andrea L. Richardson, Department of Pathology, Johns Hopkins Medicine, Sibley Memorial Hospital, 5255 Loughboro Rd. NW, Washington, DC, 20016, email: aricha58@jhmi.edu, +1-202-660-6309

Background: Platinum based therapy is an effective treatment for a subset of triple negative breast cancer and ovarian cancer patients. In order to increase response rate and decrease unnecessary use, robust biomarkers that predict response to therapy are needed.

Patients and methods: We performed an integrated genomic approach combining differential analysis of gene expression and DNA copy number in sensitive compared to resistant triple negative breast cancers in two independent neoadjuvant cisplatin treated cohorts. Functional relevance of significant hits was investigated *in vitro* by overexpression, knockdown and targeted inhibitor treatment.

Results: We identified two genes, the Bloom helicase (BLM) and Fanconi anemia complementation group I (FANCI), that have both increased DNA copy number and gene expression in the platinum sensitive cases. Increased level of expression of these two genes was also associated with platinum but not with taxane response in ovarian cancer. As a functional validation, we found that overexpression of BLM promotes DNA damage and induces sensitivity to cisplatin, but has no effect on paclitaxel sensitivity.

Conclusions: A biomarker based on the expression levels of the BLM and FANCI genes is a potential predictor of platinum sensitivity in triple negative breast cancer and ovarian cancer.

Short description

Through integrated analysis of gene expression and copy number data from two independent clinical trials in triple negative breast cancer, we identify two genes, BLM and FANCI, involved in double-strand DNA repair where increased expression is related to sensitivity to platinum induced DNA damage. Further functional validation reveals that overexpression of BLM alone promotes DNA damage.

Key words: platinum based chemotherapy, gene expression based predictor of treatment sensitivity, triple negative breast cancer, ovarian cancer,

Introduction

BRCA1 plays an important role in response to replication stress and repair of stalled or collapsed replication forks, and complete absence of functional BRCA1 leads to defective error-free homologous recombination-type double strand break repair. BRCA1^{-/-} tumors are particularly sensitive to platinum-containing chemotherapy and inhibitors of PARP1, whereas BRCA wild-type cancers showed a more limited response to these agents [1]. Platinum salts generate both interstrand and intrastrand crosslinks that slow or stall replication forks [2]. Stalled replication forks may collapse into double-strand breaks (DSBs) and/or

become sites for translesional synthesis-induced mutagenesis, potentially causing genome instability.

Platinum-sensitive triple-negative breast cancers (TNBC) and serous ovarian cancers carry extensive genomic rearrangements and allelic imbalance suggesting these cancers may share similar defects in DNA repair acquired through alternative mechanisms than through BRCA1 loss [3]. The overall level of such genomic aberrations can be characterized and quantified by a recently developed clinical measure, the “HRD score” [3-5]. These results suggest that platinum sensitivity may be related to a functional defect that occurs when BRCA1 levels are insufficient and a biomarker indicative of such defects may be predictive of sensitivity to DNA cross-linkers such as platinum agents.

To explore and define specific molecular alterations that might be associated with cisplatin sensitivity, we combined differential analysis of gene expression and DNA copy number in cisplatin sensitive compared to cisplatin resistant TNBC.

Materials and Methods

Breast cancer cohorts

This study is based on previously published clinical and molecular data from two cisplatin-treated triple negative breast cancer trials, Cisplatin 1 and Cisplatin 2 [3, 6]. SNP data is available from the National Center for Biotechnology Information Gene Expression Omnibus (GEO) at GSE28330, and RNA expression data for Cisplatin 1 is available GEO at GSE18864.

Data acquisition and data generation

Gene expression data based on the Affymetrix HGU133plus2 platform was generated from a subset of the Cisplatin 2 cohort from which sufficient material was available as described previously [6] with the exception that the samples were not subjected to double amplification. Data available from GEO at GSE103668. Gene expression data from the ovarian cancer trial OV-01 [7] based on the Affymetrix HGU133A platform was acquired from GEO at GSE15622.

Cell lines

Breast cancer cell lines MDA-MB-231, MDA-MB-436, MDA-MB-453, HCC38, HCC1143, HCC1937 and BT549 were cultured *in vitro* and subjected to DNA damage. Transfections of siRNA and plasmid DNA were performed using Lipofectamine RNAiMAX or Lipofectamine 2000 (Life Technologies) respectively.

For full experimental details, see supplementary methods.

Results

Identification of genes associated with cisplatin response

We performed a leave-one-out (LOO) differential gene expression analysis in two independent cisplatin-treated TNBC cohorts separately to identify genes significantly associated with response. Permutation testing identified 12 genes where expression was significantly associated with platinum response in both cisplatin TNBC cohorts (Table S1, Figure 1A, Fig. S1A). A similar leave-one-out analysis of the DNA copy numbers were performed. This identified

234 genes associated with cisplatin response in both cisplatin TNBC cohorts (Table S2, Figure 1B, Fig. S1B). Only two genes were identified in both analyses for association with platinum sensitivity, the Bloom helicase (*BLM*) and the Fanconi anemia complementation group I (*FANCI*) genes, both located at chromosome 15q26. The copy number of both genes was significantly higher in the cisplatin-sensitive tumors in both TNBC cohorts (BLM: cisplatin-1, $p=0.003$, cisplatin-2, $p=0.008$, FANCI: cisplatin-1, $p=0.003$, cisplatin-2, $p=0.003$, Figure S1C). Both BLM and FANCI showed DNA copy number gain in 33% of sensitive versus 0% of resistant tumors in Cisplatin-1, and gain in 44% of sensitive versus 12% of resistant tumors in Cisplatin-2. Similarly, in both cohorts cisplatin-sensitive tumors had significantly higher BLM and FANCI mRNA expression (BLM: cisplatin-1, $p = 0.0028$; cisplatin-2, $p = 0.0075$; Fig 1C-D, FANCI: cisplatin-1, $p = 0.0036$; cisplatin-2, $p = 0.0125$; Fig 1E-F).

As validation, gene expression levels of BLM and FANCI as measured by Affymetrix U133 microarray were significantly correlated with results obtained by qRT-PCR for the same samples (BLM, $r = 0.87$; FANCI, $r = 0.76$; Fig. S1D-E). Western blot analysis in protein extracts from a series of tumor samples (Figure S1H) with matched array-based mRNA measurements showed good correlation between mRNA and protein levels for BLM (Spearman $r = 0.68$, $p = 0.0023$, Figure S1F), but not for FANCI (Spearman $r = -0.01$, $P = 0.97$, Figure S1G).

We found the expression level of BRCA1 transcript as measured by qRT-PCR significantly associated with cisplatin resistance [3, 6]. When we tested association of the ratio of average of BLM and FANCI levels divided by BRCA1 levels from microarray expression versus cisplatin response, the ratio was

significantly higher in the cisplatin sensitive tumors in both cohorts (cisplatin-1, $p = 0.0230$; cisplatin-2, $p = 0.0023$; Figure 1G-H).

To further validate these findings, we acquired a publicly available gene expression data set from a serous ovarian cancer trial of either carboplatin monotherapy or paclitaxel monotherapy [7]. The average expression levels of BLM and FANCI were significantly higher in the carboplatin-sensitive ovarian cancers ($p = 0.026$ and $p = 0.036$ respectively, Figure 1I-J). The ratio of BLM+FANCI/BRCA1 was also significantly higher in carboplatin sensitive ovarian cancers ($p = 0.026$, Figure 1K). Altogether, these data indicate that the expression levels of BRCA1 are inversely correlated with those of BLM and FANCI. Moreover, the association of BLM and FANCI with paclitaxel response was not significant and the trend was in the opposite direction (Figure 1L-N), suggesting that the correlations might be cisplatin- or drug class-specific.

The performance of the ratio of BLM+FANCI/BRCA1 to predict platinum response was also compared to previously published genomic scar or gene expression based signatures in the cisplatin-1 and cisplatin-2 cohorts. This gene expression ratio performed better than the previously published predictors (Supplementary material, Figure S2-4). The BLM+FANCI/BRCA1 expression ratio also performed better in the platinum monotherapy treated serous ovarian cancer cohort [7] and a platinum treated gastric cancer cohort (supplementary material, Figure S5 and S7), but had no predictive power in non-platinum treated ovarian cancer samples (Figure S6). It did not predict response to therapy in non-platinum treated neoadjuvant triple negative breast cancer cohorts and it did not have prognostic power in non-platinum treated breast cancer cohorts (see supplementary material, figures S8-S14).

Relationship of BLM and BRCA1 expression and sensitivity to DNA damaging agents in cell lines

BLM and BRCA1 protein levels were measured in a panel of breast cancer cell lines by Western blot analysis (Figure 2A) and densitometry quantitation of expression ratios are displayed in bar plots (Figure 2B, C). Two cell lines without *BRCA1* sequence mutations (BT549, HCC1143) and the HCC38 cell line in which BRCA1 expression is suppressed by methylation displayed high BLM/actin and low BRCA1/BLM expression ratios. Two other BRCA1wt cell lines (MDA231 and MDA453) displayed relatively lower BLM/actin and higher BRCA1/BLM expression ratios. On the other hand, HCC1937 and MDA436, which are known to be BRCA1-null, displayed high levels of BLM expression normalized to actin along with very low BRCA1/BLM expression ratios. This cell line panel was evaluated for sensitivity to various treatments by colony formation assay. The three panels of Figure 2 D-F show that the pattern of sensitivity to cisplatin, UV radiation treatment, and PARP inhibitor olaparib across the panel of cell lines is associated with the pattern of relative expression of BRCA1/BLM and BLM/actin. The two sequence-mutated BRCA1 cell lines and the three cell lines with low BRCA1/BLM and high BLM/actin have greater sensitivity to all three treatments. The two cell lines with low BLM and high BRCA1/BLM (MDA231, MDA453) are relatively resistant to the same treatments. In contrast, there is no apparent relationship between BLM and BRCA1 expression with the pattern of sensitivity to the microtubule stabilizer paclitaxel (Fig 2G). These data suggest that BRCA1/BLM protein ratio may be predictive of the sensitivity to canonical DNA

damaging agents (e.g. cisplatin, UV, and olaparib) but not necessarily to agents like paclitaxel that work through distinct mechanisms.

Effect of modulating BRCA1 expression on BLM and FANCI expression

In order to elucidate the correlation between BRCA1, BLM and FANCI expression levels, we knocked down endogenous BRCA1 in U2OS cells in which the role of BRCA1 in the double-strand break repair (DSBR) response has been well studied. After one week of treatment following transfection of BRCA1-specific shRNA, the expression of BRCA1, BLM, and FANCI were measured by Western blot analysis (Figure 2I). Cells in which BRCA1 was knocked-down showed increased expression of BLM protein compared to control cells treated with shRNA specific to luciferase (Figure 2I, left panel). Given that BRCA1 is a known negative transcriptional regulator of BLM [8], it is possible that the increase in BLM protein levels in BRCA1 depleted cells is a direct consequence of this negative regulation. On the other hand, the expression levels of FANCI did not change upon knockdown of BRCA1 by shRNA (Figure 2I, right panel).

Effect of modulating BLM and FANCI expression on sensitivity to cisplatin and taxanes

Since BLM showed a good correlation between mRNA and protein expression levels and was up-regulated by down-regulation of BRCA1, further functional validation was performed. We performed knockdown experiments in BT549 breast cancer cells, which inherently express high levels of BLM. Gene-specific siRNA treatment resulted in reduced protein expression of BLM as determined by Western blot (Fig 2J). In these cells, the IC50s for cisplatin and paclitaxel were

determined by colony formation assay. The knockdown of either BLM or FANCI by siRNA resulted in greater resistance to cisplatin but no significant effect on sensitivity to paclitaxel (Fig 2K). Given that cell cycle state can influence chemosensitivity, we examined whether a change in cell cycle upon BLM and/or FANCI depletion could explain the decreased chemosensitivity to cisplatin. Neither loss of BLM (siBLM) or FANCI (siFANCI) had a significant effect on the cell cycle state (Figure S15, A & B). This further suggests a direct link between BLM and/or FANCI levels and sensitivity to DNA damaging agents.

We tested the reverse hypothesis by overexpressing BLM using an HA-tagged lentivirus expression vector in MDA231 cells, which display low levels of BLM and relative resistance to cisplatin. Overexpression was assessed by Western blot analysis for endogenous BLM or for the HA-tag (Figure 2L). In order to address the specificity of this effect, we used siRNA-mediated knockdown of BLM, which reduced the expression levels of both endogenous and HA-tagged BLM protein. We also tested the effect of adding a small molecule inhibitor of BLM (BLMi), which can suppress BLM activity without affecting BLM protein levels (Fig 2L, lane 3,4 and Fig S16). As shown in Fig. 2M, overexpression of BLM resulted in decrease in the IC50 (greater sensitivity) to cisplatin. This effect was reversed by treatment with the BLM helicase inhibitor (BLMi) and by siRNA knockdown of BLM. These results suggest that increased BLM expression levels and specifically BLM helicase activity promotes increased sensitivity to cisplatin.

BLM overexpression increases spontaneous and cisplatin-induced DNA damage

To determine if BLM expression levels affect the accumulation of DNA damage in breast cancer cells, we performed immunofluorescence for markers of DNA damage in MDA231 cells in which BLM expression levels or activity was modulated. Overexpression of BLM resulted in no detectable difference in spontaneous or cisplatin-induced BRCA1 foci, but did suppress RAD51 focus formation (Figure S17). This is consistent with reports that BLM can displace RAD51 localization at damaged replication forks [9]. Since RAD51 is required for homologous recombination-dependent and -independent DSBR mechanisms at stalled replication forks [10], such loss of RAD51 recruitment to sites of DNA damage in BLM overexpressing cells could further explain the increased sensitivity of BLM overexpressing cells to stalled fork-inducing agents like cisplatin (Fig. 2M).

Overexpression of BLM also resulted in increased spontaneous DNA damage as evidenced by increased γ -H2AX and phospho-53BP1 (53BP1-p) foci in the absence of any drug treatment (Figure S18, Figure 3A, black bars in Fig. 3C, D). This increase in DNA damage is greater in cells after 4 hours treatment with cisplatin (Figure 3B, grey bars in Fig. 3C, D). The addition of a small molecule BLM helicase inhibitor (BLMi) or siRNA to BLM (siBLM) blocks the effect of BLM overexpression on DNA damage foci. The accumulation of endogenous DNA damage (as seen by γ -H2AX and 53BP1-p foci in untreated cells) as well as cisplatin-induced DNA damage in BLM overexpressing cells, further explains the increased sensitivity of BLM overexpressing cells to cisplatin (Fig. 2M).

Discussion

In this study, we identified two neighboring genes, BLM and FANCI, with consistent copy gain and overexpression in cisplatin sensitive primary breast tumors. The association between this three-gene signature of high BLM+FANCI/BRCA1 and platinum sensitivity was confirmed in an independent cohort of serous ovarian cancer treated with single agent carboplatin. Through *in vitro* studies, we found that chronic repression of BRCA1 expression in cell lines led to an increase in BLM expression and no change in FANCI expression suggesting a possible compensatory or direct transcriptional effect on BLM in the setting of low or insufficient BRCA1. Our findings are consistent with previous reports that BRCA1 regulates transcription of BLM in prostate cancer cell lines with overexpression of BRCA1 resulting in down regulation of BLM [8].

BLM has recently been shown to play an important role at stalled replication forks [11, 12]. Unlike BRCA1, which is required for the stability of stalled replication forks, loss of BLM does not result in its degradation. Instead, BLM has been shown to help restart the stalled forks while suppressing firing of new origins in response to replication stress. An increase in spontaneous DNA damage, especially accumulation of 53BP1-p foci in BLM overexpressing cells (Fig. 3), is suggestive of degradation and/or collapse of spontaneously arising stalled replication forks.

Homologous recombination (HR) is required for repair of cisplatin-induced damage, as the loss of HR factors such as BRCA1 and BRCA2 increases cisplatin sensitivity [13, 14]. Both pro- and anti-HR roles have been described for BLM. Its function in DNA end resection and the ability to promote DNA repair synthesis in D-loops supports HR, while the disruption of RAD51 filament formation, the disruption of D-loops and the dissolution of double Holliday

junctions opposes HR [15-18]. One hypothesis to explain these results is that at increased BLM levels the anti-HR functions of the protein dominate, leading to increased cisplatin sensitivity.

Reduced HR in BLM overexpressing cells should cause impaired repair of DSBs generated at cisplatin interstrand crosslinks. However, cisplatin-induced single strand lesions are much more common [19], therefore the observed damage foci may mostly arise at replication forks that encounter single strand lesions. It has been shown previously that RAD51 is required for efficient repair and restart of stalled replication forks in an HR dependent and independent manner [10]. Disruption of RAD51 filament formation by BLM could antagonize post-replication repair promoted by BRCA1, consistent with a BLM function that requires helicase activity, and explaining the similarity of the consequences of low BRCA1 or high BLM. Given that we do not see a disruption of BRCA1 foci in BLM-overexpressing cells despite an apparent reduction in Rad51 foci formation, we cannot rule out that cisplatin sensitivity observed in BLM-overexpressing cells is at least in part related to a BRCA1-independent pathway wherein loss of RAD51 at the sites of DNA damage sensitizes the BLM overexpressing cells to cisplatin.

Funding

ALR, NT, JEG, JDI, ZCW and ZS are supported by the Breast Cancer Research Foundation. ZS is supported by the Basser Foundation, EU FP7 project PREDICT and by the NovoNordisk Foundation (ID 16854). DS is supported by Momentum

Grant LP2011-015 of the Hungarian Academy of Sciences. NJB is supported by the The Villum Kann Rasmussen Foundation. Analysis of the cisplatin-2 samples was supported by a grant to JEG from the V-Foundation.

Conflict of interest

NJB, ZCW, ZS and ALR are inventors on a patent application for the use of the ratio of BLM+FANCI/BRCA1 to predict response to chemotherapy. All remaining authors have declared no conflicts of interest.

REFERENCES

1. Basourakos SP, Li L, Aparicio AM et al. Combination Platinum-based and DNA Damage Response-targeting Cancer Therapy: Evolution and Future Directions. *Curr. Med. Chem.* 2017; 24(15):1586–1606.
2. Sokol AM, Cruet-Hennequart S, Pasero P, Carty MP. DNA polymerase η modulates replication fork progression and DNA damage responses in platinum-treated human cells. *Sci Rep* 2013; 3(1):3277.
3. Birkbak NJ, Wang ZC, Kim J-Y et al. Telomeric allelic imbalance indicates defective DNA repair and sensitivity to DNA-damaging agents. *Cancer Discov* 2012; 2(4):366–375.
4. Abkevich V, Timms KM, Hennessy BT et al. Patterns of genomic loss of heterozygosity predict homologous recombination repair defects in epithelial ovarian cancer. *Br. J. Cancer* 2012; 107(10):1776–1782.
5. Popova T, Manié E, Rieunier G et al. Ploidy and large-scale genomic instability consistently identify basal-like breast carcinomas with BRCA1/2 inactivation. *Cancer Res* 2012; 72(21):5454–5462.
6. Silver DP, Richardson AL, Eklund AC et al. Efficacy of neoadjuvant

- Cisplatin in triple-negative breast cancer. *J. Clin. Oncol.* 2010; 28(7):1145–1153.
7. Ahmed AA, Mills AD, Ibrahim AEK et al. The extracellular matrix protein TGFBI induces microtubule stabilization and sensitizes ovarian cancers to paclitaxel. *Cancer Cell* 2007; 12(6):514–527.
 8. De Luca P, Vazquez ES, Moiola CP et al. BRCA1 loss induces GADD153-mediated doxorubicin resistance in prostate cancer. *Mol. Cancer Res.* 2011; 9(8):1078–1090.
 9. Ouyang KJ, Woo LL, Zhu J et al. SUMO modification regulates BLM and RAD51 interaction at damaged replication forks. *PLoS Biol.* 2009; 7(12):e1000252.
 10. Petermann E, Orta ML, Issaeva N et al. Hydroxyurea-stalled replication forks become progressively inactivated and require two different RAD51-mediated pathways for restart and repair. *Mol. Cell* 2010; 37(4):492–502.
 11. Davies SL, North PS, Hickson ID. Role for BLM in replication-fork restart and suppression of origin firing after replicative stress. *Nat. Struct. Mol. Biol.* 2007; 14(7):677–679.
 12. Pichierri P, Franchitto A, Rosselli F. BLM and the FANC proteins collaborate in a common pathway in response to stalled replication forks. *EMBO J.* 2004; 23(15):3154–3163.
 13. Raaphorst GP, Leblanc M, Li LF. A comparison of response to cisplatin, radiation and combined treatment for cells deficient in recombination repair pathways. *Anticancer Res.* 2005; 25(1A):53–58.
 14. Qing Y, Yamazoe M, Hirota K et al. The epistatic relationship between BRCA2 and the other RAD51 mediators in homologous recombination. *PLoS Genet.* 2011; 7(7):e1002148.
 15. Cejka P, Cannavo E, Polaczek P et al. DNA end resection by Dna2-Sgs1-RPA and its stimulation by Top3-Rmi1 and Mre11-Rad50-Xrs2. *Nature* 2010; 467(7311):112–116.
 16. Nimonkar AV, Genschel J, Kinoshita E et al. BLM-DNA2-RPA-MRN and EXO1-BLM-RPA-MRN constitute two DNA end resection machineries for human DNA break repair. *Genes Dev.* 2011; 25(4):350–362.
 17. Bugreev DV, Yu X, Egelman EH, Mazin AV. Novel pro- and anti-recombination activities of the Bloom's syndrome helicase. *Genes Dev.* 2007; 21(23):3085–3094.
 18. Karow JK, Constantinou A, Li JL et al. The Bloom's syndrome gene product promotes branch migration of holliday junctions. *Proc. Natl. Acad. Sci. U.S.A.* 2000; 97(12):6504–6508.

19. Wang D, Lippard SJ. Cellular processing of platinum anticancer drugs. *Nat Rev Drug Discov* 2005; 4(4):307–320.

Figure legends

Figure 1. BLM and FANCI are significantly associated with response to cisplatin chemotherapy.

(A-B) Scatter-plots of leave-one-out analysis in Cisplatin-1 and Cisplatin-2 cohorts identifies BLM and FANCI as the only two genes that show significant association with response in both clinical trials, based on both gene expression data **(A)** and copy number aberration data **(B)**. Red dashed lines indicate significance thresholds based on permutation testing. Color intensity indicates overlapping genes. **(C-N)** Association between array expression of BLM, FANCI, and the ratio of BLM+FANCI expression to BRCA1, and response to cisplatin chemotherapy in Cisplatin-1 **(C, E, G)**, Cisplatin-2 **(D, F, H)** and OV01 carboplatin treated **(I, J, K)** and OV01 paclitaxel treated **(L, M, N)** cohorts. Red dots indicate the BRCA1 mutant cases.

Figure 2. Relationships of BLM, FANCI and BRCA1 expression levels in breast cancer cell lines and association with therapy sensitivities. **(A)**

Western blot analysis of BLM, BRCA1 and Actin protein abundance in a panel of breast cancer cell lines. BT549, HCC1143, HCC38, MDA231 and MDA453 are BRCA1 wildtype genotype. HCC1937 and MDA436 are BRCA1 mutated **(B)** Densitometry of Western blot in panel A for quantification of BLM/Actin and **(C)** BRCA1/BLM ratio. **(D-G)** Bar plots of IC50 to treatments in panel of cell lines. Breast cancer cells lines were irradiated with increasing doses of UV-C or subjected to cisplatin, olaparib or paclitaxel treatment, and four weeks later

assayed for colony formation. Error bars represent the standard deviation between three independent experiments. **(D)** IC₅₀ to cisplatin **(E)** IC₅₀ to UV-C **(F)** IC₅₀ to PARP-inhibitory olaparib **(G)** IC₅₀ to paclitaxel. **(I)** Western blot showing the effect of shRNA BRCA1 or shLuciferase control (shLuc) on expression of BLM (left side) and FANCI (right side) in U2OS cells. GAPDH is shown as a loading control. **(J)** Western blot demonstrating gene specific siRNA knockdown of BLM or FANCI expression in BT549 breast tumor cells. **(K)** Bar plots indicate the ratio of IC₅₀ for the cisplatin (black bars) and paclitaxel (grey bars) in gene-specific siRNA treated cells, relative to scramble control siRNA treated cells. **(L)** Western blot for endogenous BLM and HA-tag in MDA231 cells transfected with control vector (lane 1), HA-tag BLM cDNA (BLM, lane2), HA-tag BLM co-treated with BLM helicase small molecule inhibitory (BLM+BLMi, lane 3), and HA-tag BLM co-treated with 50 nM siRNA BLM (BLM+siBLM, lane 4). **(M)** Bar plots indicate the ratio of IC₅₀ for the cisplatin in MDA231 cells treated with control vector (black bar), HA-tag BLM cDNA (BLM, medium grey bar), HA-tag BLM co-treated with BLM helicase small molecule inhibitory (BLM+BLMi, dark grey bar), and HA-tag BLM co-treated with 50 nM siRNA BLM (BLM+siBLM, light grey bar).

Figure 3. Increased DNA damage upon BLM overexpression in MDA231 cells. MDA231 cells were infected with control vector or HA-tag BLM cDNA (BLM) and co-treated with 20 μ M BLM small molecule inhibitory (Bi) or 100nM siBLM (si). Cells were mock treated (cisplatin: 0h, panel **A**) or treated with 10 μ M cisplatin for 4 hours (cisplatin: 4h, panel **B**) and released for 24 hours.

Immunofluorescence for phospho-H2Ax and phospho-53BP1 was performed and nuclei counterstained with DAPI. **(A)** Representative immunofluorescent images for indicated markers in mock treated cells (cisplatin: 0h) indicating spontaneous DNA damage foci **(B)** Representative immunofluorescent image in cells treated with 10 μ M cisplatin for 4 hours (cisplatin: 4h) indicating drug induced damage foci. All images were obtained at the same magnification and exposure time. All images were analyzed in parallel for each experiment. **(C and D)** Cells containing foci recognized by relevant antibodies in immunofluorescence assays, were identified and counted. At least 100 cells were counted for each category of foci at each time point. Bar plots indicate percentages of cells, noted above, that contain γ H2AX-p foci **(C)** and PB53-p foci **(D)** in mock treated cells (cisplatin: 0h, black bars) and 10 μ M cisplatin treated cells (cisplatin: 4h, grey bars).

Key message

Through integrated analysis of gene expression and copy number data from two independent clinical trials in triple negative breast cancer, we identify two genes, BLM and FANCI, involved in double-strand DNA repair where increased expression is related to sensitivity to platinum induced DNA damage. Further functional validation reveals that overexpression of BLM alone promotes DNA damage.

Figure 1

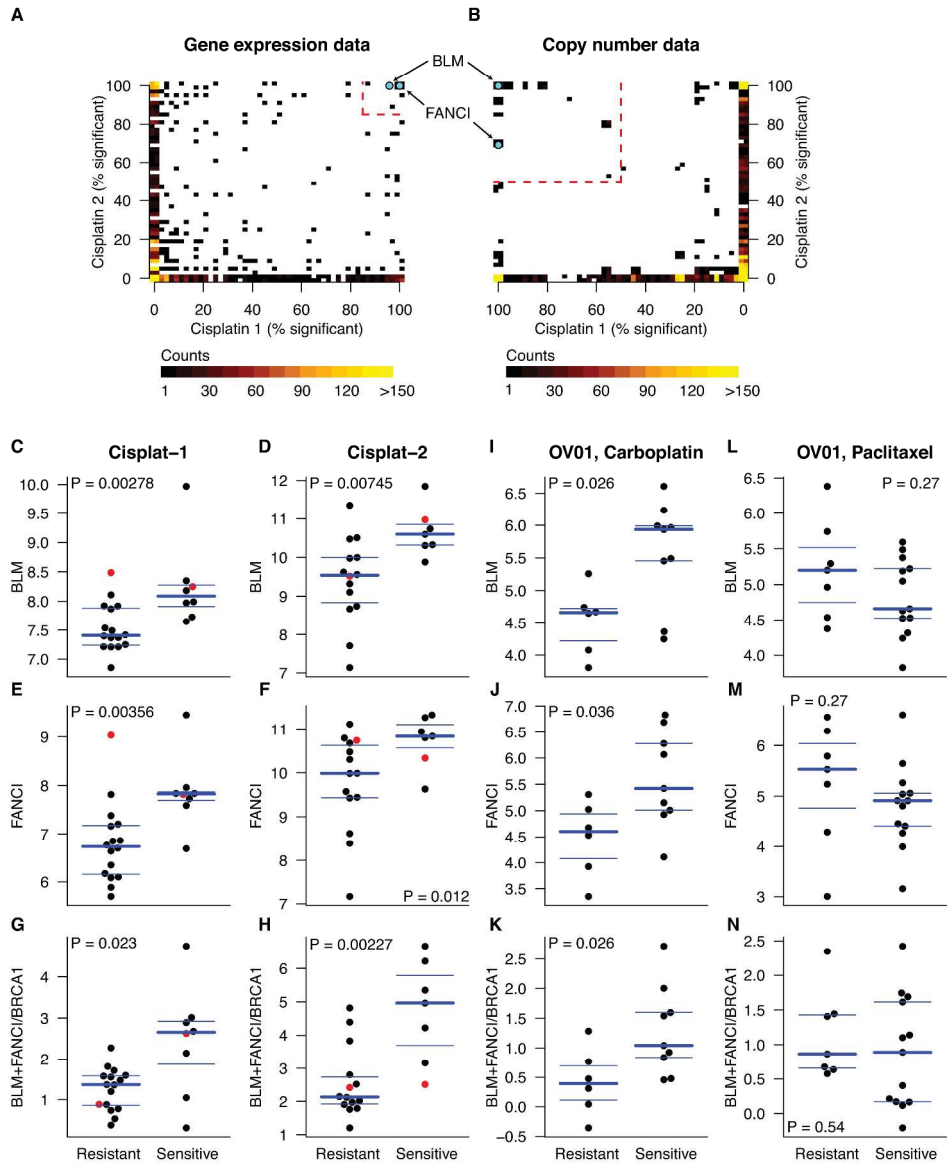


Figure 1. BLM and FANCI are significantly associated with response to cisplatin chemotherapy. (A-B) Scatter-plots of leave-one-out analysis in Cisplatin-1 and Cisplatin-2 cohorts identifies BLM and FANCI as the only two genes that show significant association with response in both clinical trials, based on both gene expression data (A) and copy number aberration data (B). Red dashed lines indicate significance thresholds based on permutation testing. Color intensity indicates overlapping genes. (C-N) Association between array expression of BLM, FANCI, and the ratio of BLM+FANCI expression to BRCA1, and response to cisplatin chemotherapy in Cisplatin-1 (C, E, G), Cisplatin-2 (D, F, H) and OV01 carboplatin treated (I, J, K) and OV01 paclitaxel treated (L, M, N) cohorts. Red dots indicate the BRCA1 mutant cases.

329x421mm (300 x 300 DPI)

Figure 2

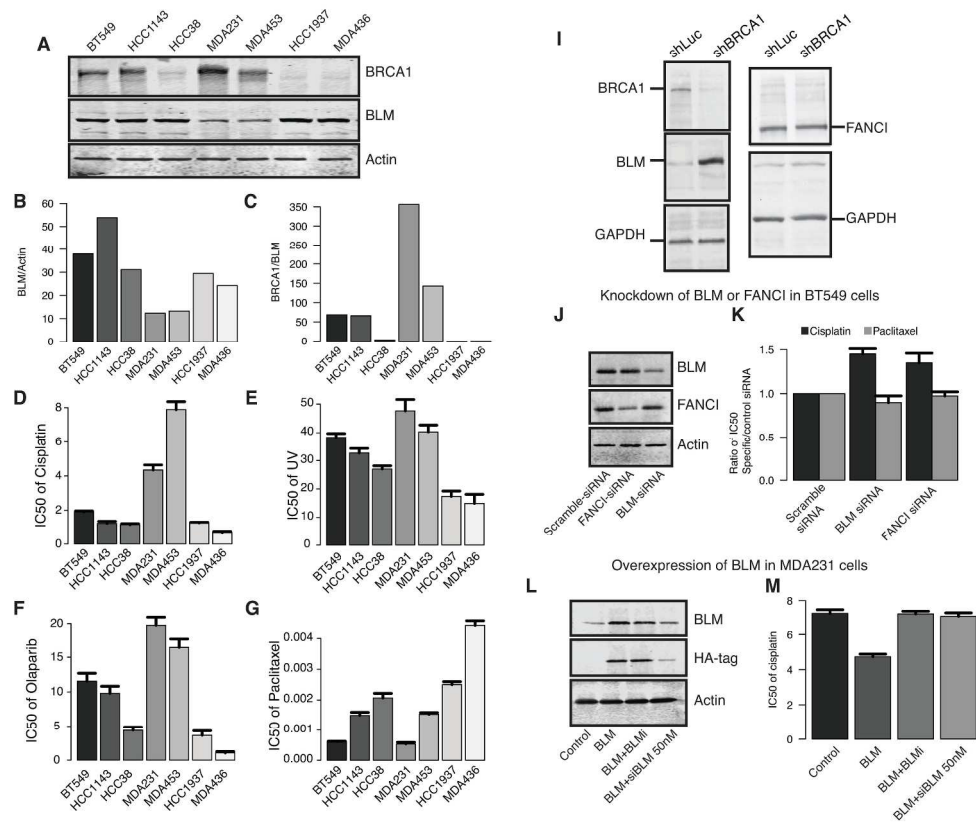


Figure 2. Relationships of BLM, FANCI and BRCA1 expression levels in breast cancer cell lines and association with therapy sensitivities. (A) Western blot analysis of BLM, BRCA1 and Actin protein abundance in a panel of breast cancer cell lines. BT549, HCC1143, HCC38, MDA231 and MDA453 are BRCA1 wildtype genotype. HCC1937 and MDA436 are BRCA1 mutated (B) Densitometry of Western blot in panel A for quantification of BLM/Actin and (C) BRCA1/BLM ratio. (D-G) Bar plots of IC50 to treatments in panel of cell lines. Breast cancer cells lines were irradiated with increasing doses of UV-C or subjected to cisplatin, olaparib or paclitaxel treatment, and four weeks later assayed for colony formation. Error bars represent the standard deviation between three independent experiments. (D) IC50 to cisplatin (E) IC50 to UV-C (F) IC50 to PARP-inhibitory olaparib (G) IC50 to paclitaxel. (I) Western blot showing the effect of shRNA BRCA1 or shLuciferase control (shLuc) on expression of BLM (left side) and FANCI (right side) in U2OS cells. GAPDH is shown as a loading control. (J) Western blot demonstrating gene specific siRNA knockdown of BLM or FANCI expression in BT549 breast tumor cells. (K) Bar plots indicate the ratio of IC50 for the cisplatin (black bars) and paclitaxel (grey bars) in gene-specific siRNA treated cells, relative to scramble control siRNA treated cells. (L) Western blot for endogenous BLM and HA-tag in MDA231 cells transfected with control vector (lane 1), HA-tag BLM cDNA (BLM, lane2), HA-tag BLM co-treated with BLM helicase small molecule inhibitory (BLM+BLMi, lane 3), and HA-tag BLM co-treated with 50 nM siRNA BLM (BLM+siBLM, lane 4). (M) Bar plots indicate the ratio of IC50 for the cisplatin in MDA231 cells treated with control vector (black bar), HA-tag BLM cDNA (BLM, medium grey bar), HA-tag BLM co-treated with BLM helicase small molecule inhibitory (BLM+BLMi, dark grey bar), and HA-tag BLM co-treated with 50 nM siRNA BLM (BLM+siBLM, light grey bar).

257x222mm (300 x 300 DPI)

Figure 3

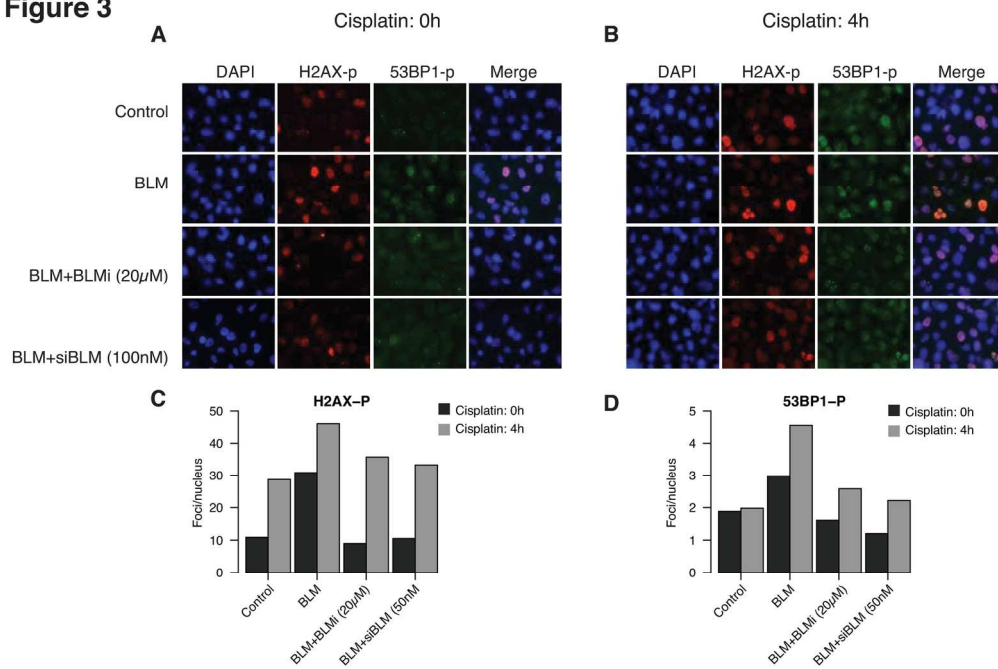


Figure 3. Increased DNA damage upon BLM overexpression in MDA231 cells. MDA231 cells were infected with control vector or HA-tag BLM cDNA (BLM) and co-treated with 20 µM BLM small molecule inhibitory (Bi) or 100nM siBLM (si). Cells were mock treated (cisplatin: 0h, panel A) or treated with 10 µM cisplatin for 4 hours (cisplatin: 4h, panel B) and released for 24 hours. Immunofluorescence for phospho-H2Ax and phospho-53BP1 was performed and nuclei counterstained with DAPI. (A) Representative immunofluorescent images for indicated markers in mock treated cells (cisplatin: 0h) indicating spontaneous DNA damage foci (B) Representative immunofluorescent image in cells treated with 10µM cisplatin for 4 hours (cisplatin: 4h) indicating drug induced damage foci. All images were obtained at the same magnification and exposure time. All images were analyzed in parallel for each experiment. (C and D) Cells containing foci recognized by relevant antibodies in immunofluorescence assays, were identified and counted. At least 100 cells were counted for each category of foci at each time point. Bar plots indicate percentages of cells, noted above, that contain γH2AX-p foci (C) and PB53-p foci (D) in mock treated cells (cisplatin: 0h, black bars) and 10µM cisplatin treated cells (cisplatin: 4h, grey bars).

172x116mm (300 x 300 DPI)